Chapter 10

UPPER EXTREMITY MOTOR REHABILITATION INTERVENTIONS

EBRSR



HEART & STROKE FOUNDATION Canadian Partnership for Stroke Recovery

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Key points

Bobath concept approaches and motor relearning programmes may not be beneficial for upper limb rehabilitation following stroke.

Brunnstrom movement therapy may be more beneficial than motor relearning programmes for upper limb function.

The literature is mixed regarding bilateral arm training for upper limb rehabilitation following stroke.

Bilateral arm training may not be beneficial compared to unilateral training for upper limb function.

Bilateral arm training in combination with other therapy approaches may not be beneficial for upper limb rehabilitation.

The literature is mixed regarding strength training and functional strength training for upper limb rehabilitation following stroke.

The literature is mixed regarding strength training and functional strength training for upper limb rehabilitation following stroke.

Task-specific training, alone or in combination with other therapy approaches, may be beneficial for some aspects of upper limb function following stroke.

Higher and lower intensity task-specific training may have similar effects on upper limb function.

Constraint-induced movement therapy may be beneficial for upper limb rehabilitation in the chronic phase following stroke.

The literature is mixed regarding constraint-induced movement therapy for upper limb rehabilitation in the subacute/acute phase following stroke.

Modified constraint-induced movement therapy may be beneficial for upper limb rehabilitation in the chronic phase following stroke.

Modified constraint-induced movement therapy may not be beneficial for upper limb rehabilitation in the subacute/acute phase following stroke.

Higher and lower intensity constraint-induced movement therapy may have similar effects on upper limb function in the chronic phase following stroke.

The literature is mixed regarding constraint-induced movement therapy in combination with other therapy approaches for upper limb rehabilitation following stoke.

Trunk restraint with reaching training or distributed constraint induced therapy may improve some aspects of upper limb function following stroke, but the effect of combining trunk restraint with constraint-induced movement therapy is less clear. Stretching programs may improve some aspects of upper limb function following stroke.

Orthotics may not be beneficial for upper limb rehabilitation following stroke.

Mirror therapy on its own or in combination with other interventions can improve many aspects of upper limb function following stroke.

Mental practice, alone or in combination with constraint-induced movement therapy, may be beneficial for upper limb rehabilitation following stroke.

Mental practice in combination with virtual reality training may not be beneficial for upper limb function.

Action observation may be beneficial for some aspects of upper limb function following stroke.

The literature is mixed regarding music therapy for upper limb rehabilitation following stroke.

The literature is mixed regarding telerehabilitation for upper limb rehabilitation following stroke.

The evidence is mixed regarding arm/shoulder end-effector robotics, alone or in combination with other therapy approaches, for upper limb rehabilitation following stroke.

The evidence is mixed regarding arm/shoulder exoskeleton, hand exoskeleton, and hand endeffector robotics for upper limb rehabilitation.

Virtual therapy alone may not be more beneficial than conventional therapy for upper limb rehabilitation following stroke, however it may be beneficial for certain aspects of upper limb function when used in combination with conventional or other therapy approaches.

The literature is mixed regarding brain-computer interface technology for upper limb motor rehabilitation following stroke, either on its own or combined with other therapies, but it may not be beneficial alone for other aspects of upper limb function.

The literature is mixed regarding EMG biofeedback alone for upper limb rehabilitation following stroke, however it may not be beneficial when combined with other therapy approaches.

The literature is mixed regrading cyclic and EMG-triggered neuromuscular electrical stimulation types, as well as functional electrical stimulation, alone or combined with other therapy approaches, for upper limb rehabilitation following stroke.

The various types of neuromuscular electrical stimulation may not be more beneficial compared to one another.

Transcutaneous electrical nerve stimulation may be beneficial for some aspects of upper limb function following stroke.

Noxious thermal stimulation may not be beneficial for upper limb rehabilitation following stroke, whereas innocuous thermal stimulation may improve some aspects of upper limb function.

Muscle vibration may be beneficial for improving upper limb function following stroke.

The literature is mixed regarding additional afferent and peripheral stimulation for upper limb rehabilitation following stroke.

The literature is mixed regarding invasive cortical and nerve stimulation for upper limb rehabilitation following stroke.

The literature is mixed regarding low frequency repetitive transcranial magnetic stimulation, alone or in combination with other therapy approaches, for upper limb rehabilitation following stroke.

High frequency repetitive transcranial magnetic stimulation, alone or in combination with other therapy approaches, may be beneficial for upper limb rehabilitation.

The literature is mixed regarding bilateral repetitive transcranial magnetic stimulation for upper limb rehabilitation.

Theta burst stimulation alone may not be beneficial for upper limb function following stroke, however it may be beneficial for certain aspects of upper limb function when used in combination with repetitive transcranial magnetic stimulation.

The literature is mixed regarding anodal, cathodal, or dual transcranial direct current stimulation, alone or in combination with other therapy approaches, for upper limb rehabilitation following stroke.

Botulinum A likely improves spasticity in the upper limb following stroke, but not range of motion or activities of daily living. The effect on general upper limb motor function is conflicting and less clear.

Botulinum toxin A in combination with other types of therapeutic approaches may be beneficial for certain aspects of upper limb function.

Botulinum toxin B has been less well studied to date in comparison to botulinum toxin A.

Steroid injections may not be beneficial for upper limb rehabilitation following stroke.

Cerebrolysin may be beneficial for aspects of upper limb function following stroke.

The evidence is mixed regarding Levodopa for upper limb rehabilitation following stroke.

The evidence is mixed regarding atorvastatin for upper limb rehabilitation following stroke.

Antidepressants may be beneficial for aspects of upper limb function following stroke.

Dexamphetamine or methylphenidate may be beneficial for aspects of upper limb function following stroke.

Methylphenidate combined with dual transcranial direct current stimulation may be beneficial for upper limb rehabilitation following stroke.

The evidence is mixed regarding acupuncture alone for upper limb rehabilitation following

stroke. Acupuncture combined with conventional or other therapy approaches may not be beneficial for upper limb function. Some forms of acupuncture may be more beneficial than others.

Electroacupuncture with neuronavigation-assisted aspiration may be beneficial for upper limb rehabilitation following stroke, however the evidence is mixed regarding electroacupuncture and transcutaneous electrical acupoint stimulation.

Both meridian acupressure and massage therapy may be beneficial for some aspects of upper limb function following stroke.

Modified Sackett Scale

Level of evidence	Study design	Description	
Level 1a	Randomized controlled trial (RCT)	More than 1 higher quality RCT (PEDro score ≥6).	
Level 1b	RCT	1 higher quality RCT (PEDro score ≥6).	
Level 2	RCT	Lower quality RCT (PEDro score <6).	
	Prospective controlled trial (PCT)	PCT (not randomized).	
	Cohort	Prospective longitudinal study using at least 2 similar groups with one exposed to a particular condition.	
Level 3	Case Control	A retrospective study comparing conditions, including historical cohorts.	
Level 4 Pre-Post A prospective and a post-te		A prospective trial with a baseline measure, intervention, and a post-test using a single group of subjects.	
	Post-test	A prospective post-test with two or more groups (intervention followed by post-test and no re-test or baseline measurement) using a single group of subjects	
	Case Series	A retrospective study usually collecting variables from a chart review.	
Level 5	Observational	Study using cross-sectional analysis to interpret relations. Expert opinion without explicit critical appraisal, or based on physiology, biomechanics or "first principles".	
	Case Report	Pre-post or case series involving one subject.	

New to the 19th edition of the Evidence-based Review of Stroke Rehabilitation

1) PICO conclusion statements

This edition of Chapter 10: Upper extremity motor rehabilitation interventions synthesizes study results from only randomized controlled trials (RCTs), all levels of evidence (LoE) and conclusion statements are now presented in the Population Intervention Comparator Outcome (PICO) format.

For example:



Outcome

New to these statements is also the use of colours where the levels of evidence are written.

Red statements like above, indicate that the majority of study results when grouped together show no significant differences between intervention and comparator groups.

Green statements indicate that the majority of study results when grouped together show a significant between group difference in favour of the intervention group.

For example:

Population: Stroke survivors

Intervention

Bilateral arm training may produce greater Meng et al.	0040
improvements in motor function than conventional therapy.	2018; 017; I. 2008; et al.

Yellow statements indicate that the study results when grouped together are mixed or conflicting, some studies show benefit in favour of the intervention group, while others show no difference between groups.

For example:

Population: Stroke survivors

Outcome Interv		rven	tion		
		DEXTERITY			
LoE		Conclusion Statement	•	RCTs	References
1a There is conflicting evidence about the effect of CIMT to improve dexterity when compared to conventional therapy or motor relearning programmes during the		4	Shah et al. 2016; Yoon et al. 2014; Boake et al. 2007; Ro et al. 2006		
	acute/subacu	ute phase poststroke.			

Comparator

2) Upper extremity rehabilitation outcome measures

For the studies reviewed, upper extremity rehabilitation outcome measures were classified into the following broad categories to allow for synthesis of results and formulation of PICO conclusion statements:

Motor function: These outcome measures covered gross motor movements and a series of general impairment measures when using the upper extremities.

Dexterity: These outcome measures assessed fine motor and manual skills through a variety of tasks, particularly with the use of a stroke survivor's hand.

Activities of daily living: These outcome measures assessed performance and level of independence in various everyday tasks.

Spasticity: These outcome measures assessed changes in muscle tone, stiffness, and contractures.

Range of motion: These outcome measures assessed a patient's ability to freely move their upper extremity through flexion, abduction, and subluxation movements for instance, both passively and actively.

Proprioception: These outcome measures assessed sensory awareness about one's body and the location of limbs.

Stroke severity: These outcome measures assessed the severity of one's stroke through a global assessment of a multitude of deficits a stroke survivor may experience.

Muscle strength: These outcome measures assessed muscle power and strength during movements and tasks.

Outcome measures that fit these categories are described in the next few pages.

Outcome measures definitions

Motor Function

Action Research Arm Test (ARAT): Is a measure of activity limitation in the paretic arm that assesses a patient's ability to handle objects differing in size, weight and shape. The test evaluates 19 tests of arm motor function, both distally and proximally. Each test is given an ordinal score of 0, 1, 2, or 3, with higher values indicating better arm motor status. The total ARAT score is the sum of the 19 tests, and thus the maximum score is 57. This measure has been shown to have good test-retest reliability and internal validity when used to assess motor function in chronic stroke patients (Ward et al. 2019; Nomikos et al. 2018)

Brunnstrom Recovery Stages (BRS): Is a measure of motor function and muscle spasticity in stroke survivors. The measure contains 35 functional movements which are done with the guidance of a clinician (e.g. should abduction, shoulder adduction, leg flexion/extension). These movements are evenly divided into 2 sections: upper extremity and lower extremity. Each movement is then rated on a 6-point scale (1=Flaccidity is present, and no movements of the limbs can be initiated, 2=Movement occurs haltingly and spasticity begins to develop, 3=Movement is almost impossible and spasticity is severe, 4=Movement starts to be regained and spasticity begins to decline, 5=More difficult movement combinations are possible as spasticity declines further. 6=Spasticity disappears, and individual joint movements become possible). This measure has been shown to have good reliability and concurrent validity (Naghdi et al. 2010; Safaz et al. 2009).

Disabilities of the Arm, Shoulder and Hand (QuickDASH): Is a shortened version of DASH – a patient-reported outcome measure intended for upper extremity disorders. It consists of 11 items from the original 30-item DASH questionnaire, where each item has 5 response options, with scaled scores ranging from 0 (no disability) to 100 (most severe disability). The measure is shown to be valid and reliable in populations with upper extremity disorders (Gummesson et al. 2006; Salaffi et al. 2018).

Fugl-Meyer Assessment (FMA): Is an impairment measure used to assess locomotor function and control of the upper and lower extremities, including balance, sensation, and joint pain in patients poststroke. It consists of 155 items, with each item rated on a three-point ordinal scale. The maximum motor performance score is 66 points for the upper extremity section, 34 points for the lower extremity section, 14 points for the balance section, 24 points for sensation section, and 44 points each for passive joint motion and joint pain section, for a maximum of 266 points that can be attained. The upper extremity section consists of four categories (Shoulder/Elbow/Forearm, Wrist, Hand/Finger, and Coordination) and includes 23 different movements which evaluate 33 items. The items are scored on a 3-point rating scale: 0=unable to perform, 1=partial ability to perform and 2=near normal ability to perform. The measure is shown to have

good reliability and construct validity (Okuyama et al. 2018; Villian-Villian et al. 2018; Nillson et al. 2001; Sanford et al. 1993).

Finger Oscillation Test (FOT): Measures motor control and speed and is used to help detect brain damage through motor dysfunction by assessing the speed of finger movement. It measures the maximal tapping speed of the index finger of each hand by requiring the patient to work the lever arm of a mechanical counter up and down as fast as he or she can. The average number of taps in a 10-second interval is determined, and the patient performs five trials. The measure is considered a reliable indicator of brain function (Prigatano et al. 2004; Eng et al. 2013).

Jebsen-Taylor Hand Function Test (JTHFT): Is a measure used to evaluate fine motor skills with weighted and non-weighted hand functions. The test is derived from hand functions required for activities of daily living and is scored as the time taken (in seconds) to complete each subtest, with a maximum of 120 seconds permitted for each subtest. The test is shown to have good test-retest reliability (Allgower et al. 2017; Stern 1992)

Manual Function Test (MFT): Is an upper-limb function assessment measure used for evaluating proximal arm movements as well as fine and gross dexterity of hemiparetic patients after stroke. The test includes 8 subtests including forward and lateral elevation of arm, grasping, pinching, and pegboard manipulations, and ratings can range from 0 (severely impaired) to 32 (full function). The measure has been shown to have good reliability and validity (Miyamoto et al. 2009; Michimata et al. 2008).

Motor Club Assessment (MCA): Is a measure of functional movement that indicates balance and movement by assessing the range of active movement for shoulder shrugging, arm lifting, forearm supination, wrist cocking, and finger extension. Each movement is rated on a 3-point scale (where 0 = no movement, and 2 = full range of movement). (Sunderland et al. 1989)

Motor Evaluation Scale for Upper Extremity in Stroke Patients (MES-UE): Is a measure that assesses the quality of arm movement performance of the hemiparetic arm and hand in stroke patients. The scale encompasses 10 arm function items with six response categories (scores 0-5), nine hand function items with three response categories (scores 0-2), and three functional tasks with three response categories (scores 0-2). The measure is shown to be valid and reliable for measuring quality of arm movement in stroke patients (Van de Winckel et al. 2006).

Motor Status Scale (MSS): Is a measure of upper limb impairment and disability following stroke. It is divided into 4 sections and assesses shoulder, elbow/forearm, wrist and hand movements on a 6-point scale (maximum score = 82 points). This clinical scale is thought to provide a more complete measurement of upper-limb motor function than the FMA, as it evaluates the complete range of motor function of the upper limb by employing a finer grading of isolated movements. The scale has been shown to have good validity and reliability (Ferraro et al. 2002; Wei et al. 2011).

Rancho Los Amigos Functional Test for the Hemiparetic Upper Extremity (RLAFT-

UE): Is a measure used to quantify functional movement ability of the hemiparetic arm in stroke patients. The test consists of a series 17 timed activities of daily living that focus on completion of everyday tasks involving the impaired limb (e.g., zipping a jacket, placing a pillow in a pillowcase). The tasks are arranged in seven levels by degree of difficulty ranging from simple single joint movements at the shoulder to complex multi-joint movements involving the hand and arm. The test has been shown to have high inter- and intra-rater reliability (Kahn et al. 2006; Wilson et al. 1984).

Rivermead Motor Assessment (RMA): Is a multi-faced measure that assesses gross motor function, leg and trunk movements and arm movements in post-stroke patients. The arm movements section consists of 15 items ranging from specific isolated movements (e.g. protracting shoulder girdle in supine position) to complex tasks (e.g. placing a string around the head and tying a bow at the back). Patients perform all movements actively, and dichotomous scores indicate either success (score 1) or failure (score 0). The measure is shown to have good test-retest reliability, content validity, and construct validity (Dong et al. 2018, Van de Winckel et al. 2007).

Sodring Motor Evaluation Scale (SMES): Is a measure of motor function and activities in patients with stroke. It is comprised of 3 subscales that evaluate the motor function of the upper and lower limb, and gross motor function. The first 2 subscales assess simple voluntary movements, while the third evaluates functional tasks including trunk movements, balance, and gait. The scale is comprised of 32 different items scored using a 5-point scale. The measure is shown to have good concurrent and construct validity, as well as good inter-rater reliability (Gor-Garcia_Fogeda et al. 2014).

Stroke Impairment Assessment Set (SIAS): Is a measure of overall motor function and visuospatial ability in stroke survivors. The measure consists of 20 functional tasks (e.g. walking, combing hair, bending, tying shoes). These tasks are then subdivided into 2 areas: tasks specific for the lower extremity and tasks specific for the upper extremity. Each task is then scored on a 6-point scale (0=cannot complete task, 5=completes task as well as the unaffected side). This measure has been shown to have good reliability and validity (Panarese et al. 2016; Seki et al. 2014).

Stroke Rehabilitation Assessment of Movement (STREAM): Is a measure of overall gross motor function in stroke survivors. The measure consists of 30 functional tasks (e.g. filling up and drinking from a cup, walking, getting into and out of the bathtub, buttoning a shirt). These tasks are then subdivided into 3 areas: upper limb, lower limb and basic mobility. Each task is then scored on a 3-point scale (0=cannot complete task, 2=completes task as well as the unaffected side). This measure has been shown to have good reliability and validity (Mateen et al. 2018).

Sollerman Hand Function Test (SHFT): Is a measure of general hand function and dexterity in stroke survivors. The measure consists of 20 functional tasks (e.g. stirring liquid, tying shoes, drinking from a cup, opening/shutting doors). Each task is then

scored on a 6-point scale (0=cannot complete task, 5=completes task as well as the unaffected side). This measure has been shown to have good inter/intra reliability and validity (Singh et al. 2015; Brogardh et al. 2007).

Stroke Upper Limb Capacity Scale (SULCS): Is a measure of basic arm capacities and overall arm strength in stroke survivors. The measure consists of 10 functional tasks (e.g. carrying a briefcase, typing on a computer, writing on a notepad). These tasks are then subdivided into 3 areas: upper limb capacity with no control from wrist and fingers, upper limb capacity with basic control from wrist and fingers, and upper limb capacity with advanced control from wrist and fingers. Each task is then scored on a 3-point scale (0=cannot complete task, 2=completes task as well as the unaffected side). This measure has been shown to have good reliability and concurrent validity (Houwink et al. 2011; Roorda et al. 2011).

University of Maryland Arm Questionnaire (UMAQ): Is a measure of gross functional dexterity in the upper arm for stroke survivors. The measure consists of 10 functional tasks (e.g. opening/closing jars, opening/closing doors, reaching and grabbing common household items). Each task is then scored on a 6-point scale (0=cannot complete task, 5=completes task as well as the unaffected side). This measure has been shown to have good reliability and validity (Beebe et al. 2009, Bovend' Eerdt et al. 2002).

Upper Extremity Function Test (UEFT): Is a measure of total upper extremity dexterity and function in stroke survivors. The measure consists of 15 functional tasks (e.g. moving a jar around, stacking coins, reaching and grabbing a cup). There are 3 subsections of the UEFT: (speed of execution, functional rating, task analysis). Each task is then measured on a 6-point scale (-3=cannot complete task, +3=completes task as well as the unaffected side). This measure has good test/re-test reliability and validity (Platz et al. 2009; Feys et al. 2002).

Wolf Motor Function Test (WMFT): Is a measure that quantifies upper extremity motor ability in stroke survivors. The measure consists of 17 tasks (e.g. lifting arm up using only shoulder abduction, picking up a pencil, picking up a paperclip). These tasks are then subdivided into 3 areas: functional tasks, measures of strength, and quality of movement. Patients are scored on a 6-point scale (1=cannot complete task, 6=completes task as well as the unaffected side. This measure has been shown to have good reliability and validity (Wolf et al. 2005; Wolf et al. 2001).

Dexterity

Box and Block Test (BBT): Is a measure of gross unilateral manual dexterity in stroke survivors. This measure consists of 1 functional task. This task involves a patient moving as many wooden blocks as possible from one end of a partitioned box to the other, in a span of 60 seconds. Patients are scored based on the number of blocks they transfer (the higher the blocks transferred, the better the outcome). The measure has been shown to have good reliability and validity. (Higgins et al. 2005; Platz et al. 2005).

Finger to Nose Test (FNT): Is a measure of overall manual dexterity in stroke survivors. This measure consists of 1 functional task. This task involves the patient touching their index finger to their nose as 10 times as fast as possible. This task is then repeated 1 additional time. Patients are scored based on the number of times they touch their nose (the faster the time the better the outcome). The measure has been shown to have good reliability and construct plus concurrent validity (Rodrigues et al. 2017)

Grating Orientation Task (GOT): Is a measure of overall tactile spatial acuity in stroke survivors. This measure consisted of 1 functional task. Patients were asked to differentiate between a smooth and grooved surface that was placed both proximally and then distally from the patient. This process is repeated 10 different times. Patients are scored based on the number of times they successfully identify the type of surface (the higher the rate of identification, the better the outcome). This measure has been shown to have good reliability and validity (Craig 1999).

Grooved Pegboard Test (GPT): Is a measure of fine motor control in stroke survivors. This measure consists of 1 functional task. Patients are asked to place 25 pegs into the grooved pegboard and are typically given 5-10 minutes to do so. The patients are then scored based on the number of pegs inserted and the time it took them to do so (the higher the insertion rate and the lower the time, the better the outcome). This measure has been shown to have good reliability and validity (Lee et al. 2016; Thompson-Butel et al. 2014).

Minnesota Manual Dexterity Test (MMDT): Is a measure of fine motor control and general dexterity in stroke survivors. The measure consists of 2 functional tasks. Patients are asked to place wooden discs instead of a cylindrical object for the first task. Then, they are asked to turn the discs clockwise 180 degrees and told to shut the lid on the cylinder. Patients are scored on the amount discs inserted and on the screwing of the lid. The higher the number of discs put in the cylinder and the faster/tighter the lid is screwed on, the better the outcome. This measure has been shown to have good reliability and validity (Wang et al. 2018; Surrey et al. 2003).

Nine Hole Peg Test (9HPT): Is a measure of overall manual dexterity in stroke survivors. The measure consists of 1 functional task. Patients are asked to take 9 pegs

out of a container and insert them into the pegboard. Once all 9 pegs are inserted they are then taken out of the pegs as quickly as possible and placed back in the container. Patients are scored on how quickly they can insert and take out the pins, so the faster the time, the better the outcome. This measure has been shown to have good reliability and concurrent validity (da Silva et al. 2017).

Purdue Pegboard Test (PPT): Is a measure of precision grip strength and speed in stroke survivors. The measure consists of 1 functional task. Patients are asked to place as many pins as they can onto the pegboard in 30 secs, and then repeat this exercise for their other hand. Patients are scored on the number of pins they can place onto the pegboard in the given amount of time. This measure has been shown to have good reliability and validity (Gonzalez et al. 2017, Wittich & Nadon, 2017).

Activities of daily living

Arm Motor Ability Test (AMAT): Is a measure of upper extremity limitation for stroke survivors in performing activities of daily living. The measure consists of 13 common unilateral and bilateral tasks (e.g. manipulating objects such as utensil and telephones; donning/doffing a piece of clothing). Each task is scored on two, 6-point ordinal scales assessing functional ability and the quality of the movement performed. The measure has been shown to have good reliability and construct validity, in its full form and in abbreviated versions for stroke survivors (Fulk et al. 2017; O'Dell 2013; O'Dell 2011).

Assessment of Motor and Process Skills (AMPS): Is a measure of processing skills and overall independence for stroke survivors in performing activities of daily living (ADL) (Ahn et al. 2016). The measure consists of 16 motor tasks (e.g. picking up/setting down a mug, donning/doffing a piece of clothing, turning doorknobs) and 20 process tasks (e.g.memory testing, matching shapes, word recall) (Ahn et al. 2016) Each task is scored on 10 item tool assessing functional ability and the accuracy/speed at which the skill(s) are completed (Lam et al. 2018). This measure has been shown to have good reliability and validity in both its full and abbreviated form (Lam et al. 2018; Ahn et al. 2016).

Barthel Index (BI): Is a measure of how well a stroke survivor can function independently and how well they can perform activities of daily living (ADL). The measure consists of a 10-item scale (e.g. feeding, grooming, dressing, bowel control). Possible total scores range from 0 to 100. This measure has been shown to have good reliability and validity in its full form (Gonzalez et al. 2018; Park et al. 2018).

ABILHAND: Is a measure of how well a stroke survivor utilizes their hands to complete various manual tasks. The measure consists of 23 common bimanual activities (e.g. hammering a nail, wrapping gifts, cutting meat, buttoning a shirt, opening mail). Each task is then scored on a 3-point scale (0=impossible, 1=difficult, 2=easy) assessing overall ability. This measure has been shown to have good reliability and validity in its full form (Ashford et al. 2008; Penta et al. 2001).

Canadian Occupational Performance Measure (COPM): Is a measure of how well a stroke survivor engages in self-care, productivity and leisure. The measure consists of 25 functional items/tasks (e.g. bathing, ability to work at least part-time, activities involved in). Each task is then scored on a single 10-point rating scale primarily measuring proficiency in each of the 3 sub-categories (self-care, productivity and leisure). This measure has been shown to have good reliability and validity in its full form. (Yang et al. 2017).

Chedoke Arm and Hand Activity Inventory (CAHAI): Is an upper limb measure that uses a 13-point quantitative scale in order to assess recovery of the arm and hand in performing activities of daily living after a stroke. It is a performance test using 13 bimanually performed real-life items, designed to encourage bilateral upper limb use.

Scores represent the patient's relative ability to independently perform stabilisation or manipulation in ADL with the impaired upper limb. The measure is shown to have good test-retest and interrater reliability, as well as good construct and concurrent validity (Ward et al. 2019; Schuster-Amft et al. 2018; Barteca et al. 2004).

Duruoz Hand Index (DHI): Is a measure used to assess hand-related activity limitation based on questions concerning activities in a person's daily life. It contains 18 activities commonly performed by the hand in the kitchen, during dressing, while performing personal hygiene, while performing office tasks, and other general items. The measure is shown to have good construct validity, test-retest reliability, and internal consistency in patients with stroke (Sezer et al. 2007).

Frenchay Arm Test (FAT): Is a measure of the upper extremity motor control that a stroke survivor possesses. The measure consists of 5 common tasks that require use of the upper extremity (e.g. stabilize a ruler/draw a line with a pencil, comb hair, clip a clothespin onto the edge of a table, grasp a cylinder, drink from a glass of water and then set it down). Each task is then scored on a 2-point scale wherein each task receives either a 0 (unsuccessful completion) or a 1 (successful completion). This measure has been shown to have good reliability and validity in its full form. (Heller et al. 1987; Parker et al. 1986)

Frenchay Activities Index (FAI): Is a measure of activities that stroke survivors have participated in recently. The measure consists of 15 items that are in turn split up into 3 subscales (domestic chores, leisure/work and outdoor activities). These items include: preparing meals, washing clothes, light/heavy housework, social outings etc. Each task is then scored on a 4-point scale with 1 being the lowest score. This measure has been shown to have good reliability and concurrent validity in its full form (Schuling et al. 1993)

Functional Activity Scale (FAS): Is a measure of functional everyday activities that stroke survivors participate in daily. The measure consists of 15 functional activities (e.g. cooking, cleaning, zipping up a coat). Each activity is then scored on a 5-point scale (0=cannot complete activity, 4=completes activity as well as the unaffected side). This measure has been shown to have good reliability and validity (Pang et al. 2006).

Functional Independence Measure (FIM): Is an 18-item outcome measure composed of both cognitive (5-items) and motor (13-items) subscales. Each item assesses the level of assistance required to complete an activity of daily living on a 7-point scale. The summation of all the item scores ranges from 18 to 126, with higher scores being indicative of greater functional independence. This measure has been shown to have excellent reliability and concurrent validity in its full form (Granger et al. 1998, Linacre et al. 1994; Granger et al. 1993).

Goal Attainment Scale (GAS): Is a measure that quantifies the progress made towards obtaining personalized rehabilitation goals. The measure consists of 5 levels of goal achievement. The items in these levels consist of various goals individual patients

would like to achieve (e.g. bathing independently, being able to do housework, walking unaided). The patient is then rated on a 4-point scale on their ability to carry out said goals (-2=far behind schedule, +2=far ahead of schedule). This measure has been found to have good reliability and validity in its full form (Hanlan et al. 2017; Krasny-Pacini et al. 2016)

Modified Barthel Index (MBI): Is a measure of how well a stroke survivor can function independently and how well they can perform activities of daily living (ADL). The measure consists of a 10-item scale (e.g. feeding, grooming, dressing, bowel control). Possible scores range from 0 to 20. This measure has been shown to have good reliability and validity in its full form. (MacIsaac et al. 2017; Ohura et al. 2017).

Motor Activity Log (MAL): Is a patient-reported measure of the use and quality of movement of the impaired arm. The measure consists of 30 functional tasks (e.g. handling utensils, buttoning a shirt, combing hair). Each task is then measured on a 6-point scale (0=complete inability to use affected arm). This measure has been shown to have good reliability and validity (Chuang et al. 2017).

Motor Assessment Scale (MAS): Is a performance-based measure that assesses everyday motor function. The measure consists of 8 motor-function based tasks (e.g. supine lying, balanced sitting, walking). Each task is then measured on a 7-point scale (0=suboptimal motor performance, 6=optimal motor performance). This measure has been shown to have good reliability and concurrent validity (Simondson et al. 2003).

Nottingham Extended Activities of Daily Life (NEADL): Is a measure of a stroke survivor's independence with regards to their performance on various activities of daily living. The measure consists of 22 functional tasks (e.g. walking, cooking, cleaning, participation in active hobbies). These tasks are then further divided into 4 distinct subscales (mobility, kitchen, domestic, and leisure activities). In turn, each task is measured on a 5-point (0=not at all, 4=on my own with no difficulty). This measure has been shown to have good reliability and validity (das Nair et al. 2011; Sahin et al. 2008).

Nottingham Stroke Dressing Assessment (NSDA): Is a measure of a stroke survivor's ability to successfully dress themselves. The measure consists of 25 functional dressing tasks (e.g. buttoning up a shirt, buckling a belt/watch, putting on pants). These tasks are then measured on a 4-point scale (0=cannot complete task, 3=completes task as well as the unaffected side). This measure has been shown to have good reliability and validity (Walker et al. 2011).

Stroke Impact Scale (SIS): Is a patient-reported measure of multi-dimensional stroke outcomes. The measure consists of 59 functional tasks (e.g. dynamometer, reach and grab, walking, reading out loud, rating emotional regulation, word recall, number of tasks completed, and shoe tying). These tasks are then divided into 8 distinct subscales which include: strength, hand function, mobility, communication, emotion, memory, participation and activities of daily living (ADL). Each task is measured on a 5-point

scale (1=an inability to complete the task, 5=not difficult at all). The measure has been shown to have good reliability and validity (Mulder et al. 2016; Richardson et al. 2016).

STAIS Stroke Questionnaire (SSQ): Is a measure of activities and participation in the physical environment for stroke survivors. The measure consists of 36 functional tasks (e.g. taking a bath or shower, ability to handle your finances, opening and closing doors). Each task is measured on a 4-point scale (1=no ability, 4=complete ability). The measure has been shown to have good reliability and concurrent validity (Bouffioulx et al. 2010 Bouffioulx et al. 2008)

Upper Limb Self-Efficacy Test (UPSET): Is a measure of a stroke survivor's confidence in their ability to carry out upper limb specific tasks with their affected side. The measure consists of 20 functional tasks (e.g. shaking hands, flipping a coin, opening/shutting doors). Each task is then measured on a 5-point scale (0=cannot complete task, 4=completes task as well as the unaffected side). The measure has been shown to have good test/retest reliability and validity (Abdullahi, 2016; Pang et al. 2007).

Spasticity

Ashworth Scale (AS): Is a measure of resistance to passive movement in stroke survivors. The measure contains 15 functional m'ovements which are done with the guidance of a trained clinician. These movements are evenly divided into 2 sections: upper extremity and lower extremity. Each movement is then rated on a 5-point scale (0=no increase in muscle tone, 1=barely discernible increase in muscle tone, 2=moderate increase in muscle tone 3=profound increase in muscle tone (movement of affected limb is difficult) 4=complete limb flexion/rigidity (nearly impossible to move affected limb)). This measure has been shown to have good reliability and validity (Merholz et al. 2005; Watkins et al. 2002).

Bhakta Finger Flexion Scale (BFFS): Is a measure of the overall finger flexion experienced by stroke survivors when completing functional tasks. This measure consists of 27 functional tasks (e.g. writing with a pen, typing, squeezing a ball). Each task is then rated on a 3-point scale (0=cannot complete task; fingers too rigid, 2=easily completes task; flexes and extends fingers). This measure has been shown to have good reliability and validity (Christina et al. 2015).

Disability Assessment Scale (DAS): Is a measure of resistance to passive movement in the upper extremity for stroke survivors. The measure consists of 20 functional tasks (e.g. brushing teeth, buttoning a shirt, gait technique & general pain). These tasks are then divided into 4 sections: hygiene, dressing, limb position and pain. Each task is then rated from: 0=no disability, 1=mild disability 2=moderate disability, 3=severe disability. This measure has been shown to have good reliability and validity (Thibaut et al. 2013; Brashear et al. 2002)

Modified Ashworth Scale (MAS): Is a measure of muscle spasticity for stroke survivors. The measure contains 20 functional movements which are done with the guidance of a trained clinician. These movements are evenly divided into 2 sections: upper extremity and lower extremity. Each movement is then rated on a 6-point scale (0=no increase in muscle tone, 1=barely discernible increase in muscle tone 1+=slight increase in muscle tone, 2=moderate increase in muscle tone 3=profound increase in muscle tone (movement of affected limb is difficult) 4=complete limb flexion/rigidity (nearly impossible to move affected limb)). This measure has been shown to have good reliability and validity (Merholz et al. 2005; Blackburn et al. 2002).

Modified Tardieu Scale (MTS): Assesses spasticity through measuring the quality and angle of muscle movements in response to stretches of different velocities. The velocities of muscle movement are as slow as possible (V1), speed of the limb falling from gravity (V2), and when the joint is moved as fast as possible (V3). The quality and angle of muscle reactions are recorded during these velocities. The quality of muscle reactions are scored as: 0 (no resistance throughout the duration of the stretch), 1 (slight resistance), 2 (clear catch occurring at a precise angle, followed by a release), 3 (fatigable clonus), 4 (infatigable clonus), 5 (joint is immovable) (Li et al. 2014b).

Resistance to Passive Movement Scale (REPAS): Is a measure of general muscle spasticity for stroke survivors. The measure contains 52 functional movements which are done with the guidance of a trained clinician. These movements are evenly divided into 2 sections: upper extremity and lower extremity. Each movement is then rated on a 5-point scale (0=no increase in muscle tone, 1=barely discernible increase in muscle tone, 2=moderate increase in muscle tone 3=profound increase in muscle tone (movement of affected limb is difficult) 4=complete limb flexion/rigidity (nearly impossible to move affected limb)). This measure has been shown to have good test/retest reliability and concurrent validity (Platz et al. 2008).

Spasm Frequency Scale (SFS): Is a measure of the amount of spasms experienced by stroke survivors in a day. The measure is only concerned with measuring the amount of spasms in a single day. The amount of spasms per day are rated based on a 5-point scale (0=No spasms. 1= One or fewer spasms per day 2=Between 1 and 5 spasms per day 3=Five to less than 10 spasms per day 4=Ten or more spasms per day, or continuous contraction). This measure has been shown to have good reliability and validity (Santamato et al. 2013; Snow et al. 1990).

Range of motion

Active Range of Motion (AROM): Is a measure of the range of motion stroke survivors possess without receiving assistance. The measure consists of 20 functional movements for both the upper and lower extremity. The movements are evenly divided into 2 sections: upper extremity and lower extremity. These movements are then rated on a 4-point ordinal scale (0=cannot complete movement, 3=completes movement as well as the unaffected side). This measure has been shown to have good reliability and validity (Beebe & Lang 2009, Dickstein et al. 1986)

Maximal Elbow Extension Angle During Reach (MEEAR): Is a measure of the amount of elbow extension undergone by a stroke survivor while they are reaching for an object. The measure consists of 1 functional movement which is when a patient reaches for an object and their rate of elbow extension is measured (the higher the rate of extension, the better the outcome). This measure has been shown to have good inter/intra reliability and concurrent validity (Murphy et al. 2011; Cristea et al. 2003).

Passive Range of Motion (PROM): Is a measure of the range of motion stroke survivors possess while receiving assistance. The measure consists of 30 functional movements for both the upper and lower extremity. The movements are evenly divided into 2 sections: upper extremity and lower extremity. These movements are then rated on a 5-point ordinal scale (0=cannot complete movement, 4=completes movement as well as the unaffected side). This measure has been shown to have good test/retest reliability and validity (Lynch et al. 2005).

Proprioception

Joint Position Sense Test (JPST): Is a measure of how well stroke survivors can perceive the position of their joints in motion and standing still. The measure consists of 1 functional task repeated several times. This task involves the patient holding 2 different shaped objects that also weigh different from each other and then told to identify which one weighs more and which one has a stranger shape. The more times the patient (s) identifies which shape is heavier/unique, then the better the outcome. This measure has been shown to have good reliability and validity (Kattenstroth et al. 2013).

Kinesthetic Visual Imagery Questionnaire (KVIQ): Is the measure of the visual acuity and muscle movement that stroke survivors possess. The measure consists of 20 functional tasks (e.g. tying shoes, reading out loud, reaching for an object, peripheral vision testing). Each task is then measured on 3-point scale (0=cannot complete task, 2=completes task as well as the unaffected side). This measure has been shown to have good reliability and validity (Salles et al. 2017; Demanboro et al. 2018).

Revised Nottingham Sensory Assessment (RNSA): Is a measure of somatosensory perception in stroke survivors. The measure consists of 1 functional task repeated with 11 different objects. The task involves patients identifying 11 different objects with their eyes closed. The higher the rate of objects identified leads to a better overall outcome. This measure is shown to have good reliability and validity (Boccuni et al. 2018; Gorst et al. 2018).

Stroke severity

Modified Rankin Scale (MRS): Is a measure of functional independence for stroke survivors. The measure contains 1 item. This item is an interview that lasts approximately 30-45 minutes and is done by a trained clinician. The clinician asks the patient questions about their overall health, their ease in carrying out ADLs (cooking, eating, dressing) and other factors about their life. At the end of the interview the patient is assessed on a 6-point scale (0=bedridden, needs assistance with basic ADLs, 5=functioning at the same level as prior to stroke). This measure has been shown to have good reliability and validity (Quinn et al. 2009; Wilson et al. 2002).

National Institutes of Health Stroke Scale (NIHSS): Is a measure of somatosensory function in stroke survivors during the acute phase of stroke. This measure contains 11 items and 2 of the 11 items are passive range of motion (PROM) assessments delivered by a clinician to the upper and lower extremity of the patient. The other 9 items are visual exams conducted by the clinician (e.g. gaze, facial palsy dysarthria, level of consciousness). Each item is then scored on a 3-point scale (0=normal, 2=minimal function/awareness). This measure has been shown to have good reliability and validity (Heldner et al. 2013; Weimar et al. 2004).

Neurological Function Deficit Scale (NFDS): Is a measure of neurological deficits experienced by stroke survivors in both the upper and lower extremities. This measure contains 40 functional movements done with the guidance of a clinician (e.g. should abduction, shoulder adduction, leg flexion/extension). These movements are evenly divided into 2 sections: upper extremity and lower extremity. Each movement is then measured on a 6-point scale (0=normal function, 5=severe stroke). This measure has been shown to have good test/retest reliability and validity (Yao & Ouyang. 2014).

Muscle strength

Hand Grip Strength (HGS): Is a measure of the overall hand grip strength in stroke survivors. The measure consists of 1 functional task. This task involves a patient squeezing the dynamometer and then receiving a hand grip strength measurement. This action is then repeated 1 additional time and the best of the two readings is used as a score. This measure has been shown to have good test/retest reliability and validity (Bertrand et al. 2015).

Isokinetic Peak Torque (IPT): Is a measure of the work capacity of specific muscle groups of a stroke survivor. The measure consists of 1 functional task. The patient performs elbow flexion/extension while attached to a machine that measures force output. The process is then repeated for the leg. The output is then compared to healthy patients that are approximately the same age and build. This measure has been shown to have good test/retest reliability (Horvat et al. 1997).

Manual Muscle Strength Test (MMST): Is a measure of how well a stroke survivor can complete various upper extremity movements while resistance is applied by a trained clinician. The measure consists of 3 functional tasks: muscle contraction, total range of motion and resistance to applied pressure. Patients are scored on a 12-point scale (0=no movement, T=trace/barely discernable movement, 10=movement carried out as well as the unaffected side). This measure has been shown to have good reliability and validity (Kristensen et al. 2017; Ada et al. 2016)

Medical Research Council Scale (MRCS): Is a measure of overall muscle strength a stroke survivor possesses. The measure consists of 33 functional tasks (e.g. opening/shutting cupboards, screwing and unscrewing lids, lifting of light objects). Each task is then rated on a 4-point scale (0=cannot complete task, 3=completes task as well as the unaffected side). This measure has been shown to have good reliability and validity (Hsieh et al. 2011; Fasoli et al. 2004).

Therapy based interventions Neurodevelopmental techniques



Adopted from: http://www.bobathconcept.eu/en/main-site/

There are several approaches that are considered to be neurodevelopmental techniques (NDT). These include the Bobath concept, Brunnstrom movement therapy and motor relearning programmes.

The Bobath concept is a comprehensive, problem-solving treatment approach that focuses on motor recovery (e.g. function, movement and tone) of an individual's affected side after a lesion in the central nervous system (Michielsen et al. 2017). Prior to its introduction in the 1950's, stroke rehabilitation largely assumed a compensatory approach towards the unaffected side for rehabilitation (Kollen et al. 2009). The Bobath concept like other neurodevelopmental techniques relies on the tenets of neuroplasticity, in that motor recovery of the affected side is possible through individualised treatment plans that focus on how tasks are completed, facilitation of movements through therapeutic handling, movement analysis, modification of the environment and appropriate use of verbal cues from therapists (Michielsen et al. 2017).

Brunnstrom movement therapy focuses on retraining motor movements through emphasis of the synergistic and reflexive muscle movements that develop during recovery from hemiplegia. The approach encourages the use of abnormal or spastic muscle movements of the flexors and extensors during early recovery to regain muscle synergies, contrary to the Bobath concept which inhibits these movements (Pandian 2012; Brunnstrom 1970).

The motor relearning programme employs practice of task-specific activities to remediate specific motor skills needed to perform that task. Motor tasks are practiced in context relevant environments to enhance sensory input and modulate performance (Pandian 2012).

A total of 9 RCTs were found that evaluated neurodevelopmental techniques for upper extremity motor rehabilitation, interventions categories are listed below.

Three RCTs compared the Bobath concept to conventional therapy (van der Lee et al. 1999; Gelber et al. 1995; Basmajian et al. 1987). Two RCTs compared motor relearning programmes to conventional therapy (Walker et al. 2012; Platz et al. 2009). Three RCTs compared motor relearning programmes to Bobath concept approaches (Langhammer and Stanghelle, 2011; Platz et al. 2005; van Vliet et al. 2005). One RCT compared Brunnstrom movement therapy to a motor relearning programme (Pandian et al. 2012).

The methodological details and results of all 9 RCTs are presented in Table 1.

Table 1. RCTs evaluating neurodevelopmental techniques for upper extremity moto	or
rehabilitation	

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end}	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Time post stroke category	Bobath concept approach compared to conve	entional therapy
<u>van der Lee et al.</u> (1999) RCT (7) N _{start} =66 N _{end=} 57 TPS=Chronic	E: Bobath concept C: Forced-use therapy Duration: 6h, 5d/wk for 2wk	Action Research Arm Test (+con)
Gelber et al. (1995) RCT (5) N _{start} =20 N _{end} =20 TPS=Acute	E: Bobath concept C: Traditional techniques Duration: <i>Not reported</i>	 Functional Independence Measure (-) Box and Block Test (-) Nine Hole Peg Test (-)
Basmajian et al. (1987) RCT (6) N _{start} =29 N _{end} =23 TPS=Sub-acute	E: Bobath concept C: Physical and behavioural therapy using EMG Duration: 45min, 3d/wk for 5wk	 Upper Extremity Performance Test for the Elderly (-) Finger Oscillation Test (-)
	Motor relearning programmes compared to con	ventional therapy
Walker et al. (2012) RCT (7) Nstart=70 NEnd=64 TPS=Acute	E: Motor relearning programme C: Dressing without a task-oriented approach Duration: 3d/wk for 6wk	 Nottingham Stroke Dressing Assessment (-) 10-hole peg transfer test (-)
Platz et al. (2009) RCT (8) Nstart=148 Nend=135 TPS=Not reported	E: Motor relearning programme E2: Passive therapy (with splints) C: Conventional therapy Duration: 45min, 5d/wk for 4wk	 Fugl-Meyer Assessment (-) Upper Extremity Performance Test for the Elderly (-)
Мо	tor relearning programme compared to Bobath	concept approaches
Langhammer & Stanghelle (2011). RCT (8) N _{start} =61 N _{end} =53 TPS=Not reported	E: Motor relearning programme E2: Bobath concept Duration: 40min, 5d/wk for 2wk	 Motor Assessment Scale (+exp) Sodring Motor Evaluation Scale (+exp) Motor Activity Log (+exp)
Platz et al. 2005 RCT (8) N _{start} =62 N _{end} =62 TPS=Subacute	E: Motor relearning programme (Arm BASIS) E2: Bobath concept C: No augmented exercise therapy time Duration: 4wk	Fugl-Meyer Assessment (-)
van Vliet et al. (2005) RCT (7) Nstart=120	E: Motor Relearning Programme E2: Bobath concept Duration: 23min, 5d/wk for 4wk	Motor Assessment Scale (-)

N _{end} =105 TPS=Acute		
	Brunnstrom movement therapy vs Motor relear	ning programme
Pandian et al. (2012) RCT (6) Nstart=30 Nend=30 TPS=Chronic	E: Brunnstrom hand manipulation treatment C: Motor relearning programme Duration: 1h, 3d/wk for 4wk	Fugl-Meyer Assessment (+exp)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about neurodevelopmental techniques

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	Bobath concept approaches may not have a difference in efficacy when compared to conventional therapy for improving motor function.	2	Van der lee et al. 1999; Basmajian et al. 1987
1b	Motor relearning programmes may not have a difference in efficacy when compared to conventional therapy for improving motor function.	1	Platz et al. 2009
1a	There is conflicting evidence about the effect of motor relearning programmes to improve motor function when compared to Bobath concept approaches .	2	Langhammer Stanghelle et al. 2011; Platz et al. 2005
1b	Brunnstrom movement therapy may produce greater improvements in motor function than motor relearning programmes.	1	Pandian et al. 2012

ACTIVITIES OF DAILY LIVING			
LoE	Conclusion Statement	RCTs	References
2	Bobath concept approaches may not have a difference in efficacy when compared to conventional therapy for improving performance of activities of daily living.	1	Gelber et al. 1995
1b	Motor relearning programmes may not have a difference in efficacy when compared to conventional therapy for improving performance of activities of daily living.	1	Walker et al. 2012
1a	There is conflicting evidence about the effect of motor relearning programmes to improve performance of activities of daily living when compared to Bobath concept approaches .	2	Langhammer Stanghelle et al. 2011; Van Vliet et al. 2005

DEXTERITY			
LoE	Conclusion Statement	RCTs	References
2	Bobath concept approaches may not have a difference in efficacy when compared to conventional therapy for improving dexterity.	1	Gelber et al. 1995
1b	Motor relearning programmes may not have a difference in efficacy when compared to conventional therapy for improving dexterity.	1	Walker et al. 2012

Key points

Bobath concept approaches and motor relearning programmes may not be beneficial for upper limb rehabilitation following stroke.

Brunnstrom movement therapy may be more beneficial than motor relearning programmes for upper limb function.

Bilateral arm training



Adopted from: https://www.newswise.com/articles/stroke-survivors-rehab-arms-with-in-home-device

Bilateral arm training is a technique whereby patients perform the same movements with both the right and left upper limbs simultaneously. The use of bilateral arm training techniques with the upper limb following stroke has been encouraged recently with the development of new theories regarding neural plasticity. Theoretically, the use of the intact limb helps to promote functional recovery of the impaired limb through facilitative coupling effects between the damaged and intact cerebral hemispheres through neural networks linked via the corpus callosum (Morris et al. 2008; Summers et al. 2007).

Interventions for bilateral arm training included: 13 RCTs evaluating bilateral arm training compared to unilateral arm training (Han and Kim, 2016; Shim et al. 2015; McCombe et al. 2014; Byl et al. 2013; Dispa et al. 2013; Kim et al. 2013; Wu et al. 2013; Morris and van Wijck, 2012; Yang et al. 2012; Lin et al. 2010; Stoykov et al. 2009; Morris et al. 2008; Summers et al. 2007). Five RCTs evaluating bilateral arm training compared to conventional rehabilitation (Meng et al. 2018; Lee et al. 2017; Lee et al. 2013; Stinear et al. 2008; Desrosiers et al. 2005). Two RCTs evaluating bilateral arm training with rhythmic auditory cueing compared to unilateral arm training or conventional rehabilitation (Whitall et al. 2011; Luft et al. 2004), and task-oriented bilateral arm training (Hsieh et al. 2016; Song et al. 2015). A single RCT looked at bilateral arm training compared to CIMT (Brunner et al. 2012; Wu et al. 2011), and another two compared bilateral arm training with rhythmic auditory cueing to modified CIMT (van Delden et al. 2015; van Delden et al. 2013).

The methodological details and results of all 29 RCTs evaluating bilateral arm training for the upper extremity motor rehabilitation are presented in Table 2.

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)	
Bilateral arm training compared to unilateral arm training			
Han & Kim (2016) RCT (5) N _{start} =25 N _{End} =25 TPS=Not reported	E: Bilateral arm training C: Unilateral arm training Duration: 5x/wk for 6wk	 Box and Block Test (-) Elbow Amplitude (-) Shoulder Amplitude (+exp) 	
Shim et al. (2015) RCT (6) N _{Start} =20 N _{End} =20 TPS=Chronic	E: Bilateral training C: Unilateral training Duration: 30min, 5x/wk for 6wk	 Motor Function Test (+exp) Functional Independence Measure (+exp) Affected hand amount of sedentary and moderate activity (+exp) 	
McCombe et al.(2014) RCT (7) N _{Start} =30 N _{End} =26 TPS=Subacute	E: Bilateral + Unilateral training C: Unilateral training Duration: 1h, 3d/wk for 12wk	 Wolf Motor Function Test (+exp) University of Maryland Arm Questionnaire (+exp) Fugl-Meyer Assessment (-) Box and Block Test (-) 	
<u>Byl et al</u> . (2013) RCT (5) N _{Start} =18 N _{End} =15 TPS=Subacute	E: Bilateral orthosis C: Unilateral orthosis Duration: 90 min for 12 sessions	Fugl-Meyer Assessment (-)	
Dispa et al. (2013) RCT (7) N _{Start} =10 N _{End} =10 TPS=Not given	E: Bilateral therapy C: Unilateral therapy Duration: 1h, 3d/wk for 4wk	 Fugl-Meyer Assessment (-) Purdue pegboard Test (-) ABILHAND scale (-) STAIS-stroke questionnaire (-) 	
Kim et al. (2013) RCT (3) N _{start} =15 N _{end} =15 TPS=Subacute	E1: Bilateral robotic training E2: Unilateral robotic training C: Usual Care Duration: 90min, 2d/wk for 6wk	Fugl-Meyer Assessment (-)	
Wu et al. (2013c) RCT (7) Nstart=53 NEnd=53 TPS=Chronic	E1: Bilateral robotic training E2: Unilateral robotic training C: Conventional therapy Duration: 90 to 105min, 1d/wk for 4wk	 Motor Activity Log (-) Wolf Motor Function Test (-) ABILHAND Scale (-) 	
Morris & van Wijck (2012) RCT (7) N _{start} =106 N _{end} =85 TPS=Not reported	E: Bilateral training C: Unilateral training Duration: 20min, 5d/wk for 6wk	 9 Hole Peg Test (+exp) Action Research Arm Test (-) 	
<u>Yang et al. (</u> 2012) RCT (7) N _{start} =21 N _{end} =21 TPS=Chronic	E1: Unilateral robot assisted training E2: Bilateral robot assisted training C: Standard training group Duration: 90min, 5d/wk for 4wk	 Fugl-Meyer Assessment (-) Medical Research Council Scale (-) 	
Lin et al. (2010) RCT (6) N _{start} =33	E: Bilateral training C: Unilateral training Duration: 2h, 5d/wk for 3wk	 Fugl Meyer Assessment (+exp) Functional Independence Measure (-) Motor Assessment Log (-) 	

Table 2. RCTs evaluating BAT interventions	for upper extremit	y motor rehabilitation
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N _{end} =33					
TPS=Chronic					
Stoykov et al. (2009)	E: Bilateral training	Motor Assessment Scale (-)			
RCT (5)	C: Unilateral training	Motor Status Scale (-)			
N _{start} =21	Duration: 1h, 3d/wk for 8wk				
N _{end=} 21					
Morris et al. (2008)	E: Bilateral training	Arm Research Arm Test (-)			
	C: Unilateral training	Rivermead Motor Assessment (-)			
N Start=100	Duration: 20min, 5d/wk for 6wk	 9 Hole Peg Test (+exp) Modified Barthol Index () 			
TPS-Chronic		• Modified Barther Hidex (-)			
Summore of al (2007)	E: Bilatoral training	Modified Motor Assessment Scale (Lovp)			
BCT (5)	C: Unilateral training	• Modified Motor Assessment Scale (Texp)			
Netart=12	Duration: Not reported				
Nend=10					
TPS=Chronic					
	Bilateral arm training compared to c	onventional rehabilitation			
Meng et al. (2018)	E: Hand-Arm Bimanual Intensive Therapy	Fugl-Meyer Assessment (+exp)			
RCT (7)	C: Conventional Rehabilitation Program	Action Research Arm Test (+exp)			
N _{start} =128	Duration: 1h (twice per d), 5d/wk for 2wk				
N _{end} =123					
TPS=Acute					
Lee et al. (2017)	E: Bilateral Arm Training	 Fugl-Meyer Assessment (+exp) 			
RCT (6)	C: Upper Extremity Training	Box and Block Test (+exp)			
N _{Start} =30	Duration: 1h, 5d/wk for 8wk	 Modified Barthel Index (+exp) 			
NEnd=30					
$\frac{\text{Lee et al.}}{\text{PCT}(6)}$	E: Bilateral training + conventional	Functional independence measure (+exp)			
	C: Conventional rehabilitation				
Nstart=20 $N_{rad}=26$	Duration: 30min 3d/wk for 4wk				
TPS=Chronic					
Stinear et al. (2008)	E: Bilateral training	Fugl Mever Assessment (+exp)			
RCT (6)	C: Self-directed motor practice	• Grip strength (-)			
N _{start} =32	Duration: 10min (three times per day).				
N _{end} =27	7d/wk for 4wk				
TPS= Chronic					
Desrosiers et al. (2005)	E: Bilateral training	Fugl Meyer Assessment (-)			
RCT (7)	C: Conventional therapy	Grip strength (-)			
N _{start} =41	Duration: 45min, 15-20 sessions	Box and Block Test (-)			
N _{end} =33		 Purdue Pegboard Test (-) 			
TPS=Subacute		Finger-to-Nose Test (-)			
		Upper Extremity Performance test for the Elderly (-)			
		Functional Independence Measure (-) The Assessment of Mature and Dependence Obility (-)			
		Ine Assessment of Motor and Process Skills (-)			
	with instantic auditory cueing compared t	o unilateral arm training or conventional renabilitation			
$\frac{\text{vinital et al.}}{\text{RCT}(6)}$		Fugi integer Assessment (-) Wolf Motor Function Test (-)			
$N_{\text{start}} = 111$	C: Dose matched unilateral therapeutic	Stroke Impact Scale (-)			
NEnd=92	exercises	Flbow extension (-)			
TPS=Chronic	Duration: 20min, 3d/wk for 6wk	Shoulder extension (-)			
	,	• Wrist extension (+exp)			
		Elbow flexion (-)			
Luft et al. (2004)	E: Bilateral arm training + rhythmic auditory	Fugl Meyer (-)			
RCT (7)	cueing	Wolf Motor Arm Test (-)			
N _{start} =26	C: Therapeutic exercises.	University of Maryland Arm Questionnaire for Stroke (-)			
N _{end} =21	Duration: 1 h, 3d/wk for 6wk	Elbow Strength (-)			
IPS=Chronic		Shoulder Strength (-)			
Task-oriented bilateral arm training compared to task-oriented training or bilateral arm training with rhythmic auditory cueing					

Hsieh et al. (2016) RCT (6) N _{Start} =31 N _{End} =31 TPS=Subacute	E: Bilateral arm priming + task-oriented training C: Task-oriented training alone Duration: 90min, 5d/wk for 4wk	 Fugl-Meyer Assessment (-) Box and Block Test (-) Grip Strength (-) Modified Rankin Scale (-) Functional Independence Measure (-) Activities of Daily Living (-) Stroke Impact Scale (+exp)
<u>Song et al</u> . (2015) RCT (5) N _{Start} =40 N _{End} =40 TPS=Chronic	E: Task-oriented bilateral arm training E2: Bilateral arm training with rhythmic auditory cueing Duration: 30min, 5d/wk for 12wk	 Box and Block Test (+exp) Jebsen Taylor Hand Function Test (+exp) Modified Barthel Index (+exp)
	Bilateral arm training com	pared to TENS
<u>Stinear et al</u> . (2014) RCT (6) N _{Start} =57 N _{End} =51 TPS=Not given	E: Bilateral training C: TENS Duration: 45min, 5d/wk for 4wk	 Modified Ashworth Scale (-) Stroke Impact Scale (-)
EMG-triggered	NMES with bilateral arm training compared	to EMG-triggered NMES with unilateral training
Singer et al.(2013) RCT (4) N _{Start} =24 N _{End} =21 TPS=Chronic	E: Bilateral training + EMG-triggered NMES C: Unilateral training + EMG-triggered NMES Duration: 30min, 7d/wk for 6wk	 Fugl-Meyer Assessment (-) Arm Motor Ability Test (-)
Cauraugh & Kim (2002) RCT (5) N _{start} =25 N _{end} =25 TPS=Chronic	E: EMG-triggered NMES + bilateral training E2: EMG-triggered NMES + unilateral trainin C: Control Duration: 90min, 4d/wk for 2wk	 E1 vs E2/C Box and Block Test: (+exp) E2 vs C Box and Block Test (+exp₂)
	Bilateral arm training com	pared to CIMT
Brunner et al. (2012) RCT (7) Nstart=30 Nend=30 TPS=Not given	E: Bilateral training C: mCIMT Duration: 4h, 7d/wk for 4wk	 Action Research Arm Test (-) 9 Hole Peg Test (-) Motor Activity Log (-)
WU et al. (2011) RCT (5) Nstart=66 Nend=58 TPS=Chronic	E: dCl1 E2: Bilateral training C: Control Duration: 2h, 5d/wk for 3wk	 E/E2 VS C Normalized Movement Unit for unilateral and bilateral task (+exp, exp2) E2 vs C Peak Velocity for unilateral and bilateral tasks (exp2) E vs C Wolf Motor Function Test (+exp) E vs E2/C Motor Activity Log (+exp) Wolf Motor Function Test (-) Peak Velocity for unilateral and bilateral tasks (-) Normalized Movement Unit for unilateral and bilateral task (-)
Modified CIM	T with unilateral training compared to rhyth	mic auditory cueing with bilateral arm training
van Delden et al. (2015) RCT (6) N _{Start} =60 N _{End} =52 TPS=Subacute	E: Modified CIMT + unilateral training E2: Rhythmic auditory cueing + bilateral training C: Dose-matched Control Duration: 1h, 3d/wk for 6wk	 E2 vs C Bimanual coordination task: (+exp₂) E vs C Unimanual reference task (+con) E vs E2 Unimanual reference task (+exp₂)
<u>van Delden et al. (2013)</u> RCT (6)	E1: Modified CIMT + unilateral training E2: Rhythmic auditory cueing + bilateral training	 Action Research Arm Test (-) Nine Hole Peg Test (-) Motricity Index (-)

N _{Start} =60	C: Dose-matched control group	•	Fugl-Meyer Assessment (-)
N _{End} =55	Duration: 1h, 3d/wk for 6wk	•	Motor Activity Log (-)
TPS=Subacute		•	Stroke Impact Scale (-)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time

post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05
Conclusions about bilateral arm training

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	Bilateral arm training may not have a difference in efficacy when compared to unilateral arm training for improving motor function.	11	Shim et al. 2015; McCombe et al. 2014; Byl et al. 2013; Dispa et al. 2013; Kim et al. 2013; Wu et al. 2013; Morris and van Wijck, 2012; Yang et al. 2012; Lin et al. 2010; Stoykov et al. 2009; Morris et al. 2008
1a	Bilateral arm training may produce greater improvements in motor function than conventional therapy.	4	Meng et al. 2018; Lee et al. 2017; Stinear et al. 2008; Desrosiers et al. 2005
1b	Bilateral arm training with rhythmic auditory cueing may not have a difference in efficacy when compared to unilateral arm training or conventional therapy for improving motor function.	2	Whiteall et al. 2011; Luft et al. 2004
1b	Task-oriented bilateral arm training may not have a difference in efficacy when compared to task-oriented training for improving motor function.	1	Hsieh et al. 2016
2	EMG-triggered NMES with bilateral arm training may not have a difference in efficacy when compared to EMG-triggered NMES with unilateral arm training for improving motor function.	1	Singer et al. 2013
1b	Bilateral arm training may not have a difference in efficacy when compared to CIMT for improving motor function.	2	Brunner et al. 2012; Wu et al. 2011
1a	There is conflicting evidence about the effect of bilateral arm training with rhythmic auditory cueing to improve motor function when compared to mCIMT .	2	Van Delden et al. 2015; Van Delden et al. 2013

SPASTICITY

LoE	Conclusion Statement	RCTs	References
1b	Bilateral arm training may not have a difference in efficacy when compared to TENS for improving spasticity.	1	Stinear et al. 2014

STROKE SEVERITY				
LoE	Conclusion Statement	RCTs	References	
1b	Task-oriented bilateral arm training may not have a difference in efficacy when compared to task-oriented training for improvements on measures of stroke severity.	1	Hsieh et al. 2016	

DEXTERITY			
LoE	Conclusion Statement	RCTs	References
1a	Bilateral arm training may not have a difference in efficacy when compared to unilateral arm training for improving dexterity.	5	Han and Kim, 2016; McCombe et al. 2014; Dispa et al. 2013; Morris and van Wijck, 2012; Morris et al. 2008
1a	There is conflicting evidence about the effect of bilateral arm training to improve dexterity when compared to conventional therapy .	2	Lee et al. 2017; Desrosiers et al. 2005
1b	Task-oriented bilateral arm training may not have a difference in efficacy when compared to task- oriented training for improving dexterity.	1	Hsieh et al. 2016
2	Task-oriented bilateral arm training may produce greater improvements in dexterity than bilateral arm training with rhythmic auditory cueing.	1	Song et al. 2015
2	EMG-triggered NMES with bilateral arm training may produce greater improvements in dexterity than EMG-triggered NMES with unilateral arm training or conventional therapy.	1	Cauraugh and Kim, 2002
1b	Bilateral arm training may not have a difference in efficacy when compared to CIMT for improving dexterity.	1	Brunner et al. 2012
1b	Bilateral arm training with rhythmic auditory cueing may not have a difference in efficacy when compared to mCIMT for improving dexterity.	1	Van Delden et al. 2013

MUSCLE STRENGTH				
LoE	Conclusion Statement	RCTs	References	
1b	There is conflicting evidence about the effect of bilateral arm training to improve muscle strength when compared to unilateral arm training .	2	Han and Kim, 2016; Yang et al. 2012	
1a	Bilateral arm training may not have a difference in efficacy when compared to conventional therapy for improving muscle strength.	2	Stinear et al. 2008; Desrosiers et al. 2005	
1b	Bilateral arm training with rhythmic auditory cueing may not have a difference in efficacy when compared to unilateral arm training or conventional therapy for improving muscle strength.	2	Whiteall et al. 2011; Luft et al. 2004	
1b	Task-oriented bilateral arm training may not have a difference in efficacy when compared to task-oriented training for improving muscle strength.	1	Hsieh et al. 2016	

ACTIVITIES OF DAILY LIVING			
LoE	Conclusion Statement	RCTs	References
1a	Bilateral arm training may not have a difference in efficacy compared to unilateral arm training for improving performance of activities of daily living.	7	Shim et al. 2015; Dispa et al. 2013; Wu et al. 2013; Lin et al. 2010; Stoykov et al. 2009; Morris et al. 2008; Summers et al. 2007
1a	There is conflicting evidence about the effect of bilateral arm training to improve performance of activities of daily living when compared to conventional therapy .	3	Lee et al. 2017; Lee et al. 2013; Desrosiers et al. 2005
1b	Bilateral arm training with rhythmic auditory cueing may not have a difference in efficacy when compared to unilateral arm training for improving performance of activities of daily living.	1	Whiteall et al. 2011
1b	There is conflicting evidence about the effect of task-oriented bilateral arm training to improve performance of activities of daily living when compared to task-oriented training .	1	Hsieh et al. 2016
2	Task-oriented bilateral arm training when compared to bilateral arm training with rhythmic auditory cueing may produce greater improvements in performance of activities of daily living.	1	Song et al. 2015
1b	Bilateral arm training may not have a difference in efficacy when compared to TENS for improving performance of activities of daily living.	1	Stinear et al. 2014
2	EMG-triggered NMES with bilateral arm training may not have a difference in efficacy when compared to EMG-triggered NMES with unilateral arm training for improving performance of activities of daily living.	1	Singer et al. 2013
1b	There is conflicting evidence about the effect of bilateral arm training to improve performance of activities of daily living when compared to CIMT .	2	Brunner et al. 2012; Wu et al. 2011
1b	Bilateral arm training with rhythmic auditory cueing may not have a difference in efficacy when compared to mCIMT for improving performance of activities of daily living.	1	Van Delden et al. 2013

Key points

The literature is mixed regarding bilateral arm training for upper limb rehabilitation following stroke.

Bilateral arm training may not be beneficial compared to unilateral training for upper limb function.

Bilateral arm training in combination with other therapy approaches may not be beneficial for upper limb rehabilitation.

Strength training



Adopted from: <u>https://www.flintrehab.com/2018/arm-exercises-for-stroke-patients/</u>

Strength training can be defined as an intervention involving repetitive and effortful muscle contractions with the goal of increasing motor unit activity (Ada et al. 2006). The interventions analyzed were classified as either traditional strength training or functional strength training. Traditional strength training involves resistance training in which individual muscles are often isolated and stabilized through protocols involving free weights or machines (Tomljenovic et al. 2011). Functional strength training is based on the principle of specific adaptations to imposed demands (SAID) in which training programs involve tasks that are modeled after common daily activities (Tomljenovic et al. 2011). These tasks often involve multiple muscle groups and require functional movements that are more applicable and may produce gains in strength in performing everyday tasks (Tomljenovic et al. 2011).

18 RCTs were found evaluating strength training for upper extremity motor rehabilitation. Nine RCTs compared strength training to conventional rehabilitation, simple joint mobilization or scapular exercises (Coroian et al. 2018; Dell'Uomo et al. 2017; Kim et al. 2017; Kim and Yim, 2017; Jeon et al. 2016; Da Silva et al. 2015; Lin et al. 2015; Winstein et al. 2004; Trombly et al. 1986). Four RCTs looked at strength training compared to task-specific training (Folkerts et al. 2017; Awad et al. 2015; Thielman et al. 2013; Corti et al. 2012). Three RCTs compared functional strength training to conventional therapy, non-functional strength training or movement performance therapy (Hunter et al. 2018; Park et al. 2017; Graef et al. 2016). Two RCTs looked at functional strength training compared to task-specific training (Agni and Kulkarni, 2017; Pattern et al. 2013).

The methodological details and results of all 18 RCTs are presented in Table 3.

Table 3.	RCTs evaluating strength training	interventions for upper	[·] extremity motor
rehabilita	ation		

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Strength training	versus conventional rehabilitation, simple joint	mobilization or scapular exercises
Coroian et al. (2018) RCT (7) N _{Start} =20 N _{End} =16 TPS=Chronic	E: Isokinetic Strengthening C: Passive Joint Mobilization Duration: 45min/d, 3d/wk for 6wk	 Fugl-Meyer Assessment (+con) Isokinetic Peak Torque (-) Box and Block Test (-) Modified Ashworth Scale (-)
Dell'Uomo et al. (2017) RCT (5) N _{Start} =28 N _{End} =28 TPS=Subacute	E: Scapulohumeral Rehabilitation C: Conventional Arm/Trunk Rehabilitation Duration: 20min/d, 5d/wk for 6wk	 Barthel Index (-) Fugl-Meyer Assessment (-) Modified Ashworth Scale (-)
<u>Kim et al.</u> (2017) RCT (5) N _{Start} =24 N _{End} =17 TPS=Chronic	E: Scapular Stabilization Exercise C: Simple Scapular Exercise Duration: 30min/d, 3d/wk for 8wk	Manual Function Test (+exp)
Kim & Yim (2017) RCT (5) N _{Start} =30 N _{End} =29 TPS=Chronic	E: Hand Training and Treadmill Weight Bearing Training C: Conventional Therapy Duration: 30min/d, 3d/wk for 6wk	Handgrip Strength (-)
<u>Jeon et al.</u> (2016) RCT (5) N _{Start} =12 N _{End} =12 TPS=Chronic	E: Repetitive bilateral and unilateral movements with strength exercises C: Conventional rehabilitation Duration: 30min/d, 3d/wk for 12wk	 Flexion and abduction range of motion (+exp)
Da Silva et al. (2015) RCT (8) Nstart=20 NEnd=20 TPS=Chronic	E: Strength training C: Standard care Duration: 30min/d, 2d/wk for 6wk	 TEMPA (+exp) Glumerohumeral flexion strength (+exp) Active shoulder Range of Motion (+exp) Fugl-Meyer Assessment (+exp)
Lin et al. (2015) RCT (7) N _{Start} =33 N _{End} =33 TPS=Chronic	E: Bilateral Isometric Handgrip Force Training with Visual Feedback C: Routine Therapy Duration: 30min/d, 3d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Wolf Motor Function Test (+exp) Motor Assessment Scale (+exp) Barthel Index (+exp)
Winstein et al. (2004) RCT (6) N _{start} =64 N _{end} =44 TPS=Acute	E1: Strength training E2: Functional task practice C: Standard care Duration: 1h/d, 5d/wk for 4wk	 <u>E1/E2 vs. C</u> Fugl Meyer Assessment: (+exp & +exp₂) Functional test of the hemiparetic upper extremity (+exp & +exp₂) Isometric torque (+exp & +exp₂)
Trombly et al. (1986) RCT (4) N _{start} =20 N _{end} =20 TPS=Chronic	E1: Resisted Grasp E2: Resisted Extension C: Ballistic Extension Duration: 7d/wk for 3wk	 Finger Extension Range of Motion (-) Speed and ability to rapidly reverse movement (-)
	Strength training versus task-specifie	c training
Folkerts et al (2017) RCT Crossover (4)	E1: Eccentric Strength Training followed by Task-Oriented Strength Training	 Action Research Arm Test (-) Shoulder, Elbow and Wrist Strength (-)

Nstart=11 NEnd=10 TPS=Chronic	E2: Task-Oriented Strength Training followed by Eccentric Strength Training Duration: 3d/wk for 4wk	
Awad et al. (2015) RCT (4) N _{Start} =30 N _{End} =23 TPS=Chronic	E: Shoulder Strength Training, Trunk Control Training, and Additional Strengthening Exercises. C: Shoulder Strength Training and Trunk Control Training. Duration: 3d/wk for 6wk	 Shoulder Abduction Peak Torque (+exp) Shoulder External Rotator Peak Torque (+exp) Supraspinatus Peak Force (+exp) Upper Trapezius Peak Force (+exp) Serratus Anterior Peak Force (+exp) Scapular Upward Rotation Angle (+exp) Spinal Lateral Deviation Angle (+exp)
Thielman et al. (2013b) RCT (6) N _{Start} =16 N _{End} =16 TPS=Chronic	E: Progressive resistive strength training C: Task-related training Duration: <i>Not reported</i>	 Activate range of motion for shoulder and elbow (+exp) Wolf Motor Function Test (+exp) Reaching (+exp)
Corti et al. (2012) RCT Crossover (7) N _{start} =14 N _{end} =14 TPS=Chronic	E1: Power Training E2: Functional Task Practice Duration: 90min/d, 3d/wk for 10wk	 Shoulder Flexion and Elbow Extension (+exp)
Functional strength tr	aining versus conventional therapy, strength tra	aining or movement performance therapy
Hunter et al. (2018) RCT (6) N _{Start} =288 N _{End} =240 TPS=Acute	E: Functional Strength Training C: Movement Performance Therapy Duration: 90min/d, 5d/wk for 6wk	 Action Research Arm Test (-) Wolf Motor Function Test (-) Grip and Pinch Force (-)
Park et al. (2017) RCT (5) N _{start} =30 N _{End} =26 TPS=Subacute	E: Boxing C: Conventional Therapy Duration: 30min/d, 3d/wk for 6wk	 Manual Function Test (+exp) Unaffected Side Hand Grip Strength (+exp)
Graef et al. (2016) RCT (8) N _{Start} =28 N _{End} =27 TPS=Chronic	E: Strength training with a functional goal C: Strength training with non-functional movements Duration: 30min/d, 3d/wk for 5wk	 Upper-Extremity Performance Test (+exp) Shoulder Strength (-) Grip Strength (-) Shoulder Active Range of Motion (-) Fugl-Meyer Assessment (-) Modified Ashworth Scale (-)
	Functional strength training versus task-s	pecific training
Agni and Kulkarni (2017) RCT (5) Nstan=45 Nend=37 TPS=Chronic	E1: Strength Training E2: Functional Task-Related Training E3: Functional Task-Related Training with Strength Training Duration: 70min/d, 3d/wk for 6wk	 E1 vs. E2: Chedoke Arm and Hand Inventory (exp₂) Manual Muscle Strength (+exp) Fugl-Meyer Assessment (-) E1 vs E3: Chedoke Arm and Hand Inventory (+exp₃) Manual Muscle Strength (+exp₃) Fugl-Meyer Assessment (-) E2 vs E3: Chedoke Arm and Hand Inventory (-) Manual Muscle Strength (+exp₃) Fugl-Meyer Assessment (-) E2 vs E3: Chedoke Arm and Hand Inventory (-) Manual Muscle Strength (+exp₃) Fugl-Meyer Assessment (-)
Patten et al. (2013) USA RCT (7) N _{start} =19 N _{end} =17	E: Functional Task Practice and Power Training C: Functional Task Practice Duration: 75min/d, 3d/wk for 4wk	 Wolf Motor Function Test (-) Ashworth Scale (-) Functional Independence Measure (+exp)

TPS=Chronic		
Abbreviations and table notes: C=c	ontrol group: D=days: E=experimental group: H=hours: Mir	=minutes: RCT=randomized controlled trial: TPS=time

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group - indicates no statistically significant between groups differences at α =0.05

Conclusions about strength training

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	Strength training may produce greater improvements in motor function than conventional therapy, simple joint mobilization or scapular exercises.	6	Coroian et al. 2018; Dell'Uomo et al. 2017; Kim et al. 2017; Da Silva et al. 2015; Lin et al. 2015; Winstein et al. 2004
1b	There is conflicting evidence about the effect of strength training to improve motor function when compared to task-specific training.	3	Agni and Kulkarni, 2017; Folkerts et al. 2017; Thielman et al. 2013
1a	Functional strength training may not have a difference in efficacy when compared to conventional therapy, strength training or movement performance therapy for improving motor function.	4	Hunter et al. 2018; Agni and Kulkarni, 2017; Park et al. 2017; Graef et al. 2016
1b	Functional strength training may not have a difference in efficacy when compared to task-specific training for improving motor function.	2	Agni and Kulkarni, 2017; Pattern et al. 2013

DEXTERITY

LoE	Conclusion Statement	RCTs	References		
1b	Strength training may not have a difference in efficacy when compared to conventional therapy, simple joint mobilization or scapular exercises for improving dexterity.	2	Corian et al. 2018; Trombly et al. 1986		

SPASTICITY				
LoE	Conclusion Statement	RCTs	References	
1b	Strength training may not have a difference in efficacy when compared to conventional therapy, simple joint mobilization or scapular exercises for improving spasticity.	2	Coroian et al. 2018; Dell'Uomo et al. 2017	
1b	Functional strength training may not have a difference in efficacy when compared to strength training for improving spasticity.	1	Graef et al. 2016	
1b	Functional strength training may not have a difference in efficacy when compared to task-specific training for improving spasticity.	1	Pattern et al. 2013	

RANGE OF MOTION					
LoE	LoE Conclusion Statement RCTs Reference				
1a	Strength training may produce greater improvements in range of motion than conventional therapy, simple joint mobilization or scapular exercises.	4	Jeon et al. 2016; Da Silva et al. 2015; Winstein et al. 2004; Trombly et al. 1986		
1a	Strength training may produce greater improvements in range of motion than task-specific training .	2	Thielman et al. 2013; Corti et al. 2012		

1b	Functional strength training may not have a difference in efficacy when compared to strength	1	Graef et al. 2016
	training for improving range of motion.		

ACTIVITIES OF DAILY LIVING				
LoE	Conclusion Statement	RCTs	References	
1b	There is conflicting evidence about the effect of strength training to improve performance of activities of daily living when compared to conventional therapy, simple joint mobilization or scapular exercises .	2	Dell'Uomo et al. 2017; Lin et al. 2015	
2	Functional strength training may produce greater improvements in performance of activities of daily living than strength training .	1	Agni and Kulkarni, 2017	
1b	There is conflicting evidence about the effect of functional strength training to improve performance of activities of daily living when compared to task-specific training .	2	Agni and Kulkarni, 2017; Pattern et al. 2013	

MUSCLE STRENGTH			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of strength training to improve muscle strength when compared to conventional therapy, simple joint mobilization or scapular exercises.	3	Coroian et al. 2018; Kim and Yim, 2017; Da Silva et al. 2015
2	Strength training may produce greater improvements in muscle strength than task-specific training.	3	Agni and Kulkarni, 2017; Folkerts et al. 2017; Awad et al. 2015
1a	There is conflicting evidence about the effect of functional strength training to improve muscle strength when compared to conventional therapy , strength training or movement performance therapy .	4	Hunter et al. 2018; Agni and Kulkarni, 2017; Park et al. 2017; Graef et al. 2016
2	Functional strength training may produce greater improvements in muscle strength than task-specific training.	1	Agni and Kulkarni, 2017

Key points

The literature is mixed regarding strength training and functional strength training for upper limb rehabilitation following stroke.

Task-specific training



Adopted from: https://www.sittoday.com/lifestyles/health-med-fit/custom-made-rehab-helps-victims-of-stroke/article_06eb5759-3291-5730-930f-725c0d436450.html

Task-specific training involves integrating tasks that are relevant to daily life (e.g. pouring a drink into a cup) into rehabilitation programs, while repetitive task training involves repeated practice of these tasks (Van Peppen et al. 2004; McCombe Waller et al. 2008; Stewart et al. 2006). Usually these consist of motor tasks that are focused on improvement of performance and function through goal-directed practice and repetition (Hubbard et al. 2009). It is well established that task-specific practice is required for motor learning to occur (Schmidt, 1991). Focal transcranial magnetic stimulation and functional magnetic resonance imaging have shown that task-specific training, in comparison to traditional stroke rehabilitation, yields long-lasting cortical reorganization specific to the corresponding areas being used (Classen et al.1998). More specifically, Karni et al. (1995), using functional magnetic resonance imaging, and Classen et al. (1998), using transcranial magnetic stimulation, both reported a slowly evolving, long-term, experience-dependent reorganization of the adult primary motor cortex following daily practice of task-specific motor activities.

Also, of interest is that task-specific sessions (i.e., thumb and hand movements), as short as 15 minutes in duration, are also effective in inducing lasting cortical representational changes (Bütefisch et al.1995; Classen et al.1998). According to Page (2003), intensity alone does not account for the differences between traditional stroke and task-specific rehabilitation. For example, Galea et al. (2001) reported that stroke patients who underwent a 3-week long program consisting of 45-minute task-specific, upper limb training showed improvements in measures of motor function, dexterity, and increased use of the more affected upper limbs. According to Page (2003), other, task-specific, low-intensity regimens designed to improve use and function of the affected limb have also reported significant improvements (Smith et al. 1999; Whitall et al. 2000; Winstein et al. 2001).

A total of 16 RCTs were found that looked task-specific training for upper extremity motor rehabilitation. 12 RCTs looked at task-specific training compared to conventional rehabilitation (Skubik-Peplaski et al. 2017; Brkic et al. 2016; Winstein et al. 2016; Kim et al. 2015; Hubbard et al. 2015; Zondervan et al. 2014; Shimodozono et al. 2013; Thielman et al. 2013; Arya et al. 2012; Thielman et al. 2012; Boyd et al. 2010; Thielman et al. 2004). Two RCTs looked at the intensity of task-specific training delivered (Waddell et al. 2017; Lang et al. 2016). One RCT looked at robotic training with task-specific training compared to robotic training (Hung et al. 2016), and another RCT looked at EMG-triggered NMES with task-specific training compared to EMG-triggered NMES (Kim et al. 2016).

The methodological details and results of all 16 RCTs are presented in Table 4.

Authors (Year) Study Design (PEDro Score) Sample Sizestart	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Sample Size _{end} Time post stroke category		
	Task-specific training compared to conventior	nal rehabilitation
Skubik-Peplaski et al. (2017) RCT (7) N _{Start} =16 N _{End} =16 TPS=Chronic	E: Repetitive Task Practice C: Occupation-Based Intervention Duration: 55min/d, 2d/wk for 4wk	 Fugl-Meyer Assessment (-) Stroke Impact Scale (-) Canadian Occupational Performance Measure (-)
Brkic et al. (2016) RCT (5) N _{Start} =24 N _{End} =22 TPS=Acute	E: Repetitive upper limb functional task practice C: Conventional rehabilitation Duration: 7d/wk for 4wk	 Action Research Arm Test (+exp) Grip Strength (+exp)
Winstein et al. (2016) ICARE Trial RCT (7) N _{start} =361 N _{End} =361 TPS=Subacute	E1: Structured, task-oriented upper extremity training E2: Dose-equivalent occupational therapy C: Monitoring-only occupational therapy Duration: 1h/d, 3d/wk for 10wk	 <u>E1/E2 vs C; E1 vs E2</u> Wolf Motor Function Test: (-) <u>E1/E2 vs C; E1 vs E2</u> Stroke Impact Scale: (-)
Kim et al. (2015) RCT (8) N _{Start} =44 N _{End} =40 TPS=Chronic	E: Target reach training with visual biofeedback, routine occupational and physical therapy C: Routine occupational and physical therapy Duration: 1h/d, 3d/wk for 4wk	 Fugl-Meyer Upper Extremity (+exp) Wolf Motor Function Test (+exp) Reaching speed (+exp) Range of Motion of the shoulder (+exp) Reach distance (-)
Hubbard et al. (2015) RCT (6) N _{Start} =23 N _{End} =23 TPS=Acute	E: Task-specific training and standard care C: Standard Care Duration: 2h/d, 5d/wk for 3wk	 Upper Limb Motor Assessment Scale (-) Modified Rankin Scale (-)
Zondervan et al. (2014) RCT (6) N _{Start} =17 N _{End} =16 TPS=Chronic	E: Self-guided, high-repetition home therapy with mechanical arm exerciser C: Conventional therapy Duration: 1h/d, 3d/wk for 3wk	 Fugl-Meyer Assessment (-) Motor Activity Log (-) Ashworth Scale (-)
Shimodozono et al. (2013) RCT (7) N _{Start} =52 N _{End} =49 TPS=Subacute	E: Repetitive functional exercise C: Conventional rehabilitation Duration: 40min/d, 5d/wk for 4wk	 Action Research Arm Test (+exp) Grasp and pinch (+exp) Fugl Meyer (+exp)
Thielman et al. (2013a) RCT (6) N _{start} =37 N _{end} =37 TPS=Chronic	E1: Task-Related Training (TRT) E2: Progressive Resistive Exercises (PRE) Duration: <i>Not reported</i>	 Motor Activity Log (+exp) Wolf Motor Function Test (+exp) Reaching Performance Scale (+exp) Fugl-Meyer Assessment (+exp)
Arya et al. (2012) MTST Trial RCT (9) N _{Start} =103 N _{End} =102 TPS=Subacute	E: Task-specific training C: Standard training using the Bobath approach Duration: 1h/d, 4-5d/wk for 4wk	 Fugl Meyer Score (+exp) Action Research Arm Test (+exp)
Thielman (2012) RCT (6) Nstart=16 NEnd=16 NEnd=16	E1: Task-Related Training E2: Resistive Exercise Training Duration: 40-45min/d, 2-3d/wk for 6wk	 Wolf Motor Function Test (-) Fugl Meyer Assessment (-)

Table 4. RCTs evaluating task-specific training for upper extremity motor rehabilitation

TPS=Chronic					
Boyd et al. (2010) RCT (5) N _{start} =18 N _{end} =18 TPS=Chronic	E: Task-specific training C: General arm training Duration: 3 sessions	Change in reaction and movement time (+exp)			
Thielman et al. (2004) RCT (4) N _{start} =12 N _{end} =12 TPS=Chronic	E: Progressive resistive exercises C: Task-related training Duration: 35min/d, 3d/wk for 4wk	 Modified Ashworth Scale (-) Rivermead Motor Assessment (-) 			
	Intensity of task-specific training	Ig			
Waddell et al. 2017 RCT (5) N _{Start} =85 N _{End} =78 TPS=Chronic	E1: 13.6 hours of task-specific training (100 repetitions/session) E2: 20 hours of task-specific training (200 repetitions/session) E3: 26.3 hours of task-specific training dose group (300 repetitions/session) Duration: 25-50min/d, 4d/wk for 8wk	Action Research Arm Test (-)			
Lang et al. (2016) RCT (5) N _{Start} =85 N _{End} =82 TPS=Chronic	E1: 3200 repetitions of task-specific upper limb training E2: 6400 repetitions of task-specific upper limb training E3: 9600 repetitions of task-specific upper limb training C: Individualized maximum repetitions Duration: 1h/d, 4d/wk for 8wk	 Action Research Arm Test (-) Stroke Impact Scale (-) Canadian Occupational Performance Measure (-) 			
	Robotic training with task-specific tr	raining			
Hung et al. (2016) RCT (8) N _{Start} =21 N _{End} =21 TPS=Chronic	E: Robotic training + task-specific training C: Robotic training + impairment-oriented training Duration: 20min/d, 3d/wk for 6wk	 Fugl-Meyer Assessment (+exp) Stroke Impairment Scale (+exp) 			
	EMG-triggered NMES with task-specific training				
Kim et al. (2016) RCT (6) N _{Start} =20 N _{End} =20 TPS=Chronic	E: EMG-triggered NMES with task-oriented training on paretic arm C: EMG-triggered NMES Duration: 30min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Box and Block Test (+exp) Jebsen-Taylor Hand Function Test (+exp) 			

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α=0.05 in favour of the control group

- indicates no statistically significant between groups differences at $\alpha{=}0.05$

Conclusions about task-specific training

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	Task-specific training may produce greater improvements in motor function than conventional therapy .	11	Skubik-Peplaski et al. 2017; Brkic et al. 2016; Winstein et al. 2016; Kim et al. 2015; Zondervan et al. 2014; Shimodozono et al. 2013; Thielman et al. 2013; Arya et al. 2012; Thielman et al. 2012; Boyd et al. 2010; Thielman et al. 2004	
2	Higher intensity task-specific training may not have a difference in efficacy when compared to lower intensity task-specific training for improving motor function.	2	Waddell et al. 2017; Lang et al. 2016	
1b	Robotic training with task-specific training may produce greater improvements in motor function than robotic training with impairment-oriented training.	1	Hung et al. 2016	
1b	EMG-triggered NMES with task-specific training may produce greater improvements in motor function than EMG-triggered NMES alone.	1	Kim et al. 2016	

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
1b	EMG-triggered NMES with task-specific training may produce greater improvements in dexterity than EMG-triggered NMES alone.	1	Kim et al. 2016	

SPASTICITY					
LoE	Conclusion Statement	RCTs	References		
1a	Task-specific training may produce greater improvements in spasticity than conventional therapy.	2	Zondervan et al. 2014; Thielman et al. 2004		

RANGE OF MOTION				
LoE	Conclusion Statement	RCTs	References	
1b	Task-specific training may produce greater improvements in range of motion than conventional therapy.	1	Kim et al. 2016	

STROKE SEVERITY				
LoE	Conclusion Statement	RCTs	References	
1b	Task-specific training may not have a difference in efficacy when compared to conventional therapy for improvements on measures of stroke severity.	1	Hubbard et al. 2015	

ACTIVITIES OF DAILY LIVING				
LoE	Conclusion Statement	RCTs	References	
1a	Task-specific training may not have a difference in efficacy when compared to conventional therapy for improving performance of activities of daily living.	5	Skubik-Peplaski et al. 2017; Winstein et al. 2016; Hubbard et al. 2015; Zondervan et al. 2014; Thielman et al. 2013	
2	Task-specific training may produce greater improvements in performance of activities of daily living than strength training .	1	Agni and Kulkarni, 2017	
2	Higher intensity task-specific training may not have a difference in efficacy when compared to lower intensity task-specific training for improving performance of activities of daily living.	1	Lang et al. 2016	
1b	Robotic training with task-specific training may produce greater improvements in performance of activities of daily living than robotic training with impairment-oriented training.	1	Hung et al. 2016	

MUSCLE STRENGTH						
LoE	LoE Conclusion Statement RCTs Reference					
1b	Task-specific training may produce greater improvements in muscle strength than conventional therapy.	2	Brkic et al. 2016; Shimodozono et al. 2013			

Key points

Task-specific training, alone or in combination with other therapy approaches, may be beneficial for some aspects of upper limb function following stroke.

Higher and lower intensity task-specific training may have similar effects on upper limb function.

Constraint-Induced Movement Therapy (CIMT)



Roughly 80% of all stroke survivors are left with motor impairments of the upper limb which affects their ability to perform activities of daily living (ADLs) (Kwakkel et al. 2016; Langhorne et al. 2009). Constraint-Induced Movement Therapy (CIMT) is a neurorehabilitation technique originally designed in the 1970s for the purpose of improving upper extremity function post-stroke (Christie et al. 2019; Morris et al. 2006). Traditional CIMT involves three key components: 1) immobilization of the non-paretic hand/arm using a mitt for 90% of waking hours, 2) high intensity task-oriented training with the paretic hand/arm, and 3) behavioural strategies to encourage use of the paretic upper limb after the patient leaves therapy, also known as a transfer package (Etoom et al. 2016).

CIMT is designed to overcome the tendency among hemiparetic patients to avoid the use of their paretic limb, a process termed "learned non-use". By constraining the non-paretic upper limb, the patient is forced to activate the muscles and neural pathways of their paretic limb, promoting neuroplasticity and use-dependent cortical reorganization (Taub et al. 1999). This form of treatment has shown promise, especially among stroke survivors with moderate upper limb disability. Modified versions of CIMT (mCIMT) have since been developed with varied dosage, timing, and composition of therapy but generally include less intense training of the paretic limb over a longer period of time (Kwakkel et al. 2016). CIMT is often compared to "forced use", or constraint only treatments, which are conceptually simpler versions of CIMT that do not apply operant training techniques.

Here we provide a review of 54 published RCTs related to CIMT for upper extremity motor rehabilitation. In order to better contextualize this body of evidence, studies were separated and classified according to the type of treatment (CIMT or mCIMT) as well as the time poststroke (acute/subacute phase (<6 months) or chronic stage (>6 months)), leading to 4 groups of RCTs. The authors' own declaration of the type of therapy (i.e. mCIMT or CIMT) was used for classification purposes.

The first two tables (Table 5, Table 6) list the summary of 12 RCTs examining CIMT in the acute/subacute phase (Seok et al. 2016; Shah et al. 2016; Song et al. 2016; Batool et al. 2015;

Thrane et al. 2015; Yoon et al. 2014; Dromerick et al. 2009; Boake et al. 2007; Ro et al. 2006; Page et al. 2005; Plougman and Corbett 2004; Dromerick et al. 2000) and 22 RCTs evaluating CIMT in the chronic phase (Souza et al. 2015; Nadeau et al. 2014; Takebayshi et al. 2013; Huseyinsinoglu et al. 2012; Khan et al. 2011; Wu et al. 2011; Lin et al. 2010; Wolf et al. 2010; Lin et al. 2009; Dahl et al. 2008; Lin et al. 2008; Sawaki et al. 2008; Wolf et al. 2008; Lin et al. 2007; Wu et al. 2007; Brogardh and Bengt, 2006; Richards et al. 2006; Underwood et al. 2006; Wolf et al. 2006; Alberts et al. 2004; Suputtitada et al. 2004; Wittenberg et al. 2003) poststroke.

The last two tables (Table 7, Table 8) list the summary of 7 RCTs examining mCIMT in the acute/subacute phase (Kwakkel et al. 2016; Liu et al. 2016; El-Helow et al. 2014; Treger et al. 2012; Brogardh et al. 2009; Hammer and Lindmark, 2009; Myint et al. 2007) and 13 RCTs in the chronic phase (Doussoulin et al. 2017; Hsieh et al. 2016; Yadav et al. 2016; Barzel et al. 2015; Smania et al. 2012; Wang et al. 2011; Hayner et al. 2010; Page et al. 2008; Lin et al. 2007; Wu et al. 2007b; Wu et al. 2007c; Page et al. 2004; Page et al. 2002).

		1		
Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)		
	E ON IT			
<u>Snan et al.</u> (2016a)		Motor Activity Log (+exp)		
RCI (5)	C: Motor Relearning Program	Nine Hole Peg Test (-)		
NStart=45	Duration: 80% of working hours	 Fugl-Meyer Assessment (-) 		
N _{End} =40				
TPS=Subacute				
Song et al. (2016a)	E: Scalp cluster acupuncture and Constraint	 Fugl-Meyer Assessment (-) 		
RCT (5)	Induced Movement Therapy			
N _{Start} =30	C: Body acupuncture and traditional rehabilitation			
N _{End} =30	therapy			
TPS=Acute	Duration: 5-6h, 6d/wk for 2wk			
Batool et al. (2015)		 Modified Ashworth Scale (+exp) 		
PCT (5)	C: Motor Poloarning Programmo	Eunctional Independence Measure (Leve)		
	Durotion 2b. Colvuls for 2wls			
INStart=42				
TPS=Subacute				
<u>Thrane et al.</u> (2015)	E: CIMT	 Wolf Motor Function Test (-) 		
RCT (7)	C: Usual Care	Stroke Impact Scale (-)		
N _{Start} =47	Duration: 3h, 1/d for 10d	 Fugl-Meyer Assessment (-) 		
N _{End} =47				
TPS=Acute				
Boake et al. (2007)	E: CIMT	Fugl Meyer Motor recovery (-)		
RCT (5)	C: Traditional rehabilitation	Grooved Pegboard test (-)		
Nstart=23	Duration: 3h. 6d/wk for 2wk	Motor Activity Log: Quality of Movement		
Nend=16		(+exp)		
TPS=Acute				
$P_{0} et al. (2006)$	E. CIMT	Grooved Rephard test (+eyp)		
PCT (6)	C: Traditional rebabilitation	Eucl-Meyer Assessment (+exp)		
	Duration: 2h. 6d/wk for 2wk	Motor Activity Log (Lovp)		
Nstart-O				
TPS=Acute				
Page et al. (2005b)	E: CIMT	Action Research Arm Test (-)		
RCT (5)	C: Regular rehabilitation	 Fugl-Meyer Assessment (-) 		
N _{start} =10	Duration: 30min, 3d/wk for 10wk	Motor Activity Log (-)		
N _{end} =10				
TPS=Subacute				
Ploughman & Corbett (2004)	E: Forced Use Therapy (Constraint without	Chedoke McMaster Impairment Inventory (-)		
RCT (5)	Shaping)	Action Research Arm Test (-)		
N _{start} =23	C: Conventional Therapy	Functional Independence Measure (-)		
N _{end} =23	Duration: 1-6h (incremental increase), 5d/wk for			
TPS=Subacute	2wk			
Dromerick et al. (2000)	E. CIMT	Action Research Arm Test (+exp)		
BCT (6)	C: Traditional upper extremity therapy	Functional Independence Measure (-)		
N23	Duration: 2h 5d/wk for 2wk	Barthel Index (-)		
High Intensity CIMT compared to CIMT				
	E4. Use interacts ODAT	F0/0 F1		
VECTORS (Study Acronym)		$\frac{EZ/UVSE1}{2}$		
Dromerick et al. (2009)	E2: Standard CIMI	• Action Research Arm Test: (+exp ₂ , +con)		
RCI (6)	C: ADL and UE bilateral training Exercises	Functional Independence Measure (-)		
N _{start} =52	Duration: 2-3h, 5d/wk for 2wk	Stroke Impact Scale (-)		
N _{end} =52				

Table 5. Summary of RCTs Evaluating CIMT in the acute/subacute (<6months) phase for upper extremity motor rehabilitation</th>

TPS=Subacute					
	CIMT combined with another intervention				
Seok et al. (2016) RCT (5) N _{Start} =32 N _{End} =30 TPS=Subacute	E1: CIMT with Visual Biofeedback E2: Visual Biofeedback C: Conventional Occupational Therapy Duration: 1h, 5d/wk for 2wk	E1 vs C Grasp Strength (+exp) Pinch Strength (+exp) Wolf Motor Function Test (+exp) Fugl-Meyer Assessment (+exp) <u>E2 vs C</u> Grasp Strength (-) Pinch Strength (-) Wolf Motor Function Test (+exp ₂) Fugl-Meyer Assessment (+exp ₂)			
Yoon et al. (2014) RCT (7) Nstart=26 NEnd=26 TPS=Subacute	E1: CIMT combined with mirror therapy E2: CIMT C: Conventional therapy Duration: 6h, 5d/wk for 2wk	E1 v E2Box and block test (+exp)Nine-hole pegboard test (+exp)Grip strength (+exp)Brunnstrom Recovery Stages (-)Wolf motor function test (-)Fugl-Meyer Assessment (-)Korean Modified Barthel Index (-) $E1 v C$ Box and block test (+exp)Nine-hole pegboard test (+exp)Grip strength (+exp)Brunnstrom Recovery Stages (-)Wolf motor function test (+exp)Fugl-Meyer Assessment (-)Korean Modified Barthel Index (+exp)Fugl-Meyer Assessment (-)Korean Modified Barthel Index (+exp)E2 vs CBox and block test (+exp2)Nine-hole pegboard test (-)Grip strength (+exp2)Brunnstrom Recovery Stages (-)Wolf motor function test (+exp2)Fugl-Meyer Assessment (-)Korean Modified Barthel Index (+exp2)Fugl-Meyer Assessment (-)Wolf motor function test (+exp2)Fugl-Meyer Assessment (-)Korean Modified Barthel Index (+exp2)			

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group - indicates no statistically significant between groups differences at α =0.05

Table 6. Summary of RCTs evaluating CIMT in the chronic (>6months) phase poststroke for upper extremity motor rehabilitation

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Huseyinsinoglu et al. (2012) RCT (6) N _{start} =24 N _{end} =21 TPS=Chronic	E: CIMT C: Bobath Duration: 3h/d for 10d	 Motor Activity Log (+exp) Wolf Motor Function Test (-) Functional Independence Measure (-)
Khan et al. (2011) RCT (6) N _{start} =44 N _{end} =39 TPS=Chronic	E1: CIMT E2: Therapeutic Climbing C: Conventional Neurological Therapy Duration: 15-20h/wk for 4wk	E1 vs E2 Wolf Motor Function Test (+exp) Motor Activity Log (-) Isometric Strength (-) Active Range of Motion (-) <u>E1 vs C</u> Wolf Motor Function Test (-) Motor Activity Log (-) Isometric Strength (-) Active Range of Motion (-)
Wu et al. (2011) RCT (5) N _{start} =66 N _{end} =65 TPS=Chronic	E1: Distributed CIMT E2: Bilateral Arm Training C: Routine Therapy Duration: 2h, 5d/wk for 3wk	 E1/E2 vs C Unilateral and Bilateral Smoothness while Reaching: (+exp, +exp₂) <u>E1 vs E2/C</u> Motor Activity Log: (+exp) <u>E1 vs E2/C</u> Wolf Motor Function Test: (+exp)
Lin et al. (2010) RCT (5) N _{start} =13 N _{end} =13 TPS=Chronic	E: Distributed CIMT C: Routine Therapy Duration: 2h, 5d/wk for 3wk	 Fugl-Meyer Assessment (+exp) Motor Activity Log (+exp)
Lin et al. (2009a) RCT (5) N _{start} =32 N _{end} =32 TPS=Chronic	E: CIMT C: Dose Matched Control Intervention Duration: 2h, 5d/wk for 3wk	 Fugl-Meyer Assessment (+exp) Stroke Impact Scale (+exp) Nottingham Extended Activities of Daily Living (+exp) Motor Activity Log (+exp)
Dahl et al. (2008) RCT (8) N _{start} =30 N _{end} =30 TPS=Chronic	E: CIMT C: Community-based rehabilitation Duration: 6h, 5d/wk for 2wk	 Wolf Motor Function Test: post (+exp), 6mo (-) Motor Activity Log (-) Functional Independence Measure (-) Stroke Impact Scale (-)
Lin et al. (2008) RCT (5) N _{start} =22 N _{end} =22 TPS=Chronic	E: CIMT C: Traditional Intervention Duration: 2h, 5d/wk for 3wk	 Fugl-Meyer Assessment (+exp) Functional Independence Measure (+exp) Motor Activity Log (-) Nottingham Extended Activities of Daily Living Scale (-), mobility subsection (+exp)
Lin et al. (2007) RCT (5) N _{start} =35 N _{end} =32 TPS=Chronic	E: CIMT C: Traditional therapy (neurodevelopmental) Duration: 2h, 5d/wk for 3wk	 Fugl-Meyer Assessment (+exp) Functional Independence Measure (+exp) Motor Activity Log (-)
Wu et al. (2007a) RCT (6)	E: CIMT C: Regular interdisciplinary rehab	 Motor Activity Log (+exp) Fugl-Meyer Assessment (-)

N _{start} =47	Duration: 2h, 5d/wk for 3wk	
TPS=Chronic		
Underwood et al. (2006) RCT (8) N _{start} =41 N _{end} =32 TPS=Chronic	E: CIMT + shaping procedure C: Usual care Duration: 6h, 5d/wk for 2wk	 Fugl-Meyer Assessment (-) Wolf Motor Function Test (-)
Wolf et al. (2006) RCT (8) EXCITE Nstart=222 Nend=201 TPS=Chronic	E: CIMT + shaping procedure C: Usual care Duration: 6h, 5d/wk for 2wk	 Wolf Motor Function Test (+exp) Motor Activity Log (+exp)
Alberts et al. (2004) RCT (6) N _{start} =10 N _{end} =10 TPS=Subacute	E: CIMT C: Conventional rehabilitation Duration: 6h, 5d/wk for 2wk	 Maximum precision grip (+exp) Wolf Motor Function Test (+exp)
Suputtitada et al. (2004) RCT (6) N _{start} =69 N _{end} =69 TPS=Chronic	E: CIMT C: Bimanual-upper-extremity training based on NDT approach Duration: 6h, 5d/wk for 2wk	 Action Research Arm Test (+exp) Pinch test (+exp)
	High compared to low intensit	ty CIMT
<u>Souza et al.</u> (2015) RCT (5) N _{start} =24 N _{end} =19 TPS=Chronic	E1: CIMT high intensity (3h) E2: CIMT low intensity (1h) Duration: 1/3h, 3-4d/wk for 4wk	 Fugl-Meyer Assessment (-) Motor Activity Log (-)
Brogårdh & Bengt (2006) RCT (7) Nstart=16 N _{end} =16 TPS=Chronic	E: CIMT and using mitt at home for another 3 months every other day C: CIMT Duration: 6h, 5d/wk for 2wk	 Modified Motor Assessment Scale (-) Sollerman Hand Function Test (-) Motor Activity Log (-)
Wittenberg et al. (2003) RCT (5) Nstart=16 Nend=16 TPS=Chronic	E: Intense CIMT (6h) C: Less intense CIMT (3h) Duration: 3/6h/d for 10d	 Motor Activity Log (+exp) Wolf Motor Function Test (-) Assessment of Motor and Process Skills (-)
High intensity C	CIMT compared to low intensity CIMT con	nbined with cyloserine (antibiotic)
Nadeau et al. (2014) RCT (7) Nstart=24 NEnd=22 TPS=Chronic	E1: CIMT-6hr + cycloserine C1: CIMT-6hr + placebo E2: CIMT-2hr + cycloserine C2: CIMT-2hr + placebo Duration: 2/6h, 3-5d/wk for 10wk Early compared to delayed	 Fugl-Meyer Assessment (-) Wolf Motor Function Test (-) Motor Activity Log (-)
Wolf et al. (2010)	E1: CIMT early (3-9 months' post stroke)	Wolf Motor Function Test (+exp)
RCT (8) N _{start} =226 N _{end} =192 TPS=Chronic	E2: CIMT delayed (15 to 21 months post stroke) Duration: 90% of waking time for 2wk	 Motor Activity Log (+exp) Stroke Impact Scale (+exp)
<u>Sawaki et al. (2008)</u> RCT (8) N _{start} =30 N _{end} =30 TPS=Chronic	E: Early CIMT C: Delayed CIMT (4mo after randomization) Duration: 90% of d for 2wk	 Grip strength (+exp) Wolf Motor Function Test (-)
Wolf et al. (2008)	E1: CIMT early (3-9 months' post stroke)	Wolf Motor Function Test (+exp)

RCT (8) N _{start} =98 N _{end} =70 TPS=Chronic	E2: CIMT delayed (15 to 21 months post stroke) Duration: 90% of waking time for 2wk	 Motor Activity Log (+exp) Functional Independence Measure (+exp) Stroke Impact Scale (+exp)
	CIMT with transfer packa	ge
Takebayashi et al. (2013) RCT (5) Nstart=23 NEnd=21 TPS=Chronic Taub et al. (2013) RCT (5) Nstart=45 NEnd=40 TPS=Chronic	E: CIMT + transfer package (train affected arm) C: CIMT Duration: 4.5h spread over 2wk E1: Shaping training + CIMT transfer package (TP) E2: Repetitive task practice + TP E3: Repetitive task practice C: Shaping training	 Fugl-Meyer Assessment (-) Motor Activity Log (+exp) <u>E1/E2 vs. E3/C</u> Motor Activity Log (+exp, +exp₂) <u>E1/E2 vs. E3/C</u> Wolf Motor Function Test (+exp, +exp₂)
CIM	Γ combined with rTMS or donepezil (chol	inesterase inhibitor)
Richards et al. (2006) Secondary analyses of two parallel RCTs (7) N _{start} =39 N _{end} =35 TPS=Chronic	E1: Traditional CIMT (6h) + donepezil C1: Traditional CIMT (6h) + placebo E2: Shortened CIMT (1h) + repetitive transcranial magnetic stimulation (rTMS) C2: Shortened CIMT (1h) + sham rTMS Duration:1/6h, 5d/wk for 2wk	 E1 vs C1 Motor Activity Log (+exp) Wolf Motor Function Test: (-) E2 vs C2 Motor Activity Log (-) Wolf Motor Function Test: (-)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group - indicates no statistically significant between groups differences at α =0.05

phase for upper extremi		
Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Kwakkel et al. (2016) RCT (7) Nstart=159 NEnd=159 TPS=Subacute	 E1: Electromyographic Neuromuscular Stimulation on finger extensors E2: Modified Constraint Induced Movement Therapy C1: Unfavourable prognosis based on voluntary finger extension. Received usual care. C2: Favourable prognosis based on voluntary finger extension. Received usual care. Duration: 3h, 5d/wk for 3wk 	E2 vs C2; E1 vs C1 Action Research Arm Test: (+exp2) Fugl-Meyer Assessment: (-) Wolf Motor Function Test (-) Motricity Index (-) Erasmus Modified Nottingham Sensory Assessment (-) Nine-Hole Peg Test (-) Frenchay Arm Test (-) Motor Activity Log (-) Stroke Impact Scale-Hand (+exp2) E1 vs C Action Research Arm Test (-)
RCT (6) N _{Start} =90 N _{End} =86 TPS=Subacute	Therapy E2: Self-Regulated Modified Constraint Induced Movement Therapy C: Conventional Therapy Duration: 1h, 5d/wk for 2wk	 Action Research Arm Test (+exp) Fugl-Meyer Assessment (+exp) Lawton Instrumental Activities of Daily Living (+exp) Motor Activity Log (+exp) <u>E2 vs C</u> Action Research Arm Test (+exp₂) Fugl-Meyer Assessment (+exp₂) Lawton Instrumental Activities of Daily Living (-) Motor Activity Log (+exp₂) <u>E1 vs E2</u> Action Research Arm Test (-) Fugl-Meyer Assessment (+exp₂) Lawton Instrumental Activities of Daily Living (-) Motor Activity Log (+exp₂) E1 vs E2 Action Research Arm Test (-) Fugl-Meyer Assessment (+exp₂) Lawton Instrumental Activities of Daily Living (+exp₂) Motor Activity Log (+exp₂)
EI-Helow et al. (2014) RCT (6) N _{start} =60 N _{end} =60 TPS=Acute	E: Modified Constraint Induced Movement Therapy C: Conventional Rehabilitation Duration: 6h/d for 2wk	 Fugl-Meyer Assessment (+exp) Action Research Arm Test (+exp)
Treger et al. (2012) RCT (7) N _{start} =28 N _{end} =28 TPS=Subacute	E: mCIMT C: Traditional rehabilitation Duration: 4h, 2d/wk for 2wk	 Functional Independence Measure (-) Manual Function Test (-)
Brogårdh et al. (2009) RCT (5) N _{start} =24 N _{end} =24 TPS=Subacute	E: Shortened CIMT (mitt use) C: No mitt use Duration: 90% of waking time for 12d	 Motor Assessment Scale (-) Sollerman Hand Function Tst (-) 2-Point Discrimination Test (-) Motor Activity Log Test (-)

Table 7. Summary of RCTs Evaluating Modified CIMT in the acute/Subacute (<6 months) phase for upper extremity motor rehabilitation

Hammer & Lindmark (2009)	E: Restraining sling and Standard	Fugl-Meyer Assessment (-)
RCT (6)	Rehabilitation	Action Research Arm Test (-)
N _{Start} =30	C: Standard Rehabilitation	 Motor Assessment Scale (-)
N _{End} =26	Duration: 6h, 5d/wk for 2wk	 16-Hole Peg Test (-)
TPS=Subacute		 Grip strength ratio (-)
		Modified Ashworth Scale (-)
Myint et al. (2007)	E: mCIMT	Action Research Arm Test (+exp)
RCT (7)	C: Traditional rehabilitation	 Motor Activity Log (+exp)
N _{start} =43	Duration: 4h/d for 10d	
N _{end} =43		
TPS=Subacute		

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp2 indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Table 8. Summary of RCTs Evaluating Modified CIMT in the Chronic (>6 months) phase for upper extremity motor rehabilitation

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Hsieh et al. (2016) RCT (7) Nstart=34 Nend=34 TPS=Chronic	E: mCIMT C: Regular Therapy Duration: 105min, 5d/wk for 4wk	 Wolf Motor Function Test (+exp) Nottingham Extended Activities of Daily Living (+exp) Functional Independence Measure (-)
Yadav et al. (2016) RCT (5) Nstart=65 Nend=60 TPS=Chronic	E: mCIMT C: Conventional rehabilitation Duration: 3h, 3d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Motor Activity Log (+exp)
Barzel et al. (2015) RCT (6) N _{start} =156 N _{end} =156 TPS=Chronic	E: Home CIMT C: Standard Therapy Duration: 5h/wk for 4wk	 Motor Activity Log (+exp) Wolf Motor Function Test (-) Nine Hole Peg Test (-) Stroke Impact Scale (-) Barthel Index (-) Instrumental Activities of Daily Living (-)
<u>Smania et al.</u> (2012) RCT (8) N _{start} =66 N _{end} =40 TPS=Chronic	E: mCIMT C: Dose-match task-specific therapy Duration: 2h, 5d/wk for 2wk	 Wolf Motor Function Test (+exp) Motor Activity Log (+exp)
Wang et al. (2011) RCT (4) Nstart=30 Nend=30 TPS=Chronic	E1: mCIMT E2: Intensive conventional therapy C: Conventional therapy Duration: 3h, 5d/wk for 4wk	Wolf Motor Function Test (+exp)
Hayner et al. (2010) RCT (4) N _{start} =12 N _{end} =12 TPS=Chronic	E: mCIMT C: Bilateral training Duration: 6h/d for 10d	 Wolf Motor Function Test (-) COPM (-)
Page et al. (2008) RCT (5) Nstart=35 N _{end} =35 TPS=Chronic	E1: mCIMT + physical and occupational therapy E2: Traditional rehab C: No therapy Duration: 5h, 5d/wk for 10wk	 Fugl-Meyer Assessment (-) Action Research Arm Test (+exp)
Lin et al. (2007) RCT (7) N _{start} =34 N _{end} =31 TPS=Chronic	E: mCIMT C: Traditional rehab Duration: 6h, 5d/wk for 3wk	 Motor Activity Log (+exp) Functional Independence Measure (+exp)
Wu et al. (2007b) RCT (5) Nstart=26 Nend=26 TPS=Chronic	E: mCIMT + a restraining mitt on the unaffected hand C: Traditional therapy Duration: 2h, 5d/wk for 3wk	 Fugl-Meyer Assessment (+exp) Functional Independence Measure (+exp) Motor Activity Log (+exp) Stroke Impact Scale (+exp)
<u>Wu et al. (2007c)</u> RCT (6) N _{start} =30	E: mCIMT C: Regular occupational therapy Duration: 2h, 5d/wk for 3wk	 Motor Activity Log (+exp) Functional Independence Measure (+exp)

Nerd-30		
TDO Chronie		
Page et al. (2004)	E: mCIMT	<u>E vs C1:</u>
RCT (6)	C1: Traditional Rehabilitation	 Fugl-Meyer Assessment (+exp)
Nstart=17	C2: No Therapy	 Action Research Arm Test (-)
Nend=17	Duration: 5h. 5d/wk for 10wk	E1 vs C2:
TPS-Chronic		 Fugl-Meyer Assessment (+exp)
		Action Research Arm Test (+exp)
		C1 vs C2:
		Fugl-Mever Assessment (-)
		 Action Research Arm Test (+con1)
		•
Page et al. (2002)	E1: mCIMT + physical and	Fugl-Mever Assessment (+exp)
BCT (5)	occupational therapy	Action Research Arm Test (+exp)
	E2: Traditional rebab	
N -14	C: No therepy	
TPS=Chronic	Duration: 30min, 3d/wk for 10wk	
	mCIMT in group or individual settin	g
Doussoulin et al. (2017)	E1: mCIMT group therapy	Motor Activity Log (+exp)
RCT (5)	E2: mCIMT individual therapy	Action Research Arm Test (+exp)
N _{Start} =36	Duration: 3h/d for 10d	Functional Independence Measure (+exp)
Nr		1 - (- 17
1		

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at $\alpha\text{=}0.05$ in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about CIMT and mCIMT

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	CIMT may not have a difference in efficacy when compared to conventional therapy or motor relearning programmes for improving motor function during the acute/subacute phase poststroke.	8	Shah et al. 2016; Song et al. 2016; Thrane et al. 2015; Yoon et al. 2014; Dromerick et al. 2009; Boake et al. 2007; Page et al. 2005; Plougman and Corbett 2004; Dromerick et al. 2000
2	CIMT combined with visual biofeedback may produce greater improvements in motor function than conventional therapy on its own during the acute/subacute phase poststroke.	1	Seok et al. 2016
1b	CIMT combined with mirror therapy may not have a difference in efficacy when compared to CIMT on its own for improving motor function during the acute/subacute phase poststroke.	1	Yoon et al. 2014
1a	CIMT may produce greater improvements in motor function than conventional therapy or neurodevelopmental techniques during the chronic phase poststroke.	13	Huseyinsinoglu et al. 2012; Khan et al. 2011; Wu et al. 2011; Lin et al. 2010; Lin et al. 2009; Dahl et al. 2008; Lin et al. 2008; Lin et al. 2007; Wu et al. 2007; Underwood et al. 2006; Wolf et al. 2006; Alberts et al. 2004; Suputtitada et al. 2004
1b	High intensity CIMT may not have a difference in efficacy when compared to low intensity CIMT on its own for improving motor function during the chronic phase poststroke.	3	Souza et al. 2015; Brogardh and Bengt, 2006; Wittenberg et al. 2003
1b	High intensity CIMT with/without cycloserine may not have a difference in efficacy when compared to low intensity CIMT with/without cycloserine for improving motor function during the chronic phase poststroke.	1	Nadeau et al. 2014
1a	Early CIMT may produce greater improvements in motor function than delayed CIMT during the chronic phase poststroke.	3	Wolf et al. 2010; Sawaki et al. 2008; Wolf et al. 2008
2	There is conflicting evidence about the of CIMT with the transfer package protocol when compared to traditional CIMT for improving motor function during the chronic phase poststroke.	2	Takebayashi et al. 2013; Taub et al. 2013
1a	There is conflicting evidence about the effect of mCIMT to improve motor function when compared to conventional therapy or bilateral arm training during the acute/subacute phase poststroke.	7	Kwakkel et al. 2016; Liu et al. 2016; El-Helow et al. 2014; Treger et al. 2012; Brogardh et al. 2009; Hammer and Lindmark, 2009; Myint et al. 2007
1a	mCIMT may produce greater improvements in motor function than conventional therapy or bilateral arm training during the chronic phase poststroke.	10	Hsieh et al. 2016; Yadav et al. 2016; Barzel et al. 2015; Smania et al. 2012; Wang et al. 2011; Hayner et al. 2010; Page et al. 2008; Wu et al. 2007b; Page et al. 2004; Page et al. 2002
2	Group based mCIMT may produce greater improvements in motor function than one on one mCIMT sessions during the chronic phase poststroke.	1	Doussoulin et al. 2017

DEXTERITY			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of CIMT to improve dexterity when compared to conventional therapy or motor relearning programmes during the	4	Shah et al. 2016; Yoon et al. 2014; Boake et al. 2007; Ro et al. 2006
	acute/subacute phase poststroke.		
1b	CIMT combined with mirror therapy may produce greater improvements in dexterity than CIMT on its own during the acute/subacute phase poststroke.	1	Yoon et al. 2014
1b	mCIMT not have a difference in efficacy when compared to conventional therapy or bilateral arm training for improving dexterity during the acute/subacute phase poststroke.	1	Kwakkel et al. 2016
1b	mCIMT not have a difference in efficacy when compared to conventional therapy or bilateral arm training for improving dexterity during the chronic phase poststroke.	1	Barzel et al. 2015

SPASTICITY

LoE	Conclusion Statement	RCTs	References
2	CIMT may produce greater improvements in spasticity than conventional therapy or motor relearning programmes during the acute/subacute phase poststroke.	1	Batool et al. 2015
1b	mCIMT not have a difference in efficacy when compared to conventional therapy or bilateral arm training for improving spasticity during the acute/subacute phase poststroke.	1	Hammer and Lindmark, 2009

RANGE OF MOTION					
LoE	LoE Conclusion Statement RCTs References				
1b	CIMT not have a difference in efficacy when compared to conventional therapy or neurodevelopmental techniques for improving range of motion during the chronic phase poststroke.	1	Khan et al. 2011		

PROPRIOCEPTION				
LoE	Conclusion Statement	RCTs	References	
1b	mCIMT not have a difference in efficacy when compared to conventional therapy or bilateral arm training for improving proprioception during the acute/subacute phase poststroke.	2	Kwakkel et al. 2016; Brogardh et al. 2009	

MUSCLE STRENGTH			
LoE	Conclusion Statement	RCTs	References
1b	CIMT may produce greater improvements in muscle strength than conventional therapy or motor relearning programmes during the acute/subacute phase poststroke.	1	Yoon et al. 2014
2	CIMT combined with visual biofeedback may produce greater improvements in muscle strength than conventional therapy or motor relearning programmes during the acute/subacute phase poststroke.	1	Seok et al. 2016
1b	CIMT combined with mirror therapy may produce greater improvements in muscle strength than CIMT on its own during the acute/subacute phase poststroke.	1	Yoon et al. 2014
1a	CIMT may produce greater improvements in muscle strength than conventional therapy or neurodevelopmental techniques during the chronic phase poststroke.	2	Alberts et al. 2004; Suputtitada et al. 2004
1b	Early CIMT may produce greater improvements in muscle strength than delayed CIMT during the chronic phase poststroke.	1	Sawaki et al. 2008
1a	mCIMT not have a difference in efficacy when compared to conventional therapy or bilateral arm training for improving muscle strength during the acute/subacute phase poststroke.	2	Kwakkel et al. 2016; Hammer and Lindmark, 2009

ACTIVITIES OF DAILY LIVING			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of CIMT to improve performance of activities of daily living when compared to conventional therapy or motor relearning programmes during the acute/subacute phase poststroke.	8	Shah et al. 2016; Batool et al. 2015; Thrane et al. 2015; Yoon et al. 2014; Boake et al. 2007; Ro et al. 2006; Page et al. 2005; Dromerick et al. 2000
1b	CIMT combined with mirror therapy may not have a difference in efficacy when compared to CIMT on its own for improving performance of activities of daily living during the acute/subacute phase poststroke.	1	Yoon et al. 2014
1a	CIMT may produce greater improvements in performance of activities of daily living than conventional therapy or neurodevelopmental techniques during the chronic phase poststroke.	10	Huseyinsinoglu et al. 2012; Khan et al. 2011; Wu et al. 2011; Lin et al. 2010; Lin et al. 2009; Dahl et al. 2008; Lin et al. 2008; Lin et al. 2007; Wolf et al. 2006
1b	High intensity CIMT may not have a difference in efficacy when compared to low intensity CIMT on its	3	Souza et al. 2015; Brogardh and Bengt, 2006; Wittenberg et al. 2003

	own for improving performance of activities of daily living during the chronic phase poststroke.		
1b	High intensity CIMT with/without cycloserine may not have a difference in efficacy when compared to low intensity CIMT with/without cycloserine for improving performance of activities of daily living during the chronic phase poststroke.	1	Nadeau et al. 2014
1a	Early CIMT may produce greater improvements in performance of activities of daily living than delayed CIMT during the chronic phase poststroke.	2	Wolf et al. 2010; Wolf et al. 2008
2	CIMT with the transfer package protocol may not have a difference in efficacy when compared to traditional CIMT for performance of activities of daily living during the chronic phase poststroke.	1	Takebayashi et al. 2013
1a	mCIMT not have a difference in efficacy when compared to conventional therapy or bilateral arm training for improving performance of activities of daily living during the acute/subacute phase poststroke.	6	Kwakkel et al. 2016; Liu et al. 2016; Treger et al. 2012; Brogardh et al. 2009; Hammer and Lindmark, 2009; Myint et al. 2007
1a	mCIMT may produce greater improvements in performance of activities of daily living than conventional therapy or bilateral arm training during the chronic phase poststroke.	8	Hsieh et al. 2016; Yadav et al. 2016; Barzel et al. 2015; Smania et al. 2012; Hayner et al. 2010; Lin et al. 2007; Wu et al. 2007b; Wu et al. 2007c
2	Group based mCIMT may produce greater improvements in performance of activities of daily living than one on one mCIMT sessions during the chronic phase poststroke.	1	Doussoulin et al. 2017

Key points

Constraint-induced movement therapy may be beneficial for upper limb rehabilitation in the chronic phase following stroke.
The literature is mixed regarding constraint-induced movement therapy for upper limb rehabilitation in the subacute/acute phase following stroke.
Modified constraint-induced movement therapy may be beneficial for upper limb rehabilitation in the chronic phase following stroke.
Modified constraint-induced movement therapy may not be beneficial for upper limb rehabilitation in the subacute/acute phase following stroke.
Higher and lower intensity constraint-induced movement therapy may have similar effects on upper limb function in the chronic phase following stroke.
The literature is mixed regarding constraint-induced movement therapy in combination with other therapy approaches for upper limb rehabilitation following stoke.

Trunk restraint



Adopted from: https://www.ortopedia-almirall.com/en/producto/cinturon-sujecion-tronco-y-pelvis-cierre-magnetico/

Reaching movements performed with the affected arm poststroke are often accompanied by compensatory trunk or shoulder girdle movements, which overextend the reach of the arm (Michaelsen et al. 2001). Restriction of compensatory trunk movements may encourage recovery of "normal" reaching patterns in the hemiparetic arm when reaching for objects placed within arm's length (Michaelsen & Levin, 2004). Eight RCTs (Bang et al. 2015; Lima et al. 2014; Wu et al. 2012a; Wu et al. 2012b; Thielman et al. 2010; Woodbury et al. 2009; Michaelsen et al. 2006; Michaelsen and Levin, 2004) have evaluated the effectiveness of trunk restraint combined with other training to improve the movement quality of reaching tasks. Their methodological details and results are presented in Table 9.

Authors (Year)						
Study Design (PEDro Score)	Duration: Session length, frequency per	Result (direction of effect)				
Sample Sizestart	week for total number of weeks					
Time post stroke category						
mCIMT + trunk restraint training						
Bang et al. (2015)	E: mCIMT + trunk resistant training	Action Research Arm Test (+exp)				
RCT (9)	C: mCIMT	Fugl-Meyer Assessment (+exp)				
Nstart=18	Duration: 30 min, 5 d/wk, for 4 wk	Modified Barthel Index (+exp)				
TPS=Subacute		• Motor Activity Log (+exp)				
Lima et al. (2014)	E: mCIMT + trunk resistant training	Motor Activity Log (-)				
RCT (8)	C: mCIMT	Bilateral Activity Assessment Scale (-)				
N _{Start} =22	Duration: Not Reported	Wolf Motor Function Test (-)				
NEnd=15		Global strength (-)				
Woodbury et al. (2009)	E: mCIMT + trunk restraint	Hand path trajectories (+exp)				
RCT (5)	C: mCIMT	······································				
N _{start} =11	Duration: 6 hr, 5d/wk for 2 wk					
Nend=11						
	Distributed CIT + trunk restraint tra	ining				
<u>Wu et al.</u> (2012a)	E1: Distributed constraint-induced therapy	<u>E1/E2 vs. C</u>				
RUT (5) Notart=57		 Action Research Arm Test (+exp, exp₂) Frenchav Activities Index (+exp, exp₂) 				
Nend=57	C: Usual care (neurodevelopmental treatment	 Motor Activity Log (+exp, exp2) 				
TPS=Chronic	techniques)	Stroke Impact Scale (+exp, exp ₂)				
	Duration: 2hr, 5d/wk for 3 wk	5 /52 0				
<u>Wu et al. (</u> 2012b)	E1: Distributed constraint-induced therapy	<u>E1/E2 vs. C</u> Motor Activity Log (Lovp. Lovp.)				
Not (5) N _{start} =45	E2: dCIT	 Fual-Mever Assessment (+exp) 				
N _{end} =45	C: Dose-matched control intervention	·				
TPS=Chronic	(neurodevelopmental treatment techniques)					
	Duration: 2hr, 3d/wk for 3 wk					
Thielman (2010)	E: Auditory feedback about trunk position	Reaching Performance Scale Near Target				
RCT (4)	C: Trunk restraint with external device	(+exp)				
N _{start} =16	Duration: 45 min, 3d/wk for 4 wk	Reaching Performance Scale Far Target				
N _{end} =16		(-)				
IPS=Unronic						
Reach to grasp training with trunk restraint						
RCT (7)	restraint	 Fugl-Mever Assessment (+exp) 				
Nstart=30	C: Unrestrained reach-to-grasp training	Box and Block Test (-)				
N _{End} =10	Duration: 40 min, 3d/wk for 5 wk					
IPS=Chronic	E. Deach to see the fair a structure start of	Obevildes herizontal - dducting ()				
RCT (5)	E: Reach-to-grasp training + trunk restraint	 Shoulder norizontal adduction (-) Shoulder flexion (-) 				
N _{start} =28	Duration: 60 sessions over 8 weeks	 Elbow Extension (+exp) 				
N _{end} =28						
TPS=Chronic						

Table 9. RCTs evaluating trunk restraint training for upper extremity motor rehabilitation

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α=0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at $\alpha\text{=}0.05$

Conclusions about trunk restraint training

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of trunk restraint combined with mCIMT to improve motor function when compared to mCIMT .	3	Bang et al. 2015; Lima et al. 2014; Woodbury et al. 2009
2	Trunk restraint combined with distributed CIT may produce greater improvements in motor function than conventional rehabilitation.	2	Wu et al. 2012a; Wu et al. 2012b
2	There is conflicting evidence about the effect of auditory feedback regarding trunk position to improve motor function when compared to trunk restraint training.	1	Thielman 2010
1b	Trunk restraint combined with reaching training may produce greater improvements in motor function than reaching training alone .	2	Michaelsen & Levin 2004; Michaelsen et al. 2006

LoE	Conclusion Statement	RCTs	References	
1h	Trunk restraint combined with reaching training	1	Michaelsen et al. 2006	
	difference in efficacy for dexterity.			

ACTIVITIES OF DAILY LIVING			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of trunk restraint combined with mCIMT to improve performance of activities of daily living when compared to mCIMT .	3	Bang et al. 2015; Lima et al. 2014; Woodbury et al. 2009
2	Trunk restraint combined with distributed CIMT may produce greater improvements in performance of activities of daily living than conventional rehabilitation.	2	Wu et al. 2012a; Wu et al. 2012b

Key points

Trunk restraint with reaching training or distributed constraint induced therapy may improve some aspects of upper limb function following stroke, but the effect of combining trunk restraint with constraint-induced movement therapy is less clear.

Stretching programs



Adopted from: http://advrehabnj.com/2014/10/08/trigger-finger-occupational-therapy/

Spasticity following stroke relates to hypertonicity or increased active tension of the muscle. Contracture may also occur as a result of spasticity and atrophic changes in the mechanical properties of muscles. Since surgery is the only treatment option once a contracture has developed, prevention is encouraged. Stretching may help to prevent contracture formation and, although well-accepted as a treatment strategy, although the evidence base is extremely limited for this intervention.

The methodological details and results of two RCTs evaluating stretching for upper extremity motor rehabilitation are presented in Table 10.
Table 10. RCTs evaluating stretching interventions for upper extremity motor rehabilitation

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
You et al. (2014) RCT (5) N _{Start} =45 N _{End} =41 TPS=Chronic	E1: Stretching program + joint stabilizing exercise (combo) E2: Stretching program C: Traditional exercise therapy Duration: 30min/d, 5d/wk for 8wk	 E1 vs C Muscle thickness (+exp) Motor assessment scale (+exp) E2 vs C Muscle thickness (+exp₂) Motor assessment scale (+exp₂) E1 vs E2 Muscle thickness (-) Motor assessment scale (-)
Tseng et al. (2007) RCT (7) Nstart=59 Nend=59 TPS=Chronic	E1: Nurse assisted range of motion exercise program E2: Nurse supervised range of motion exercise program C: Usual care Duration: 20-40min/d, 6d/wk for 4wk	 <u>E1/E2 vs C</u> Joint angles (+exp, +exp₂) FIM (+exp, +exp₂)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

- indicates no statistically significant between groups differences at d=0.05

Conclusions about stretching programs

SPASTICITY

LoE	Conclusion Statement	RCTs	References	
2	Stretching programs may produce greater improvements in spasticity than conventional therapy.	1	You et al. 2014	

RANGE OF MOTION				
LoE	Conclusion Statement	RCTs	References	
1b	Stretching programs may produce greater improvements in range of motion than conventional therapy.	1	Tseng et al. 2007	

ACTIVITIES OF DAILY LIVING				
LoE	LoE Conclusion Statement RCTs References			
1b	Stretching programs may produce greater improvements in performance of activities of daily living than conventional therapy .	2	You et al. 2014; Tseng et al. 2007	

Key points

Stretching programs may improve some aspects of upper limb function following stroke.

Orthotics



Adopted from: https://www.amazon.com/Soft-Resting-Hand-Splint-Left/dp/B007G4TVIK

Upper limb orthotic devices such as splints or kinesthetic tape are generally used to minimize or prevent contractures, reduce spasticity and pain, and prevent edema poststroke (Lannin & Herbert, 2003). Arm weighted support rehabilitation through orthic devices can facilitate recovery of hand movements through performing semiautonomous rehabilitation programs (Bartolo et al. 2014).

14 RCTs were found that used orthotic devices for upper extremity motor rehabilitation (Choi et al. 2016a; Choi et al. 2016b; Lannin et al. 2016; Kim et al. 2015; Bartolo et al. 2014; Page et al. 2013; Barry et al. 2012; Basaran et al. 2012; Jung et al. 2011; Lannin et al. 2007; Lannin et al. 2003; Langlois et al. 1991; Poole et al. 1990; Rose et al. 1987), the methodological details and results of these RCTs are presented in Table 11.

Authors (Year)	Interventions	Outcome Measures
Sample Sizestart	week for total number of weeks	Result (direction of enect)
Sample Size _{end} Time post stroke category		
<u>Choi et al.</u> (2016a) RCT (5)	E: Hand Splints and a General Rehabilitation Program	Modified Ashworth Scale (-)
N _{Start} =30 N _{End} =30	C: General Rehabilitation Program Duration: 30min/d, 5d/wk for 12wk	
<u>Choi et al.</u> (2016b) RCT (4)	E: Dorsal Resting Hand Splint C: Volar Resting Hand Splint	 Modified Ashworth Scale (+exp) Active Range of Motion (+exp)
N _{start} =52 N _{End} =52 TPS-Chronic	Duration: 30min/d, 5dwk for 8wk	
Lannin et al. (2016) RCT (5)	E: Task-specific training + training with the Saebo-Flex device	Motor Assessment Scale (-) Box and Block Test (-)
N _{Start} =9 N _{End} =6 TPS=Acute	C: Task-specific training Duration: 45-60min/session, 1-3sessions/d, 5- 7d/wk for 4-12wk	Grip Strength (-)
Kim et al. (2015) RCT (7) Nstart=30 NEnd=30 TPS=Subacute	E: Taping C: No taping Duration: 30min/d, 3d/wk for 28wk	 Manual Function Test (+) Modified Motor Assessment Scale (+exp)
Bartolo et al. (2014)	E: Arm orthosis	Arm abduction (+exp) Arm adduction (+exp)
NStart=28	Duration: 30min/d, 6d/wk for 2wk	 Arm flexion (+exp) Arm extension (+exp)
TPS=Acute		 Normalized jerk (+exp) Fugl Meyer Assessment (-) Modified Ashworth Scale (-)
Page et al. (2013) RCT (6) Nstart=16 Nscat=16	E: Myomo brace C: Repetitive task practice Duration: 30min/d, 3d/wk for 8wk	 Fugl Meyer Assessment (-) Canadian Occupational Performance Measure (-) Stroke Impact Scale (-)
TPS=Chronic		
Barry et al. (2012) RCT (7)	E: Dynamic hand orthosis C: Manual assisted therapy	 Grip strength (-) Action Research Arm Test (-)
Nend=19 TPS=Subacute		Stroke Impact Scale (-)
Basaran et al.(2012) RCT (6)	E1: Volar splint E2: Dorsal splint	E1 vs E2 vs C Modified Ashworth Scale (-)
N _{start} =39	C: No splint Duration: up to 10h/d for 5wk	Passive range of motion (-)
TPS=Chronic		
<u>Jung et al.</u> (2011) RCT (4)	E: Hand stretching/splint device C: No splint	Modified Ashworth Scale (+exp)
N _{Start} =21	Duration: 40min/d, 6d/wk for 3wk	
TPS=Chronic		
Lannin et al. (2007)	E1: Extension splint	Wrist contracture (-)
RCT (7) Nator=63	E2: Neutral splint	
N _{end} =63	Duration: 9-12h/d for 4wk	
TPS=Acute		
Lannin et al. (2003)	E: Hand splint	Wrist flexor (-)

Table 11. RCTs evaluating orthotic devices for upper extremity motor rehabilitation

RCT (8) N _{start} =28 N _{finish} =27 TPS=Subacute	C: No hand splint Duration: up to 12h/d, 5d/wk for 4wk	Finger flexor (-)
Langlois et al. (1991) RCT (3) N _{start} =9 N _{end} =9 TPS=Chronic	E1: Spint 22hr/d E2: Splint 12hr/d E3: Splint 6hr/d Duration: 6, 12, or 22h/d for 4wk	Spasticity (-)
Poole et al. (1990) RCT (5) N _{start} =18 N _{end} =18 TPS=Acute	E: Splint C: No splint Duration: 30min/d, 5d/wk for 3wk	Fugl Meyer Assessment (-)
Rose et al. (1987) RCT (4) N=30	E1: Dorsal orthosis E2: Volar orthosis C: No orthosis Duration: 2h	E1/E2 vs C Passive range of motion (+exp) E1 vs C Spontaneous flexion (+exp) E2 vs C Spontaneous flexion (-)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about orthotic devices

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	Orthotic devices may not have a difference in efficacy when compared to conventional therapy, repetitive task practice, or no orthotic device for improving motor function.	5	Kim et al. 2015; Bartolo et al. 2014; Page et al. 2013; Barry et al. 2012; Poole et al. 1990	

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
1b	Orthotic devices may not have a difference in efficacy when compared to conventional therapy, repetitive task practice, or no orthotic device for improving dexterity.	2	Lannin et al. 2016; Barry et al. 2012	

SPASTICITY				
LoE	Conclusion Statement	RCTs	References	
1b	Orthotic devices may not have a difference in efficacy when compared to conventional therapy, repetitive task practice, or no orthotic device for improving spasticity.	7	Choi et al. 2016a; Choi et al. 2016b; Bartolo et al. 2014; Basran et al. 2012; Jung et al. 2011; Lannin et al. 2007; Langlois et al. 1991	

RANGE OF MOTION				
LoE	Conclusion Statement	RCTs	References	
1a	Orthotic devices may produce greater improvements in range of motion than conventional therapy, repetitive task practice, or no orthotic device.	5	Choi et al. 2016b; Bartolo et al. 2014; Basran et al. 2012; Lannin et al. 2003; Rose et al. 198	

ACTIVITIES OF DAILY LIVING			
LoE	Conclusion Statement	RCTs	References
1a	Orthotic devices may not have a difference in efficacy when compared to conventional therapy, repetitive task practice, or no orthotic device for improving performance of activities of daily living.	4	Lannin et al. 2016; Kim et al. 2015; Page et al. 2013; Barry et al. 2012

MUSCLE STRENGTH				
LoE	Conclusion Statement	RCTs	References	
1b	Orthotic devices may not have a difference in efficacy when compared to conventional therapy, repetitive task practice, or no orthotic device for improving muscle strength.	2	Lannin et al. 2016; Barry et al. 2012	

Key points

Orthotics may not be beneficial for upper limb rehabilitation following stroke.

Mirror Therapy



Adopted from: https://www.saebo.com/shop/saebo-mirror-box/

In mirror therapy, a mirror is placed beside the unaffected limb, blocking view of the affected limb and creating an illusion of two limbs as if they are both functioning normally. Mirror therapy functions through a process known as mirror visual feedback wherein the movement of one limb is perceived as movement from the other limb (Deconinck et al. 2015). In the brain, mirror therapy is thought to induce neuroplastic changes that promote recovery by increasing excitability of the ipsilateral motor cortex which projects to the paretic limb (Deconinck et al. 2015). Ramachandran et al. (1995) first used this method to understand the effect of vision on phantom sensation and pain in arm amputees. This method has since been adapted from its original use as a means to enhance upper-limb function following stroke (Sathian et al. 2000).

A total of 25 RCTs were found that evaluated mirror therapy for upper extremity rehabilitation poststroke. Of these 18 RCTs looked at mirror therapy compared to conventional rehabilitation or the Bobath concept approach (Radajewska et al. 2017; Colomer et al. 2016; Gurbuz et al. 2016; Kim et al. 2016; Lim et al. 2016; Pervane Vural et al. 2016; Arya et al. 2015; Cristina et al. 2015; Park et al. 2015; Invernizzi et al. 2013; Radajewska et al. 2013; Timmerman et al. 2013; Wu et al. 2013a; Lee et al. 2012; Michielsen et al. 2011; Dohle et al. 2009; Yavuzer et al. 2008; Altschuler et al. 1999). Two RCTs looked at mirror therapy with bilateral arm training (Rodrigues et al. 2016; Samuelkamaleshkumar et al. 2014), mirror therapy combined with: transcranial direct current stimulation (Cho et al. 2015), functional electrical stimulation (Kim et al. 2015), neuromuscular electrical stimulation (Yun et al. 2011), rTMS (Ji et al. 2014), and in a group or individual setting (Thieme et al. 2012).

The methodological details and results of these 25 RCTs are presented in Table 12.

Table 12. Summary of RCTs evaluating mirror therapy for the upper extremity motor rehabilitation

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
M	irror therapy compared to conventiona	al rehabilitation
Radajewska et al. (2017) RCT (5) N _{Start} =60 N _{End} =60 TPS=Subacute	E: Mirror therapy C: Conventional rehabilitation Duration: 30min/d, 5d/wk for 3wk	 Frenchay Arm Test (+exp)
Colomer et al. (2016) RCT (7) N _{Start} =34 N _{End} =31 TPS=Chronic	E: Mirror Therapy C: Passive Mobilization Duration: 45min/d, 3d/wk for 8wk	 Nottingham Sensory Assessment (+exp) Fugl-Meyer Assessment (-)
Gurbuz et al. (2016) RCT (6) N _{Start} =31 N _{End} =31 TPS=Subacute	E: Mirror Therapy C: Conventional Therapy Duration: 60-120min/d, 5d/wk for 4wk	 Brunnstrom Recovery Stage (-) Fugl-Meyer Assessment (+exp) Function Independence Measure (-)
<u>Kim et al.</u> (2016) RCT (5) N _{Start} =25 N _{End} =25 TPS=Chronic	E: Mirror Therapy C: Conventional Therapy Duration: 30min/d, 5dwk for 4wk	 Action Research Arm Test (+exp) Fugl-Meyer Assessment (+exp) Box and Block Test (+exp) Functional Independence Measure (+exp)
Lim et al. (2016) RCT (5) N _{Start} =60 N _{End} =60 TPS=?	E: Mirror Therapy C: Sham Therapy Duration: 20min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Modified Barthel Index (+exp) Brunnstrom Recovery Stage (-)
Pervane Vural et al. (2016) RCT (6) N _{Start} =30 N _{End} =30 TPS=Subacute	E: Mirror Therapy C: Conventional rehabilitation Duration: 4h/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Visual Analog Scale (+exp) Brunnstrom Recovery Stage (+exp) Functional Independence Measure (+exp) Modified Ashworth Scale (+exp)
Arya et al. (2015) RCT (8) Nstart=33 NEnd=32 TPS=Chronic	E: Task-based mirror therapy C: Standard Rehabilitation Duration: 90min/d, 5d/wk for 8wk	 Fugl-Meyer Assessment (+exp)
Cristina et al. (2015) RCT (6) N _{Start} =15 N _{End} =15 TPS=Subacute	E: Mirror therapy C: Conventional therapy Duration: 30min/d, 5d/wk for 6wk	 Modified Ashworth Scale: writ (+exp) Bhakta finger flexion scale (+exp)
Park et al. (2015) RCT (6) Nstart=30 NEnd=30 TPS=Chronic	E: Mirror therapy C: Non-reflecting mirror Duration: 5d/wk for 6wk	 Manual Function Test (+exp) FIM (+exp)
Invernizzi et al. (2013) RCT (7) N _{Start} =26 N _{End} =25 TPS=Acute	E: Mirror therapy C: Conventional therapy Duration: 30-60min/d, 5d/wk for 4wk	 Action Research Arm Test (+exp) Motricity Index (+exp) Fugl-Meyer Assessments (+exp)
Radajewska et al. (2013)	E: Mirror therapy	Frenchay Arm Test (+exp)

RCT (3) Nstart=60 N _{End} =60 TPS=2	C: Conventional therapy Duration: 30min/d, 5d/wk for 3wk			
Timmerman et al. (2013) RCT (7) N _{Start} =42 N _{End} =42 TPS=Subacute	E: Mirror therapy C: Bobath concept Duration: 30min/d, 3d/wk for 6wk	 Frenchay Arm Test (-) Functional Assessment Scale (-) Wolf Motor Function Test (-) 		
<u>Wu et al.</u> (2013a) RCT (6) N _{Start} =33 N _{End} =21 TPS=Chronic	E: Mirror therapy C: Conventional therapy Duration: 1.5h/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Modified Ashworth Scale (-) ABILHAND (-) 		
Lee et al. (2012) RCT (5) N _{start} =28 N _{end} =26 TPS=Subacute	E: Mirror therapy C: Standard care Duration: 50min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Brunnstrom recovery stages (+exp) Manual Function Test (+exp) 		
Michielsen et al. (2011) RCT (7) N _{Start} =40 N _{End} =40 TPS=Chronic	E: Mirror therapy C: Control therapy Duration: 1h/d, 5d/wk for 6wk	 Action Research Arm Test (-) ABILHAND (-) Grip force (-) Tardieu Scale (-) Fugl-Meyer Assessment (+exp) 		
Dohle et al. (2009) RCT (7) Nstart=36 NEnd=36 TPS=Acute	E: Mirror therapy C: Control therapy Duration: 30min/d, 5d/wk for 6wk	 Fugl-Meyer Assessment (-) 		
<u>Yavuzer et al. (</u> 2008) RCT (7) N _{Start} =40 N _{End} =40 TPS=Subacute	E: Mirror Therapy C: Sham Therapy Duration: 2-5h/d, 5d/wk for 4wk	 Brunnstrom Recovery Stages (+exp) Funtional Indepence Measure (+exp) Modified Ashworth Scale (-) 		
Altschuler et al. (1999) RCT (7) N _{Start} =40 N _{End} =40 TPS=Chronic	E: Mirror therapy C: Sham therapy Duration: 30min/d, 6d/wk for 4wk	 Brunnstrom Recovery Stage (+exp) Fugl Meyer self-care Score (+exp) Modified Ashworth Scale (-) 		
	Mirror therapy combined with bilatera	I arm training		
Rodrigues et al. (2016) RCT (7) N _{start} =16 N _{End} =16 TPS=Chronic	E: Mirror therapy and Bilateral Training C: Bilateral Training Duration: 1h/d, 3d/wk for 4wk	Upper extremity function test (-)		
Samuelkamaleshkumar et al. (2014) RCT (7) N _{Start} =20 N _{End} =20 TPS=Subacute	E: Mirror therapy + bilateral arm training C: Control group Duration: 6h/d, 5d/wk for 3wk	 Fugl-Meyer Assessment (+exp) Brunnstrom Recovery Stage (+exp) Box and Block Test (+exp) Modified Ashworth Scale (-) 		
	Mirror therapy combined with	tDCS		
<u>Cho et al.</u> (2015) RCT (5) N _{Start} =30 N _{End} =27 TPS=Chronic	E: Mirror therapy + tDCS C: Sham mirror therapy + tDCS Duration: 45min/d, 3d/wk for 6wk	 Box and Block Test (+exp) Grip strength (+exp) Jebsen Taylor Hand Function (-) Fugl-Meyer Assessment (-) 		
Mirror therapy combined with functional electrical stimulation				
Kim et al. (2015)	E: FES + mirror therapy	Box and Block Test (-)		

RCT (6)	C: FES + sham mirror therapy	Fugl-Meyer Assessment (+exp)
Nstart=28	Duration: 30min/d, 5d/wk for 4wk	Brunnstrom Recovery Stage (-)
N _{End} =23		Manual Function Test (+exp)
TPS=Chronic		
Mirror t	herapy combined with neuromuscular	electrical stimulation
Yun et al. (2011)	E1: Cyclic NMES + mirror therapy	E1 vs. E2/E3
RCT (4)	E2: Cvclic NMES	 Fugl-Mever Assessment (+exp)
N=60	E3: Mirror therapy	Hand flexion (-)
TPS=Acute	Duration: 30min/d. 5d/wk for 3wk	Wrist flexion (-)
		• Wrist extension (-)
		•
	Mirror therapy combined with	rTMS
Ji et al. (2014)	E1: Mirror therapy + rTMS	E1 vs. E2
RCT (7)	E2: Mirror therapy	 Fugl-Meyer Assessment (+exp)
Nstart=35	C: Sham therapy	Box and Block Test (+exp)
N _{End} =35	Duration: 30min/d. 5d/wk for 4wk	E2 vs. C
TPS=Chronic		 Fugl-Mever Assessment (+exp₂)
		 Box and Block Test (+exp₂)
	Group vs individual mirror the	erapy
Thisms at al. (2012)	E4. Individual minutes the second	Action Desservels Arm Test ()
$\frac{1}{1}$	E1: Individual mirror therapy	Action Research Arm Test (-)
RCT (6)	E2: Group mirror therapy	Fugi-Meyer Assessment (-)
NStart=0U	C: Snam mirror therapy	Bartnei Index (-)
NEnd=49	Duration: 30min/d, 4dwk for 5wk	• Stroke Impact Scale (-)
IPS=Subacute		<u>E1 vs. E2</u>
		 Modified Ashworth Scale (+exp)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about mirror therapy

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	Mirror therapy may produce greater improvements in motor function than conventional therapy or Bobath concept approaches .	15	Colomer et al. 2016; Gurbuz et al. 2016; Kim et al. 2016; Lim et al. 2016; Pervane Vural et al. 2016; Arya et al. 2015; Park et al. 2015; Ji et al. 2014; Invernizzi et al. 2013; Wu et al. 2013a; Lee et al. 2012; Michielsen et al. 2011; Dohle et al. 2009; Altschuler et al. 1999
1a	There is conflicting evidence about the effect of mirror therapy combined with bilateral arm training to improve motor function when compared to bilateral arm training or conventional therapy .	2	Rodrigues et al. 2016; Samuelkamaleshkumar et al. 2014
2	Mirror therapy combined with tDCS may not have a difference in efficacy compared to sham mirror therapy combined with tDCS for improving motor function.	1	Cho et al. 2015
1b	Mirror therapy combined with high frequency rTMS may produce greater improvements in motor function than mirror therapy on its own or sham stimulation.	1	Ji et al. 2014
1b	Mirror therapy combined with FES may produce greater improvements in motor function than sham mirror therapy with FES.	1	Kim et al. 2015
2	Mirror therapy combined with cyclic NMES may produce greater improvements in motor function than mirror therapy or cyclic NMES on their own.	1	Yun et al. 2011
1b	Mirror therapy provided in a group setting may not have a difference in efficacy when compared to mirror therapy in a one on one setting to improve motor function.	1	Thieme et al. 2012

DEXTERITY			
LoE	Conclusion Statement	RCTs	References
1b	Mirror therapy may produce greater improvements in dexterity than conventional therapy or Bobath concept approaches .	2	Kim et al. 2016; Ji et al. 2014
1b	Mirror therapy combined with bilateral arm training may produce greater improvements in dexterity than bilateral arm training or conventional therapy.	1	Samuelkamaleshkumar et al. 2014
2	Mirror therapy combined with tDCS may produce greater improvements in dexterity than sham mirror therapy combined with tDCS.	1	Cho et al. 2015
1b	Mirror therapy combined with FES may not have a difference in efficacy compared to sham mirror therapy with FES for improving dexterity.	1	Kim et al. 2015

SPASTICITY			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of mirror therapy to improve spasticity when compared to conventional therapy or Bobath concept approaches .	6	Pervane Vural et al. 2016; Cristina et al. 2015; Wu et al. 2013a; Michielsen et al. 2011; Yavuzer et al. 2008; Altschuler et al. 1999
1b	Mirror therapy combined with bilateral arm training may produce greater improvements in spasticity than bilateral arm training or conventional therapy.	1	Samuelkamaleshkumar et al. 2014
1b	Mirror therapy provided in a group setting may produce greater improvements in spasticity than mirror therapy administered in a one on one setting.	1	Thieme et al. 2012

RANGE OF MOTION			
LoE	Conclusion Statement	RCTs	References
2	Mirror therapy combined with cyclic NMES may not have a difference in efficacy when compared to cyclic NMES or mirror therapy on their own for improving range of motion.	1	Yun et al. 2011

PROPRIOCEPTION			
LoE	Conclusion Statement	RCTs	References
1b	Mirror therapy may produce greater improvements in proprioception than conventional therapy or Bobath concept approaches .	1	Colomer et al. 2016

ACTIVITIES OF DAILY LIVING			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of mirror therapy to improve performance of activities of daily living when compared to conventional therapy or Bobath concept approaches .	11	Radajewska et al. 2017; Gurbuz et al. 2016; Kim et al. 2016; Lim et al. 2016; Pervane Vural et al. 2016; Park et al. 2015; Radajewska et al. 2013; Timmerman et al. 2013; Wu et al. 2013; Muchielsen et al. 2011; Yavuzer et al. 2008
1b	Mirror therapy in a group setting may not have a difference in efficacy compared to mirror therapy in a one on one setting to improve performance of activities of daily living.	1	Thieme et al. 2012

MUSCLE STRENGTH			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of mirror therapy to improve muscle strength when compared to conventional therapy or Bobath concept approaches .	2	Invernizzi et al. 2013; Michielsen et al. 2011
2	Mirror therapy combined with tDCS may produce greater improvements in muscle strength than sham mirror therapy combined with tDCS.	1	Cho et al. 2015

Key points

Mirror therapy on its own or in combination with other interventions can improve many aspects of upper limb function following stroke.

Mental practice



Adopted from: https://www.ucbmsh.com/motor-imagery-for-improvement-of-gait-in-stroke-patient/

Mental practice as the name suggests, involves cognitively rehearsing a specific task by repetitively imagining oneself performing the precise movements involved in the task in the absence of performing the physical movement (Page et al. 2014). Mental practice is speculated to be effective because of its ability to use the same motor schema as when physically practicing the same task through the activation of similar neural regions and networks during mental practice (Page et al. 2014). The use of mental practice was adapted from the field of sports psychology where the technique has been shown to improve athletic performance, when used as an adjunct to standard training methods (Page et al. 2014). The technique is believed to be advantageous in stroke survivors because certain motor skills may be difficult to physically practice; stroke survivors spend a majority of their time inactive and alone; and repetitive task-specific practice is a prerequisite for cortical plasticity and subsequent motor changes (Page et al. 2014). Mental practice can be used to supplement conventional therapy and can be used at any stage of recovery.

17 RCTs evaluated mental practice compared to conventional rehabilitation or a sham intervention for upper extremity motor rehabilitation (Oh et al. 2016; Park et al. 2015b; Liu et al. 2014; Mihara et al. 2013; Oostra et al. 2013; Lee et al. 2012; Letswaart et al. 2011; Page et al. 2011; Bovend'Eerdt et al. 2010; Riccio et al. 2010; Liu et al. 2009b; Muller et al. 2007; Page et al. 2007; Page et al. 2005; Liu et al. 2004; Page et al. 2001; Page et al. 2000). Two RCTs combined mental practice with modified constraint induced movement therapy (mCIMT) compared to mCIMT on its own (Park et al. 2015a; Page et al. 2009). Another RCT combined mental practice with Nintendo Wii virtual reality interactive game training compared to Nintendo Wii training on its own (Park et al. 2016).

The methodological details and results of all 20 RCTs evaluating mental practice interventions for upper extremity motor rehabilitation are presented in Table 13.

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Oh et al. (2016) RCT Crossover (7) N _{Start} =10 N _{End} =10 TPS=Chronic	E: Mental Practice C: Conventional Therapy Duration: 20min/d, 3d/wk for 3wk	 Fugl-Meyer Assessment (-) Motor Activity Log (-)
Park et al. (2015b) RCT (6) N _{Start} =29 N _{End} =29 TPS=Chronic	E: Mental practice C: Physical therapy Duration: 10min/d, 5d/wk for 2wk	 Fugl-Meyer Assessment (+exp) Action Research Arm Test (+exp) Modified Barthel Index (+exp)
Liu et al. (2014) RCT (7) N _{Start} =20 N _{End} =20 TPS=Subacute	E: Motor imagery + mental practice of affected hand C: Motor imagery + mental practice of unaffected hand Duration: 45min/d, 5d/wk for 4wk	Action Research Arm test (+exp)
Mihara et al. (2013) RCT (9) N _{Start} =20 N _{End} =20 TPS=Chronic	E: Mental practice C: Sham intervention Duration: 20min/d, 3d/wk for 2wk	 Fugl-Meyer Assessment (+exp) Action Research Arm test (-)
Oostra et al. (2013) RCT (8) N _{Start} =20 N _{End} =20 TPS=Chronic	E: Mental practice C: Physical training Duration: 30min/d, 5d/wk for 6wk	Action Research Arm Test (+exp)
Ietswaart et al. (2011) RCT (7) Nstart=121 Nend=101 TPS=Subacute	E1: Motor imagery E2: Attention placebo C: Usual care Duration: 45min/d, 3d/wk for 4wk	Action Research Arm Test (-)
Page et al. (2011) RCT (6) N _{start} =32 N _{end=} 29 TPS=Subacute	E: Audiotaped mental practice C: Audiotaped sham intervention Duration: 30min/d, 3d/wk for 10wk	 Fugl-Meyer Assessment (-) Action Research Arm Test (-)
Bovend'Eerdt et al. (2010) RCT (8) N _{start} =50 N _{end} =48 TPS=Chronic	E: Mental practice C: Conventional therapy Duration: 30min/d, 2-3d/wk for 5wk	 Barthel Index (-) Nottingham Extended ADL (-) Action Research Arm Test (-)
Riccio et al. (2010) RCT Crossover (5) N _{start} =36 N _{end} =36 TPS=Chronic	E: Mental practice C: Conventional rehabilitation Duration: 1h/d, 5d/wk for 3wk	 Motricity Index (+exp) Arm Function Test (+exp)
Liu et al. (2009b) RCT (5) N _{start} =35 N _{end} =35 TPS=Subacute	E: Mental Imagery C: Conventional Functional Rehabilitation Duration: 1h, 5d/wk for 3wk	Improvement in Trained Tasks (+exp)
Müller et al. (2007)	E1: Mental practice	<u>E1/E2 vs. C</u>
RCT (4)	E2: Motor practice	• Jebsen Hand Function Test: (+exp ₁ , +exp ₂)

Table 13. RCTs evaluating mental practice interventions for upper extremity motor rehabilitation

N _{start} =17 N _{end} =17 TPS=Acute	C: Conventional therapy Duration: 30min/d, 5d/wk for 4wk	•	Pinch grip: (+exp1, +exp2)	
Page et al. (2007) RCT (6) Nstart=32 Nend=32 TPS=Chronic	E: Mental Practice C: Sham Relaxation Exercise Intervention Duration: 30min/d, 2d/wk for 6wk	•	Fugl-Meyer Assessment (+exp) Action Research Arm Test (+exp)	
Page et al. (2005a) RCT (6) N _{start} =11 N _{end} =8 TPS=Chronic	E: Mental practice C: Relaxation techniques Duration: 30min/d, 2d/wk for 6wk	•	Action Research Arm Test (+exp) Motor Activity Log: Amount of Use (+exp), Quality of Movement (+exp)	
Liu et al. (2004) RCT (4) N _{start} =49 N _{end} =46 TPS=Acute	E: Mental Imagery C: Functional training Duration: 1h/d, 5d/wk for 3wk	•	Fugl-Meyer Assessment (-)	
Page et al. (2001) RCT (5) N _{start} =13 N _{end} =13 TPS=Subacute	E: Imagery training C: Occupational therapy Duration: 10min/d, 4d/wk for 6wk	•	Fugl-Meyer Assessment (+exp) Action Research Arm Test (+exp)	
Page et al. (2000) RCT (4) N _{start} =16 N _{end} =13 TPS=Chronic	E: Imagery training C: Occupational therapy Duration: 30min/d, 3d/wk for 4wk	•	Fugl-Meyer Assessment (+exp)	
	Mental practice combined with I	mC	IMT	
Park et al. (2015a) RCT (7) N _{Start} =26 N _{End} =26 TPS=Chronic	E: Mental practice + mCIMT C: mCIMT Duration: 30min/d, 5d/wk for 6wk	•	Fugl-Meyer Assessment (+exp) Action Research Arm Test (+exp) Modified Barthel Index (+exp)	
Page et al. (2009) RCT (4) N _{start} =10 N _{end} =10 TPS=Chronic	E: Mental practice + Modified Constraint Induced Movement Therapy C: Modified Constraint Induced Movement Therapy Duration: 30min/d, 5d/wk for 10wk	•	Action Research Arm Test (+exp) Fugl-Meyer Assessment (+exp)	
Nintendo Wii combined with mental practice				
Park et al. (2016) RCT (7) N _{Start} =30 N _{End} =30 TPS=Chronic	E: Nintendo Wii + mental practice C: Nintendo Wii Duration: 5min/d, 5d/wk for 4wk	•	Fugl-Meyer Assessment (-) Motor Activity Log (-)	

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about mental practice

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	Mental practice may produce greater improvements in motor function than conventional rehabilitation or a sham intervention .	15	Oh et al. 2016; Park et al. 2015b; Liu et al. 2014; Mihara et al. 2013; Oostra et al. 2013; Lee et al. 2012; Page et al. 2011; Bovend'Eerdt et al. 2010; Riccio et al. 2010; Muller et al. 2007; Page et al. 2007; Page et al. 2005; Liu et al. 2004; Page et al. 2001; Page et al. 2000
1b	Mental practice combined with mCIMT may produce greater improvements in motor function than mCIMT on its own.	2	Park et al. 2015a; Page et al. 2009
1b	Mental practice combined with Nintendo Wii training may not have a difference in efficacy compared to Nintendo Wii training on its own for improving motor function.	1	Park et al. 2016

ACTIVITIES OF DAILY LIVING				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of mental practice to improve performance of activities of daily living when compared to conventional rehabilitation or a sham intervention .	6	Oh et al. 2016; Park et al. 2015b; Rajeesh et al. 2015; Bovend'Eerdt et al. 2010; Liu et al. 2009b; Page et al. 2005	
1b	Mental practice combined with mCIMT may produce greater improvements in performance of activities of daily living than mCIMT on its own .	1	Park et al. 2015a	
1b	Mental practice combined with Nintendo Wii training may not have a difference in efficacy compared to Nintendo Wii training on its own for improving performance of activities of daily living.	1	Park et al. 2016	

MUSCLE STRENGTH					
LoE	LoE Conclusion Statement RCTs References				
2	Mental practice may produce greater improvements in muscle strength than conventional rehabilitation or a sham intervention.	2	Riccio et al. 2010; Muller et al. 2007		

Key points

Mental practice, alone or in combination with constraint-induced movement therapy, may be beneficial for upper limb rehabilitation following stroke.

Mental practice in combination with virtual reality training may not be beneficial for upper limb function.

Action observation



Adopted from: <u>https://www.youtube.com/watch?v=QE3CUhmKi7U</u>

Action observation is a form of therapy whereby an individual observes another individual performing a motor task, either on a video or a real demonstration, and then may attempt to perform the same task themselves. For example, the patient may be instructed to watch a video showing an adult stretching out his hand to pick up a cup, bringing the cup to his mouth, and then returning the cup to its initial position - the act of drinking. After observing the video sequence for a time, the participants may or may not be asked to perform the same action (Borges et al. 2018).

The therapy is considered a multisensory approach designed to increase cortical excitability in the primary motor cortex by activating central representations of actions through the mirror neuron system (Kim and Kim, 2015). Although action observation has been evaluated mainly in healthy volunteers, a few studies have evaluated its benefit in motor relearning following stroke.

Seven RCTs were found that evaluated action observation techniques compared to conventional rehabilitation or sham action observation for upper extremity motor rehabilitation (Kuk et al. 2016; Kim and Kim, 2015; Zhu et al. 2015; Sale et al. 2014; Cowles et al. 2013; Franceschini et al. 2012; Ertelt et al. 2007); their methodological details and results are presented in Table *x*.

Table 14. RCTs evaluating action observation interventions for upper extremity motor rehabilitation

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)	
$\frac{Fu \text{ et al. 2017}}{RCT (5)}$ N _{start} =70 N _{End} =53 TPS=Subacute	E: Video clip of 30 actions relating to sjhoulder, elbow, wrist, forearm and hand movements. C: Conventional therapy Duration: 20min, 6x/wk for 8 wk	 Fugl-Meyer Assessment (-) Wolf motor function test (-) Modified Barthel Index (-) 	
<u>Kuk et al.</u> (2016) RCT (5) N _{Start} =22 N _{End} =20	E: Video clip of a motor task followed by execution of the same motor task C: Pictures of landscapes followed by execution of the motor task Duration: 1min/d for 5d	Box and Block Test (+exp)	
$\label{eq:constant} \begin{array}{c} \frac{Kim \; and \; Kim}{RCT \; (6)} & (2015) \\ N_{Start} = 12 \\ N_{End} = 12 \end{array}$	E: Action observation + occupational therapy C: Placebo observation + occupational therapy Duration: 30min/d, 5d/wk for 6wk	Wolf Motor Function Test (-)	
Zhu et al. (2015) RCT (5) N _{Start} =70 N _{End} =61	E: Upper Limb Action Observation Therapy C: Conventional Rehabilitation Therapy Duration: 30min/d, 6d/wk for 8wk	 Fugl-Meyer Assessment (+exp) Barthel Index (+exp) Modified Ashworth Scale (+exp) 	
<u>Sale et al.</u> (2014) RCT (7) N _{Start} =67 N _{End} =67	E: Action observation C: Standard rehabilitation Duration: 3min/d, 5d/wk for 4wk	 Box and Block Test (+exp) Fugl Meyer Assessment (+exp) 	
Cowles et al. (2013) RCT (7) N=29	E: Action observation C: Conventional therapy Duration: 1h/d, 5d/wk for 3wk	 Motricity Index (-) Action Research Arm Test (+con) 	
Franceschini et al. (2012) RCT (8) N=102	E: Video footage C: Static images Duration: 15min/d, 5d/wk for 4wk	 Box and Block Test (+exp) Frenchay Arm Test (-) Modified Ashworth Scale (-) FIM (-) 	
Ertelt et al. (2007) RCT (5) N=15	E: Action observation therapy C: Traditional therapy Duration: 12min/d, 5d/wk for 18d	 Frenchay Arm Test (+exp) Wolf Motor Function Test (+exp) Stroke Impact Scale (+exp) 	
Action observation compared to task-oriented training			
Kim and Bang, 2016 RCT (5) N _{Start} =22 N _{End} =22 TPS=Subacute	E: Action observation C: Task-oriented training Duration: 40min, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Box and block test (+exp) Modified Barthel Index (+exp) Modified Ashworth Scale (-) 	

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α=0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group - indicates no statistically significant between groups differences at α =0.05

Conclusions about action observation

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of action observation interventions to improve motor function when compared to conventional rehabilitation or sham action observation .	6	Fu et al. 2017; Kim and Kim, 2015; Zhu et al. 2015; Sale et al. 2014; Cowles et al. 2013; Ertelt et al. 2007	
2	Action observation may produce greater improvements in motor function than task-oriented training.	1	Kim and Bang, 2016	

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
1a	Action observation may produce greater improvements in dexterity than sham stimulation or conventional therapy.	3	Kuk et al. 2016; Sale et al. 2014; Franceschini et al. 2012	
2	Action observation may produce greater improvements in dexterity than task-oriented training.	1	Kim and Bang, 2016	

ACTIVITIES OF DAILY LIVING					
LoE	LoE Conclusion Statement RCTs References				
1b	There is conflicting evidence about the effect of action observation interventions to improve activities of daily living when compared to sham stimulation or conventional therapy .	4	Fu et al. 2017; Zhu et al. 2015; Franceschini et al. 2012; Ertelt et al. 2007		
2	Action observation may produce greater improvements in activities of daily living than task-oriented training.	1	Kim and Bang, 2016		

MUSCLE STRENGTH				
LoE	References			
	Action observation may not have a difference in	1	Cowles et al. 2017	
1b	efficacy when compared to sham stimulation or	1		
	conventional therapy for improving muscle strength.			

SPASTICITY				
LoE	Conclusion Statement	RCTs	References	
2	Action observation may produce greater improvements in spasticity than sham stimulation or conventional therapy.	1	Zhu et al. 2015	
2	Action observation may not have a difference in efficacy when compared to task-oriented training for improving spasticity.	1	Kim and Bang, 2016	

Key points

Action observation may be beneficial for some aspects of upper limb function following stroke.

Music therapy



Adopted from: https://steinhardt.nyu.edu/site/ataglance/2017/03/music-therapy-helps-with-recovery-post-stroke.htm

Music therapy is defined as listening, singing, and creating music with/without rhythm and percussion instruments, and is based on four rehabilitation principles: extended repetition of simple finger and arm movements, auditory-motor coupling to reinforce motor learning due to instant auditory feedback, individualized training, and emotional/motivational support due to the emotions invoked by music and the acquisition of a new skill (Zhang et al. 2016). As such it involves many components of conventional upper limb rehabilitation interventions including repetitive task practice, finger individualization, as well as tactile and auditory feedback (van Wijck et al. 2012). The rehabilitation program can also be shaped by increasing the tempo of the songs or incorporating more difficult music pieces based on individual performance (Jun et al. 2013).

Six RCTs (Scholz et al. 2016; Tong et al. 2015; Thielbar et al. 2014; Van Vugt et al. 2014; Jun et al. 2013; Altenmuller et al. 2009) evaluated the effects of music therapy compared to conventional therapy, task-oriented therapy and sham interventions on improving upper extremity motor rehabilitation.

The methodological details and results of all six RCTs are presented in Table 15.

Table 15. RCTs evaluating music therapy interventions for upper extremity motor rehabilitation

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Scholz et al. (2016) RCT (4) N _{Start} =25 N _{End} =25 TPS=Acute	E: Music Sonification Therapy C: Sham Movement Training Duration: 30min/d, 5d/wk for 2wk	 Fugl-Meyer Assessment (-) Action Research Arm Test (-) Nine Hold Peg Test (-) Stroke Impact Scale (-)
Tong et al. (2015) RCT (5) Nstart=33 NEnd=30 TPS=Chronic	E: Audible Music Instrumental Training C: Mute Music Instrumental Training Duration: 30min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (-) Wolf Motor Function Test (+exp)
Thielbar et al. (2014) RCT (6) N _{Start} =14 N _{End} =14 TPS=Chronic	E: Virtual keyboard music playing C: High intensity, task oriented occupational therapy Duration: 1hr/d, 3d/wk for 6wk	 Action Research Arm Test (-) Fugl Meyer Assessment (+exp) Jebsen Taylor Hand Function Test (+exp) Grip strength (-) Pinch strength (-)
Van Vugt et al. (2014) RCT (4) N _{Start} =36 N _{End} =28 TPS=Subacute	E: Playing piano together C: Playing piano sequentially Duration: 30min/d, 5d/wk for 2wk	Nine Hole Peg Test (-)
<u>Jun et al.</u> (2013) RCT (4) N _{Start} =40 N _{End} =30 TPS=Acute	E: Music movement therapy C: Routine intervention Duration: 1hr/d, 3d/wk for 8wk	 Shoulder and elbow flexion (+exp) Arm strength (-) Modified Barthel Index (-)
Altenmüller et al. (2009) RCT (5) Nstart=62 NEnd=62 TPS=Acute	E: MIDI piano and electronic drum training + conventional therapy C: Conventional therapy only Duration: 1hr/d, 5d/wk for 3wk	 Box and Block Test (+exp) Nine Hole Pegboard Test (+exp) Action Research Arm Test (+exp) Finger/Hand tapping (+exp)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group

 $+exp_2$ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at $\alpha{=}0.05$

Conclusions about music therapy

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1b	There is conflicting evidence about the effect of music therapy to improve motor function when compared to conventional therapy, task-oriented therapy and sham interventions .	4	Scholz et al. 2016; Tong et al. 2015; Thielbar et al. 2014; Altenmuller et al. 2009	

ACTIVITIES OF DAILY LIVING					
LoE	LoE Conclusion Statement RCTs References				
2	Music therapy may not have a difference in efficacy when compared to conventional therapy, task- oriented therapy and sham interventions for improving performance of activities of daily living.	1	Scholz et al. 2016, Jun et al. 2013		

MUSCLE STRENGTH				
LoE	Conclusion Statement	RCTs	References	
2	Music therapy may not have a difference in efficacy when compared to conventional therapy, task- oriented therapy and sham interventions for improving muscle strength.	2	Thielbar et al. 2014, Jun et al. 2013	

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
2	There is conflicting evidence about the effect of music therapy to improve dexterity when compared to conventional therapy, task-oriented therapy and sham interventions .	3	Scholz et al. 2016, Van Vugt et al. 2014, Altenmuller et al. 2009	

RANGE OF MOTION				
LoE	Conclusion Statement	RCTs	References	
2	Music therapy may produce greater improvements in range of motion than conventional therapy, task-	1	Jun et al. 2013	
2	Music therapy may produce greater improvements in range of motion than conventional therapy, task- oriented therapy and sham interventions.	1	Jun et al. 2013	

Key points

The literature is mixed regarding music therapy for upper limb rehabilitation following stroke.

Technology based interventions Telerehabilitation



Adopted from: http://www.telereadaptation.com/en/projet/telerehabilitation-in-speech-therapy/

Telerehabilitation is the process of providing rehabilitation services remotely through information and communication technologies (e.g. a kiosk, telephone and computer) (Dodakian et al. 2017; Emmerson et al. 2017). This rehabilitation method is particularly useful for patients who cannot access a rehabilitation center (Benvenuti et al. 2014). Additionally, this intervention can be delivered for a longer duration and at a reduced cost when compared to therapies provided in the inpatient rehabilitation setting (Benvenuti et al. 2014).

Only two RCTs looked at upper limb rehabilitation using telerehabilitation (Emerson et al. 2017; Wolg et al. 2015), though several RCT protocols and observational studies have been published. In one RCT the intervention group was a home exercise program delivered through a tablet (Emerson et al. 2017), while the other RCT delivered a home exercise program through a novel hand robot system (Wolf et al. 2015). Both RCTs were compared to home exercise programs on their own,

The methodological details and results of the two RCTs evaluating telerehabilitation for the upper extremity motor rehabilitation are presented in Table 16.

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Emmerson et al. (2017) RCT (7) N _{Start} =62 N _{End} =58 TPS=Chronic	E: Home exercise program using an electronic tablet with automated reminders C: Paper-based home exercise program Duration: 45min/d, 5d/wk for 4wk	 Wolf Motor Function Test (-) Grip Strength (-)
Wolf et al. (2015) RCT (7) Nstart=99 NEnd=92 TPS=Subacute	E: Telerehabilitation through an upper extremity hand robot with home exercise program C: Home exercise program only Duration: 3h/d, 5d/wk for 8-12wk	 Fugl Meyer Assessment (-) Action Research Arm Test (-) Wolf Motor Function Test (+exp)

Table 16. RCTs evaluating telerehabilitation for upper extremity motor rehabilitation

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group

 $+exp_2$ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at $\alpha\text{=}0.05$

Conclusions about telerehabilitation

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of telerehabilitation to improve motor function when compared to conventional therapy, task-oriented therapy and sham interventions .	2	Emmerson et al. 2017; Wolf et al. 2015	

MUSCLE STRENGTH				
LoE	Conclusion Statement	RCTs	References	
1b	Telerehabilitation may not have a difference in efficacy compared to home exercise programs for	1	Emmerson et al. 2017	
	improving muscle strength.			

Key points

The literature is mixed regarding telerehabilitation for upper limb rehabilitation following stroke.

Robotics



Adopted from: https://www.strokengine.ca/wp-content/uploads/2015/05/robotics_ARMin-300x226.jpg http://www.gentle.rdg.ac.uk/103-0325_IMG.JPG: https://cpmsales.net/wp-content/uploads/CENTURA.jpg: http://img.medicalexpo.com/images_me/photo-g/74722-10591286.jpg

Robotic devices can be used to help facilitate passive range of motion, to help maintain range and flexibility, to temporarily reduce hypertonia, and to provide resistance during passive movement. Assistance can also be provided during active movements when a patient cannot complete a movement independently. Robotics may be most appropriate for patients with dense hemiplegia, although robotics can be used with higher-level patients who wish to increase strength by providing resistance during the movement. According to Lum et al. (2002) robotic devices may be the most beneficial in severely impaired patients where unassisted movement is not possible, and especially during the acute phase of recovery during which spontaneous recovery occurs. Krebs et al. (2003) noted that robotic devices rely on the repetition of specific movements to improve functional outcomes.

Upper limb robotic devices can be classified based on the type of robot, the actuation method, the form of transmission, and the sensor used (Yue et al. 2017). The type of robot is based on the alignment of the device and the use and includes end-effectors and exoskeletons (Yue et al. 2017). End-effectors are external to the patient and are connected at a single distal point, whereas exoskeletons are worn by the patient and include mechanical joints that align to the human limb joints (Sicuri et al. 2014; Yue et al. 2017). Actuation of the robot refers to the way in which the energy is produced and includes use of an electric motor, hydraulics, pneumatics, or human muscle (Yue et al. 2017). Transmission refers to the way in which the robot transfers the motion of the actuator to that of the arm, and includes linkages and cables (Yue et al. 2017). Lastly, sensors detect the force and position of the upper limb to provide feedback in response,

and these include physical or bioelectrical signals such as through an electroencephalogram or an electromyogram (Yue et al. 2017).

A table of various robotic devices used in stroke rehabilitation is outlined below (Table 17).

Robotic Devices	Description			
Arm/Shoulder End- Effectors	MIT-Manus was one of the first robotic devices to be developed and is the most commonly used end-effector (Sicuri et al. 2014). It is a 2-degree-of-freedom robot manipulator that assists in goal-directed shoulder and elbow movements within the horizontal plane, while			
MIT-Manus (InMotion)	providing visual, auditory and tactile feedback (Masiero et al. 2007). A commercially available unit (InMotion ²) of this device is also available.			
 GENTLE/S (Haptic Master) MIME (Mirror Image Movement Enhancer) Neuro-X Arm Assist 	GENTLE/S or the Haptic Master is a 3-degree-of-freedom haptic interface arm with a wrist attachment mechanism, two embedded computers, a monitor and speakers and an overhead arm support system (Coote et al. 2008). The affected arm is de-weighted through a free moving elbow splint attached to the overhead frame (Coote et al. 2008). The subject is connected to the device by a wrist splint and feedback is provided during task-oriented training (Coote et al. 2008)			
 Bi-Manu-Track Arm Guide NeReBot Armeo Boom Continuous Passive 	MIME is a 6-degree-of-freedom robotic manipulator that is attached at the forearm through a splint. It provides bimanual movements as well as unilateral passive, active-assisted, and resisted movements of the hemiparetic upper extremity (Kahn et al. 2006; Burgar et al. 2011). More force is applied to the more affected forearm during goal-directed movements.			
Motion Devices (CYBEX and NORM, Shoulder 600)	Neuro-X is a 2-degree-of-freedom upper limb rehabilitation robot that assists in performing shoulder abduction-adduction and elbow flexion-extension movements in a horizontal plane. Feedback is provided through use of a monitor on which tasks are performed (Lee et al, 2016).			
	Arm Assist is a low-cost robotic system for rehabilitation of the shoulder and elbow post- stroke. The arm is supported through a device while playing interactive games (Tomic et al. 2017).			
	Bi-Manu-Track is a 1 degree-of-freedom device that enables bilateral and passive/active practice of forearm and wrist movement (Van Delden et al. 2012).			
	The ARM Guide offers 3 degrees of freedom and uses a motor and chain drive to move the user's hand along a linear rail, which assists reaching in a straight-line trajectory (Kahn et al. 2006).			
	The NeReBot is a 3-degrees-of-freedom, cable-driven device that produces sensorimotor stimulation and spatial movements of the shoulder and elbow. It is portable and can be used when the patient is either prone or sitting (Rosati et al. 2007; Masiero et al. 2007).			
	Armeo Boom is a 3-degree-of-freedom cable-driven manipulator (Sicuri et al. 2014).			
	A continuous passive motion device mobilizes a joint through supporting repetitive and reproducible movements (Hu et al. 2009).			
Arm/Shoulder Exoskeletons	ARMin is 7-degree-of-freedom exoskeleton robot that provides intensive and task-specific training to target improvements in motor function (Klamroth-Marganska et al. 2014).			
ARMin	Pneu-WREX is 4-degree-of-freedom pneumatically actuated upper extremity orthosis that provides robot assisted movement rehabilitation (Reinkensmeyer et al. 2012).			
Pneu-WREX Armeo Spring	Armeo Spring is 5-degree-of-freedom exoskeleton robot with an adjustable suspension system (Gijbels et al. 2011). Auditory and visual feedback are provided through the virtual reality system while various functional tasks are performed (Gijbels et al. 2011).			
Hand End-Effectors	The Amadeo assists in hand rehabilitation, having an end-effecter design. It helps with finger movements to allow for synchronization (Sale et al. 2014).			
Amadeo				
Hand Exoskeletons	The Music Glove is used with a game that promotes specific pinching movements to match musical notes displayed on a screen (Zondervan et al. 2016).			

Table 17. Robotic devices used for upper limb rehabilitation post-stroke

•	Music GloveGloreha (HAnd	The Gloreha hand rehabilitation glove provides repetitive and passive mobilization of the fingers with multisensory feedback through a computing device (Vanoglio et al. 2017).			
•	REhabilitation GLOve) RAPAEL Smart Glove FINGER Robot Modified Hand	The RAPAEL Smart Glove provides a 9-axis movement and position sensors along with acceleration channels, angular rate channels, magnetic field channels to assess wrist movement, and bending sensors to assess finger movement (Shin et al. 2016). The glove is worn during video games that are specifically designed to encourage specific rehabilitation exercises within the wrist and fingers (Shin et al. 2016).			
•	Exoskeleton Robot Hand Mentor 	The FINGER robotic exoskeleton provides assistance with flexion and extension of the finger while playing a musical computer game (Rowe et al. 2017).			
		The modified hand exoskeleton robot enables individual finger control through joint movement sensing (Susanto et al., 2015). The robot is used to assist with gestures such as hand grasping/opening as well as finger pinching/opening (Susanto et al. 2015).			
		The Hand Mentor robotic device facilitates and assists in movement of the wrist and fingers. While the arm unit stabilizes the forearm, movement in the wrist and fingers are isolated. Visual and auditory feedback are provided through a computer control box (Linder et al. 2015).			

54 RCTs were found that evaluated upper limb robotics for motor rehabilitation.

40 RCTs evaluated arm/shoulder end-effectors (Ellis et al. 2018; Hsieh et al. 2017; Kim et al. 2017; Tomic et al. 2017; Fan et al. 2016; Lee et al. 2016; McCabe et al. 2015; Prange et al. 2015; Ang et al. 2014; Hesse et al. 2014; Hsieh et al. 2014; Lemmens et al. 2014; Masiero et al. 2014; Timmermans et al. 2014; Sale et al. 2014; Hsieh et al. 2012; Liao et al. 2012; Abdullah et al. 2011; Burgar et al. 2011; Conroy et al. 2011; Hsieh et al. 2011; Masiero et al. 2011; Lo et al. 2010; Hu et al. 2009; Coote et al. 2008; Hesse et al. 2008; Rabadi et al. 2008; Volpe et al. 2008; Masiero et al. 2007; Kahn et al. 2006; Lum et al. 2006; Masiero et al. 2006; Hesse et al. 2005; Fasoli et al. 2004; Stein et al. 2004; Volpe et al. 2004; Lum et al. 2002; Burgar et al. 2000; Volpe et al. 2000; V

Four RCTs evaluated arm/shoulder exoskeletons (Daunoraviciene et al. 2018; Brokaw et al. 2014; Klamroth-Marganska et al. 2014; Reinkensmeyer et al. 2012). Two RCTs evaluated hand end-effectors (Sale et al. 2014; Hwang et al. 2012).

Nine RCTs evaluated hand exoskeletons (Rowe et al. 2017; Vanoglio et al. 2017; Shin et al. 2016; Zondervan et al. 2016; Linder et al. 2015; Susanto et al. 2015; Wolf et al. 2015; Friedman et al. 2014; Kutner et al. 2010). One RCT evaluated miscellaneous robotic devices (Bustamante Valles et al. 2016).

The methodological details and results of all 56 RCTs are presented in Table 18.

Table To. NOTS evalua	ting robotics for upper extremi	
Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
····· poor on one one gory	Arm/Shoulder End-Effect	hors
Ellis et al. 2018 RCT (8) N _{Start} =32 N _{End} =32 TPS=Chronic	E: Progressive Abduction Loading Therapy and Horizontal-Plane Viscous Resistance using Robotic Device (Haptic Master) C: Progressive Abduction Loading Therapy Duration: 30min/d, 3d/wk for 8wk	 Maximum Reaching Distance (+exp) Elbow Extension and Rotation (+exp) Shoulder Extension, Abduction (+exp) Fugl-Meyer Assessment (-) Motor Activity Log (-) Quality of Movement (-) Rancho Los Amigos Functional Test for the Hemiparetic Upper Extremity (-)
Hsieh et al. (2017) RCT (6) N _{Start} =31 N _{End} =21 TPS=Subacute	E: Bilateral priming robot-aided (Bi-Manu- Track) therapy with task-oriented therapy C: Task-oriented therapy Duration: 90min/d, 5d/wk for 4wk	 Stroke Impacto Spper Externity () Stroke Impact Scale (-) Fugl-Meyer Assessment (-) Box and Block Test (-) Grip Strength (-) Modified Rankin Scale (-) Functional Independence Measure (-)
Kim et al. (2017) RCT (5) N _{Start} =33 N _{End} =30 TPS=Chronic	E: External Focus with Robotic Arm (InMotion ARM) C: Internal Focus with Robotic Arm Duration: 45min/d, 3d/wk for 4wk	 Joint Independence (-) Fugl-Meyer Assessment (-) Wolf Motor Function Test (-)
Tomic et al. (2017) RCT (7) N _{Start} =26 N _{End} =26 TPS=Subacute	E: ArmAssist Robot C: Conventional Therapy Duration: 30min/d, 5d/wk for 3wk	 Fugl-Meyer Assessment (+exp) Wolf Motor Function Test (+exp) Barthel Index (-)
Fan et al. (2016) RCT (4) N _{Start} =6 N _{End} =6 TPS=Chronic	E: Robot-assisted bilateral arm therapy (Bi- Manu-Track) C: Dose-matched control therapy Duration: 45min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (-) Wolf Motor Function Test (-)
Lee et al. (2016) RCT (4) N _{Start} =58 N _{End} =44 TPS=Acute	E: Robotic-assisted therapy (Neuro-X) C: Conventional rehabilitation Duration: 1hr/d, 5d/wk for 2wk	 Manual Muscle Test (-) Manual Function Test (-) Modified Barthel Index (-)
McCabe et al. (2015) RCT (6) N _{Start} =39 N _{End} =35 TPS=Chronic	E1: Robotic training (InMotion ARM) + motor learning E2: Motor learning + functional electrical stimulation C: Motor learning Duration: 5hr/d, 5d/wk for 12wk	 Arm Motor Ability Test (-) Fugl-Meyer Assessment (-)
Prange et al. (2015) RCT (7) N _{Start} =70 N _{End} =68 TPS=Acute	E: Arm training with robot (ArmeoBoom) C : Conventional training Duration : 30min/d, 4d/wk for 6wk	 Stroke Upper Limb Capacity Scale (-) Reaching Distance (-) Fugl-Meyer Assessment (-)
Ang et al. (2014) RCT (7) N _{Start} =26 N _{End} =25 TPS=Chronic	E: Brain Computer Interface Coupled with MIT-Manus shoulder-elbow robotic feedback C: Training with the MIT-Manus Duration: 2hr/d, 3d/wk for 4wk	Fugl-Meyer Assessment (-)
<u>Hesse et al.</u> (2014) RCT (8)	E: Group robot therapy (Bi-Manu-Track) + individual arm therapy	 Box and Block Test (-) Action Research Arm Test (-)

Table 18. RCTs evaluating robotics for upper extremity motor rehabilitation

N _{Start} =50 N _{End} =46 TPS=Acute	C: Individual arm therapy Duration: 30min/d, 5d/wk for 6wk	
Hsieh et al. (2014) RCT (8) Nstart=48 NEnd=48 TPS=Chronic	E1: Robotic training (Bi-Manu-Track) + dCIT (distributed constraint induced therapy) E2: Robotic therapy C: Conventional therapy Duration: 1hr/d, 5d/wk for 5wk	E1 vs E2 • Fugl-Meyer Assessment (+exp) • Wolf Motor Function Test (+exp) <u>E1 vs C</u> • Fugl-Meyer Assessment (+exp) • Wolf Motor Function Test (+exp) <u>E2 vs C</u> • Fugl-Meyer Assessment: (+exp ₂) • Wolf Motor Function Test (+exp ₂) • E1 vs E2, E1 vs C & E2 vs C • Motor Activity Log (-)
Lemmens et al. (2014) RCT (7) Nstart=16 NEnd=16 TPS=Chronic	E: Robotic therapy (Haptic Master) C: No robotic therapy Duration: 30min (2x/d), 4d/wk for 8wk	 Fugl-Meyer Assessment (-) Action Research Arm Test (-) Motor Activity Log (-)
Masiero et al. (2014a) RCT (7) N _{Start} =34 N _{End} =30 TPS=Chronic	E: Robotic therapy (NeReBot) C: Standard therapy Duration: 2hr/d, 5d/wk for 5wk	 Fugl-Meyer Assessment (-) Box and Block test (-) Frenchay Arm Test (-) Medical Research Council Scale (-) Functional Independence Measure (-)
Timmermans et al. (2014) RCT (8) N _{Start} =22 N _{End} =22 TPS=Chronic	E: Robotic arm training (Haptic Master) C: Task oriented arm training Duration: 30min (2x/d), 4d/wk for 8wk	 Fugl-Meyer Assessment (-) Action Research Arm test (-) Motor Activity Log (-)
<u>Sale et al.</u> (2014) RCT (6) N _{Start} =53 N _{End} =53 TPS=Acute	E: Robot aided therapy (MIT-Manus) + reaching tasks C: Reaching tasks Duration: 1hr/d, 2d/wk for 10wk	 Fugl-Meyer Assessment (+exp) Motricity Index (+exp)
Hsieh et al. (2012) RCT (7) Nstart=54 N _{end} =53 TPS=Chronic	E1: High intensity robotic therapy (Bi- Manu-Track) E2: Low intensity robotic therapy C: Conventional therapy Duration: 90min/d, 5d/wk for 3wk	E1 vs E2 Fugl-Meyer Assessment: (+exp) E1 vs C Fugl-Meyer Assessment: (+exp) E1 vs E2 & E1 vs C Medical Research Council Scale (-) Motor Activity Log (-) Stroke Impact Scale (-)
Liao et al. (2012) RCT (7) N _{start} =20 N _{end} =20 TPS=Chronic	E: Robotic therapy (Bi-Manu-Track) C: Dose-matched conventional therapy Duration: 100min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Motor Activity Log (+exp) ABILHAND (+exp)
Abdullah et al. (2011) RCT (5) N _{start} =20 N _{end} =20 TPS=Acute	E: Robot assisted therapy C: Dose-matched conventional therapy Duration: <i>Not Specified</i>	Chedoke Arm and Hand Activity Inventory (-)
Burgar et al. (2011) RCT (5) N=54 TPS=Acute	E1: High intensity robotic therapy (MIME) E2: Low intensity robotic therapy C: Conventional therapy Duration: 1hr/d, 3d/wk for 8wk	E1 vs C • Functional Independence Measure (+exp) • Modified Ashworth Scale (-) • Fugl-Meyer Assessment (-)
<u>Conroy et al. (2011)</u> RCT (6) N _{start} =62	E1: Robot-assisted (InMotion ARM) planar reaching	Fugl-Meyer Assessment (-)

N _{end} =54 TPS=Chronic	E2: Robot-assisted planar and vertical reaching C: Intensive conventional arm therapy Duration: 1hr/d. 3d/wk for 6wk	
Hsieh et al. (2011) RCT (8) N _{start} =18 N _{end} =18 TPS=Chronic	E1: High intensity robot-assisted therapy (Bi-Manu-Track) E2: Low intensity robot-assisted therapy C: Conventional therapy Duration: 45min/d, 3d/wk for 6wk	 E1 vs E2 Fugl-Meyer Assessment: (+exp) E2 vs. C Fugl-Meyer Assessment (-) E1 vs C Motor Activity Log (+exp) E1 vs E2/C Motor Activity Log (-) ABILHAND (-) Medical Research Council Scale (-)
Masiero et al. (2011) RCT (5) Nstart=21 Nend=21 TPS=Acute	E: Robotic arm therapy (NeReBot) C: Conventional therapy Duration: <i>Not Specified</i>	 Medical Research Council Scale (+exp) Fugl-Meyer Assessment (-) Functional Independence Measure (-) Modified Ashworth Scale (-) Frenchay Arm Test (-) Box and Block Test (-)
Lo et al. (2010) RCT (7) N _{start} =127 N _{end} =127 TPS=Chronic	E1: Intensive robot assisted therapy (MIT- Manus) E2: Intensive comparison therapy C: Usual care Duration: 1hr/d, 3d/wk for 12wk	E1 vs C Fugl-Meyer Assessment (-) Wolf Motor Function Test (-) Stroke Impact Scale (+exp) Modified Ashworth Scale (-) E1 vs E2 Fugl-Meyer Assessment (-) Wolf Motor Function Test (-) Stroke Impact Scale (-) Modified Ashworth Scale (-)
Hu et al. (2009) RCT (5) N _{start} =27 N _{end} =27 TPS=Chronic	E: EMG-driven robot (CYBEX and NORM Continuous Passive Motion) C: Passive motion device Duration: 20min/d, 5d/wk for 7wk	 Fugl-Meyer Assessment (+exp) Modified Ashworth Scale (+exp)
Coote et al. (2008) RCT (6) Nstart=23 Nend=20 TPS=Chronic	E: Robot-mediated therapy (GENTLE/s) C: Sling suspension phase Duration: 30min/d, 3d/wk for 9wk	Fugl-Meyer Assessment (+exp)
Hesse et al. (2008) RCT (8) Nstart=54 Nend=47 TPS=Subacute	E: Computerized arm trainer (Reha-Slide Mechanical Arm Trainer) C: Electrical stimulation Duration: 25min/d, 5d/wk for 6wk	 Fugl-Meyer Assessment (-) Barthel Index (-)
Rabadi et al. (2008) RCT (5) Nstart=30 Nend=30 TPS=Acute	E1: Robot (MIT-Manus)-unilateral group E2: Ergometer (bilateral) group C: Conventional therapy Duration: 3hr/d, 3d/wk for 4wk	E1 vs E2/C Fugl-Meyer Assessment (-)
<u>Volpe et al.</u> (2008) RCT (5) N _{start} =21 N _{end} =21 TPS=Chronic	E: Sensorimotor arm training delivered by robotic device (MIT-Manus) C: Sensorimotor arm training delivered by a therapist Duration: 1hr/d, 3d/wk for 6wk	 Fugl-Meyer Assessment (-) Motor Power Scale (-)
<u>Masiero et al.</u> (2007) RCT (5) N _{start} =20 N _{end} =20	E: Robotic Training (NeReBot) C: Exposure to robotic device Duration: 1hr/d, 4d/wk for 5wk	 Fugl-Meyer Assessment (+exp) Medical Research Council (+exp) Functional Independence Measure (+exp) Modified Ashworth Scale (-)

TPS=Acute		Γ	
Kahn et al. (2006)	E: Active-assistive reaching exercise using	•	Rango Los Amigos Functional Test (-)
RCT (4)	a robotic device (Arm Guide)		
N _{start} =19	C: Task-matched amount of reaching		
N _{end} =19	without assistance		
TPS=Chronic	Duration: 40min/d, 6d/wk for 4wk		
Lum et al. (2006)	E1: Robot-unilateral (MIME)	Г	E3 vs C
RCT (4)	E2: Robot-bilateral	•	Fugl-Meyer Assessment (+exp ₃)
N _{start} =30	E3: Robot-combined	•	Motor Status Score (+exp ₃)
Nend=23	C: Conventional therapy	•	Functional Independence Measure (-)
TPS=Subacute	Duration: 30min/d, 3d/wk for 4wk	•	Motor power examination (-)
		•	Modified Ashworth Scale (-)
			<u>E3 vs E1</u>
		•	Fugl-Meyer Assessment (-)
		•	Motor Status Score (-)
		•	Functional Independence Measure (-)
		•	Motor power examination (-)
		•	Modified Ashworth Scale (-)
Masiero et al. (2006)	E: Additional sensorimotor robotic training	•	Fugl-Meyer Assessment (+exp)
RCT (5)	(NeReBot)	•	Motricity Index (+exp)
N _{start} =35	C: Exposure to robotic device with no	•	Functional Independence Measure (+exp)
N _{end} =35	training		Medical Research Council Scale (-)
TPS=Acute	Duration: 1hr/d, 4d/wk for 8wk		
<u>Hesse et al.</u> (2005)	E: Computerized arm training enabling		Fugl-Meyer Assessment (+exp)
RCT (8)	repetitive practice (Bi-Manu-Track)		
N _{start} =44	C: Electrical stimulation		
N _{end} =39	Duration: 20min/d, 5d/wk for 6wk		
TPS=Subacute		╞	
Fasoli et al. (2004)	E: Robot assisted (MIT-Manus) movement	•	Fugl-Meyer Assessment (+exp)
RCT (6)	training	•	Motor status score (-)
N _{start} =56	C: Robot exposure	•	Medical Research Council score (-)
Nend=56	Duration: 90min/d, 2d/wk for 12wk		
1PS=Acute		┝	
<u>Stein et al. (</u> 2004)	E1: Robot-aided (InMotion ARM)	•	Fugl-Meyer Assessment (-)
RCT (5)	progressive resistance training	•	Strength (-)
N _{start} =47	E2: Active-assisted robot-aided exercise		
Nend=40	Duration: 1nr/d, 4d/wk for 3wk		
	E. Orationary Descine Mating Device	┢	
	E: Continuous Passive Motion Device	•	Fugi-Meyer Assessment (-)
Not (4) Note=32			Modified Ashworth Scale ()
N -22	Durotion : 20min/d. 2d/wk for 4wk	•	Modified Astiworth Scale (-)
TPS-Acute			
Lum et al. (2002)	E: Robot (MIME)-assisted movement		Fugl-Meyer Assessment (+eyp)
RCT (6)	training		Strength under extremity (±exp)
Netort=30	C: Conventional therapy		Reach upper extremity (+exp)
Nend=27	Duration: 1hr/d 5d/wk for 6wk		Functional Independence Measure (-)
TPS=Chronic			
Burgar et al. (2000)	E: Robotic (MIME) device therapy	•	Fugl-Mever Assessment (-)
RCT (5)	C: Conventional care (physical therapy)		Functional Independence Measure (-)
N _{start} =21	Duration: 2hr/d. 3d/wk for 10wk		Barthel Index (-)
N _{end} =21			
TPS=Chronic			
Volpe et al. (2000a)	E: Robotic training (MIT-Manus)	•	Motor Power score: shoulder and elbow (+exp),
RCT (6)	C: Exposure to the robotic device without		wrist and hand (-)
N _{start} =56	training	•	Motor Status score: shoulder and elbow (+exp),
N _{end} =56	Duration: 30min/d, 3d/wk for 4wk		wrist and hand (-)
TPS=Acute		•	Functional Independence Measurer (+exp)
		•	Fugl-Meyer Assessment: (-)
Volpe et al. (1999)	E: Robot (MIT-Manus)	•	Motor Status score (+exp)

RCT (6) N _{start} =20 N _{end} =12	C: Sham treatment Duration: 45min/d, 5d/wk for 6wk	 Motor Status score (-) Motor Power score (+exp) Fugl-Meyer Assessment (-) 	
TPS=Acute			
Arm/Shoulder Exoskeletons			
Daunoraviciene et al. (2018) Lithuania RCT (5) N _{Start} =34 N _{End} =34 TPS= Subacute	E: Robot-assisted Training (Armeo Spring) C: Conventional Therapy Duration: 1hr/d, 4d/wk for 5wk	 Functional Independence Measure (-) Fugl-Meyer Assessment (-) Shoulder Flexion, Abduction, Adduction, and Internal Rotation (+exp) Elbow Flexion, Supination, and Pronation (+exp) Wrist Range of Motion (-) 	
Brokaw et al. (2014) RCT (3) N _{Start} =12 N _{End} =10 TPS=Chronic	E: Robotic therapy (ARMin) C: Conventional therapy Duration: 90min/d, 3d/wk for 4wk	 Fugl-Meyer Assessment (-) Action Research Arm Test (+exp) Box and Bock Test (-) 	
Klamroth-Marganska et al. (2014) RCT (8) N _{Start} =77 N _{End} =73 TPS= Chronic	E: Robotic therapy (ARMin) C: Conventional treatment Duration: 1hr/d, 3d/wk for 8wk	 Fugl-Meyer Assessment (+exp) Strength (+exp) Motor Activity Log (-) Modified Ashworth Scale (-) Wolf Motor Function test (-) 	
Reinkensmeyer et al. (2012) RCT (7) N _{start} =26 N _{end} =26 TPS=Chronic	E: Robotic training (Pneu-WREX) C: Conventional tabletop therapy Duration: 1hr/d, 3d/wk for 8wk	 Fugl-Meyer Assessment (-) Nottingham sensory test (-) Grip strength (-) Box and Block Test (-) 	
	Hand End-Effectors		
<u>Sale et al.</u> (2014) RCT (7) N _{Start} =20 N _{End} =20 TPS=Acute	E: Amadeo robotic therapy + physiotherapy C: Occupational therapy Duration : 1hr/d, 3d/wk for 4wk	 Box and Block Test (+exp) Fugl-Meyer Assessment (+exp) 	
Hwang et al. (2012) RCT (6) N _{start} =17 N _{end} =17 TPS=Chronic	E: Active Amadeo robot training C: Early passive therapy Duration : 45min/d, 5d/wk for 2wk	 Jebsen-Taylor Hand Function (-) Fugl-Meyer Assessment (-) Ashworth Scale (-) Nine Hole Peg Test (-) Stroke Impact Scale (-) 	
Hand Exoskeletons			
Rowe et al. (2017) RCT (7) Nstart =30 NEnd =30 TPS=Chronic	E: High Robotic Assistance Finger Training (FINGER robot) C: Low Robotic Assistance Finger Training Duration: 90min/d, 5d/wk for 8wk	 Fugl-Meyer Assessment (+exp) Box and Block Test (-) Action Research Arm Test (-) Nine-Hole Peg Test (-) Finger Tapping (-) Motor Activity Log (-) National Institutes of Health Stroke Scale (-) 	
Vanoglio et al. (2017) RCT (7) Nstart =30 NEnd =27 TPS=Acute	E: Robotic Glove with Multisensory Feedback (Gloreha hand rehab glove) C: Conventional Therapy Duration: 30min/d, 3d/wk for 5wk	 Motricity Index (+exp) Nine Hole Peg Test (+exp) Grip Strength (+exp) Pinch Test (+exp) Quick Version of Disabilities of the Arm, Shoulder, and Hand Questionnaire (+exp) 	
Shin et al. (2016) RCT (8) Nstart=46 NEnd=46 TPS=Chronic	E: RAPAEL SmartGlove virtual reality task training C: Conventional therapy Duration: 40min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Jebsen Taylor Hand Function Test (+exp) Stroke Impact Scale (+exp) Purdue Pegboard Test (-) 	
<u>Zonuervan et al.</u> (2016)	E. DOME-DASED TRAINING WITH A MUSICE OVE	 initial Activity Log (+exp) 	

RCT (6) N _{Start} =18 N _{End} =17 TPS=Chronic	C: Conventional tabletop exercise Duration: 25min/d, 6d/wk for 4wk	 Box and Block Test (-) 9-Hole Peg Test (-) Action Research Arm Test (-) 	
Linder et al. (2015) RCT (5) N _{Start} =99 N _{End} =99 TPS=Acute	E: Robot-assisted therapy program + home exercise program (Hand Mentor) C: Home exercise program Duration: 30min/d, 2d/wk for 5wk	Stroke Impact Scale (+exp)	
Susanto et al. (2015) RCT (7) N _{Start} =19 N _{End} =19 TPS=Chronic	E: Robotic paretic hand therapy (exoskeleton device) C: Task therapy without robotic aid Duration: 1hr/d, 5d/wk for 4wk	Wolf Motor Function Test (+exp)	
Wolf et al. (2015) RCT (6) Nstart=99 NEnd=92 TPS=Acute	E: Telemonitored robotic assisted home exercise therapy program (Hand Mentor) C: Dose-matched usual care home program Duration; 3hr/d, 5d/wk for 8wk	 Wolf Motor Function Test (+exp) Fugl-Meyer Assessment (-) Action Research Arm Test (-) 	
Friedman et al. (2014) RCT (6) N _{Start} =12 N _{End} =12 TPS=Chronic	E1: IsoTrainer E2: Music glove training C: Control Duration: 1hr/d, 3d/wk for 2wk	 Wolf Motor Function Test (-) Fugl-Meyer Assessment (-) Action Research Arm Test (-) <u>E2 vs C</u> Box and Block Test: (+exp₂) Nine Hole Peg Test: (+exp₂) <u>E1 vs E2</u> Box and Block Test: (-) Nine Hole Peg Test: (-) <u>E1 vs C</u> Box and Block Test: (-) Nine Hole Peg Test: (-) Nine Hole Peg Test: (-) 	
Kutner et al. (2010) RCT (7) N _{start} =30 N _{end} =26 TPS=Subacute/Chronic	E: Robot therapy (Hand Mentor) C: Conventional therapy Duration: 1hr/d, 5d/wk for 6wk	Stroke Impact Scale (+exp)	
Other			
Bustamante Valles et al. (2016) RCT (3) N _{Start} =27 N _{End} =20 TPS=Chronic	E: Rehabilitation using a technology- assisted rehabilitation gymnasium (circuit with various robots) C: Traditional therapy Duration: 2hr/d, 4d/wk for 6wk	 Fugl-Meyer Assessment (-) Box and Block Test (-) 	

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α=0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group - indicates no statistically significant between groups differences at α =0.05
Conclusions about robotics

	MOTOR FUNCTION		
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of various arm/shoulder end-effectors (MIME, Neuro-X, GENTLE/s/Haptic Master, MIT-Manus/In Motion, NeReBot, ArmeoBoom, continuous passive motion) to improve motor function when compared to conventional therapy.	17	Tomic et al. 2017; Lee et al. 2016; Prange et al. 2015; Lemmens et al. 2014; Masiero et al. 2014; Sale et al. 2014; Abdullah et al. 2011; Burgar et al. 2011; Conroy et al. 2011; Masiero et al. 2011; Lo et al. 2010; Rabadi et al. 2008; Kahn et al. 2006; Lum et al. 2006; Volpe et al. 2004; Lum et al. 2002; Burgar et al. 2000
1a	A specific arm/shoulder end-effector (Bi-Manu-Track) may produce greater improvements in motor function than conventional therapy .	5	Fan et al. 2016; Hsieh et al. 2014; Hsieh et al. 2012; Liao et al. 2012; Hsieh et al. 2011
1a	Arm/shoulder end-effectors (Bi-Manu-Track, Haptic Master) may not have a difference in efficacy when compared to task-oriented therapy for improving motor function.	2	Hsieh et al. 2017; Timmermans et al. 2014
1a	Arm/shoulder end-effectors (Haptic Master, MIT- Manus/InMotion) may not have a difference in efficacy when compared to active control therapies (progressive abduction loading therapy, motor learning, or sensorimotor arm training) for improving motor function.	3	Ellis et al. 2018; McCabe et al. 2015; Volpe et al. 2008
1a	Arm/shoulder end-effectors (GENTLE, MIT- Manus/InMotion, and continuous passive motion devices) may produce greater improvements in motor function than passive control therapies (a sling suspension phase, passive motion, and sham robotics).	3	Hu et al. 2009; Coote et al. 2008; Volpe et al. 1999.
1b	Arm/shoulder end-effectors (MIT-Manus/InMotion, NeReBot) may produce greater improvements in motor function than exposure to the robotic device without active assistance.	4	Masiero et al. 2007; Masiero et al. 2006; Fasoli et al. 2004; Volpe et al. 2000a
1b	Arm/shoulder end-effectors (Bi-Manu-Track, MIT- Manus/InMotion) provided in a group setting may not have a difference in efficacy when compared to arm/shoulder end-effectors provided in a one on one setting for improving motor function.	2	Kim et al. 2017; Hesse et al. 2014
1b	There is conflicting evidence about the effect of a specific arm/shoulder end-effector (Bi-Manu-Track) to improve motor function when compared to cyclic NMES .	2	Hesse et al. 2008; Hesse et al. 2005
1b	Arm/shoulder end-effectors (MIT-Manus/InMotion) combined with additional therapies may not have a difference in efficacy when compared to using these robotics alone for improving motor function.	2	Ang et al. 2014; Stein et al. 2004
1a	There is conflicting evidence about the effect of arm/shoulder exoskeletons (ARMin, Armeo Spring	4	Daunoraviciene et al. 2018; Brokaw et al. 2014; Klamroth-

	or Pneu-WREX) to improve motor function when compared to conventional therapy.		Marganska et al. 2014; Reinkensmeyer et al. 2012
1a	There is conflicting evidence about the effect of hand end-effectors (Amadeo hand robot) to improve motor function when compared to conventional therapy.	2	Sale et al. 2014; Hwang et al. 2012
1a	There is conflicting evidence about the effect of hand exoskeletons (Glohera, SmartGlove, Hand Mentor, Music Glove) to improve motor function when compared to conventional therapy.	6	Rowe et al. 2017; Shin et al. 2016; Zondervan et al. 2016; Wolf et al. 2015; Susanto et al. 2015; Friedman et al. 2014

MUSCLE STRENGTH				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of arm/shoulder end-effectors (Bi-Manu-Track, Neuro- X, MIME, NeReBot, and MIT-Manus) to improve muscle strength when compared to conventional therapy.	9	Lee et al. 2016; Masiero et al. 2014; Sale et al. 2014; Hsieh et al. 2012; Hsieh et al. 2011; Masiero et al. 2011; Rabadi et al. 2008; Lum et al. 2006; Lum et al. 2002	
1b	A specific arm/shoulder end-effector (Bi-Manu- Track) may not have a difference in efficacy when compared to task-oriented training for improving muscle strength.	1	Hsieh et al. 2007	
2	Arm/shoulder end-effectors (MIT-Manus//InMotion) may not have a difference in efficacy when compared to active control therapies (sensorimotor arm training, progressive resistance training) for improving muscle strength.	2	Volpe et al. 2008; Stein et al. 2004	
1b	There is conflicting evidence about the effect of arm/shoulder exoskeletons (ARMin or Pneu-WREX) to improve muscle strength when compared to conventional therapy.	2	Klamroth-Marganska et al. 2014; Reinkensmeyer et al. 2012	
1b	Hand exoskeletons (Gloreha) may produce greater improvements in muscle strength than conventional therapy.	1	Vanoglio et al. 2017	

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
1b	Arm/shoulder end-effectors (MIME, MIT-Manus, NeReBot, and continuous passive motion devices) may not have a difference in efficacy when compared to conventional therapy for improving dexterity.	6	Masiero et al. 2014; Burgar et al. 2011; Masiero et al. 2011; Lo et al. 2010; Lum et al. 2006; Volpe et al. 2004	
1b	A specific arm/shoulder end-effector (Bi-Manu- Track) may not have a difference in efficacy when compared to task-oriented training for improving dexterity.	1	Hsieh et al. 2017	
2	Arm/shoulder end-effectors (Bi-Manu-Track, MIT- Manus/InMotion) provided in a group setting may not have a difference in efficacy when compared to arm/shoulder end-effectors provided in a one on one setting for improving dexterity.	1	Hesse et al. 2014	
1b	Arm/shoulder exoskeletons (ARMin, Pneu-WREX) may not have a difference in efficacy when compared to conventional therapy for improving dexterity.	2	Brokaw et al. 2014; Reinkensmeyer et al. 2012	
1a	There is conflicting evidence about the effect of hand end-effectors (Amadeo hand robot) to improve dexterity when compared to conventional therapy.	2	Sale et al. 2014; Hwang et al. 2012	
1a	There is conflicting evidence about the effect of hand exoskeletons (Glohera, SmartGlove, Music Glove) to improve dexterity when compared to conventional therapy.	4	Vanoglio et al. 2017; Shin et al. 2016; Zondervan et al. 2016; Friedman et al. 2014	

RANGE OF MOTION				
LoE	Conclusion Statement	RCTs	References	
	A specific arm/shoulder end-effector (Haptic Master)		Ellis et al. 2018	
1h	combined with progressive resistance training may	1		
	produce greater improvements in range of motion than			
	progressive resistance training alone.			
•	External focus while using an arm/shoulder end-		Kim et al. 2017	
2	effector (MIT-Manus/InMotion) may not have a	1		
	difference in efficacy when compared to internal	1		
	focus while using an arm/shoulder end-effector			
	(MIT-Manus/InMotion) for improving range of motion.			
	A specific arm/hand exoskeleton (Armeo Spring)	1	Daunoraviciene et al.	
2	may produce greater improvements in range of motion	I	2018	
	than conventional therapy.			

	ACTIVITIES OF DAILY LIV	ING	
LoE	Conclusion Statement	RCTs	References
1a	Arm/shoulder end-effectors (MIME, Bi-Manu-Track, Neuro-X, Haptic Master, NeReBot, ArmAssist) may not have a difference in efficacy when compared to conventional therapy or task-oriented training for improving performance of activities of daily living.	16	Hsieh et al. 2017; Tomic et al. 2017; Lee et al. 2016; Hsieh et al. 2014; Lemmens et al. 2014; Masiero et al. 2014; Timmermans et al. 2014; Hsieh et al. 2012; Liao et al. 2012; Burgar et al. 2011; Hsieh et al. 2011; Masiero et al. 2011; Lo et al. 2010; Lum et al. 2006; Lum et al. 2002; Burgar et al. 2000
1a	Arm/shoulder end-effectors (Haptic Master, MIT- Manus/InMotion) may not have a difference in efficacy when compared to active control therapies (progressive abduction loading therapy or motor learning) for improving performance of activities of daily living.	2	Ellis et al. 2018; McCabe et al. 2015
1b	A specific arm/shoulder end-effector (Bi-Manu- Track) may not have a difference in efficacy when compared to cyclic NMES for improving performance of activities of daily living.	1	Hesse et al. 2008
1b	Arm/shoulder exoskeletons (ARMin, Armeo Spring) may not have a difference in efficacy when compared to conventional therapy for improving performance of activities of daily living.	2	Daunoraviciene et al. 2018; Klamroth- Marganska et al. 2014
1b	Hand end-effector (Amadeo hand robot) may not have a difference in efficacy when compared to early passive training for improving performance of activities of daily living.	1	Hwang et al. 2012
1a	Hand exoskeletons (MusicGlove, SmartGlove, Hand Mentor) may produce greater improvements in performance of activities of daily living than conventional therapy.	4	Zondervan et al. 2016; Shin et al. 2016; Linder et al. 2015; Kutner et al. 2010
1b	High assistance FINGER exoskeleton robotic use may not have a difference in efficacy when compared to low assistance FINGER exoskeleton robotic use for improving performance of activities of daily living.	1	Rowe et al. 2017

	PROPRIOCEPTION					
LoE	Conclusion Statement	RCTs	References			
1b	Arm/shoulder exoskeletons (Pneu-WREX) may produce greater improvements in proprioception than conventional therapy.	1	Reinkensmeyer et al. 2012			

SPASTICITY				
LoE	Conclusion Statement	RCTs	References	
1b	Arm/shoulder end-effectors (MIME, MIT- Manus/InMotion, NeReBot, and continuous passive motion devices) may not have a difference in efficacy when compared to conventional therapy for improving spasticity.	6	Masiero et al. 2014; Burgar et al. 2011; Masiero et al. 2011; Lo et al. 2010; Lum et al. 2006; Volpe et al. 2004	
2	A specific arm/shoulder end-effector (NeReBot) may not have a difference in efficacy when compared to exposure to the robotic device without active assistance for improving spasticity.	1	Masiero et al. 2007	
1b	Arm/shoulder exoskeletons (Armin) may not have a difference in efficacy when compared to conventional therapy for improving spasticity.	1	Klamroth-Marganska et al. 2014	
1b	Hand end-effector (Amadeo hand robot) may not have a difference in efficacy when compared to early passive training for improving spasticity.	1	Hwang et al. 2012	
1b	Hand exoskeletons (Gloreha) may produce greater improvements in spasticity than conventional therapy.	1	Vanoglio et al. 2017	

Key points

The evidence is mixed regarding arm/shoulder end-effector robotics, alone or in combination with other therapy approaches, for upper limb rehabilitation following stroke.

The evidence is mixed regarding arm/shoulder exoskeleton, hand exoskeleton, and hand end-effector robotics for upper limb rehabilitation.

Virtual reality



Adopted from: https://philadelphia.cbslocal.com/2016/05/15/virtual-reality-stroke-rehab/

Virtual reality interventions are described as the use of immersive multimedia created through computer programs that allows users to engage in simulated environments representative of both real-world and imagined places and objects (Iruthayarajah et al. 2017; Laver et al. 2017). These virtual reality interventions are presented typically as games with haptic feedback, that allow for the creation of a multisensory experience. Virtual reality interventions meet as the four guiding principles of rehabilitation: intensity, task-specific training, biofeedback and motivation (Dias et al. 2019). Research on the use of virtual reality for stroke rehabilitation is increasing as technology becomes more accessible and affordable. This includes using existing gaming consoles (e.g. Nintendo Wii, Xbox Kinect, Playstation Eyetoy) for therapeutic purposes or designing new systems specifically for rehabilitation (Laver et al. 2017).

A total of 36 RCTs evaluating virtual reality interventions for upper extremity motor rehabilitation were found, the characteristics of these interventions are described below.

30 RCTs evaluated virtual reality interventions compared to conventional therapy, recreational therapy or sham interventions (Askin et al. 2018; Faria et al. 2018; Kim et al. 2018; Kiper et al. 2018; Lee et al. 2018; Adie et al. 2017; Ballester et al. 2017; Brunner et al. 2017; Rand et al. 2017; Standen et al. 2017; Turkbey et al. 2017; Choi et al. 2016; Givon et al. 2016; Kong et al. 2016; Lee et al. 2016a; Lee et al. 2016c; Saposnik et al. 2016; da Silva Ribeiro et al. 2015; Shin et al. 2015; Choi et al. 2014; Fan et al. 2014; Kiper et al. 2014; Shin et al. 2014; Thielbar et al. 2014; Duff et al. 2013; Lee et al. 2013; Sin and Lee, 2013; Crosbie et al. 2012; Saposnik et al. 2010; Yavuzer et al. 2008). One RCT compared virtual reality bilateral arm training to bilateral arm training (Lee et al. 2018), and another combined virtual reality training with tDCS (Lee and Chun, 2014). One RCT compared virtual reality training with tDCS (Lee and Chun, 2014). One RCT compared asymmetric training with virtual reality training to mCIMT (McNulty et al. 2015). One RCT compared asymmetric training with virtual reality to symmetric training (Lee et al. 2014). One RCT compared virtual reality training with virtual reality to symmetric training (Lee et al. 2014). One RCT compared virtual reality training to no training (Jang et al. 2005).

The methodological details and results of all 36 RCTs are presented in Table 19.

Table 19. RCTs evaluating	virtual reality	interventions for	upper extremity	motor
rehabilitation	-			

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Virtual	reality training compared to conventional th	erapy, recreational therapy or sham
Askin et al. (2018) RCT (6) N _{Start} =40 N _{End} =38 TPS=Chronic	E: Xbox Kinect-based virtual reality training + physical therapy C: Physical therapy Duration: 1h/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Motricity Index (+exp) Active range of motion (+exp) Brunnstrom Recovery Stages (-) Modified Ashworth Scale (-) Box and Block Test (-)
Faria et al. (2018) RCT (4) Nstart = 32 NEnd = 24 TPS=Chronic	E: Virtual reality (Reh@Task) C: Time-matched standard occupational therapy Duration: 45min/d, 3d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Chedoke Arm and Hand Activity Inventory (-) Barthel Index (-) Motricity Index (-) Modified Ashworth Scale (-)
Kim et al. (2018) RCT (8) N _{Start} =23 N _{End} =19 TPS=Chronic	E: Kinect-based virtual reality C: Sham virtual reality Duration: 30min/d, 5d/wk for 2wk	 Fugl-Meyer Assessment (-) Brunnstrom Stage: Arm and Hand (-) Box and Block Test (-) Korean Modified Barthel Index (-)
Kiper et al. (2018) RCT (7) N _{Start} =139 N _{End} =136 TPS=Subacute	E: Reinforced feedback in virtual environment + conventional rehabilitation C: Conventional rehabilitation Duration: 1h/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Functional Independence Measure (+exp) National Institute of Health Stroke Scale (+exp)
Lee et al. (2018) RCT (6) N _{Start} =31 N _{End} =30 TPS=Subacute	E: Virtual reality canoe paddle training + conventional therapy C: Conventional therapy Duration: 30min/d, 3d/wk for 5wk	 Manual Function Test (+exp)
Adie et al. (2017) RCT (7) N _{Start} =235 N _{End} =209 TPS=Chronic	E: Wii arm exercises C: Home-based arm exercises Duration: 45min/d for 6wk	 Action Research Arm Test (-) Stroke Impact Questionnaire (-) Canadian Occupational Performance Measure (-) Motor Activity Log (-)
$\frac{\text{Ballester et al.}}{\text{RCT (5)}}$ $\frac{\text{N}_{\text{Start}} = 39}{\text{N}_{\text{End}} = 35}$ TPS=Chronic	E: Home-based virtual reality C: Home-based occupational therapy Duration: 30min/d, 5d/wk, 3wk	 Fugl-Meyer Assessment (-) Chedoke Arm and Hand Activity Inventory (+exp) Barthel Index (-) Medical Research Council Scale (-) Ashworth Scale (-) Grip force (-)
$\label{eq:start} \begin{array}{l} \frac{Brunner \ et \ al.}{RCT} (2017) \\ RCT \ (5) \\ N_{Start} = 120 \\ N_{End} = 102 \\ TPS = Subacute \end{array}$	E: Virtual reality training C: Conventional training Duration: 60min/d, 4-5d/wk for 4wk	 Action Research Arm Test (-) Box and Block Test (-) Functional Independence Measure (-)
Rand et al. (2017) RCT (7) Nstart = 24 N _{End} =21 TPS=Chronic	E: Video games self-training C: Traditional self-training Duration: 60min/d, 6d/wk for 5wk	 Action Research Arm Test (-) Motor Activity Log (-) Box and Block Test (-)

Standen et al. (2017) RCT (6) Nstart =27 NEnd =18 TPS=Subacute Turkbey et al. (2017) RCT (7) Nstart =20 NEnd =19 19	E: Home-based virtual reality C: Conventional therapy Duration: up to 60min/d, 7d/wk for 8wk E: Xbox Kinect virtual reality training + conventional rehabilitation C: Conventional rehabilitation Duration: 60min/d, 5d/wk for 4wk	 Motor Activity Log (+exp) Wolf Motor Function Test (-) Wolf Grip Strength (+exp) Nine-Hole Peg Test (-) Nottingham Extended Activities of Daily Living (-) Box and Block Test (+exp) Wolf Motor Function Test (+exp) Brunnstrom Motor Recovery Stage (+exp) Functional Independence Measure (-)
TPS=Subacute <u>Choi et al</u> . (2016) RCT (6) N _{Start} =24 N _{End} =24	E: Virtual reality rehabilitation program + conventional occupational therapy C: Conventional occupational therapy Duration: 30min/d, 5d/wk for 2wk	 Fugl-Meyer Assessment (+exp) Brunnstrom Stage (+exp) Manual Muscle Test (+exp) Modified Barthel Index (-)
Givon et al. (2016) RCT (6) N _{Start} =47 N _{End} =43 TPS=Chronic	E: Virtual reality video game therapy C: Traditional therapy Duration: 60min/d, 2d/wk for 12wk	 Action Research Arm Test (-) Grip strength (-)
<u>Kong et al.</u> (2016) RCT (7) N _{Start} =105 N _{End} =97 TPS=Acute	E: Nintendo Wii virtual reality training C: Conventional therapy Duration: 60min/d, 4d/wk for 3wk	 Fugl-Meyer Assessment (-) Action Research Arm Test (-) Stroke Impact Scale (-) Functional Independence Measure (-)
Lee et al. (2016a) RCT (7) N _{Start} =26 N _{End} =26 TPS=Chronic	E: Virtual reality-based rehabilitation C: Group-based rehabilitation Duration: 30min/d, 3d/wk for 8wk	 Fugl-Meyer Assessment (+exp) Manual Function Test (+exp) Box and Block Test (-) Modified Barthel Index (-)
Lee et al. (2016c) RCT (5) N _{Start} =14 N _{End} =10 TPS=Acute	E: Canoe game-based virtual reality training + conventional rehabilitation C: Conventional rehabilitation Duration: 30min/d, 3d/wk for 4wk	 Fugl-Meyer Assessment (+exp)
Saposnik et al. (2016) RCT (6) N _{Start} =141 N _{End} =121 TPS=Acute	E: Virtual reality training using Nintendo Wii C: Recreational activities Duration: 60min/d, 5d/wk for 2wk	 Wolf Motor Function Test (-) Box and Block Test (+con) Stroke Impact Scale (-) Barthel Index (-) Functional Independence Measure (-) Grip Strength (-)
$\label{eq:start} \begin{array}{c} \underline{da \; Silva \; Ribeiro \; et \; al.}}{RCT \; (7)} \\ RCT \; (7) \\ N_{Start} = 30 \\ N_{End} = 30 \\ TPS = Chronic \end{array}$	E: Virtual reality training using Nintendo Wii C: Conventional physical therapy Duration: 20min/d, 3d/wk for 12wk	Fugl-Meyer Assessment (-)
Shin et al. (2015) RCT (6) Nstart=35 NEnd=32 TPS=Chronic	E: Virtual reality + conventional occupational therapy C: Conventional occupational therapy Duration: 30min/d, 5d/wk for 4wk	Fugl-Meyer Assessment (-)
Choi et al. (2014) RCT (8) Nstart=20 NEnd=20 TPS=Chronic	E: Virtual reality therapy using Nintendo Wii C: Conventional occupational therapy Duration: 30min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (-) Box and Block Test (-) Manual Function Test (-) Grip strength (-) Modified Barthel Index (-)
Fan et al. (2014) RCT (7)	E1: Virtual reality E2: Conventional therapy	E1 vs E2 vs E3 vs C • Jebsen-Taylor Hand Function Test (-)

NEnd=20 C: No treatment TPS=Chronic Duration: 60min/d, 3d/wk for 3wk Kiper et al. (2014) E: Reinforced feedback in virtual	
TPS=Chronic Duration: 60min/d, 3d/wk for 3wk Kiper et al. (2014) E: Reinforced feedback in virtual 	
Kiper et al. (2014) E: Reinforced feedback in virtual • Fudl-Mever Assessment (+exp)	
RCT (6) environment + traditional rehabilitation • Functional Independence Measure (+exp)	
N _{Start} =46 C: Traditional rehabilitation	
N _{End} =44 Duration: 2h/d, 5d/wk for 4wk	
TPS=Chronic	
Shin et al. (2014) E: Virtual reality training + conventional • Fugl-Meyer Assessment (-)	
RCT (5) occupational therapy • Modified Barthel Index (-)	
Nstart=16 C: Occupational therapy Medical Research Council Score (-)	
N _{End} =16 Duration: 30min/d, 5d/wk for 2wk • Range of Motion (-)	
TPS=Chronic	
Thielbar et al. (2014)E: Virtual reality keypad/glove• Action Research Arm Test (+exp)	
RCT (6) C: Occupational therapy • Jebsen-Taylor Hand Function Test (-)	
Nstart=14 Duration: 18h/d, 3d/wk for 6wk • Fugl-Meyer Assessment (-)	
Chip Strength (-) Dinch Strength (-)	
TPS=Chronic	
Duff et al. (2013) E: Adaptive mixed reality rehabilitation Fuel-Meyer Assessment (+con)	
BCT (5) C: Traditional therapy	
Nstart=25 Duration: 60min/d. 3d/wk for 4wk • Stroke Impact Scale (-)	
• Motor Activity Log (-)	
TPS=Chronic	
Lee et al. (2013) E: Xbox Kinect-based virtual reality + • Manual Muscle Test (-)	
RCT (6) Conventional occupational therapy • Modified Ashworth Scale (-)	
N _{Start} =14 C: Conventional occupational therapy • Functional Independence Measure (-)	
N _{End} =14 Duration: 60min/d, 3d/wk for 6wk	
TPS=Chronic	
Sin & Lee (2013) E: Xbox Kinect-based virtual reality training Range of Motion (+exp)	
RCT (5) + conventional occupational therapy • Fugl-Meyer Assessment (+exp)	
Nstart=40 C: Conventional occupational therapy • Box and Block Test (+exp)	
N _{End} =35 Duration: 30min/d, 3d/wk for 6wk	
TPS=Chronic	
Crosbie et al. (2012) E: Virtual reality training • Motricity Index (-)	
RCT (8) C: Conventional physiotherapy • Action Research Arm Test (-)	
Nstart=18 Duration: 30-45min/d, 3d/wk for 3wk	
N _{end} =17	
TPS=Chronic	
Saposnik et al. (2010) E: Virtual reality training using Nintendo Wii • Wolf Motor Function Test (+exp)	
RCT (7) C: Recreational therapy • Grip strength (-)	
N _{start} =22 Duration: 60min/d, 4d/wk for 2wk • Box and Block Test (-)	
• Stroke impact Scale (-)	
IPS=Acute	
Yavuzer et al. (2008) E: Playstation EyeToy games + • Brunnstrom Recovery Stages (-)	
Conventional therapy Functional Independence Measure (+exp)	
C: Snam therapy + conventional therapy	
TDS-Subcouto	
virtual reality bilateral arm training compared to bilateral arm training	
E: VITUAI reality-based bilateral arm training • Jebsen Laylor Hand Function Lest (+exp)	
No. O Dilateral anni training • Dox and block rest (+exp) No. -20 • <	
Nstart-20 Duration. comminut, survivi or 4wk (• Citooved regional rest (+exp)	
Virtual reality with EES compared to EES	

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Lee et al. (2018) RCT (7) N _{Start} =48 N _{End} =41	E: Virtual reality + functional electrical stimulation C: Functional electrical stimulation Duration: 30min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Wolf Motor Function Test (-) Box and Block Test (-) Jebsen-Taylor Hand Function Test (-) 		
TPS=Chronic		Stroke Impact Scale (-)		
Virtual reality compared to and combined with cathodal tDCS				
Lee & Chun (2014) RCT (7) N _{Start} =64 N _{End} =59 TPS=Subacute	E1: Cathodal transcranial direct current stimulation (tDCS) E2: Virtual reality E3: Cathodal tDCS + virtual reality Duration: 30min/d, 5d/wk for 3wk	E3 vs E2/E1 Manual Function Test (+exp ₃) Fugl-Meyer Assessment (+exp ₃) Manual Muscle Test (-) Box and Block Test (-) Modified Ashworth Scale (-) Modified Barthel Index (-) E2 vs E1 Manual Function Test (+exp ₂) Fugl-Meyer Assessment (+exp ₂) Manual Muscle Test (-)		
		 Manual Muscle Test (-) Box and Block Test (-) Modified Ashworth Scale (-) 		
		Modified Barthel Index (-)		
	Virtual reality with a hand orthosis compa	red to conventional therapy		
Nijenhuis et al. (2017) RCT (6) N _{Start} =20 N _{End} =19 TPS=Chronic	E: Hand orthosis + computerised gaming exercises C: Conventional exercise Duration: 30min/d, 6d/wk for 6wk	 Action Research Arm Test (-) Fugl-Meyer Assessment (-) Grip Strength (-) Box and Block Test (-) Motor Activity Log (-) Stroke Impact Scale (-) 		
	Virtual reality compared to	n mCIMT		
McNulty et al. (2015) RCT (7) N _{Start} =41 N _{End} =40 TPS=Chronic	E: Nintendo Wii-based movement therapy C: Modified constraint-induced movement therapy Duration: 60min/d, 5d/wk for 2wk	 Wolf Motor Function Test (-) Motor Activity Log (-) Fugl-Meyer Assessment (-) Modified Ashworth Scale (-) Box and Block Test (-) Grooved Pegboard Test (-) Range of motion (-) 		
A	symmetric training with virtual reality comp	pared to symmetric training		
Lee et al. (2014) RCT (5) N _{Start} =30 N _{End} =24 TPS=Chronic	E: Asymmetric training using virtual reality + conventional physical therapy C: Symmetric training + conventional physical therapy Duration: 60min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Box and Block test (+exp) Grip strength (+exp) Modified Ashworth Scale (-) Range of motion (+exp) 		
	Virtual reality training compared	to no training		
<u>Jang et al</u> . (2005) RCT (5) N _{start} =10 N _{end} = 10 TPS=Chronic	E: Virtual reality training C: No treatment Duration: 60min/d, 5d/wk for 4wk	 Box and Block test (+exp) Fugl-Meyer Assessment (+exp) Manual Function Test (+exp) 		

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α=0.05

Conclusions about virtual reality

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of virtual reality interventions to improve motor function when compared to conventional therapy, recreational therapy or sham interventions.	30	Askin et al. 2018; Faria et al. 2018; Kim et al. 2018; Kiper et al. 2018; Lee et al. 2018; Adie et al. 2017; Brunner et al. 2017; Rand et al. 2017; Standen et al. 2017; Turkbey et al. 2017; Choi et al. 2016; Givon et al. 2016; Kong et al. 2016; Lee et al. 2016a; Lee et al. 2016c; Saposnik et al. 2015; Shin et al. 2015; Choi et al. 2014; Fan et al. 2014; Kiper et al. 2014; Shin et al. 2014; Thielbar et al. 2014; Mipf et al. 2013; Crosbie et al. 2012; Chosbie et al. 2014; Chosbie et al. 2014; Thielbar et al. 2014; Duff et al. 2013; Crosbie et al. 2012; Saposnik et al. 2010
1b	Virtual reality bilateral arm training may produce greater improvements in motor function than bilateral arm training.	1	Lee et al. 2016b
1b	Virtual reality interventions combined with FES may not have a difference in efficacy when compared to FES alone for improving motor function.	1	Lee et al. 2018
1b	Virtual reality interventions on their own or combined with cathodal tDCS may produce greater improvements in motor function than cathodal tDCS.	1	Lee and Chun, 2014
1b	Virtual reality training with a hand orthosis may not have a difference in efficacy when compared to conventional therapy for improving motor function.	1	Nijenhuis et al. 2017
1b	Virtual reality training may not have a difference in efficacy when compared to mCIMT for improving motor function.	1	McNulty et al. 2015
2	Asymmetric virtual reality training may produce greater improvements in motor function than symmetric conventional training.	1	Lee at al. 2014
2	Virtual reality training may produce greater improvements in motor function than no training .	1	Jang at al. 2005

STROKE SEVERITY				
LoE	Conclusion Statement	RCTs	References	
1b	Virtual reality interventions may produce greater improvements on measures of stroke severity than conventional therapy.	1	Kiper et al. 2018	

RANGE OF MOTION			
LoE	Conclusion Statement	RCTs	References
2	There is conflicting evidence about the effect of virtual reality interventions to improve range of motion when compared to conventional therapy, recreational therapy or sham interventions.	2	Shin et al. 2014; Sin and Lee, 2013
1b	Virtual reality training may not have a difference in efficacy when compared to mCIMT for improving range of motion.	1	McNulty et al. 2015
2	Asymmetric virtual reality training may produce greater improvements in spasticity than symmetric conventional training.	1	Lee at al. 2014

DEXTERITY			
LoE	Conclusion Statement	RCTs	References
1a	Virtual reality interventions may not have a difference in efficacy when compared to conventional therapy, recreational therapy or sham interventions for improving dexterity.	10	Askin et al. 2018; Kim et al. 2018; Brunner et al. 2017; Rand et al. 2017; Standen et al. 2017; Turkbey et al. 2017; Lee et al. 2016a; Choi et al. 2014; Sin and Lee, 2013; Saposnik et al. 2010
1b	Virtual reality bilateral arm training may produce greater improvements in dexterity than bilateral arm training.	1	Lee et al. 2016b
1b	Virtual reality interventions combined with FES may not have a difference in efficacy when compared to FES alone for improving dexterity.	1	Lee et al. 2018
1b	Virtual reality interventions on their own or combined with cathodal tDCS may not have a difference in efficacy when compared to cathodal tDCS for improving dexterity.	1	Lee and Chun, 2014
1b	Virtual reality training with a hand orthosis may not have a difference in efficacy when compared to conventional therapy for improving dexterity.	1	Nijenhuis et al. 2017
1b	Virtual reality training may not have a difference in efficacy when compared to mCIMT for improving dexterity.	1	McNulty et al. 2015
2	Asymmetric virtual reality training may produce greater improvements in dexterity than symmetric conventional training.	1	Lee at al. 2014
2	Virtual reality training may produce greater improvements in dexterity than no training .	1	Jang at al. 2005

SPASTICITY			
LoE	Conclusion Statement	RCTs	References
1a	Virtual reality interventions may not have a difference in efficacy when compared to conventional therapy, recreational therapy or sham interventions for improving spasticity.	4	Askin et al. 2018; Faria et al. 2018; Ballester et al. 2017; Lee et al. 2013
1b	Virtual reality interventions on their own or combined with cathodal tDCS may not have a difference in efficacy when compared to cathodal tDCS for improving spasticity.	1	Lee and Chun, 2014
1b	Virtual reality training may not have a difference in efficacy when compared to mCIMT for improving spasticity.	1	McNulty et al. 2015
2	Asymmetric virtual reality training may not have a difference in efficacy when compared to symmetric conventional training for improving spasticity.	1	Lee at al. 2014

MUSCLE STRENGTH			
LoE	Conclusion Statement	RCTs	References
1a	Virtual reality interventions may not have a difference in efficacy when compared to conventional therapy, recreational therapy or sham interventions for improving muscle strength.	12	Askin et al. 2018; Faria et al. 2018; Ballester et al. 2017; Standen et al. 2017; Choi et al. 2016; Givon et al. 2016; Saposnik et al. 2016; Choi et al. 2014; Shin et al. 2014; Lee et al. 2013; Crosbie et al. 2012; Saposnik et al. 2010
1b	Virtual reality bilateral arm training may produce greater improvements in muscle strength than bilateral arm training.	1	Lee et al. 2016b
1b	Virtual reality interventions on their own or combined with cathodal tDCS may not have a difference in efficacy when compared to cathodal tDCS for improving muscle strength.	1	Lee and Chun, 2014
1b	Virtual reality training with a hand orthosis may not have a difference in efficacy when compared to conventional therapy for improving muscle strength.	1	Nijenhuis et al. 2017
2	Asymmetric virtual reality training may produce greater improvements in muscle strength than symmetric conventional training.	1	Lee at al. 2014

ACTIVITIES OF DAILY LIVING			
LoE	Conclusion Statement	RCTs	References
1b	Virtual reality interventions combined with FES may not have a difference in efficacy when compared to FES alone for improving performance of activities of daily living.	1	Lee et al. 2018
1b	Virtual reality interventions on their own or combined with cathodal tDCS may not have a difference in efficacy when compared to cathodal tDCS for improving performance of activities of daily living.	1	Lee and Chun, 2014
1b	Virtual reality training with a hand orthosis may not have a difference in efficacy when compared to conventional therapy for improving performance of activities of daily living.	1	Nijenhuis et al. 2017
1b	Virtual reality training may not have a difference in efficacy when compared to mCIMT for improving performance of activities of daily living.	1	McNulty et al. 2015

Key points

Virtual therapy alone may not be more beneficial than conventional therapy for upper limb rehabilitation following stroke, however it may be beneficial for certain aspects of upper limb function when used in combination with conventional or other therapy approaches.

Brain computer interfaces



Adopted from: http://www.tech-faq.com/brain-computer-interface.html

Brain-computer interface (BCI) technology has only recently emerged as a potential rehabilitative treatment option following stroke. BCI records and decodes local brain activity during the performance of a motor movement (Van Dokkum et al. 2015). The decoded brain signals can be configured into visual, auditory or haptic feedback, and even for the control of external devices to help facilitate movement (Van Dokkum et al. 2015). BCI promotes the recruitment of brain areas involved in motor planning and execution and facilitates neural plasticity of neural networks using these areas, helping patients learn to generate normal brain activity or use brain activity to operate training devices (Van Dokkum et al. 2015). The evidence base for this intervention is still however in its infancy.

The methodological details and results of 5 RCTs evaluating BCI for the upper extremity motor rehabilitation in chronic stroke survivors are presented in Table 20.

Table 20. RCTs evaluating brain computer interfaces interventions for upper extremity motor rehabilitation

Authors (Year) Study Design (PEDro Score)	Interventions Duration: Session length, frequency	Outcome Measures Result (direction of effect)
Sample Sizestart	per week for total number of weeks	
Sample Sizeend		
Lin et al. (2018)	E1: Motion tracking device+ VR game	E2 vs C
$\frac{\operatorname{Ent}\operatorname{et}\operatorname{al.}\left(2010\right)}{\operatorname{RCT}\left(6\right)}$	E1: Motion tracking device + brain-	 Fugl-Mever Assessment (exp₂)
Not (0) Nstart =15	computer interface attention-monitoring	F1 vs C
NEnd =15	electroencephalogram device + VR	 Fugl-Mever Assessment (-)
TPS=Chronic	game	
	C: Conventional therapy	
	Duration: 35min/d, 3d/wk for 4wk	
Young et al. (2016)	E: Brain computer interface training	Stroke Impact Scale (-)
RCT (5)	C: No training	Action Research Arm Test (-)
N _{Start} =19	Duration: 120min/d for 9-15d	9 Hole Peg Test (-)
N _{End} =10		
TPS=Chronic		
Ang et al. (2015)	E: Brain computer interface + MIT-	 Fugl-Meyer Assessment (-)
RCT (7)	Manus robotic training	
N _{Start} =26	C: MIT-Manus robotic training	
N _{End} =25	Duration: 90min/d, 3d/wk for 4wk	
TPS=Chronic		
<u>Ang et al</u> . (2014)	E1: Brain-computer interface + haptic	<u>E1 vs C</u>
RCT (8)	knob (HK) robot	Fugl-Meyer Assessment (-)
N _{Start} =22	E2: HK robot	<u>E2 vs C</u>
N _{End} =21	C: Standard Arm Therapy (SAT)	Fugl-Meyer Assessment (-)
TPS=Chronic	Duration: 90min/d, 3d/wk for 6wk	$\frac{E1 \text{ VS E2}}{E \text{ usl Mover Assessment ()}}$
Ramos-Murguialday et al. (2013)	E: Brain machine interface (BMI)	Fugl Mever Assessment (+exp)
RCT (8)	C: Sham BMI	Motor Activity Log (-)
Noter 32	Duration: 5d/wk for 4wk	Goal Assessment Scale (-)
NEnd=30		Ashworth Scale (-)
TPS=Chronic		

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about brain computer interfaces

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
2	Brain computer interfaces may not have a difference in efficacy compared to no training for improving motor function.	1	Young et al. 2016
1a	Brain computer interfaces combined with robotic training may not have a difference in efficacy compared to robotic training alone for improving motor function.	2	Ang et al. 2015; Ang et al. 2014
1b	Brain computer interfaces may produce greater improvements in motor function than sham training.	1	Ramos-Murguialday et al. 2013
1b	Brain computer interfaces combined with motion tracking, electroencepholography and virtual reality may produce greater improvements in motor function than conventional therapy.	1	Lin et al. 2018

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
2	Brain computer interfaces may not have a difference in efficacy compared to no training for improving	1	Young et al. 2016	
	dexterity.			

SPASTICITY				
LoE	Conclusion Statement	RCTs	References	
1b	Brain computer interfaces may not have a difference in efficacy compared to sham training for improving spasticity.	1	Ramos-Murguialday et al. 2013	

ACTIVITIES OF DAILY LIVING				
LoE	E Conclusion Statement RCTs References			
2	Brain computer interfaces may not have a difference in efficacy compared to no training for improving activities of daily living.	1	Young et al. 2016	
1b	Brain computer interfaces may not have a difference in efficacy compared to sham training for improving activities of daily living.	1	Ramos-Murguialday et al. 2013	

Key points

The literature is mixed regarding brain-computer interface technology for upper limb motor rehabilitation following stroke, either on its own or combined with other therapies, but it may not be beneficial alone for other aspects of upper limb function.

EMG biofeedback



Adopted from: http://www.udbhavphysiotherapy.com/services/emg-biofeedback/10

EMG biofeedback for the treatment of hemiparesis after stroke is performed through the application of electrodes onto specific muscle groups important for a desired motor movement to monitor electrical activity during muscle contraction (Nelson, 2007). It then provides feedback of muscle activity back to the patient by conversion of myoelectrical activity into visual and/or auditory information to increase patient awareness and facilitate motor movement (Sturma et al. 2018). EMG biofeedback is particularly useful for small muscle contractions that are otherwise unnoticeable kinaesthetically or visually in the earlier stages of stroke recovery or in cases of severe paresis (Nelson, 2007).

The methodological details and results of 17 RCTs evaluating EMG biofeedback for the upper extremity motor rehabilitation are presented in Table 21.

Table 21. RCTs evaluating EMG biofeedback interventions for upper extremity motor rehabilitation

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Thielbar et al. (2017) RCT (6) N _{Start} =23 N _{End} =22 TPS=Chronic	E: EMG-driven actuated glove + conventional occupational therapy C: Occupational therapy Duration: 1 hr/d, 3d/wk for 6wk	 Hand Aperture (+exp) Action Research Arm Test (-) Wolf Motor Function Test (-) Fugl-Meyer Assessment (-) Chedoke McMaster Stroke Assessment (-) Grip/Pinch Strength (-)
<u>Kim et al.</u> (2017) RCT (2) N _{start} =30 N _{End} =30 TPS=Chronic	E: EMG Biofeedback and Conventional Therapy C: Conventional Therapy	 Fugl-Meyer Assessment (-) Manual Function Test (-) Functional Independence Measure (-)
Garrido-Montenegro et al. (2016) RCT (8) N _{Start} =14 N _{End} =14 TPS=Chronic	E: EMG/Biofeedback + conventional occupational therapy C: Occupational therapy Duration: 1 hr/d, 4d/wk for 4wk	 Barthel Index (+exp) Instrumental Activities of Daily Living (+exp) Action Research Arm Test (+exp) Motor Activity Log (+exp)
$\frac{Chang-Yong et al.}{RCT (7)}$ $\frac{N_{Start}=44}{N_{End}=40}$ TPS=Chronic	E: Target reaching training with biofeedback + routine therapy C: Routine therapy Duration: <i>Not Specified</i>	 Fugl-Meyer Assessment (+exp) Wolf Motor Function Test (+exp) Reach speed (+exp) Reaching angle (+exp) Maximum reach distance (-)
Rayegani et al. (2014) RCT (5) N _{start} =46 N _{End} =30 TPS=Chronic	E: OT + EMG + biofeedback E2: OT + neurofeedback C: OT Duration: 40 min/d, 5d/wk for 2wk	Jebsen Taylor Hand Test (-)
Armagan et al.(2003) RCT (7) N _{start} =27 N _{end} =27 TPS=Subacute	E: EMG/Biofeedback Therapy C: Sham EMG/biofeedback Duration: 45 min/d, 2d/wk for 5wk	 Active range of motion (+exp) Changes in EMG surface potentials (+exp) Brunnstrom stages (-) Complex movement (-)
<u>Crow et al</u> . (1989) RCT (8) N _{start} =40 N _{end} =40 TPS=Subacute	E: EMG/Biofeedback Therapy C: Sham EMG/biofeedback Duration: <i>Not Specified</i>	Action Research Arm test (+exp)
Basmajian et al. (1987) RCT (6) N _{start} =30 N _{end} =29 TPS=Chronic	E: EMG/Biofeedback Therapy C: Physical Therapy using neuro- facilitatory Duration: <i>Not Specified</i>	 Upper extremity function test (-) Finger Oscillation test (-)
Inglis et al. (1984) RCT (5) N _{start} =30 TPS=Chronic	E: EMG/Biofeedback+ physiotherapy C: Physiotherapy Duration: <i>Not Specified</i>	 Active range of motion (+exp) Brunnstrom (+exp) Muscle strength (+exp)
Basmajian et al.(1982) RCT (6) N _{start} =37 N _{end} =37 TPS=Chronic	E: EMG/Biofeedback Therapy C: Physical Therapy using neuro- physiological approach Duration: <i>Not Specified</i>	 Upper extremity function test (-) Min rate of manipulation test (-) 9-hole peg test (-)

Prevo et al. (1982)	E: EMG/Biofeedback Therapy	Proximal and distal agonists (-)
RCT (3)	C: Conventional Therapy	
N=28	Duration: 30 min/d, 2d/wk for 6wk	
Greenberg & Fowler (1980)	E: EMG/Biofeedback Therapy	 Active elbow extension (-)
RCT (5)	C: Conventional Occupational Therapy	
N=20	Duration: Not Specified	
Hurd et al. (1980)	E: Actual myofeedback	 Active range of motion (-)
RCT (6)	C: Simulated myofeedback	 Muscle activity (-)
N=24	Duration: Not Specified	
TPS=Chronic		
Mroczek et al. (1978)	E: EMG biofeedback	 Range of Motion (-)
RCT (5)	C: Physical therapy	
N=9	Duration: Not Specified	
Lee et al. (1976)	E1: True myofeedback	 Peak amplitude (-)
RCT (4)	E2: Placebo myofeedback	
N=18	C: No myofeedback with conventional	
TPS=Acute	training.	
	Duration: Not Specified	
EMG b	ofeedback combined with additional inte	erventions
Cordo et al. (2013)	E1: AMES robot + torque biofeedback	 Fugl Meyer Score (-)
RCT (6)	E2: AMES robot + EMG biofeedback	 Flexion torque strength (+exp)
N _{Start} =46	Duration: 30 min/d, 3d/wk for 10 wk	 Extension strength (-)
N _{End} =43		 Box and Block Test (-)
TPS=Chronic		 Stroke Impact Scale (-)
Hemmen & Seelen (2007)	E: EMG biofeedback + mental practice	Fugl-Meyer Score (-)
RCT (7)	C: Conventional electrostimulation	 Action Research Arm test (-)
N _{start} =27	Duration: 30 min/d, 5d/wk for 3 mo	
N _{end} =27		
TPS=Subacute		
Abbreviations and table material control many		

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α=0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group - indicates no statistically significant between groups differences at α =0.05

Conclusions about EMG biofeedback

MOTOR FUNCTION				
LoE	LoE Conclusion Statement RCTs Reference			
1a	EMG biofeedback may not have a difference in efficacy when compared to sham feedback or conventional therapy for improving motor function.	8	Thielbar et al. 2017; Kim et al. 2017; Garrido- Montenegro et al. 2016; Chang-Yong et al. 2015; Rayegani et al. 2014; Crow et al. 1989; Basmajian et al. 1987; Basmajian et al. 1982	
1b	EMG biofeedback combined with arm robotics may not have a difference in efficacy when compared to torque biofeedback combined with arm robotics for improving motor function.	1	Cordo et al. 2013	
1b	EMG biofeedback combined with mental practice may not have a difference in efficacy when compared to conventional electrostimulation for improving motor function.	1	Hemmen and Seelen, 2007	

DEXTERITY			
LoE	Conclusion Statement	RCTs	References
1b	EMG biofeedback may not have a difference in efficacy when compared to sham feedback or conventional therapy for improving dexterity.	1	Basmajian et al. 1982
1b	EMG biofeedback combined with arm robotics may not have a difference in efficacy when compared to torque biofeedback combined with arm robotics for improving dexterity.	1	Cordo et al. 2013

SPASTICITY				
LoE	Conclusion Statement	RCTs	References	
2	EMG biofeedback may not have a difference in efficacy compared to sham feedback or conventional therapy for improving spasticity.	2	Prevo et al.1982; Greenberg and Fowler, 1980	

RANGE OF MOTION				
LoE Conclusion Statement RCTs Reference				
1b	There is conflicting evidence about the effect of EMG biofeedback to improve range of motion when compared to sham feedback or conventional therapy .	4	Armagan et al. 2003; Inglis et al. 1984; Greenberg and Fowler, 1980; Mroczek et al. 1978	

STROKE SEVERITY				
LoE	Conclusion Statement	RCTs	References	
1b	There is conflicting evidence about the effect of EMG biofeedback to improve performance on measures of stroke severity when compared to sham feedback or conventional therapy .	2	Armagan et al. 2003; Inglis et al. 1984	

ACTIVITIES OF DAILY LIVING				
LoE	LOE Conclusion Statement RCTs Reference			
1a	There is conflicting evidence about the effect of EMG biofeedback to improve performance of activities of daily living when compared to sham feedback or conventional therapy .	3	Thielbar et al. 2017; Kim et al. 2017; Garrido-Montenegro et al. 2016	
1b	EMG biofeedback combined with arm robotics may not have a difference in efficacy when compared to torque biofeedback combined with arm robotics to improve performance of activities of daily living.	1	Cordo et al. 2013	

MUSCLE STRENGTH			
LoE	Conclusion Statement	RCTs	References
1b	There is conflicting evidence about the effect of EMG biofeedback to improve muscle strength when compared to sham feedback or conventional therapy .	2	Thielbar et al. 2017; Inglis et al. 1984
1b	There is conflicting evidence about the effect of torque biofeedback combined with arm robotics to improve muscle strength when compared to EMG biofeedback combined with arm robotics .	1	Cordo et al. 2013

Key points

The literature is mixed regarding EMG biofeedback alone for upper limb rehabilitation following stroke, however it may not be beneficial when combined with other therapy approaches.

Sensorimotor stimulation Neuromuscular electrical stimulation (NMES)



Adopted from: http://fescenter.org/patient-resources/current-clinical-trials/stroke-programs/hand-function-control-2/hand-function-control/

Neuromuscular electrical stimulation (NMES) is a technique used to generate muscle contractions in regions affected by hemiparesis by stimulating lower motor neurons involved in muscle movement through transcutaneous application of electrical currents (Monte-Silva et al. 2019; Allen & Goodman 2014). Three forms of NMES are available:

- 1. Cyclic NMES in which a muscle is repetitively stimulated at near maximum contraction on a pre-set schedule and patient participation is passive (Nascimento et al. 2013);
- Electromyography (EMG) triggered NMES, a target muscle is directly controlled or triggered by volitional EMG activity from the target or a different muscle to elicit a desired stimulation (Monte-Silva at al. 2019);
- 3. Functional electrical stimulation (FES), which refers to the application of NMES to assist voluntary during a functional task (Eraifej et al. 2017).

A total of 67 unique RCTs were found for using NMES to enhance upper extremity motor rehabilitation.

Interventions in 11 RCTs were cyclic NMES compared to sham stimulation or conventional rehabilitation (Tilkici et al. 2017; Baygutalp et al. 2014; De Jong et al. 2013; Malhotra et al. 2013; Sahin et al. 2012; Lin and Yan, 2011; Mann et al. 2005; Powell et al. 1999; Chae et al. 1998; King et al. 1996; Faghri et al. 1994). RCTs also looked at the combination of cyclic NMES with: robotics (Barker et al. 2017; Miyasaka et al. 2016; Lee et al. 2015; Hayward et al. 2013), and repetitive task training (Gharib et al. 2014).

10 RCTs looked at EMG-triggered NMES to sham stimulation or conventional rehabilitation (Park et al. 2017; Kwakkel et al. 2016; Dorsch et al. 2014; Bhatt et al. 2007; Gabr et al. 2005; Kimberley et al. 2004; Cauraugh and Kim, 2003; Cauraugh et al. 2000; Francisco et al. 1998; Bowman et al. 1979). RCTs looked at the combination of EMG-triggered NMES with: robotics (Qian et al. 2017; Hu et al. 2015; Barker et al. 2008), mirror therapy (Schick et al. 2017; Kojima et al. 2014), or a splint (Shindo et al. 2011).

14 RCTs looked at the effects of FES compared to sham stimulation or conventional rehabilitation (Demir et al. 2018; Pan et al. 2018; Carda et al. 2017; Marquez-Chin et al. 2017; Yuzer et al. 2017; Shimodozono et al. 2014; Karakus et al. 2013; Mangold et al. 2009; Hara et al. 2008; Thrasher et al. 2008; Hara et al. 2006; Ring and Rosenthal, 2005; Popovic et al. 2003; Faghri and Rodgers, 1997). RCTs looked at the combination of FES with: mirror therapy (Mathieson et al. 2018; Kim et al. 2015), botulinum toxin (Weber et al. 2010), action observation paired with brain computer interface (Kim et al. 2016), bilateral arm training (Chan et al. 2009), and task-oriented therapy (Jonsdottir et al. 2017).

Nine RCTs looked at the effect of different NMES techniques compared to each other (Jeon et al. 2017; Knutson et al. 2016; Wilson et al. 2016; Boyaci et al. 2013; You et al. 2013; Knutson et al. 2012; Chae et al. 2009; De Kroon and Ijzerman, 2008; Hemmen and Seelen, 2007)

Three RCTs looked at differing intensity of NMES (Page et al. 2012; Hsu et al. 2010; Kowalczewski et al. 2007), high versus low frequency cyclic NMES (Doucet and Griffin, 2013), and early versus delayed FES (Popovic et al. 2004).

The methodological details and results of all 67 RCTs are presented in table 22.

Table 22. Summary of RCTS evaluating NMES for upper extremity motor renabilitation				
Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)		
	Cyclic NMES versus conv	entional therapy		
Tilkici et al. (2017) RCT (6) N _{Start} =40 N _{End} =40 TPS=Chronic	E: Neuromuscular Electrical Stimulation C: Conventional Therapy Duration: 30min/d, 5d/wk for 3wk	 Modified Ashworth Scale (+exp) Wrist Extension (+exp) Brunnstrom's Recovery Stages (-) Modified Ashworth Scale (-) Fugl-Meyer Assessment (-) Duruoz Hand Index (-) Functional Independence Measure (-) 		
Baygutalp et al. (2014) RCT (5) N _{start} =30 N _{End} =30 TPS=Chronic	E: NMES + conventional therapy C: Conventional therapy Duration: 60min/d, 5d/wk for 3wk	 Modified Ashworth Scale (-) Barthel Index (-) Brunnstrom's Recovery Stages (-) 		
De Jong et al. (2013) RCT (8) N _{start} =46 N _{end} =46 TPS=Subacute	E: Arm stretch positioning + NMES C: Sham stretch positioning + Sham NMES Duration: 45 min (2x/d), 5d/wk, for 8 wk	Modified Ashworth Scale (-)		
Malhotra et al. (2013) RCT (5) N _{Start} =90 N _{End} =65 TPS=Acute	E: NMES C: Conventional therapy Duration: 30 min (2x/d), 5d/wk for 6 wk	Passive Range of Motion (-)		
Sahin et al. (2012) RCT (5) N _{start} =42 N _{end} =38 TPS=Chronic	E: Stretching + NMES C: Stretching Duration: 5d/wk for 4wk	Modified Ashworth Scale (+exp)		
Lin & Yan (2011) RCT (6) N _{stat} =46 N _{end} =37 TPS=Acute	E: Cyclic NMES + standard rehabilitation C: Standard rehabilitation Duration: 30 min/d, 5d/wk for 3 wk	 Fugl-Meyer Assessment (+exp) Barthel Index (+exp) 		
Mann et al. (2005) 5 (RCT) N _{start} =22 N _{end} =22 TPS=Chronic	E: Neuromuscular Electrical Stimulation C: Passive Extension Exercises Duration: 10-30min (2x per day) for 12wk	Action Research Arm Test (+exp)		
Powell et al. (1999) RCT (7) N _{start} =60 N _{end} =48 TPS=Subacute	E: Cyclic electrical stimulation + standard rehabilitation C: Standard rehabilitation Duration: 30 min (3x per day), 3d/wk for 8 wk	 Action Research Arm test (+exp) 		
Chae et al. (1998) RCT (6) N _{start} =46 N _{end} =28 TPS=Subacute	E: Cyclic NMES C: Sham stimulation + routine rehabilitation Duration: 1 hr/d, 5d/wk for 3 wk	 Fugl-Meyer Assessment (+exp) 		
King et al. (1996) RCT (4) N _{start} =21 N _{end} =NR TPS=Chronic	E: NMES C: Passive stretch Duration: <i>Not reported</i>	Tone reduction (+exp)		

Table 22. Summary of RCTs evaluating NMES for upper extremity motor rehabilitation

Faghri et al. (1994) RCT (4) N _{start} =26 N _{end} =NR TPS=NR	E: Cyclic NMES + conventional therapy C: Conventional Therapy Duration: 1.5-6h/d for 6wk	Arm tone (+exp)
11 0-111	Cyclic NMES combined w	ith robotics
Barker et al. (2017) RCT (7) N _{Start} =50 N _{End} =38 TPS=Subacute	E1: SMART Arm Training + Outcome- Triggered Electrical Stimulation + Conventional Therapy E2: SMART Arm Training + Conventional Therapy C: Conventional Therapy Duration: 60min/d, 5d/wk for 4wk	E1 vs E2 vs C • Motor Assessment Scale (-) • Modified Ashworth Scale (-) • Triceps Strength (-)
<u>Miyasaka et al.</u> (2016) RCT (5) N _{Start} =30 N _{End} =30 TPS=Subacute	E: NMES + robotic training C: Robotic training Duration: 1 hr/d, 5d/wk for 2 wk	 Fugl-Meyer Assessment (-) Range of Motion (-)
Lee et al. (2015) RCT (8) N _{start} =39 N _{End} =39 TPS=Chronic	E: NMES + robotic therapy C: Sham NMES + robotic therapy Duration: 90-100min/d, 5d/wk for 4wk	 Modified Ashworth Scale (+exp) Wolf Motor Function Test (+exp) Stroke Impairment Scale (+exp) Fugl-Meyer Assessment (-) Motor Activity Log (-)
Hayward et al. (2013) RCT (6) N _{start} =8 N _{end} =8 TPS=Acute	E: SensoriMotor Active Rehabilitation Training (SMART) with outcome trigger electrical stimulation (OT-stim) C: SensoriMotor Active Rehabilitation Training (SMART) Duration: 1 hr/d, 5d/wk for 4 wk	 Motor Assessment Scale (-) Upper Arm Function (-)
	Cyclic NMES with repetitiv	ve task training
Gharib et al. (2014) RCT (9) N _{Start} =40 N _{End} =40 TPS=Chronic	E: Cyclic NMES (20Hz) + repetitive task training C: Sham electrical stimulation + repetitive task practice Duration: 1 hr/d, 4d/wk for 8 wk	 Modified Ashworth Scale (+exp) Jebsen Taylor Hand Function Test (+exp) Range of Motion (+exp)
	EMG-triggered NMES compare	d to sham stimulation
Park et al. 2017 RCT (2) N _{Start} =40 N _{End} =32 TPS=NR	E: Mental Practice combined with Electromyography-Triggered Electrical Stimulation C: Conventional Rehabilitation Program Duration: 30min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Motor Activity Log (+exp)
Kwakkel et al. (2016) RCT (7) N _{Start} =159 N _{End} =159 TPS=Acute	E1: EMG-NMES (unfavourable prognosis) E2: Modified constraint-induced movement therapy (favourable prognosis) C1: Unfavourable prognosis based on preservation or return of voluntary finger extension early after stroke (received usual care) C2: Favourable prognosis based on preservation or return of voluntary finger extension early after stroke (received usual care) Duration: 1 hr/d, 3d/wk for 3 wk	E1 vs C1 Action Research Arm Test: (-) Fugl-Meyer Assessment: (-) Motricity Index: (-) Stroke Impact Scale: (-) Wolf Motor Function Test: (-) Motor Activity Log: (-) E2 vs C2 Action Research Arm Test: (+exp ₂) Fugl-Meyer Assessment: (-) Motricity Index: (-) Stroke Impact Scale: (+exp ₂) Wolf Motor Function Test: (-) Motor Activity Log: (-)
Dorsch et al. (2014) RCT (7) N _{Start} =33 N _{End} =30	E: EMG-triggered NMES C: Usual therapy Duration: 30 min/d, 6d/wk for 8wk	 Modified Ashworth Scale (-) Manual Muscle Test (-)

TPS=Acute		
Bhatt et al. (2007)	E1: EMG-triggered NMES	E1 vs E2 vs E3
RCT (3)	E2: Tracking training	Jebson Taylor Hand Function Test (-)
N _{start} =20	E3: EMG-triggered NMES + tracking	Box & Block Test (-)
N _{end} =18	training	
TPS=Chronic	Duration: 1 hr/d, 5d/wk, for 2 wk	
Gabr et al. (2005)	E: EMG-triggered NMES	Fual Mever Score (+exp)
RCT (4)	C: Home exercise	Action Research Arm Test (-)
N _{start} =12	Duration: 45 min/d. 3d/wk for 4 wk	
Nend=12		
TPS=Chronic		
Kimberley et al. (2004)	E: EMG-triggered NMES	Box & Block test (+exp)
RCT (7)	C: Sham	Motor Activity Log (+exp)
N _{start} =16	Duration: 3hr/d 5d/wk for 3 wk	 Jebsen Taylor Hand Function Test (+exp)
Nend= 16		
TPS=Chronic		
Cauraugh and Kim (2003)	E1: EMG-triggered NMES + blocked	F1/F2 vs C
BCT (5)	practice	• Box and Block Test (+exp ₁ +exp ₂)
Notort=34	E2: EMG-triggered NMES + random	 Sustained wrist/finger contraction (+exp1 +exp2)
Nand-31	practice	F1 vs F2
TPS-Chronic	C: Conventional therapy	Box and Block Test (-)
	Duration : 00 min/d 2d/wk (24br in	 Bustained wrist/finger contraction (-)
	between) for 2 wk	
Cauraudh et al. (2000)	E: EMG-triggered NMES + passive	Box and Block test (+exp)
	range of motion L stratching evercises	• Motor Accossment scale ()
	C: Dessive range of motion + stretching	• Wold Assessment scale (-)
		• Fugi-meyer upper extremity (-)
TPS Chronic	Durotion 20 min/d 4d/uk for 2 wk	
Francisco et al. (1998)	E: EMG-triggered NMES + standard	• Fugl-Meyer Assessment (+exp)
RUT (3)	therapy	• Functional Independence Measure (+exp)
Nstart=9	C: Conventional Therapy	
Nend=9	Duration: 30 min (2x per day), 5d/wk for	
IPS=Acute	4 WK	
Heckman et al. (1997)	E: EMG-triggered ES + standard	Hand extension (+exp)
RCT(4)	therapy	Muscle tone (+exp)
N _{start} =28	C: Standard therapy	
N _{end} =28	Duration: 5d/wk for 4wk	
IPS=Subacute		
Bowman et al. (1979)	E: Conventional therapy + positional	Range of motion (+exp)
RCT (3)	teedback electrical stimulation therapy	
Nstart=30	C: Conventional Therapy	
	Duration: 30min (2x per day), 5d/wk for	
IPS=NK	4WK	
	EMG-triggered NMES comb	ined with robotics
Qian et al. (2017)	E: Electromyography-Driven	Fugl-Meyer Assessment (+exp)
RCT (6)	Neuromuscular Electrical Stimulation-	Modified Ashworth Scale (+exp)
N _{Start} =24	Robot Arm	Action Research Arm Test (-)
NEnd =24	C: Conventional Therapy	Functional Independence Measure (-)
TPS=Acute-Subacute	Duration: 40min, 5d/wk for 4wk	
Hu et al. (2015)	E: EMG-driven NMES robot	 Fugl-Meyer Assessment (+exp)
RCT (6)	C: EMG-driven robot	Action Research Arm Test (+exp)
N _{Start} =26	Duration: 30 min/d, 4d/wk for 5 wk	Modified Ashworth Scale (-)
N _{End} =26		
TPS=Chronic		
Barker et al. (2008)	E1: SMART Arm + EMG-triggered	E1/E2 vs C
RCT (7)	NMES	Modified Ashworth Scale: (+exp ₁ , +exp ₂)
N _{start} =33	E2: SMART Arm	
N _{end} =30	C: Conventional therapy	

TPS=Chronic	Duration: 1 hr/d, 3d/wk for 4 wk	
	EMG-triggered NMES wit	h mirror therapy
<u>Schick et al.</u> (2017) RCT (7) N _{Start} =33 N _{End} =32 TPS=Subacute	E: Bilateral Electromyography- Neuromuscular Electrical Stimulation with Mirror Therapy C: Electromyography-Neuromuscular Electrical Stimulation	 Fugl-Meyer Assessment (-) Rivermead Assessment of Somatosensory Performance (-) Box and Block Test (-) Barthel Index (-)
Kojima et al. (2014) RCT crossover (7) N _{Start} =13 N _{End} =13 TPS=Subacute	E: Mirror therapy + EMG-triggered NMES first C: Mirror therapy + EMG-triggered NMES delayed Duration: 30 min/d, 4d/wk for 8 wk	 Fugl-Meyer Assessment (+exp) Hand range of Motion (+exp)
	EMG-triggered NMES	with splint
Shindo et al. (2011) RCT (6) N _{start} =24 N _{end} =20 TPS=Subacute	E: EMG-triggered NMES + splint C: Splint Duration: 45 min/d, 3d/wk for 3 wk	 Fugl-Meyer Assessment (+exp) Motor Activity Log (-) Action Research Arm Test (+exp)
	FES versus convention	onal therapy
Demir et al. 2018 RCT (4) Nstart =29 N _{End} =17 TPS=Chronic	E: Functional Electrical Stimulation and Conventional Physiotherapy C: Conventional Physiotherapy Duration: 15-45min (2x per day), 5d/wk for 8wk	 Fugl-Meyer Assessment (-) Modified Ashworth Scale (-) Motor Activity Log-28 (-) Jebsen-Taylor Hand Function Test (-)
Pan et al. 2018 RCT (6) N _{Start} =12 N _{End} =12 TPS=Subacute	E: Fuinctional Electrical Stimulation C: Sham Electrical Stimulation Duration: 40min/d, 2d/wk for 8wk	Fugl-Meyer Assessment (-)
Carda et al. (2017) RCT-Crossover (7) Nstart =11 NEnd =11 TPS=Chronic 1	E: Functional Electrical Stimulation C: Conventional Therapy Duration: 90min/d, 5d/wk for 2wk	 Fugl-Meyer Assessment (+exp) Motor Activity Log (+exp) Wolf Motor Function Test (-) Resistance to Passive Movement Scale (-)
Marquez-Chin et al. (2017) RCT (7) Secondary Analysis N _{start} =21 N _{End} =21 TPS=Subacute	E: Functional Electrical Stimulation C: Conventional Therapy Duration: 1h/d, 5d/wk for 8wk	 Functional Independence Measure (+exp) Fugl-Meyer Assessment (+exp)
Yuzer at al. 2017 RCT (6) Nstart =30 NEnd =30 TPS=Subacute	E: Functional Electrical Stimulation and Conventional Therapy C: Conventional Therapy Duration: 30min/d, 5d/wk for 4wk	 Barthel Index (+exp) Brunnstrom Stages (-) Upper Extremity Performance Test (-)
Shimodozono et al. (2014) RCT (8) N _{Start} =27 N _{End} =24 TPS= Subacute	E1: Continuous NMES + repetitive facilitative exercise E2 Repetitive facilitative exercise C: Conventional therapy Duration: 40 min/d, 5d/wk for 4 wk	 Fugl-Meyer Assessment (+exp₂) Elbow extension (+exp₂) Shoulder flexion (-) Wrist flexion (-)
Karakus et al. (2013) RCT (8) N _{Start} =28 N _{End} =28 TPS= Subacute	E: FES + standard rehabilitation C: Standard rehabilitation Duration: 30min/d, 5d/wk for 2wk	 Brunnstrom recovery stages (+exp) Motricity Index (+exp) Modified Ashworth Scale (-)
<u>Mangold et al</u> . (2009) RCT (5) N _{start} =23	E: FES C: Conventional therapy Duration: 1 hr/d, 3d/wk for 4 wk	 Barthel Index (-) Chedoke McMaster Stroke Assessment (-)

N _{end} =23		
Hara et al. (2008) RCT (5) N _{start} =20 N _{end} =20 TPS=Chronic	E: FES C: Conventional therapy Duration: 45 min/d, 6d/wk for 4 wk	 Range of motion (+exp) Modified Ashworth Scale (+exp)
Thrasher et al. (2008) RCT (5) N _{start} =21 N _{end} =19 TPS=Subacute	E: FES + conventional therapy C: Conventional therapy Duration: 30 min/d, 4d/wk for 12 wk	 Rehabilitation Engineering Laboratory Hand Function Test (+exp)
<u>Hara et al</u> . (2006) RCT (4) N _{start} =14 N _{end} =14 TPS=Chronic	E: FES C: Conventional therapy Duration: 1 hr/d, 2d/wk for 4 mo	 Modified Ashworth Scale (-) Range of Motion (+exp)
Ring & Rosenthal (2005) RCT(6) Nstart=22 Nend=NR TPS=Subacute	E: Neuroprosthetic FES C: Conventional therapy Duration: 25 min/d, 3d/wk for 5 wk	 Modified Ashworth Scores (+exp) Box & Block test (+exp) Jebsen Taylor Hand Function test (+exp)
Popovic et al. (2003) RCT (6) N _{start} =28 N _{end} =28 TPS=Subacute	E: FES C: Standard therapy Duration: 30 min/d, 7d/wk for 3 wk	Upper extremity performance test (+exp)
Faghri & Rodgers (1997) RCT (4) N _{start} =26 N _{end} =26 TPS=Acute	E: FES + conventional therapy C: Conventional therapy Duration: 6 hr/d, 6d/wk for 6 wk	 Range of motion (+exp) Shoulder muscle tone (+exp)
	FES combined with addition	onal therapies
Mathieson et al. (2018) RCT (8) N _{Start} =50 N _{End} =47 TPS=Acute	E1: Functional Electrical Stimulation E2: Mirror Therapy E3: Functional Electrical Stimulation with Mirror Therapy Duration: 30min (2x per day), 5d/wk for 3wk	E1 vs E2 • Action Research Arm Test (+exp) • Fugl-Meyer Assessment (+exp) • Nottingham Extended Activities of Daily Living Test (-) • Functional Independence Measure (-)
Jonsdottir et al. 2017 RCT (5) N _{Start} =82 N _{End} =45 TPS=Subacute	E: Myoelectric Continuous Control of Functional Electrical Stimulation Task- Oriented Therapy C: Task Oriented Therapy Duration: 45min/d, 5d/wk for 5-6wk	 Action Research Arm Test (-) Fugl-Meyer Assessment (-) Disability of the Arm, Shoulder, and Hand Questionnaire (-)
Kim et al. (2016) RCT (7) N _{Start} =34 N _{End} =30 TPS=Chronic	E: FES with Action observation training and brain computer interface C: Conventional training Duration: 30min, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Motor Activity Log (+exp) Modified Barthel Index (+exp) Wrist Flexion (+exp)
<u>Kim et al</u> . (2015) RCT (5) N _{Start} =33 N _{End} =29 TPS=Chronic	E1: FES with biofeedback + mirror therapy E2: FES + mirror therapy C: Conventional rehabilitation Duration: 30 min/d, 5d/wk for 4 wk	E1 vs C Functional Independence Measure (+exp) Jebsen Taylor Hand test (+exp) Manual Muscle Test (+exp) Box and Block Test (+exp) Wrist Extension (+exp) Grip strength (-) Modified Ashworth Scale (-) E1 vs E2 Functional Independence Measure (-) Jebsen Taylor Hand test (+exp)

		 Manual Muscle Test (+exp) Box and Block Test (+exp) Wrist Extension (+exp) Grip strength (+exp) Modified Ashworth Scale (-)
Weber et al. (2010) RCT (7) Nstart=23 Nend=23 TPS=Chronic	E: FES + botulinum toxin-A + home based exercise program C: Botulinum toxin-A + home-based exercise program Duration: 1 hr/d, 5d/wk for 4 wk	 Motor Activity Log (-) Action Research Arm Test (-)
Chan et al. (2009) RCT (7) Nstart=20 N _{end} =20 TPS=Chronic	E: Bilateral arm training + FES C: Bilateral arm training + sham FES Duration: 70 min/d, 3d/wk for 5 wk	 Fugl-Meyer Assessment (+exp) Functional test for the Hemiplegic Upper Extremity (+exp) Modified Ashworth Scale (-)
Alon et al. (2007) RCT (5) N _{start} =15 N _{end} =15 TPS=Subacute	E: FES + task specific training C: Task specific training Duration: 30 min(2x/d), 5d/wk for 12 wk	 Box and Block Test (+exp) Jebsen-Taylor light object lift (+exp) Fugl-Meyer Assessment (+exp)
	NMES techniques vers	us each other
<u>Jeon et al.</u> (2017) RCT (5) N _{Start} =20 N _{End} =20 TPS=Subacute	E: EMG-triggered NMES C: FES Duration: 30min, 5d/wk for 4wk	Fugl-Meyer Assessment (-)
Knutson et al. (2016) RCT (5) N _{Start} =80 N _{End} =64 TPS=Chronic	E1: Functional Electrical Stimulation E2: Cyclic NMES Duration: 2hrs, 7d/wk for 6 wk	 Fugl-Meyer Assessment (-) Arm Motor Abilities Test (-) Box and Block Test (+exp)
Wilson et al. (2016) RCT (6) N _{Start} =122 N _{End} =96 TPS=Subacute	E1: Cyclic Neuromuscular Electrical Stimulation E2: Electromyographically-triggered Neuromuscular Electrical Stimulation E3: Sensory Stimulation Duration: 40 min (2x/d), 5d/wk for 8 wk	 Fugl-Meyer Assessment (-) Modified Arm Motor Ability Task (-)
Boyaci et al. (2013) RCT (7) Nstart=31 NEnd=31 TPS=Chronic	E1: EMG-triggered NMES E2: Cyclic NMES C: Control Duration: 45 min/d, 5d/wk for 3 wk	 E1 vs C Fugl-Meyer Assessment (+exp) Motor Activity Log (+exp) Spasticity in wrist flexor (-) Spasticity in finger flexor (-) Range of Motion in active wrist extension (+exp) Range of Motion in active metacarpophalangeal joint extension (+exp) Grip strength (+exp) E2 vs C Fugl-Meyer Assessment (+exp2) Motor Activity Log (-) Spasticity in finger flexor (-) Range of Motion in active wrist extension (+exp2) Spasticity in finger flexor (-) Range of Motion in active wrist extension (+exp2) Range of Motion in active metacarpophalangeal joint extension (-) Grip strength (-) E1 vs E2 Fugl-Meyer Assessment (-) Motor Activity Log (-) Spasticity in wrist flexor (-)

		 Range of Motion in active wrist extension (-) Range of Motion in active metacarpophalangeal joint extension (-)
<u>You et al</u> . (2013) RCT (7) N _{Start} =18 N _{End} =16 TPS=Chronic	E: Mental training + EMG stimulation C: FES Duration: 40 min/d, 2d/wk for 4wk	 Grip strength (-) Range of Motion (-) Modified Ashworth Scale (-) Fugl-Meyer Assessment (+exp) Motor Activity Log (-) Barthel Index (-)
Knutson et al. (2012) RCT (6) N _{start} =21 N _{end} =21 TPS=Subacute	E1: Contralaterally controlled FES E2: Cyclic NMES Duration: 90 min/d, 3d/wk for 4 wk	 Maximum finger extension angle (-) Tracking error (% of AROM) (-) Fugl-Meyer Assessment (-) Box and Block Test (-) Arm Motor Abilities Test Score (-)
Chae et al. (2009) RCT (8) N _{start} =26 N _{end} =26 TPS=Chronic	E1: EMG-triggered NMES E2: Cyclic NMES Duration: 1 hr/d, 7d/wk for 6 wk	Arm Motor Ability Test (-)
De Kroon & Ijzerman (2008) RCT (7) N _{start} =22 N _{end} =22 TPS=Chronic	E1: EMG-triggered NMES E2: Cyclic NMES Duration: 30 min/d, 3d/wk for 6 wk	 Action Research Arm test (-) Grip Strength (-) Fugl-Meyer Score (-) Motricity Index (-)
Hemmen & Seelen (2007) RCT (7) N _{start} =27 N _{end} =27 TPS=Subacute	E1: EMG-triggered NMES E2: Cyclic NMES Duration: 30 min/d, 5d/wk for 3mo	 Fugl-Meyer Assessment (-) Action Research Arm test (-)
	Low versus high intensity N	MES studies
Page et al. (2012) RCT (7) N _{start} =32 N _{end} =32 TPS=Chronic	E1: 30 minutes of electrical stimulation therapy with repetitive task specific practice E2: 60 minutes of electrical stimulation therapy with repetitive task specific practice E3: 120 minutes of electrical stimulation therapy with repetitive task specific practice Duration: 30 min OR 60 min OR 120 min, 5d/wk for 8 wk.	E3 vs. E2/E1 Fugl-Meyer Assessment (+exp ₃) Arm Motor Ability Test (+exp ₃) Action Research Arm Test (+exp ₃)
Hsu et al. (2010) RCT (6) N _{start} =66 N _{end} =66 TPS=Acute	E1: High intensity cyclic NMES (60 min) E2: Low intensity cyclic NMES (30 min) C: No treatment Duration: 30/60 min, 5d/wk for 4 wk	
Kowalczewski et al. (2007) RCT (6) N _{start} =19 N _{end} =18 TPS=Subacute	E1: High intensity FES exercise therapy (60 min) E2: Low intensity FES exercise therapy (15 min) Duration: 15/60 min, 5d/wk for 3 wk	 Wolf Motor Function Test (+exp1) Motor Activity Log (-) Fugl-Meyer Assessment (-)

High versus low frequency cyclic NMES				
Doucet and Griffin (2013)	E1: High frequency cyclic NMES (40Hz)	 Lateral pinch strength (+exp) 		
RCT (5)	E2: Low frequency cyclic NMES (20Hz)	 Minnesota Manual Dexterity Test (+exp) 		
N _{Start} =16	Duration: 1 hr/d, 4d/wk for 4 wk	Endurance of thumb adduction (+exp)		
N _{End} =16				
TPS=Chronic				
	Early versus delayed FES			
Popovic et al. (2004)	E: Early (acute) FES	Upper extremity performance test (+exp)		
RCT (6)	C: Delayed (chronic) FES			
N _{start} =41	Duration: 30 min/d, 7d/wk for 3 wk			
N _{end} =32				
TPS=Acute				

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

 $+exp_2$ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group - indicates no statistically significant between groups differences at α =0.05

Conclusions about NMES

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	Cyclic NMES may produce greater improvements in motor function than sham stimulation or conventional therapy .	7	Tilkici et al. 2017; Baygutalp et al. 2014; Inobe et al. 2013; Lin and Yan 2011; Mann et al. 2005; Powell et al. 1999; Chae et al. 1998
1a	Cyclic NMES combined with arm robotics may not have a difference in efficacy when compared to arm robotics on their own or conventional therapy for improving motor function.	3	Miyasaka et al. 2016; Lee et al. 2015; Hayward et al. 2013
1b	Cyclic NMES combined with repetitive task training may produce greater improvements in motor function than repetitive task training alone .	1	Gharib et al. 2014
1a	EMG-triggered NMES may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improving motor function.	7	Park et al. 2017; Kwakkel et al. 2016; Bhatt et al. 2007; Gabr et al. 2005; Kimberley et al. 2004; Cauraugh et al. 2000; Francisco et al. 1998
1a	EMG-triggered NMES combined with arm robotics may produce greater improvements in motor function than arm robotics on their own or conventional therapy.	2	Qian et al. 2017; Hu et al. 2015
1a	There is conflicting evidence about the effect of EMG- triggered NMES combined with mirror therapy to improve motor function when compared to mirror therapy on its own.	2	Schick et al. 2017; Kojima et al. 2014
1b	EMG-triggered NMES combined with splints may produce greater improvements in motor function than splints on their own.	1	Shindo et al. 2011
1a	There is conflicting evidence about the effect of FES to improve motor function when compared to sham stimulation or conventional therapy .	11	Demir et al. 2018; Pan et al. 2018; Carda et al. 2017; Maquez-Chin et al. 2017; Yuzer et al. 2017; Shimodozono et al. 2014; Karakus et al. 2013; Mangold et al. 2009; Thrasher et al. 2008; Ring and Rosenthal, 2003; Popovic et al. 2003
1b	FES may produce greater improvements in motor function than mirror therapy .	1	Mathieson et al. 2018
2	There is conflicting evidence about the effect of FES combined with task-specific training to improve motor function when compared to task-specific training .	2	Jonsdottir et al. 2017; Alon et al. 2007
2	FES combined with biofeedback and mirror therapy may produce greater improvements in motor function than FES combined with mirror therapy or conventional therapy.	1	Kim et al. 2015
1b	FES combined with botulinum toxin A and a home exercise program may not have a difference in	1	Weber et al. 2010

	efficacy when compared to botulinum toxin A combined with a home exercise program for improving motor function.		
1b	Bilateral arm training combined with FES may produce greater improvements in motor function than bilateral arm training combined with sham FES.	1	Chan et al. 2009
1a	EMG-triggered NMES may not have a difference in efficacy when compared to cyclic NMES for improving motor function.	3	Wilson et al. 2016; Boyaci et al. 2013; De Kroon et al. 2008
2	EMG-triggered NMES may not have a difference in efficacy when compared to FES for improving motor function.	1	Jeon et al. 2013
1b	FES may not have a difference in efficacy when compared to cyclic NMES for improving motor function.	2	Knutson et al. 2016; Knutson et al. 2012

DEXTERITY			
LoE	Conclusion Statement	RCTs	References
1b	EMG-triggered NMES may produce greater improvements in dexterity than sham stimulation or conventional therapy.	4	Bhatt et al. 2007; Kimberley et al. 2004; Cauraugh and Kim 2003; Cauraugh et al. 2000
1b	EMG-triggered NMES combined with mirror therapy may not have a difference in efficacy when compared to mirror therapy on its own for improving dexterity.	1	Schick et al. 2017
1b	FES may produce greater improvements in dexterity than sham stimulation or conventional therapy.	1	Ring and Rosenthal, 2005
2	FES combined with task-specific training may produce greater improvements in dexterity than task- specific training.	1	Alon et al. 2007
2	FES combined with biofeedback and mirror therapy may produce greater improvements in dexterity than FES combined with mirror therapy or conventional therapy.	1	Kim et al. 2015
1b	There is conflicting evidence about the effect of FES to improve dexterity when compared to cyclic NMES .	2	Knutson et al. 2016; Knutson et al. 2012

PROPRIOCEPTION				
LoE	Conclusion Statement	RCTs	References	
1b	EMG-triggered NMES combined with mirror therapy may not have a difference in efficacy when compared to mirror therapy on its own for improving proprioception.	1	Schick et al. 2017	

SPASTICITY				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of cyclic NMES to improve spasticity when compared to sham stimulation or conventional therapy .	6	Tilkici et al. 2017; Baygutalp et al. 2014; De Jong et al. 2013; Sahin et al. 2012; King et al. 1996; Faghri et al. 1994	
1a	There is conflicting evidence about the effect of cyclic NMES combined with arm robotics to improve spasticity when compared to arm robotics on their own or conventional therapy .	2	Barker et al. 2017; Lee et al. 2015	
1b	Cyclic NMES combined with repetitive task training may produce greater improvements in spasticity than repetitive task training alone.	1	Gharib et al. 2014	
2	EMG-triggered NMES may produce greater improvements in spasticity than sham stimulation or conventional therapy.	1	Cauraugh and Kim, 2003	
1a	There is conflicting evidence about the effect of EMG- triggered NMES combined with arm robotics to improve spasticity when compared to arm robotics on their own or conventional therapy.	3	Qian et al. 2017; Hu et al. 2015; Barker et al. 2008	
1a	There is conflicting evidence about the effect of FES to improve spasticity when compared to sham stimulation or conventional therapy .	8	Demir et al. 2018; Carda et al. 2017; Yuzer et al. 2017; Karakus et al. 2013; Hara et al. 2008; Hara et al. 2006; Ring and Rosenthal, 2005; Faghri and Rodgers, 1997	
2	FES combined with biofeedback and mirror therapy may not have a difference in efficacy when compared to FES combined with mirror therapy or conventional therapy for improving spasticity.	1	Kim et al. 2015	
1a	EMG-triggered NMES may not have a difference in efficacy when compared to cyclic NMES for improving spasticity.	1	Boyaci et al. 2013	
RANGE OF MOTION				
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LoE	Conclusion Statement	RCTs	References	
1b	There is conflicting evidence about the effect of cyclic NMES to improve range of motion when compared to sham stimulation or conventional therapy .	2	Tilkici et al. 2017; Malhotra et al. 2013	
2	Cyclic NMES combined with arm robotics may not have a difference in efficacy when compared to arm robotics on their own or conventional therapy for improving range of motion.	1	Miyasaka et al. 2016	
1b	Cyclic NMES combined with repetitive task training may produce greater improvements in range of motion than repetitive task training alone.	1	Gharib et al. 2014	
2	EMG-triggered NMES may produce greater improvements in range of motion than sham stimulation or conventional therapy.	2	Heckman et al. 1997; Bowman et al. 1979	
1b	EMG-triggered NMES combined with mirror therapy may produce greater improvements in range of motion than mirror therapy on its own.	1	Kojima et al. 2014	
1b	There is conflicting evidence about the effect of FES to improve range of motion when compared to sham stimulation or conventional therapy .	4	Shimodozono et al. 2014; Hara et al. 2008; Hara et al. 2006; Faghri and Rodgers, 1997	
1a	EMG-triggered NMES may not have a difference in efficacy when compared to cyclic NMES for improving range of motion.	1	Boyaci et al. 2013	
1b	FES may not have a difference in efficacy when compared to cyclic NMES for improving range of motion.	2	Knutson et al. 2016; Knutson et al. 2012	

ACTIVITIES OF DAILY LIVING			
LoE	Conclusion Statement	RCTs	References
1a	Cyclic NMES may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improving performance of activities of daily living.	3	Tilkici et al. 2017; Baygutalp et al. 2014; Lin and Yan 2011
1a	Cyclic NMES combined with arm robotics may not have a difference in efficacy when compared to arm robotics on their own or conventional therapy for improving performance of activities of daily living.	3	Barker et al. 2017; Lee et al. 2015; Hayward et al. 2013
1a	There is conflicting evidence about the effect of EMG- triggered NMES to improve performance of activities of daily living when compared to sham stimulation or conventional therapy .	5	Kwakkel et al. 2016; Dorsch et al. 2014; Kimberely et al. 2004; Cauraugh et al. 2000; Francisco et al. 1998
1a	EMG-triggered NMES when combined with arm robotics, mirror therapy or splints may not have a difference in efficacy when compared to these	3	Barker et al. 2017; Qian et al. 2017; Schick et al. 2017

	additional interventions on their own for improving performance of activities of daily living.			
1b	EMG-triggered NMES combined with splints may not have a difference in efficacy when compared to splints on their own for improving performance of activities of daily living.	1	Shindo et al. 2011	
1a	There is conflicting evidence about the effect of FES to improve performance of activities of daily living when compared to sham stimulation or conventional5therapy.5		Demir et al. 2018; Carda et al. 2017; Marquez-Chin et al. 2017; Yuzer et al. 2017; Mangold et al. 2009	
1b	FES may not have a difference in efficacy when compared to mirror therapy for improving 1			
2	FES combined with biofeedback and mirror therapy may produce greater improvements in performance of activities of daily living than FES combined with mirror therapy or conventional therapy.	1	Kim et al. 2015	
1b	FES combined with biofeedback and mirror therapy may not have a difference in efficacy when compared to FES combined with mirror therapy or conventional therapy for improving performance of activities of daily living.	1	Kim et al. 2015	
1b	Bilateral arm training combined with FES may not have a difference in efficacy when compared to bilateral arm training combined with sham FES for improving performance of activities of daily living.	1	Chan et al. 2009	
1a	EMG-triggered NMES may not have a difference in efficacy when compared to cyclic NMES for improving performance of activities of daily living.	3	Wilson et al. 2016; Boyaci et al. 2013; Chae et al. 2009	
1b	FES may not have a difference in efficacy when compared to cyclic NMES for improving performance of activities of daily living.	2	Knutson et al. 2016; Knutson et al. 2012	

MUSCLE STRENGTH				
LoE	Conclusion Statement	RCTs	References	
1a	EMG-triggered NMES may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improving muscle strength.	2	Dorsch et al. 2014; Kwakkel et al. 2016	
1b	EMG-triggered NMES combined with arm robotics may not have a difference in efficacy when compared to arm robotics on their own or conventional therapy for improving muscle strength.	1	Barker et al. 2017	
1b	FES may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improving muscle strength.	1	Karakus et al. 2013	
2	There is conflicting evidence about the effect of FES combined with biofeedback and mirror therapy to improve muscle strength when compared to FES combined with mirror therapy or conventional therapy.	1	Kim et al. 2015	
1a	EMG-triggered NMES may not have a difference in efficacy when compared to cyclic NMES for improving muscle strength.	2	Boyaci et al. 2013; De Kroon et al. 2008	

Key points

The literature is mixed regrading cyclic and EMG-triggered neuromuscular electrical stimulation types, as well as functional electrical stimulation, alone or combined with other therapy approaches, for upper limb rehabilitation following stroke.

The various types of neuromuscular electrical stimulation may not be more beneficial compared to one another.

Transcutaneous electrical nerve stimulation (TENS)



Adopted from: http://www.massageprocedures.com/complementary-modalities/tens/

Transcutaneous electrical nerve stimulation (TENS) involves the application of electrical current through surface electrodes on the skin to facilitate activation of nerves (Teoli et al. 2019). Stimulation can be applied at a low frequency (<10Hz) to produce muscle contractions or at a high (>50Hz) frequency primarily used to produce paresthesia without muscle contractions (Teoli et al. 2019). TENS units are often small, portable, battery-operated devices (Teoli et al. 2019). The application of afferent electrical stimulation at the sensory level may help to enhance neuroplasticity of the brain, through increased activation and recruitment of cortical networks involving contralesional primary sensory cortex, supplementary motor area, dorsal premotor cortex, posterior parietal cortex, and secondary sensory cortices (Veldman et al. 2015; Sonde et al. 1998).

A total of 15 RCTs were found that evaluated the use of TENS for upper extremity motor rehabilitation poststroke (Capone et al. 2017; Chuang et al. 2017; Jung et al. 2017; Fleming et al. 2015; dos Santos-Fontes et al. 2013; Kim et al. 2013; Ikuno et al. 2012; Klaliput et al. 2009; Celnik et al. 2007; McDonnell et al. 2007; Wu et al. 2006; Conforto et al. 2002; Sonde et al. 1998; Tekeoglu et al. 1998; Butefisch et al. 1995). Of these one RCT was a multimodal intervention combining TENS with electromyography and bilateral arm traing (Chuang et al. 2017). The rest evaluated TENS compared to sham stimulation, task specific therapy and conventional rehabilitation.

The methodological details and results of all 15 RCTs are presented in Table 23.

Authors (Vear)	Interventions	
Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Duration: Session length, frequency per week for total number of weeks	Result (direction of effect)
Capone et al. (2017) Quasi-RCT (8) N _{Start} =14 N _{End} =12 TPS= Chronic	E: Robot-Assisted Therapy with Transcutaneous Stimulation of Vagus Nerve (tVNS) C: Robot-Assisted Therapy with Sham-tVNS Duration: 1h, 1d/wk for 10d	 Fugl-Meyer Assessment (+exp)
$\frac{Jung et al. 2017}{RCT (7)}$ $N_{Start} = 46$ $N_{End} = 46$ $TPS = Chronic$	E: Transcutaneous Electrical Nerve Stimulation and Task-Related Training C: Sham Transcutaneous Electrical Nerve Stimulation and Task-Related Training Duration: 1h, 5d/wk for 4wk	 Manual muscle test (+exp) Active Range of Motion (+exp) Fugl-Meyer Assessment (+exp)
Fleming et al. (2015) RCT (7) N _{Start} =33 N _{End} =30 TPS=Chronic	E: Active Somatosensory Stimulation C: Sham Somatosensory Stimulation Duration: 30min/d, 3d/wk for 4wk	 Action Research Arm Test (+exp) Fugl-Meyer Assessment (-) Motor Activity Log (-)
dos Santos-Fontes et al. (2013) RCT (8) N _{start} =20 N _{End} =20 TPS=Chronic	E: Peripheral nerve stimulation C: Sham nerve stimulation Duration: 2h/d, 7d/wk for 4wk	 Jebsen Taylor Hand Function Test (+exp)
<u>Kim et al</u> . (2013a) RCT (7) N _{start} =34 N _{End} =30 TPS=Chronic	E: TENS + task related training C: Placebo + Task related training Duration: 30 min, 5d/wk, for 4 wk	 Fugl-Meyer Assessment (+exp) Manual Function Test (+exp) Box and Block Test (+exp) Modified Ashworth Scale (-)
Ikuno et al. (2012) RCT (8) N _{start} =22 N _{end} =22 TPS=Subacute	E: Peripheral sensory nerve stimulation + task-specific therapy C: Task-specific therapy Duration: 6d/wk for 2wk	 Wolf Motor Function Test (-) Box and Block Test (-) Pinch Strength (-) Grip Strength (-)
Klaiput et al. (2009) RCT (8) N _{start} =20 N _{end} =20 TPS=Subacute	E: Peripheral nerve stimulation C: Sham stimulation Duration: 2h session	 Pinch Strength (+exp)
Celnik et al. (2007) RCT (6) N _{start} =9 N _{end} =9 TPS=Chronic	E1: Single session of peripheral nerve stimulation E2: No stimulation C: Asynchronous nerve stimulation Duration: 2h session	E1 vs E2/C • Jebsen-Taylor Hand Function Test (+exp)
McDonnell et al. (2007) RCT (7) N _{start} =20 N _{end} =20 TPS=Subacute	E: Task-specific training with TENS C: Task-specific training without TENS Duration: 1h/d, 3d/wk for 3wk	 Fugl-Meyer Assessment (-) Action Research Arm Test (-) Grip lift task (+exp)
Wu et al. (2006) RCT (6) N _{start} =9	E: Single session of peripheral nerve (somatosensory) stimulation	Jebsen Taylor Hand Function Test (+exp)

Table 23. RCTs evaluating TENS interventions for upper extremity motor rehabilitation

N _{end} =9 TPS=Chronic	C: No stimulation Duration: 2h session			
Conforto et al. (2002) RCT (6) Nstart=8 Nend=8 TPS=Chronic	E: Single session of medial nerve (somatosensory) stimulation C: Sham stimulation Duration: 2h session	Pinch muscle strength (+exp)		
Sonde et al. (1998) RCT (5) Nstart=44 N _{end} =44 TPS=Chronic	E: TENS + physiotherapy C: Physiotherapy Duration: 60min/d, 5d/wk for 12wk	 Fugl-Meyer Assessment (+exp) Barthel Index (-) 		
Tekeoglu et al. (1998) RCT (6) N _{start} =60 N _{end} =60 TPS=Subacute	E: Rehabilitation + TENS C: Rehabilitation Duration: 30min/d, 5d/wk for 8wk	Barthel Index (+exp)		
Bütefisch et al. (1995) RCT (3) N _{start} =27 N _{end} =24 TPS=Subacute	E: Enhanced specific therapy + TENS C: Enhanced non-specific therapy Duration: 15min (2x per day) for 2wk	Grip strength (-)		
EMG-triggered NMES with BAT versus EMG-TENS with BAT				
Chuang et al. (2017) RCT (7) Nstart=38 NEnd=38 TPS=Chronic	E: Electromyography-Neuromuscular Electric Stimulation with Bilateral Arm Training C: Electromyography-Transcutaneous Electrical Nerve Stimulation with Bilateral Arm Training Duration: 40min, 3d/wk for 4wk	Fugl-Meyer Assessment (-)		

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about TENS

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	TENS may produce greater improvements in motor function than sham stimulation , task-specific therapy or conventional therapy .	10	Capone et al. 2017; Jung et al. 2017; Fleming et al. 2015; dos Santos-Fontes et al. 2013; Kim et al. 2013; Ikuno et al. 2012; Celnik et al. 2007; McDonnell et al. 2007; Wu et al. 2006; Sonde et al. 1998	
1b	TENS combined with EMG and bilateral training may not have a difference in efficacy when compared to EMG-triggered NMES and bilateral training for improving motor function.	1	Chuang et al. 2017	

MUSCLE STRENGTH				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of TENS to improve muscle strength when compared to sham stimulation , task-specific therapy or conventional therapy .	5	Jung et al. 2017; Ikuno et al. 2012; Klaliput et al. 2009; Conforto et al. 2002; Butefisch et al. 1995	

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of TENS to improve dexterity when compared to sham stimulation and task-specific therapy .	2	Kim et al. 2013; Ikuno et al. 2012	

ACTIVITIES OF DAILY LIVING				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of TENS to improve performance of activities of daily living when compared to sham stimulation , task-specific therapy or conventional therapy .	3	Fleming et al. 2015; Sonde et al. 1998; Tekeoglu et al. 1998	

Key points

Transcutaneous electrical nerve stimulation may be beneficial for some aspects of upper limb function following stroke.

Thermal stimulation



Adopted from: https://beautisecrets.com/paraffin-waxtreatment

Thermal stimulation is another method used to facilitate sensorimotor function, thermal stimulation applied either in a noxious or innocuous form have different effects on sensory receptors in the body (Lin et al. 2017). The perception of pain from nociceptors produced by noxious heat (>43°C) and cold (<8°C) activates brain regions such as the second somatosensory cortex, posterior insular cortex and the premotor area that would not be activated by warm and cold receptors from innocuous heat (40-43°C) and cold (20-28°C) temperatures (Lin et al. 2017). Innocuous thermal stimulation has also been found to induce greater corticomotor excitability, and as such has been suggested to influence cortical reorganization and neuroplasticity (Lin et al. 2017).

A total of 4 RCTs were found that evaluated the use of thermal stimulation for upper extremity motor rehabilitation poststroke (Lin et al. 2017; Wang et al. 2017; Wu et al. 2010; Chen et al. 2005). Noxious thermal stimulation was used in 3 RCTs with comparator groups including innocuous thermal stimulation (Lin et al. 2017), thermal stimulation on the lower extremities (Wu et al. 2010a), and conventional rehabilitation (Chen et al. 2005). Innocuous thermal stimulation through paraffin wax compared to a placebo wax was used in a single study (Wang et al. 2017).

The methodological details and results of all 4 RCTs are presented in Table 24.

rehabilitation	5	-	
Authors (Year)	Interventions		Outcome Measures
Study Design (PEDro Score)	tudy Design (PEDro Score) Duration: Session length, frequency per		Result (direction of effect)
Sample Sizestart	week for total number of weeks		
Sample Sizeend			
Time post stroke category			
Noxious versus innocuous	thermal stimulation, lower extremity ther	nal	stimulation and conventional rehabilitation
Lin et al. (2017)	E: Noxious thermal stimulation	•	Fugl-Meyer Assessment (-)
RCT (7)	(Heat: 46-47°C; cold: 7-8°C)	•	Action Research Arm Test (-)
N _{Start} =79	C: Innocuous thermal stimulation	•	Motricity Index (-)
N _{End} =61	(Heat: 40-41°C; cold: 20-21°C)	•	Barthel Index (-)
TPS= Acute	Duration: 30min/d, for a total of 20-24	•	Modified Ashworth Scale (-)

•

•

(+exp)

• Grasping (-)

Stroke Rehabilitation Assessment of Movement

Action Research Arm Test (+exp)

• Modified Ashworth Scale (-)

Brunnstrom Recovery Stages (+exp)

Table 24, RCTs evaluating thermal stimulation interventions for upper extremity motor

N _{End} =29 TPS=Acute			
	Innocuous thermal stimulation	ı ve	rsus placebo
Wang et al. (2017)	E: Paraffin wax thermal stimulation	•	Modified Ashworth Scale (+exp)
RCT (8)	(Heat: 40-42°C)	•	Brunnstrom Recovery Stages (-)
N _{Start} =52	C: Placebo paraffin thermal stimulation		
N _{End} =52			
TPS= Subacute			

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group

+exp2 indicates a statistically significant between groups difference at α=0.05 in favour of the second experimental group

E: Thermal stimulation on upper extremity

C: Thermal stimulation on lower extremity

(Heat: 46-47°C; cold: 7-8°C)

(Heat: 46-47°C; cold: 7-8°C)

(Heat: 46-47°C; cold: 7-8°C)

C: Conventional rehabilitation

E: Thermal stimulation

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

sessions

<u>Wu et al</u>. (2010a)

TPS=Subacute

Chen et al. (2005)

RCT (6)

Nstart=23

Nend=23

RCT (7)

Nstart=46

Conclusions about thermal stimulation

MOTOR FUNCTION					
LoE	Conclusion Statement	RCTs	References		
1a	There is conflicting evidence about the effect of Noxious thermal stimulation to improve motor function when compared to innocuous thermal stimulation, thermal stimulation on the lower extremities and conventional rehabilitation.	3	Lin et al. 2017; Wu et al. 2010; Chen et al. 2005		

MUSCLE STRENGTH					
LoE	Conclusion Statement	RCTs	References		
1b	Noxious thermal stimulation may not have a difference in efficacy when compared to innocuous thermal stimulation for improving muscle strength.	1	Lin et al. 2017		

SPASTICITY

LoE	Conclusion Statement	RCTs	References		
1a	Noxious thermal stimulation may not have a difference in efficacy when compared to innocuous thermal stimulation, and conventional rehabilitation for improving spasticity.	2	Lin et al. 2017; Chen et al. 2005		
1b	Innocuous thermal stimulation may produce greater improvements on spasticity than placebo .	1	Wang et al. 2017		

Key points

Noxious thermal stimulation may not be beneficial for upper limb rehabilitation following stroke, whereas innocuous thermal stimulation may improve some aspects of upper limb function.

Muscle vibration



Adopted from: <u>https://www.humanlocomotion.org/products/focal-vibration-motors</u>

Various forms of muscle vibration applications exist including: focal muscle vibration, whole body vibration, and stochastic resonance stimulation. Whole body vibration involves standing, sitting, or performing various tasks/movements on a vibration platform with the purpose of improving muscle strength and function (Liao et al. 2015; Park et al. 2018). Focal muscle vibration is a new therapeutic approach that involves the application of low-amplitude/high-frequency vibratory stimulation to a specific muscle through small portable devices (Celletti et al. 2017). Lastly, stochastic resonance stimulation involves the application of electrical or mechanical vibration below the sensory threshold to lower the threshold of sensation of the tactile and proprioceptive systems (Stein et al. 2010).

A total of 9 RCTs were found that evaluated the use of muscle vibration therapies for upper extremity motor rehabilitation poststroke (Calabro et al. 2017; Costantino et al. 2017; Jung-Sun et al. 2016; Paoloni et al. 2014; Tavernese et al. 2013; Caliandro et al. 2012; Stein et al. 2010).

The methodological details and results of all 9 RCTs are presented in Table 25.

Table 25. RCTs evaluating muscle vibration interventions for upper extremity mot	tor
rehabilitation	

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
	Vibration Therapy	
Calabro et al. (2017) RCT (7) N _{Start} =20 N _{End} =19 TPS=Subacute-Chronic	E: Focal Muscle Vibration C: Sham Muscle Vibration Duration: 30min/d, 5d/wk for 6wk	 Modified Ashworth Scale (+exp) Functional Independence Measure (+exp) Fugl-Meyer Assessment (+exp)
Costantino et al. (2017) RCT (7) N _{Start} =32 N _{End} =32 TPS=Chronic	E: 300 Hz vibrations on the upper limbs C: Sham vibrations Duration: 30min/d, 3d/wk for 4wk	 Hand Grip Strength (+exp) Modified Ashworth Scale (+exp) Disabilities of the Arm, Shoulder and Hand Score (+exp) Functional Independence Measure (+exp) Fugl-Meyer Assessment (+exp) Jebsen Taylor Hand Function Test (+exp)
Lee et al. (2016) RCT (6) N _{Start} =45 N _{End} =45 TPS=Chronic	E1: Whole-body vibration and task- related training E2: Whole-body vibration C: Conventional Therapy Duration: 30min/d, 3d/wk for 4wk	 E1/E2 vs C Fugl-Meyer Assessment (+exp, +exp₂) Grip Strength (+exp, +exp₂) E1 vs E2 Grip Strength (+exp, +exp₂) E1 vs E2/C Wolf Motor Function Test (+exp) Modified Ashworth Scale (+exp)
Paoloni et al. (2014) RCT (8) N _{Start} =22 N _{End} =22 TPS=Chronic	E: Segmental muscle vibration + conventional therapy C: Conventional therapy Duration: 30min/d, 5d/wk for 2wk	 Muscle modulation of anterior deltoid (+exp) Muscle modulation of biceps brachii (+exp)
Tavernese et al. (2013) RCT (8) N _{Start} =44 N _{End} =44 TPS=Chronic	E: Segmental muscle vibration + standard therapy C: Standard therapy Duration: 30min/d, 5d/wk for 2wk	 Angular velocity at shoulder (+exp) Movement duration (+exp) Normalized jerk (+exp) Elbow angle (-) Shoulder angle (-) Shoulder abduction (-)
Caliandro et al. (2012) RCT (7) N _{Start} =49 N _{End} =36 TPS=Chronic	E: Focal muscle vibration C: Sham Duration: 30min/d, for 3d	Wolf Motor Function Test (+exp)
Stein et al. (2010) RCT (10) Nstart=30 Nend=30 TPS=Chronic	E: Stochastic resonance stimulation (combination of subthreshold electrical stimulation and vibration) C: Sham stimulation Duration: 3d/wk for 4wk	 Fugl-Meyer Assessment (-) Motor Activity Log (-) Action Research Arm Test (-)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group - indicates a statistically significant between groups difference at α =0.05 in favour of the control group - indicates no statistically significant between groups difference at α =0.05

Conclusions about muscle vibration

MOTOR FUNCTION					
LoE	Conclusion Statement	RCTs	References		
1a	Muscle vibration therapies may produce greater improvements in motor function than sham vibration or conventional therapy.	5	Calabro et al. 2017; Costantino et al. 2017; Lee et al. 2016; Caliandro et al. 2012; Stein et al. 2010		

MUSCLE STRENGTH					
LoE	Conclusion Statement	RCTs	References		
1a	Muscle vibration therapies may produce greater improvements in muscle strength than sham vibration or conventional therapy.	3	Costantino et al. 2017; Lee et al. 2016; Paoloni et al. 2014		

ACTIVITIES OF DAILY LIVING					
LoE	LoE Conclusion Statement RCTs References				
1a	There is conflicting evidence about the effect of muscle vibration therapies to improve performance of activities of daily living when compared to sham vibration or conventional therapy .	3	Calabro et al. 2017; Costantino et al. 2017; Stein et al. 2010		

SPASTICITY					
LoE	Conclusion Statement	RCTs	References		
1a	Muscle vibration therapies may produce greater improvements in spasticity than sham vibration or conventional therapy.	2	Calabro et al. 2017; Lee et al. 2016		

Key points

Muscle vibration may be beneficial for improving upper limb function following stroke.

Additional afferent and peripheral stimulation methods

Adopted from: https://www.saebo.com/saebostim-micro/

Additional sensory stimulation methods evaluated for motor rehabilitation included short wave therapy, repetitive peripheral magnetic stimulation, and intermittent pneumatic compression. Short-wave therapy is a non-invasive intervention in which electromagnetic radiation is applied to the region of the body typically at 27.12MHz in a continuous or pulse fashion (Wang et al. 2017). In repetitive peripheral magnetic stimulation coils are placed over paralysed muscles that generates a magnetic field that passes through the skin, and in turn can depolarize neurons to allow a muscle contraction (Momosaki et al. 2017). Repetitive peripheral magnetic stimulation compression is the application of inflatable splints where pressure is applied intermittently to increase sensory input (Cambier et al. 2003).

Additionally, a few studies looked at the effects of mirror therapy combined with the Mesh Glove, a novel form of technology that can apply sensory stimulation of varying intensities throughout the hand (Lee et al. 2015; Lin et al. 2014a; Lin et al. 2014b).

A total of 9 RCTs were found that evaluated the use of afferent and peripheral stimulation for upper extremity motor rehabilitation poststroke (Kattenstroth et al. 2018; Lee et al. 2015; Krewer et al. 2014; Lin et al. 2014a; Lin et al. 2014b; Hunter et al. 2011; Cambier et al. 2003; Feys et al. 1998; Jongbloed et al. 1989).

The methodological details and results of all 9 RCTs are presented in Table 26.

	Interventions			
Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Duration: Session length, frequency per week for total number of weeks	Result (direction of effect)		
Additional methods for Sensorv/Afferent Stimulation				
Kattenstroth et al. (2018) RCT (4) N _{Start} =71 N _{End} =48 TPS= Acute	E: Repetitive Sensory Stimulation C: Sham Repetitive Sensory Stimulation Duration: 45min/d, 5d/wk for 2wk	 Tactile Discrimination (+exp) Grating Orientation Task (+exp) Grip Strength (+exp) 9 Hole Peg Test (-) Jebsen Taylor Hand Function Test (-) Joint Position Sense Test (-) 		
Krewer et al. (2014) RCT (9) N _{Start} =63 N _{End} =44 TPS=Chronic	E: Repetitive peripheral magnetic stimulation C: Sham stimulation Duration: 20min/d, 2d/wk for 2wk	 Modified Tardieu Scale (-) Fugl-Meyer Assessment (-) Barthel Index (-) 		
<u>Hunter et al</u> . (2011) RCT (7) N _{start} =76 N _{end} =75 TPS= Acute	E: Mobilization and Tactile Stimulation (3 dose levels) C: Conventional therapy Duration: 30-120min (3x per day), 5d/wk for 2wk	 Motricity Index (-) Action Research Arm Test (-) 		
Cambier et al. (2003) RCT (7) N _{start} =23 N _{end} =23 TPS=Subacute	E: Intermittent pneumatic compression C: Sham short-wave therapy Duration: 30min/d, 5d/wk for 4wk	 Nottingham Sensory Assessment (+exp) Fugl-Meyer Assessment (+exp) Ashworth Scale (-) 		
<u>Feys et al</u> . (1998) RCT (6) N _{start} =100 N _{end} =100 TPS=Acute	E: Short-wave therapy stimulation with splints C: Sham stimulation Duration: 30min/d, 5d/wk for 6wk	 Fugl-Meyer Assessment (-) Action Research Arm Test (-) Barthel Index (-) 		
Jongbloed et al. (1989) RCT (5) N _{start} =90 N _{end} =87 TPS=Subacute	E: Sensorimotor integrative approach C: Functional approach Duration: <i>Not reported</i>	 Barthel Index (-) Sensorimotor Integration Test (-) 		
Mirror t	herapy with Mesh sensory stimulation gloves	versus mirror therapy		
Lee et al. (2015) RCT (7) N _{Start} =48 N _{End} =47 TPS=Chronic	E1: Mirror Therapy with Mesh Glove Afferent Stimulation E2: Mirror Therapy C: Mirror Therapy with Sham Stimulation Duration: 90min/d, 5d/wk for 4wk	 E1 vs E2/C Extensor Digitorum Muscle Tone (+exp) E1/C vs E2 Box and Block Test: (+exp, +con) Muscle stiffness on the flexor carpi radialis (+exp, +con) Functional Independence Measure (+exp, +con) Fugl-Meyer Assessment (-) Revised Nottingham Sensory Assessment (-) Modified Ashworth Scale (-) 		
Lin et al. (2014a) RCT (7) Nstart=16 NEnd=16 TPS=Chronic	C: Mirror therapy Duration: 90min/d, 5d/wk for 4wk	 Modified Astronom Scale (-) Box and Block Test (+exp) Functional Independence Measure (-) Action Research Arm Test (+exp) 		
Lin et al. (2014b) RCT (7)	E1: Mirror therapy + Mesh glove E2: Mirror therapy C: Therapeutic exercises	E1 vs C • Fugl-Meyer Assessment (+exp) E1 vs E2 & E1 vs C		

Table 26. RCTs evaluating afferent and peripheral stimulation interventions for upper extremity motor rehabilitation

N _{Start} =43	Duration: 90min/d, 5d/wk for 4wk	•	Box and Block Test (+exp)
N _{End} =42			<u>E1 vs E2</u>
TPS=Chronic		•	Wolf Motor Function Test (-)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about additional afferent and peripheral stimulation

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	Sensory stimulation methods such as short-wave therapy, repetitive magnetic stimulation and intermittent pneumatic compression may not have a difference in efficacy when compared to sham stimulation, conventional therapy and functional approaches for improving motor function.	6	Kattenstroth et al. 2018; Krewer et al. 2014; Hunter et al. 2011; McDonnell et al. 2007; Cambier et al. 2003; Feys et al. 1998	
1a	There is conflicting evidence about the effect of mirror therapy combined with Mesh Gloves to improve motor function when compared to mirror therapy on its own .	3	Lee et al. 2015; Lin et al. 2014a; Lin et al. 2014b	

MUSCLE STRENGTH				
LoE	Conclusion Statement	RCTs	References	
1b	There is conflicting evidence about the effect of sensory stimulation methods such as repetitive sensory stimulation and tactile stimulation to improve muscle strength when compared to sham stimulation and conventional therapy.	2	Kattenstroth et al. 2018; Hunter et al. 2011	

DEXTERITY			
LoE	Conclusion Statement	RCTs	References
1b	There is conflicting evidence about the effect of sensory stimulation methods such as short-wave therapy, repetitive magnetic stimulation and intermittent pneumatic compression to improve dexterity when compared to sham stimulation, conventional therapy and functional approaches.	2	Kattenstroth et al. 2018; McDonell et al. 2007
1a	Mirror therapy combined with Mesh Gloves may produce greater improvements in dexterity than mirror therapy on its own.	3	Lee et al. 2015; Lin et al. 2014a; Lin et al. 2014b

ACTIVITIES OF DAILY LIVING				
LoE	LoE Conclusion Statement RCTs Reference			
1a	Sensory stimulation methods such as short-wave therapy, repetitive magnetic stimulation and intermittent pneumatic compression may not have a difference in efficacy when compared to sham stimulation, conventional therapy and functional approaches for improving performance of activities of daily living.	3	Krewer et al. 2014; Feys et al. 1998; Jongbloed et al. 1989	
1a	There is conflicting evidence about the effect of mirror therapy combined with Mesh Gloves to improve	2	Lee et al. 2015; Lin et al. 2014a	

performance of activities of daily living when compared	
to mirror therapy on its own.	

SPASTICITY			
LoE	Conclusion Statement	RCTs	References
1a	Sensory stimulation methods such as short-wave therapy, repetitive magnetic stimulation and intermittent pneumatic compression may not have a difference in efficacy when compared to sham stimulation, conventional therapy and functional approaches for improving spasticity.	3	Krewer et al. 2014; Cambier et al. 2003; Jongbloed et al. 1989
1b	Mirror therapy when combined with Mesh Gloves may not have a difference in efficacy when compared to mirror therapy on its own for improving spasticity.	1	Lin et al. 2014a

PROPRIOCEPTION				
LoE	Conclusion Statement	RCTs	References	
1b	There is conflicting evidence about the effect of sensory stimulation methods such as short-wave therapy, repetitive magnetic stimulation and intermittent pneumatic compression to improve proprioception when compared to sham stimulation, conventional therapy and functional approaches.	2	Kattenstroth et al. 2018; Cambier et al. 2003	

Key points

The literature is mixed regarding additional afferent and peripheral stimulation for upper limb rehabilitation following stroke.

Invasive central nervous system stimulation Invasive cortical and nerve electrode implant stimulation



Adopted from: https://www.medgadget.com/2008/01/brain_stimulation_device_for_stroke_victims_fails_clinical_trial.html

Cortical stimulation in the motor cortex was traditionally used for the management of neuropathic pain, but preclinical evidence from animal models and clinical observations of pain patients showing motor improvements using this technique led to its adoption as an intervention for motor rehabilitation in stroke survivors (Levy et al. 2008; Tsubokawa et al. 1991). The neurosurgical procedure is performed through an extradural craniotomy where the stimulation electrode is placed on the dura matter of the motor cortex in a region predetermined from stereotaxic neuronavigation and functional magnetic resonance imaging (Levy et al. 2016; Brown et al. 2006). The frequency of stimulation is typically at 50Hz, and stimulation parameters remain consistent for the length of the intervention (Levy et al. 2016; Huang et al. 2008).

However, due to the invasive nature of this procedure and potential for adverse events, RCTs mainly investigating this technique for stroke rehabilitation were feasibility studies (Brown et al. 2006; Huang et al. 2008; Levy et al. 2008), and only recently a phase III clinical trial (Levy et al. 2016).

Vagus nerve stimulation has been shown in preclinical evidence from animal models to influence neuroplasticity, as stimulation can lead to increased acetylcholine and norepinephrine release, both of which are involved in the reorganization of cortical networks (Dawson et al. 2016). As well as pairing upper limb rehabilitation with vagus nerve stimulation has been shown to further promote plasticity in preclinical settings (Hays et al. 2016). Only one study has looked at vagus nerve stimulation with upper limb rehabilitation in stroke survivors (Dawson et al. 2016).

The methodological details and results of 5 RCTs (Levy et al. 2016; Dawson et al. 2016; Huang et al. 2008; Levy et al. 2008; Brown et al. 2006) that have evaluated the use of invasive cortical and nerve stimulation methods for improving motor function post stroke are presented in Table 27.

Table 27. RCTs evaluating invasive brain stimulation interventions for upper extremity motor rehabilitation

Authors (Year)	Interventions	Outcome Measures
Study Design (PEDro Score)	Duration: Session length frequency	Result (direction of effect)
Sample Sizestart	per week for total number of weeks	
Sample Sizeend		
Time post stroke category		
	Motor cortex stimulation	
Levy et al. (2016)	E: Cortical implant with epidural 6-	Arm Motor Ability Test (-)
RCT (6)	contact lead perpendicular to the	 Fugl-Meyer Assessment (-)
Nstart=164	primary motor cortex and a pulse	
N _{End} =128	generator	
TPS=Chronic	C: Conventional rehabilitation	
	Duration: Not Specified	
Huang et al. (2008)	E1: Motor cortex stimulation (50Hz)	• Fugl Meyer Score (+exp, +exp ₂)
RCT (5)	C1: Conventional rehabilitation	• Box and Block Test (+exp, +exp ₂)
N _{start} =24	E2: Motor cortex stimulation (101Hz)	Stroke Impact Scale (-)
N _{end} =24	C2: Conventional rehabilitation	Arm Motor Ability Test (-)
TPS=Chronic	Duration: 2.5hr/d, 5d/wk for 4 wk	Grip strength (-)
Levy et al. (2008)	E: Motor cortex stimulation	 Fugl Meyer Score (+exp)
RCT (5)	C: Conventional rehabilitation	 Arm Motor Ability Test (+exp)
N _{start} =24	Duration: Not Specified	
N _{end} =24		
TPS=Chronic		
<u>Brown et al</u> . (2006)	E: Motor cortex stimulation	 Fugl Meyer Scale (+exp)
RCT (6)	C: Conventional rehabilitation	 Stroke Impact Scale (+exp)
N _{start} =10	Duration: 30min/d, 5d/wk for 3 wk	
N _{end} =10		
TPS=Chronic		
	Vagus perve stimulation	
Dawson et al. (2016)	E: Impanted vagus nerve stimulation	• Fugl-Mever Assessment (+exp)
RCT (7)	C: Conventional rehabilitation	Action Research Arm Test (-)
Nstart=20	Duration: 20min/d, 4 d/wk for 8 wk	Grip Strength (-)
Nord=20	,	Nine Hole Peg Test (-)
TPS=Chronic		Box and Block Test (-)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about invasive cortical and nerve stimulation

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of motor cortex stimulation to improve motor function when compared to conventional therapy .	4	Levy et al. 2016; Huang et al. 2008; Levy et al. 2008; Brown et al. 2006
1b	There is conflicting evidence about the effect of vagus nerve stimulation to improve motor function when compared to conventional therapy .	1	Dawson et al. 2016

MUSCLE STRENGTH			
LoE	Conclusion Statement	RCTs	References
2	Motor cortex stimulation may not have a difference in efficacy when compared to conventional therapy for improving muscle strength.	1	Huang et al. 2008
1b	Vagus nerve stimulation may not have a difference in efficacy when compared to conventional therapy for improving muscle strength.	1	Dawson et al. 2016

DEXTERITY			
LoE	Conclusion Statement	RCTs	References
2	Motor cortex stimulation may produce greater improvements in dexterity than conventional therapy.	1	Huang et al. 2008
1b	Vagus nerve stimulation may not have a difference in efficacy when compared to conventional therapy for improving dexterity.	1	Dawson et al. 2016

ACTIVITIES OF DAILY LIVING				
1a	1aThere is conflicting evidence about the effect of motor cortex stimulation to improve performance of activities of daily living when compared to conventional therapy.Levy et al. 2016; Huang et al. 2008; Brown et al. 2006			

Key points

The literature is mixed regarding invasive cortical and nerve stimulation for upper limb rehabilitation following stroke.

Non-invasive brain stimulation

Repetitive Transcranial Magnetic Stimulation (rTMS)



Adopted from: https://www.rtmscentre.co.uk/rtms-treatment-in-the-uk/

Transcranial magnetic stimulation is a painless and non-invasive method of affecting neural activity through the exogenous generation of an electromagnetic field through a coil placed on the scalp, that consequently induces a change in the electrical fields of the brain (Peterchev et al. 2012). The voltage and current of the electromagnetic field generated are dependent on the parameters of the stimulation device, which is not distorted by the biological tissues in which it is applied in (Peterchev et al. 2012). The neuromodulatory effects of transcranial magnetic stimulation are attributed largely to neural membrane polarization shifts that can lead to changes in neuron activity, synaptic transmission, and activation of neural networks (Peterchev et al. 2012). Repetitive transcranial magnetic stimulation (rTMS) is the application of repetitive trans of transcranial magnetic stimulation at regular intervals.

After a stroke, interhemispheric competition is altered; with cortical excitability increasing in the unaffected hemisphere increasing and decreasing in the affected hemisphere (Zhang et al. 2017). rTMS can be used to help modulate this interhemispheric competition, with low stimulation frequencies (\leq 1Hz) decreasing cortical excitability and inhibiting activity of the contralesional hemisphere, while high frequency (>1Hz) stimulation increases excitability and have a facilitatory effect on activity of the ipsilesional hemisphere (Dionisio et al. 2018).

A growing number of studies have investigated the effects of rTMS on improving upper extremity motor rehabilitation after a stroke. Low frequency rTMS versus sham stimulation or conventional therapy was assessed in 28 RCTs (Long et al. 2018; Tarri et al. 2018; Watanabe et al. 2018; Askin et al. 2017; Meng and Song, 2017; Ozkeskin et al. 2017; Yang et al. 2017; Du et al. 2016; Li et al. 2016; Cassidy et al. 2015; Ludermann-Podubecka et al. 2015; Abo et al. 2014; Barros Galvao et al. 2014; Rose et al. 2014; Wang et al. 2014; Etoh et al. 2013; Higgins et al. 2013; Saskai et al. 2013; Conforto et al. 2012; Seniow et al. 2012; Emara et al. 2010; Khedr et al. 2009; Takeuchi et al. 2008; Liepert et al. 2007; Pomeroy et al. 2007; Fregni et al. 2006; Mansur et al. 2005; Takeuchi et al. 2005), while high frequency rTMS versus sham stimulation or conventional therapy was assessed in 13 RCTs (Guan et al. 2017; Du et al. 2016; Hosomi et al. 2016; Li et al. 2016; Cassidy et al. 2015; Kim et al. 2014; Saskai et al. 2013; Chang et al. 2010; Emara et al. 2010; Khedr et al. 2010; Khedr et al. 2009; Malcom et al. 2007; Khedr et al. 2005). RCTs looking at multimodal interventions with rTMS were limited, and combinations included bilateral stimulation (both high and low frequency rTMS; (Long et al. 2018; Takeuchi et al. 2009)), mirror therapy (Ji et al. 2014), virtual reality (Zheng et al. 2015), sensory cueing (Yang et al. 2017) and cyclic NMES (Tosun et al. 2017). The methodological details and results of all 39 RCTs evaluating rTMS for the upper extremity motor rehabilitation are presented in Table 28.

Authors (Voar)	Interventions	Outcome Measures
Authors (Tear)	Bunstian Ossaian launth framman	
Study Design (PEDro Score)	Duration: Session length, frequency	Result (direction of effect)
Sample Sizestart	per week for total number of weeks	
Sample Sizeend		
Time post stroke category		
Low frequency	(1Hz) rTMS vs sham stimulation or conve	entional therapy
Long et al. 2018	E1: Low Frequency (1Hz) combined	F2 vs C
RCT (7)	with High Frequency (10Hz) Repetitive	 Eugl-Meyer Assessment (+eyp₂)
	Transpranial Magnetic Stimulation	• Tugi-Meyer Assessment (+exp2)
NStart =02	Transcranial Magnetic Stimulation	• Wolf Motor Function Test (-)
NEnd =62	E2: Low Frequency (1Hz) Repetitive	
TPS=Acute	Transcranial Magnetic Stimulation	
	C: Sham Repetitive Transcranial	
	Magnetic Stimulation	
	Duration: Not specified	
Tarri et al. 2018	E. Paired associative stimulation	 Eugl-Meyer Assessment (-)
PCT (6)	(electrical stimulation + low frequency	· · · · · · · · · · · · · · · · · · ·
	$(4 \Pi^2) \times TMC$	
NStart =24		
NEnd =24	C. Snam Stimulation	
TPS=Subacute	Duration: Not specified	
Watanabe et al. (2018)	E1: Intermittent Theta-Burst Stimulation	<u>E2 vs C</u>
RCT (5)	E2: Low Frequency (1Hz) Repetitive	 Fugl-Meyer Assessment: (-)
Nstart =21	Transcranial Magnetic Stimulation	Stroke Impairment Assessment Set (-)
NEpd =21	C: Sham Stimulation	 Modified Ashworth Scale (+exp₂)
TPS-Acute	Duration: Not specified	Grin Strength (-)
Askin at al. 2017	E: Low froquency (1Hz) Popotitivo	- Poy and Plack Test (Lovn)
	E. Low nequency (112) Repetitive	• Box and block rest (texp)
	Transcranial Magnetic Stimulation to	Functional independence measure
N _{Start} =40	unaffected nemisphere	(+exp)
N _{End} =40	C: Conventional Physical Therapy	 Brunnstrom Recovery Stages (-)
TPS=Chronic	Duration: 1 hr/d, 5d/wk, for 2wk	 Fugl-Meyer Assessment (-)
		 Modified Ashworth Scale (-)
Meng & Song 2017	E: Low Frequency (1Hz) Repetitive	 National Institute of Health Stroke Scale
RCT (6)	Transcranial Magnetic Stimulation to	(+exp)
Nstart = 20	unaffected hemisphere	 Barthel Index (+exp)
Notat =20	C: Sham Repetitive Transcranial	 Eugl-Meyer Assessment (+eyn)
	Magnetic Stimulation	
IF S=NK	Duration: 20 min/d. Zd/uk for 2006	
	Duration: 30 min/d, 7d/wk for 2wk	
Ozkeskin et al. 2017	E: Low frequency (1Hz) Repetitive	 Brunnstrom Recovery Stages (-)
RCT (9)	Transcranial Magnetic Stimulation to	 Finger Touch Localization (-)
N _{Start} =21	unaffected hemisphere	 Modified Ashworth Scale (-)
N _{End} =21	C: Sham Repetitive Transcranial	 Wrist Proprioceptive Evaluations (+exp)
TPS=Chronic	Magnetic Stimulation	
	Duration: 90 min/d 5d/wk for 2wk	
Du et al. (2016)	E1: High frequency (2Hz) rTMC	F2 va C
$\frac{DU \in [a]}{PCT(7)}$	E_1 . The frequency ($3\Pi Z$) The Z_2	$\frac{L2 V3 U}{E u g Mover Accessment (Lown)}$
	E2. LOW frequency (THZ) FINIS	rugi-ivieyer Assessment (+exp ₂)
N _{Start} =69	C: Snam rIMS	Medical Research Council Score
NEnd =59	Duration: 30min/d, 5d/wk for 1wk	(+exp ₂)
TPS=Acute		 National Institute of Health Stroke
		Scale (+exp ₂)
		 Modified Rankin Scale (+exp₂)
		Barthel Index (+exp ₂)
Lietal (2016)	E1: Low frequency (1Hz) rTMS	E1 vs C
	E2: Ligh fraguency (10Uz) *TMC	Eugl Mover Accessment (Leve)
		Fugi-ivieyer Assessment (+exp)
NStart = 127		• vvoir iviotor Function Test (-)
N _{End} =12/	Duration: 40min/d, 5d/wk for 2wk	
TPS=Subacute		

Table 28. RCTs evaluating rTMS interventions for upper extremity motor rehabilitation

Ludemann-Podubecka et al. (2016)	E: Low frequency (1Hz) rTMS	Jebsen Taylor Hand Function Test
RCT (7)	C: Sham	(+exp)
Nstart =10	Duration: 30min/d, 5d/wk for 6wk	Box and Block Test (-)
TPS=Subacute		
Cassidy et al. (2015)	E1: High frequency (6Hz) rTMS	E2 vs. C
RCT (7)	E2: Low frequency (1Hz) rTMS	 Box and Block Test (+exp₂)
N _{Start} =11	C: Sham	
N _{End} =11	Duration: 1hr/d, 3d/wk for 5wk	
TPS=Chronic		
Ludemann-Podubecka et al. (2015)	E: Low frequency (1Hz) rTMS	Wolf Motor Function Test (+exp)
	C: Sham	Motor Evaluation Scale (+exp) Einger Tenning ()
$N_{\text{Start}} = 40$	Duration. Somin/d, Sd/wk for 6 wk	Finger rapping (-)
TPS=Chronic		
Abo et al. (2014)	E: Low frequency (1Hz) rTMS + OT	 Fugl-Meyer Assessment (+exp)
RCT (7)	training (NEURO)	 Wolf Motor Function Test (+exp)
N _{Start} =66	C: CIMT	
NEnd=66	Duration: 20min rTMS & 120min OT	
IPS=Chronic	(2x/d), 6d/wk tor 4wk	
Barros Galvao et al. (2014)	E: Low frequency (1Hz) r1MS	Involutied Ashworth Scale (-)
κυι (δ) Νομα-20	U. SildIII Duration: 1hr/d. 5d/wk for 2wk	 rugi-ivieyer Assessment (-) Eunctional Independence Measure (-)
N _{start} =20	Duration. mi/d, 50/wk for 2wk	 Functional independence measure (-)
TPS=Chronic		 Wrist range of motion (-)
Rose et al. (2014)	E: Low frequency (1Hz) rTMS +	Wolf Motor Function Test (-)
RCT (5)	functional task practice (FTP)	• Pinch strength (lateral and palmar) (-
Nstart=22	C: Sham + FTP)
N _{End} =19	Duration: 1.5hr/d, 4d/wk, 4wk	 Fugl-Meyer Assessment (-)
TPS=Chronic		 Action Research Arm Test (-)
		Modified Ashworth Scale (-)
		Motor Activity Log (-)
Wang et al. (2014)	E1: Low frequency (1Hz) rTMS applied	F1 vs C
RCT (9)	to primary motor cortex	 Wolf Motor Function Test (+exp)
N _{Start} =44	E2: Low frequency (1Hz) rTMS applied	 Fugl-Meyer Assessment (+exp)
N _{End} =44	to premotor area	Medical Research Council Scale
TPS=Chronic	C: Sham	(+exp)
	Duration: Not Specified	<u>E2 vs C</u>
		• Wolf Motor Function Test: (+exp ₂)
		Fugl-Meyer Assessment: (+exp ₂)
		Iniedical Research Council Scale (Lovp.)
		(τσλμ2) F1 vs F2
		 Wolf Motor Function Test (+exp)
		 Fugl-Meyer Assessment (+exp)
		Medical Research Council Scale
		(+exp)
Etoh et al. 2013	E1: Low frequency (1Hz) rTMS	Action Research Arm Test (+exp)
	U. SHAIN FIND Duration: Amin Ed/wk for Swk	Fugi Meyer Assessment (-) Simple test for evaluating hand
Nend=18	Duration. 4min, 50/WK IOF ZWK	function (-)
TPS=Chronic		 Modified Ashworth scale (-)
Higgins et al. (2013)	E: Low frequency (1Hz) rTMS	Box and Block Test (-)
RCT (7)	C: Sham	 Motor Acitivity Log (-)
Nstart=11	Duration: 90min/d, 4d/wk for 4wk	Wolf Motor Function Test (-)
N _{End} =11		
IPS=Chronic		F 2 va C
<u>Sasaki et al</u> . (2013)	E1: High frequency (10Hz) r1MS	$\underline{E2 VS U}$
	EZ. TEZ ETINO NON-lesioned nemisphere	• Grip strengtri (-)

N _{Start} =29	C: Sham	Tapping frequency (-)
N _{End} =29	Duration: 45min/d, 2d/wk for 6wk	
IPS=Acute	Et Low frequency (14-) rTMC	Jahaan Taylor Hand Eurotian toot
<u>Comorto et al</u> . (2012) RCT (6)	C: Sham	• Jebsen-Taylor Hand Function test (+exp)
Netart=29	Duration: 25min/d. 5d/wk for 4wk	 Pinch Force (+exp)
N _{end} =28		 Fugl-Meyer Assessment (+exp)
TPS=Acute		Modified Ashworth Scale (-)
Seniów et al. (2012)	E: Low frequency (1Hz) rTMS + PT	Wolf Motor Function Test (-)
RCT (8)	C: Sham + PT	 Fugl-Meyer Assessment (-)
N _{start} =40	Duration: 75min/d, 5d/wk for 3wk	
Nend=33		
Emara et al. (2010)	E1: High frequency (5Hz) rTMS	F2 vs C
RCT (7)	E2: Low frequency (1Hz) rTMS	 Finger tapping test (+exp₂)
N _{start} =60	C: Sham	 Frenchay Activities Index (+exp₂)
N _{end} =60	Duration: 30min/d, 5d/wk for 4wk	Modified Rankin Scale (+exp ₂)
TPS=Subacute		
<u>Khedr et al.</u> (2009)	E1: Low frequency (1Hz) rTMS	<u>E1 vs C</u>
KUI (δ) N = -26	E∠: High frequency (3HZ) rTMS	Grip strength (+exp) Burdue Begbeerd teek (+exp)
N _{start} =30	C. Sham Duration: 30min/d. 3d/wk for 4wk	 Purdue Pegboard task (+exp) Barthel Index (+exp)
TPS=Acute	Duradon. Johnin/u, Ju/wk lor 4wk	NIHSS (+exp)
Takeuchi et al. (2008)	E: Low frequency (1Hz) rTMS + pinch	Pinch force (+exp)
RCT (7)	force motor training	· · · /
N _{start} =20	C: Sham + pinch force motor training	
N _{end} =20	Duration: Not Specified	
TPS=Chronic		
Liepert et al. (2007)	E: Low frequency (1Hz) rTMS	Grip strength (-)
RUT (7) Num=12	C: Sham Duration: 3br/d. 3d/wk for 4wk	• 9-hole peg test (+exp)
Nend=12	Duration. Shi/u, Su/wk for 4wk	
TPS=Acute		
Pomeroy et al. (2007)	E1: Low frequency (0.5Hz) rTMS +	Flexion/extension torque (-)
RCT (8)	voluntary muscle contraction (VMC)	Action Research Arm Test (-)
N _{start} =27	E2: Low frequency (0.5Hz) rTMS +	
Nend=24	placebo VMC	
1PS=Chronic	E3: Sham FIMS + VMC C: Sham rTMS + placebo VMC	
	Duration: Not Specified	
Fregni et al. (2006)	E: Low frequency (1Hz) rTMS	 Jebsen-Taylor Hand Function test
RCT (7)	C: Sham	(+exp)
N _{start} =15	Duration: 20min/d, 5d/wk for 6wk	
N _{end} =15		
IPS=Chronic		Finger tenning test ()
$\frac{\text{Mansur et al.}}{\text{RCT}(4)}$	E: LOW TREQUENCY (1HZ) TIMS	 Finger tapping test (-) Perdue Perducard test (Lovo)
Not (4) Netort=10	Duration: Not Specified	• renue regularu lest (+exp)
Nend=10		
TPS=Chronic		
Takeuchi et al. (2005)	E: Low frequency (1Hz) rTMS	Hand and pinch force (-)
RCT (6)	C: Sham	
N _{start} =20	Duration: 25min/d, 3d/wk for 5wk	
Nend=20		
IFS=UNIONIC High free	 uency (>1Hz) rTMS vs Sham or conventiv	onal therapy
Guan et al. 2017	E: High frequency (5Hz) Repetitive	National Institutes of Health Stroke
RCT (5)	Transcranial Magnetic Stimulation	Scale (+exp)
NStart =42	C: Sham Repetitive Transcranial	Barthel Index (+exp)
$N_{End} = 27$	Wagnetic Stimulation	Fugi-Meyer Assessment (+exp) Modified Bankin Score ()
IFS=Acule	Duration. 25 min/u, 40/WK for 6WK	Iviouilleu Rahkin Score (-)

Du et al. (2016) RCT (7) N _{Start} =69 N _{End} =55 TPS=Acute	E1: High frequency (3Hz) rTMS E2: Low frequency (1Hz) rTMS C: Sham rTMS Duration: 30min/d, 5d/wk for 1wk	 <u>E1 vs C</u> Fugl-Meyer Assessment (-) Medical Research Council Score (-) National Institute of Health Stroke Scale (+exp) Modified Rankin Scale (+exp) Barthel Index (+exp)
Hosomi et al. (2016) RCT (8) N _{start} =41 N _{End} =39 TPS=Subacute Li et al. (2016)	E: High frequency (5Hz) rTMS C: Sham Duration: 1hr/d, 5d/wk for 2wk E1: Low frequency (1Hz) rTMS	 Brunnstorm Recovery Stages (-) Fugl-Meyer Assessment (-) National institute for Health Stroke Scale (-) Grip Power (-) E2 vs C
$ \frac{1}{RCT (7)} $ $ \frac{1}{N_{Start} = 127} $ $ \frac{1}{N_{End} = 127} $ $ \frac{1}{TPS = Subacute} $	E2: High frequency (10Hz) rTMS C: Sham Duration: 40min/d, 5d/wk for 2wk	 Fugl-Meyer Assessment (+exp₂) Wolf Motor Function Test (-)
Kim (2014) RCT (6) Nstart=31 NEnd=31 TPS=Chronic 1	E: High frequency (10Hz) rTMS C: Sham Duration: 10min/d, 5d/wk for 4wk	Manual Function Test (+exp)
<u>Sasaki et al</u> . (2013) RCT (8) N _{start} =29 N _{End} =29 TPS=Acute	E1: 10Hz rTMS lesioned hemisphere E2: 1Hz rTMS non-lesioned hemisphere C: Sham Duration: 45min/d, 2d/wk for 6wk	E1 vs C Grip strength (+exp) Tapping frequency (+exp)
Chang et al. (2010) RCT (5) N _{start} =28 N _{end} =28 TPS=Subacute	E: High frequency (10Hz) rTMS C: Sham Duration: 2min, 5d/wk for 2wk	 Motricity Index (+exp) Fugl-Meyer Assessment (-)
Emara et al. (2010) RCT (7) N _{start} =60 N _{end} =60 TPS=Subacute	E1: 5Hz rTMS E2: 1Hz rTMS C: Sham Duration: 30min/d, 5d/wk for 4wk	 <u>E1 vs C</u> Finger tapping test (+exp) Frenchay Activities Index (+exp) Modified Rankin Scale (+exp)
Khedr et al. (2010) RCT (8) N _{start} =48 N _{end} =38 TPS=Acute	E1: 3Hz rTMS E2: 10Hz rTMS C: Sham Duration: 30min/d, 3d/wk for 4wk	 <u>E1/E2 vs C</u> Grip strength (+exp, +exp₂) NIHSS (+exp, +exp₂) Modified Rankin Scale (+exp, +exp₂) <u>E1 vs E2</u> Grip strength (-) NIHSS (-) Modified Rankin Scale (-)
Khedr et al. (2009) RCT (8) Nstart=36 Nend=36 TPS=Acute	E1: 1Hz rTMS E2: 3Hz rTMS C: Sham Duration: 30min/d, 3d/wk for 4wk	E2 vs C Grip strength (+exp ₂) Purdue Pegboard task (+exp ₂) Barthel Index (+exp ₂) NIHSS (+exp ₂)
<u>Malcolm et al</u> . (2007) RCT (6) N _{start} =19 N _{end} =19 TPS=Chronic	E: High frequency (20Hz) rTMS C: Sham Duration: 40min/d, 6d/wk for 5wk	 Wolf Motor Function Test (-) Motor Activity Log (-)
<u>Khedr et al</u> . (2005) RCT (6) N _{start} =52 N _{end} =52	E: High frequency (3Hz) rTMS C: Sham Duration: 45min/d, 5d/wk, 2wk	 Barthel Index (+exp) NIHSS (+exp) Scandinavian Stroke Impact Scale (+exp)

TPS=Acute					
Low frequency combined w	Low frequency combined with high frequency rTMS or low frequency versus high frequency rTMS				
Long et al. 2018 RCT (7) N _{Start} =62 N _{End} =62 TPS=Acute	E1: Low Frequency Combined with High Frequency Repetitive Transcranial Magnetic Stimulation E2: Low Frequency Repetitive Transcranial Magnetic Stimulation C: Sham Repetitive Transcranial Magnetic Stimulation Duration: Not Specified	E1 vs C • Fugl-Meyer Assessment (+exp) • Wolf Motor Function Test (+exp) E1 vs E2 • Fugl-Meyer Assessment (-) • Wolf Motor Function Test (+exp)			
Takeuchi et al. (2009) RCT (6) N _{start} =30 N _{end} =30 TPS=Chronic	E1: Bilateral (dual) rTMS (1Hz and 10Hz) E2: 10Hz rTMS E3: 1Hz rTMS Duration: 15min/d, 3d/wk for 5wk	E1 vs E2 Pinch force (+exp) E1 vs E3 Pinch force (+exp)			
	rTMS plus an additional intervention				
Tosun et al. 2017 RCT (7) Nstart =25 NEnd =25 TPS=Subacute Yang et al. (2017) RCT (8) Nstart =60 NEnd =60 TPS=Subacute	E1: Low Frequency (1Hz) Repetitive Transcranial Magnetic Stimulation E2: Low Frequency Repetitive Transcranial with Cyclic NMES C: Physical Therapy Duration: 1 hr/d, 5d/wk for 4wk E1: Low frequency (1Hz) Repetitive Transcranial Magnetic Stimulation with Sensory Cueing E2: Low frequency (1Hz) Repetitive Transcranial Magnetic Stimulation C: Conventional Therapy Duration: 45 min/d, 5d/wk for 4wk	E1/E2 vs C; E1 vs E2 Fugl-Meyer Assessment (-) Motricity Index (-) Brunnstrom Recovery Stages (-) Modified Ashworth Scale (-) Barthel Index (-) E1 vs C Fugl-Meyer Assessment (-) Action Research Arm Test (-) Modified Barthel Index (-) E2 vs C Fugl-Meyer Assessment (-) Action Research Arm Test (-) Modified Barthel Index (-) E1 vs E2 Fugl-Meyer Assessment (-) Action Research Arm Test (-) Modified Barthel Index (-) E1 vs E2 Fugl-Meyer Assessment (-) Action Research Arm Test (-) Modified Barthel Index (-)			
Zheng et al. (2015) RCT (7) N _{Start} =112 N _{End} =108 TPS=Chronic	E: Low frequency (1Hz) rTMS + virtual reality (VR) training C: Sham + VR training Duration: 45min/d, 6d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Wolf Motor Function Test (+exp) Modified Barthel Index (+exp) 			
<u>Ji et al</u> . (2014) RCT (7) N _{Start} =35 N _{End} =35 TPS=Chronic	E1: Mirror therapy + high frequency (10Hz) rTMS E2: Mirror therapy C: Sham Duration: 15 min/d, 6d/wk for 4wk	 <u>E1 vs E2</u> Fugl-Meyer Assessment (+exp) Box and Block Test (+exp) <u>E1 vs C</u> Fugl-Meyer Assessment (+exp) Box and Block Test (+exp) 			

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α=0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group - indicates no statistically significant between groups differences at α =0.05

Conclusions about rTMS

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of low frequency rTMS to improve motor function when compared to sham stimulation or conventional therapy .	20	Long et al. 2018; Tarri et al. 2018; Watanabe et al. 2018; Askin et al. 2017; Meng and Song, 2017; Ozkesin et al. 2017; Yang et al. 2017; Du et al. 2016; Li et al. 2016; Ludermann-Podubecka et al. 2015; Abo et al. 2014; Barros Galvao et al. 2014; Rose et al. 2014; Wang et al. 2014; Etoh et al. 2013; Higgins et al. 2013; Conforto et al. 2012; Seniow et al. 2012; Pomeroy et al. 2007; Fregni et al. 2006
1a	High frequency rTMS may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improving motor function.	7	Guan et al. 2017; Du et al. 2016; Hosomi et al. 2016; Li et al. 2016; Kim et al. 2014; Chang et al. 2010; Malcom et al. 2007
1b	There is conflicting evidence about the effect of bilateral rTMS stimulation (both high and low frequency) to improve motor function when compared to sham stimulation or conventional therapy.	1	Long et al. 2018
1b	Low frequency rTMS with sensory cueing may not have a difference in efficacy when compared to low frequency rTMS or sham stimulation for improving motor function.	1	Yang et al. 2017
1b	Low frequency rTMS combined with virtual reality training may produce greater improvements in motor function than virtual reality training on its own or sham stimulation combined with virtual reality.	1	Zheng et al. 2015
1b	Mirror therapy combined with high frequency rTMS may produce greater improvements in motor function than mirror therapy on its own or sham stimulation.	1	Ji et al. 2014
1b	Low frequency rTMS with cyclic NMES may not have a difference in efficacy when compared to low frequency rTMS or conventional therapy for improving motor function.	1	Tosun et al. 2017

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of low frequency rTMS to improve dexterity when compared to sham stimulation or conventional therapy .	10	Askin et al. 2017; Ozkeskin et al. 2017; Cassidy et al. 2015; Ludermann- Podubecka et al. 2015; Higgins et al. 2013; Saskai et al. 2013; Emara et al. 2010; Khedr et al. 2009; Liepert et al. 2007; Mansur et al. 2005	
1a	High frequency rTMS may produce greater improvements in dexterity than sham stimulation or conventional therapy.	4	Cassidy et al. 2015; Saskai et al. 2013; Emara et al. 2010; Khedr et al. 2009	
1b	Mirror therapy combined with high frequency rTMS may produce greater improvements in dexterity	1	Ji et al. 2014	

than mirror therapy on its own or sham	
stimulation.	

SPASTICITY				
LoE	Conclusion Statement	RCTs	References	
1a	Low frequency rTMS may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improving spasticity.	7	Watanabe et al. 2018; Askin et al. 2017; Ozkeskin et al. 2017; Barros Galvao et al. 2014; Rose et al. 2014; Etoh et al. 2013; Conforto et al. 2012	
1b	Low frequency rTMS with cyclic NMES may not have a difference in efficacy when compared to low frequency rTMS or conventional therapy for improving spasticity.	1	Tosun et al. 2017	

RANGE OF MOTION				
LoE	Conclusion Statement	RCTs	References	
1a	Low frequency rTMS may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improving range of motion.	2	Barros Galvao et al. 2014; Pomeroy et al. 2007	

	PROPRIOCEPTION		
LoE	Conclusion Statement	RCTs	References
1h	Low frequency rTMS may produce greater improvements in proprioception than sham	1	Ozkeskin et al. 2017
	stimulation or conventional therapy.	•	

STROKE SEVERITY				
LoE	Conclusion Statement	RCTs	References	
1a	Low frequency rTMS may produce greater improvements on measures of stroke severity than sham stimulation or conventional therapy.	5	Askin et al. 2017; Meng and Song, 2017; Du et al. 2016; Emara et al. 2010; Khedr et al. 2009	
1a	High frequency rTMS may produce greater improvements on measures of stroke severity than sham stimulation or conventional therapy.	6	Guan et al. 2017; Du et al. 2016; Hosomi et al. 2016; Emara et al. 2010; Khedr et al. 2010; Khedr et al. 2009	
1b	Low frequency rTMS with cyclic NMES may not have a difference in efficacy when compared to low frequency rTMS or conventional therapy for improvements on measures of stroke severity.	1	Tosun et al. 2017	

ACTIVITIES OF DAILY LIVING			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of low frequency rTMS to improve performance of activities of daily living when compared to sham stimulation or conventional therapy .	9	Askin et al. 2017; Meng and Song, 2017; Yang et al. 2017; Du et al. 2016; Barros Galvao et al. 2014; Rose et al. 2014; Higgins et al. 2013; Emara et al. 2010; Khedr et al. 2009
1a	High frequency rTMS may produce greater improvements in performance of activities of daily living than sham stimulation or conventional therapy.	6	Guan et al. 2017; Du et al. 2016; Emara et al. 2010; Khedr et al. 2009; Malcom et al. 2007; Khedr et al. 2005
1b	Low frequency rTMS with sensory cueing may not have a difference in efficacy when compared to low frequency rTMS or sham stimulation for improving performance of activities of daily living.	1	Yang et al. 2017
1b	Low frequency rTMS combined with virtual reality training may produce greater improvements in performance of activities of daily living than virtual reality training on its own or sham stimulation combined with virtual reality.	1	Zheng et al. 2015
1b	Low frequency rTMS with cyclic NMES may not have a difference in efficacy when compared to low frequency rTMS or conventional therapy for improving performance of activities of daily living.	1	Tosun et al. 2017

MUSCLE STRENGTH			
LoE	Conclusion Statement	RCTs	References
1a	There is conflicting evidence about the effect of low frequency rTMS to improve muscle strength when compared to sham stimulation or conventional therapy .	10	Watanabe et al. 2018; Du et al. 2016; Rose et al. 2014; Wang et al. 2014; Saskai et al. 2013; Conforto et al. 2012; Khedr et al. 2009; Takeuchi et al. 2008; Liepert et al. 2007; Takeucchi et al. 2005
1a	High frequency rTMS may produce greater improvements in muscle strength than sham stimulation or conventional therapy.	6	Du et al. 2016; Hosomi et al. 2016; Saskai et al. 2013; Chang et al. 2010; Khedr et al. 2010; Khedr et al. 2009
1a	Bilateral rTMS stimulation (both high and low frequency) may produce greater improvements in muscle strength than low frequency rTMS.	1	Takeuchi et al. 2009
1a	Bilateral rTMS stimulation (both high and low frequency) may produce greater improvements in muscle strength than high frequency rTMS.	1	Takeuchi et al. 2009
1b	Low frequency rTMS with cyclic NMES may not have a difference in efficacy when compared to low frequency rTMS or conventional therapy for improving muscle strength.	1	Tosun et al. 2017

Key points

The literature is mixed regarding low frequency repetitive transcranial magnetic stimulation, alone or in combination with other therapy approaches, for upper limb rehabilitation following stroke.

High frequency repetitive transcranial magnetic stimulation, alone or in combination with other therapy approaches, may be beneficial for upper limb rehabilitation.

The literature is mixed regarding bilateral repetitive transcranial magnetic stimulation for upper limb rehabilitation.

Theta burst stimulation (TBS)



Adopted from: https://www.psychiatryadvisor.com/home/depression-advisor/intermittent-theta-burst-stimulation-for-major-depressive-disorder-treatment/

Theta Burst Stimulation (TBS) is an emerging treatment modality that is a patterned form of rTMS where stimulation pulses are delivered in triplets or bursts at a high frequency (50Hz), and in a short interval (200ms), intending to mimic naturally occurring theta brain oscillations (Schwippel et al. 2019). TBS can also be used to adjust interhemispheric rivalry after a stroke and promote motor recovery through the delivery of continuous TBS (cTBS) to reduce cortical excitability in the contralesional hemisphere (600 pulses over 40 seconds); or intermittent TBS (iTBS) to increase cortical excitability in the ipsilesional hemisphere (600 pulses over 190 seconds) (Schwippel et al. 2019; Cotoi et al. 2019).

A total of 9 RCTs were found that evaluated the use of TBS for upper extremity motor rehabilitation poststroke (Watanabe et al. 2018; Ackerley et al. 2016; Di Lazzaro et al. 2016; Volz et al. 2016; Kim et al. 2015; Di Lazzaro et al. 2014; Hsu et al. 2013; Sung et al. 2013; Talelli et al. 2012). Six RCTs evaluated the effects of iTBS (Watanabe et al. 2018; Ackerley et al. 2016; Volz et al. 2016; Kim et al. 2015; Hsu et al. 2013; Talelli et al. 2012), and three RCTs the effects of cTBS (Di Lazzaro et al. 2016; Di Lazzaro et al. 2016; Di Lazzaro et al. 2014; Talelli et al. 2012), both compared to sham TBS for improving upper extremity motor rehabilitation outcomes. Additionally, one RCT evaluated the effects of iTBS combined with low frequency rTMS compared to sham TBS/rTMS for improving upper extremity motor rehabilitation outcomes.

The methodological details and results of all 10 RCTs are presented in Table 29.

Authors (Year) Study Design (PEDro Score) Sample Size _{start}	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)		
Sample Size _{end}				
Intermittent TBS versus sham stimulation				
eq:statestatestatestatestatestatestatestat	E1: Intermittent Theta-Burst Stimulation E2: Low Frequency Repetitive Transcranial Magnetic Stimulation C: Sham Stimulation Duration: <i>Not Specified</i>	E1 vs C: Fugl-Meyer Assessment (-) Stroke Impairment Assessment Set (-) Modified Ashworth Scale (-) Grip Strength (-)		
Ackerley et al. (2016) RCT (8) N _{Start} =18 N _{End} =18 TPS=Chronic	E: iTBS C: Sham TBS Duration: 45min/d, 5d/wk for 2wk	 Action Research Arm Test (+exp) Fugl-Meyer Assessment (-) 		
Volz et al. (2016) RCT (5) N _{start} =26 N _{End} =17 TPS=Acute	E: iTBS C: Sham TBS Duration: <i>Not Specified</i>	 Grip Strength (+exp) Jebsen Taylor Hand Function Test (-) 		
Kim et al. (2015) RCT (8) N _{Start} =15 N _{End} =15 TPS=Chronic	E: iTBS C: Sham TBS Duration: 30min/d, 3d/wk for 4wk	 Modified Tardieu Scale (+exp) Peak torque (+exp) Modified Ashworth Scale (+exp) 		
Hsu et al. (2013) RCT (7) N _{start} =12 N _{end} =12 TPS=Subacute	E: iTBS C: Sham Duration: 30min/d, 3d/wk for 3wk	 Fugl-Meyer Assessment (+exp) Action Research Arm Test (-) 		
Talelli et al. (2012) RCT (7) Nstart=41 Nend=41 TPS=Chronic	E: iTBS C: Sham iTBS Duration: 1hr/d, 5d/wk for 2wk	 Nine Hole Peg Test (-) Jebsen Taylor Hand test (-) 		
Intermitte	nt TBS + low frequency rTMS versus sham TB	S and/or sham rTMS		
Sung et al. (2013) RCT (6) N _{Start} =54 N _{End} =54 TPS= Chronic	E1: Low frequency (1Hz) rTMS + iTBS E2: Sham rTMS + iTBS E3: Low frequency (1Hz) rTMS + sham iTBS C: Sham rTMS + sham Itbs Duration: 45min/d, 5d/wk for 4wk	 E1/E2/E3 vs C Wolf Motor Function test (+exp, +exp₂, +exp₃) Fugl-Meyer Assessment (+exp, +exp₂, +exp₃) Medical Research Council Scale (+exp, +exp₂, +exp₃) Functional Independence Measure (-) E1 vs E2 Wolf Motor Function test (+exp) 		
		 Fugl-Meyer Assessment (+exp) Medical Research Council Scale (+exp) Functional Independence Measure (-) <u>E1 vs E3</u> Wolf Motor Function test (+exp) Fugl-Meyer Assessment (-) Medical Research Council Scale (+exp) 		

Table 29. RCTs evaluating TBS interventions for upper extremity motor rehabilitation

		 Functional Independence Measure (-) E2 vs E3 Wolf Motor Function test (-) Fugl-Meyer Assessment (+exp₃) Medical Research Council Scale (+exp₃) Functional Independence Measure (-) 	
Continuous TBS versus sham stimulation			
Di Lazzaro et al. (2016) RCT (7) N _{Start} =20 N _{End} =17 TPS=Chronic	E: cTBS + robotic therapy C: Sham TBS + robotic therapy Duration: 1hr/d, 5d/wk for 2wk	Fugl-Meyer Assessment (-)	
Di Lazzaro et al. (2014) RCT (6) N _{Start} =12 N _{End} =12 TPS=Chronic	E: cTBS C: Sham Duration: 40min/d, 5d/wk for 2wk	 Action Research Arm Test (-) Nine Hole Peg Test (-) Jebsen Taylor hand test (-) Grasp strength (-) Pinch strength (-) 	
Talelli et al. (2012) RCT (7) Nstart=41 Nend=41 TPS=Chronic	E: cTBS C: Sham cTBS Duration: 1hr/d, 5d/wk for 2wk	 Nine Hole Peg Test (-) Jebsen Taylor Hand test (-) 	

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at $\alpha{=}0.05$

Conclusions about TBS

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1a	iTBS may not have a difference in efficacy when compared to sham stimulation for improving motor function.	6	Watanabe et al. 2018; Ackerley et al. 2016; Volz et al. 2016; Kim et al. 2015; Hsu et al. 2013; Talelli et al. 2012
1a	cTBS may not have a difference in efficacy when compared to sham stimulation for improving motor function.	3	Di Larazzo et al. 2016; Di Larazzo et al. 2014; Talelli et al. 2012
1b	iTBS combined with low frequency rTMS may produce greater improvements in motor function than sham stimulation with or without iTBS.	1	Sung et al. 2013
1b	There is conflicting evidence about the effect of iTBS combined with low frequency rTMS to improve motor function when compared to sham stimulation with low frequency rTMS .	1	Sung et al. 2013

MUSCLE STRENGTH			
LoE	Conclusion Statement	RCTs	References
1a	iTBS may produce greater improvements in muscle strength than sham stimulation .	4	Watanabe et al. 2018; Volz et al. 2016; Kim et al. 2015; Sung et al. 2013
1b	cTBS may not have a difference in efficacy when compared to sham stimulation for improving muscle strength.	1	Di Larazzo et al. 2014
1b	iTBS combined with low frequency rTMS may produce greater improvements in muscle strength than sham stimulation with or without iTBS.	1	Sung et al. 2013

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
1b	iTBS may not have a difference in efficacy when compared to sham stimulation for dexterity.	1	Talelli et al. 2012	
1a	cTBS may not have a difference in efficacy when compared to sham stimulation for dexterity.	2	Di Lazzero et al. 2014; Talelli et al. 2012	
ACTIVITIES OF DAILY LIVING				
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LoE	Conclusion Statement	RCTs	References	
1b	iTBS may not have a difference in efficacy when compared to sham stimulation for improving performance of activities of daily living.	2	Watanabe et al. 2018; Sung et al. 2013	
1b	iTBS combined with low frequency rTMS may not have a difference in efficacy when compared to sham stimulation or low frequency rTMS and iTBS on their own for improving performance of activities of daily living.	1	Sung et al. 2013	

SPASTICITY

LoE	Conclusion Statement	RCTs	References
1b, 2	There is conflicting evidence about the effect of iTBS to improve spasticity when compared to sham stimulation .	2	Watanabe et al. 2018; Kim et al. 2015

Key points

Theta burst stimulation alone may not be beneficial for upper limb function following stroke, however it may be beneficial for certain aspects of upper limb function when used in combination with repetitive transcranial magnetic stimulation.

Transcranial Direct Current Stimulation (tDCS)



Another form of non-invasive brain stimulation is transcranial direct-current stimulation (tDCS). This procedure involves the application of mild electrical currents (1-2 mA) conducted through two saline-soaked, surface electrodes applied to the scalp, overlaying the area of interest and the contralateral forehead above the orbit. Anodal stimulation is performed over the affected hemisphere and increases cortical excitability, while cathodal stimulation is performed over the unaffected hemisphere and decreases cortical excitability (Alonso-Alonso et al., 2007). Additionally, tDCS can be applied on both hemispheres concurrently, this is known as dual tDCS. In contrast to TMS, tDCS does not induce action potentials, but instead modulates the resting membrane potential of the neurons (Alonso-Alonso et al., 2007).

45 RCTs were found that evaluated tDCS interventions for upper extremity motor rehabilitation (Dehem et al. 2018; Shaheiwola et al. 2018; Andrade et al. 2017; Del Felice et al. 2017; Hong et al. 2017; Koh et al. 2017; Marquez et al. 2017; Mazzoleni et al. 2017; Pavlova et al. 2017; Rabadi et al. 2017; Takebayshi et al. 2017; Allman et al. 2016; Figlewski et al. 2016; Goodwill et al. 2016; Ilic et al. 2016; Mortensen et al. 2016; Powell et al. 2016; Rocha et al. 2016; Straudi et al. 2016; Ang et al. 2015; Cunningham et al. 2015; Lee et al. 2015; Sattler et al. 2015; Sik et al. 2015; Triccas et al. 2015; Au Yeung et al. 2014; Cha et al. 2014; Fusco et al. 2014; Hendy et al. 2013; Lefebvre et al. 2013; Wu et al. 2013; Stagg et al. 2012; Zimmerman et al. 2012; Hesse et al. 2011; Tanaka et al. 2011; Kim et al. 2010; Lindenberg et al. 2010; Kim et al. 2009; Boggio et al. 2007; Fregni et al. 2005).

17 RCTs compared anodal tDCS to sham stimulation (Andrade et al. 2017; Marquez et al. 2017; Pavlova et al. 2017; Allman et al. 2016; Ilic et al. 2016; Mortensen et al. 2016; Sik et al. 2015; Au Yeung et al. 2014; Fusco et al. 2013; Khedr et al. 2013; Stagg et al. 2012; Hesse et al. 2011; Tanaka et al. 2011; Kim et al. 2010; Kim et al. 2009; Boggio et al. 2007; Fregni et al. 2005).

14 RCTs compared cathodal tDCS to sham stimulation or conventional therapy (Marquez et al. 2017; Rabadi et al. 2017; Lee et al. 2015; Au Yeung et al. 2014; Fusco et

2013; Khedr et al. 2013; Wu et al. 2013; Stagg et al. 2012; Zimmerman et al. 2012; Hesse et al. 2011; Kim et al. 2010; Boggio et al. 2007; Fregni et al. 2005).

Eight RCTs compared dual tDCS to sham stimulation or conventional therapy (Koh et al. 2017; Goodwill et al. 2016; Sik et al. 2015; Cha et al. 2014; Lefebvre et al. 2014; Fusco et al. 2013; Lefebvre et al. 2013; Lindenberg et al. 2010).

Five RCTs compared anodal tDCS versus cathodal tDCS (Khedr et al. 2013; Stagg et al. 2012; Hesse et al. 2011; Boggio et al. 2007; Fregni et al. 2005). One RCT compared cathodal tDCS to dual tDCS (Del Felice et al. 2017). One RCT combined anodal tDCS with strength training (Hendy et al. 2014). Three RCTs compared anodal or cathodal tDCS with CIMT to sham stimulation with CIMT (Figlewski et al. 2016; Rocha et al. 2016; Cunningham et al. 2015). One RCT combined dual tDCS with cyclic NMES and CIMT (Takebayshi et al. 2017).

Four RCTs compared dual or anodal tDCS with robotics compared to sham stimulation with robotics or robotics alone (Dehem et al. 2018; Mazzoleni et al. 2017; Straudi et al. 2016; Triccas et al. 2015). One RCT compared anodal tDCS with robotics to cathodal tDCS with robotics (Ochi et al. 2013). Two RCTs compared anodal or dual tDCS with brain computer interfaces to sham stimulation with brain computer interfaces (Hong et al. 2017; Ang et al. 2015). One RCT compared dual tDCS with functional electrical stimulation to sham tDCS with functional electrical stimulation (Shaheiwola et al. 2018). Two RCTs compared anodal tDCS with or without peripheral nerve stimulation to peripheral nerve stimulation (Powell et al. 2016; Sattler et al. 2015). One RCT compared dual tDCS with low frequency rTMS and mirror therapy to sham tDCS and mirror therapy.

Two RCTs compared anodal or cathodal tDCS with virtual reality to virtual reality interventions with or without sham stimulation (Lee et al. 2014; Viana et al. 2014).

The methodological details and results of all 45 RCTs are presented in Table 30.

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)			
Time post stroke category	Anodel tDCS versus sham stimulation				
Andrade et al. (2017) RCT (9) N _{start} =60 N _{End} =60 TPS=Subacute	E1: Anodal Transcranial Direct Current Stimulation in Ipsilesional M1 and Constraint Induced Movement Therapy E2: Anodal Transcranial Direct Current Stimulation in Ipsilesional PMC and Constraint Induced Movement Therapy C: Sham Stimulation and Constraint Induced Movement Therapy Duration: 30min/d, 5d/wk for 6wk	E2 vs E1/C Fugl-Meyer Assessment (+exp ₂) Modified Ashworth Scale (+exp ₂) Box and Block Test (+exp ₂) Medical Research Council (+exp ₂) Barthel Index (+exp ₂)			
$\label{eq:states} \begin{array}{c} \displaystyle \frac{\text{Pavlova et al. 2017}}{\text{RCT}(7)} \\ \displaystyle \text{N}_{\text{Start}} = 11 \\ \displaystyle \text{N}_{\text{End}} = 11 \\ \displaystyle \text{TPS=Chronic} \end{array}$	E: Anodal tDCS C: Sham tDCS Duration: 20min (2x/d), 5d/wk for 4wk	 Fugl-Meyer Assessment (-) Wolf Motor Function Test (-) Box and Block Test (-) 			
Allman et al. (2016) RCT (7) N _{Start} =26 N _{End} =24 TPS=Chronic	E: Anodal tDCS C: Sham tDCS Duration: 1hr/d, for 9d	 Action Research Arm Test (+exp) Wolf Motor Function Test (+exp) Fugl-Meyer Assessment (-) 			
llic et al. (2016) RCT (8) N _{Start} =26 N _{End} =25 TPS=Chronic	E: Anodal tDCS + occupational therapy C: Sham tDCS + occupational therapy Duration: 45min/d, 5d/wk for 2wk	 Jebsen-Taylor Hand Function Test (+exp) Fugl-Meyer Assessment (-) Grip Strength (-) 			
Mortensen et al. (2016) RCT (7) N _{Start} =16 N _{End} =15 TPS=Chronic	E: Anodal tDCS + occupational therapy C: Sham tDCS + occupational therapy Duration: 30min/d for 5d	 Grip Strength (+exp) Stroke Impact Scale (-) Jebsen-Taylor Hand Function Test (-) 			
Tanaka et al. (2011) RCT (6) N _{start} =8 N _{end} =8 TPS=Subacute	E: Anodal tDCS C: Sham Duration: 30min/d, 4d/wk for 5wk	Grip strength (-)			
<u>Kim et al</u> . (2009) RCT (7) N _{start} =10 N _{end} =10 TPS=Subacute	E: Anodal tDCS C: Sham Duration: 20min/d, 5d/wk for 6wk	 Box & Block Test (+exp) Finger acceleration (+exp) 			
	Cathodal tDCS versus sham stimulation or conv	ventional therapy			
Rabadi et al. 2017 RCT (7) N _{Start} =16 N _{End} =12 TPS=Acute	E: Cathodal tDCS C: Sham tDCS Duration: 30min/d, 5d/wk for 2wk	Action Research Arm Test (-)			
Lee et al. (2015) RCT (6) N _{Start} =24 N _{End} =24 TPS=Chronic	E: Cathodal tDCS + physical therapy C: Physical therapy Duration: 30min/d, 5d/wk for 4wk	Fugl-Meyer Assessment (+exp)			
Fusco et al. (2014) RCT (6)	E: Cathodal tDCS + active electrode C: Sham tDCS	 Canadian Neurologic Scale (-) Nine Hole Peg Test (-) 			

Table 30. RCTs evaluating tDCS interventions for upper extremity motor rehabilitation

Nstart=14 NEnd=11 TPS=Subacute	Duration: 45min/d, 5d/wk for 2wk	 Barthel Index (-) Fugl-Meyer Assessment (-) 	
Wu et al.(2013) RCT (9) Nstart=90 NEnd=90 TPS=Chronic	E: Cathodal tDCS C: Sham tDCS Duration: 20min/d, 5d/wk for 4wk	Modified Ashworth Scale (+exp)	
Zimerman et al. (2012) RCT (6) N _{start} =12 N _{end} =12 TPS=Chronic	E: Cathodal tDCS C: Sham tDCS Duration: <i>Not Specified</i>	Grip strength (-)	
Hummel et al. (2005) RCT (6) N _{start} =6 N _{end} =6 TPS=Chronic	E: Cathodal tDCS C: Sham tDCS Duration: 20min/d, 3d/wk for 4wk	Jebsen-Taylor Hand Function test (+exp)	
	Dual tDCS versus sham stimulation or conve	entional therapy	
Koh et al. (2017) RCT (8) N _{Start} =25 N _{End} =18 TPS=Chronic	E: Dual tDCS with Sensory Modulation C: Sham tDCS with Sensory Modulation Duration: 30min/d, 5d/wk for 8wk	 Fugl-Meyer Assessment (-) Modified Ashworth Scale (-) Action Research Arm Test (-) Barthel Index (-) 	
<u>Goodwill et al.</u> (2016) RCT (7) N _{Start} =16 N _{End} =15 TPS=Chronic	E: Dual tDCS + upper limb training C: Sham tDCS + upper limb training Duration: 30min/d, 5d/wk for 3wk	 Tardieu Scale (-) Grip Strength (-) 	
Lefebvre et al. (2015) RCT Crossover (5) N _{Start} =19 N _{End} =19 TPS=Chronic	E: Dual tDCS C: Sham tDCS Duration: 30min/d, 5d/wk for 3wk	Purdue Pegboard Test (+exp)	
<u>Cha et al</u> . (2014) RCT (6) N _{Start} =20 N _{End} =20 TPS=Chronic	E: Dual tDCS C: Conventional training Duration: 30min/d, 5d/wk for 4wk	 Fugl-Meyer Assessment (+exp) Box and Block Test (+exp) 	
Lefebvre et al. (2014) RCT (8) N _{Start} =19 N _{End} =19 TPS=Chronic	E: Dual tDCS C: Sham Duration: 20min/d, 5d/wk for 2wk	 Purdue Pegboard Test (+exp) Precision grip (+exp) 	
Lefebvre et al. (2013) RCT (8) N _{Start} =18 N _{End} =18 TPS=Chronic	E: Dual tDCS C: Sham Duration: 30min/d, 4d/wk for 3wk	 Purdue Pegboard Test (+exp) Maximal hand grip force (+exp) 	
Lindenberg et al. (2010) RCT (4) N _{start} =20 N _{end} =20 TPS=Chronic	E: Dual tDCS C: Sham Duration: 30min/d, 5d/wk for 3wk	 Fugl-Meyer Assessment (+exp) Wolf Motor Function Test (+exp) 	
	Anodal or cathodal tDCS versus sham stimulation		
Marquez et al. 2017 RCT Crossover (8) N _{Start} =25 N _{End} =25 TPS=Chronic	E1: Anodal tDCS E2: Cathodal tDCS C: Sham tDCS Duration: 20min/d for 6d	 <u>E1/E2 vs C</u> Jebsen-Taylor Hand Function test (-) Grip Strength (-) 	

Au-Yeung et al. (2014) RCT Crossover (8) N _{Start} =10 N _{End} =10 TPS=Chronic	E1: Anodal tDCS E2: Cathodal tDCS C: Sham Duration: <i>Not Specified</i>	E1/E2 vs C • Purdue Pegboard Test (-) • Pinch strength (-)			
Khedr et al. (2013) RCT (9) Nstart=40 NEnd=40 TPS= Chronic	E1: Anodal tDCS E2: Cathodal tDCS C: Sham Duration: 25min/d for 6d	 <u>E1/E2 vs C</u> Orgogozo MCA scale (+exp, +exp₂) National Institute of Health Stroke Scale (-) Barthel Index (+exp, +exp₂) Medical Research Council Scale (-) <u>E1 vs E2</u> Orgogozo MCA scale (-) National Institute of Health Stroke Scale (-) Barthel Index (-) Medical Research Council Scale (-) 			
<u>Stagg et al</u> . (2012) RCT (6) N _{start} =13 N _{end} =13 TPS=Chronic	E1: Anodal tDCS E2: Cathodal tDCS C: Sham Duration: 80min/d, 3d/wk for 4wk	E1/E2 vs C Grip strength (+exp, +exp ₂) E1 vs E2 Grip strength (-)			
Hesse et al. (2011) RCT (10) Nstart=96 Nend=85 TPS=Chronic	E1: Anodal tDCS E2: Cathodal tDCS C: Sham Duration: 20min/d, 5d/wk for 6wk	E1/E2 vs C • Fugl-Meyer Assessment (-) E1 vs E2 • Fugl-Meyer Assessment (-)			
<u>Kim et al</u> . (2010) RCT (7) N _{start} =18 N _{end} =16 TPS=Subacute	E1: Anodal tDCS E2: Cathodal tDCS C: Sham Duration: Not Specified	E2 vs C • Fugl-Meyer Assessment (+exp ₂) • Barthel Index (-) <u>E1 vs C</u> • Fugl-Meyer Assessment (-) • Barthel Index (-)			
Boggio et al. (2007) RCT (6) N _{start} =4 N _{end} =4 TPS=Chronic	E1: Anodal tDCS E2: Cathodal tDCS C: Sham Duration: 20min, 1x/wk for 4wk	 <u>E1/E2 vs C</u> Jebsen-Taylor Hand Function test (+exp, +exp₂) <u>E1 vs E2</u> Jebsen-Taylor Hand Function test (-) 			
Fregni et al. (2005) RCT (7) N _{start} =6 N _{end} =6 TPS= Chronic	E1: Anodal tDCS E2: Cathodal tDCS C: Sham Duration: <i>Not Specified</i>	E1/E2 vs C Jebsen Taylor Hand Function test: (+exp, +exp ₂) E1 vs E2 Jebsen Taylor Hand Function test: (-)			
Sik at al. (2015)	Anodal, cathodal or dual tDCS versus sham	stimulation			
<u>SIK et al.</u> (2015) RCT (6) N _{start} =36 N _{end} =31 TPS=Subacute	E1: Anodal tDCS + P1 + O1 E2: Dual tDCS + PT + OT C: Sham tDCS + PT + OT Duration: <i>Not Specified</i>	 E1/E2 VS C Wolf Motor Function Test (+exp, +exp2) Jebsen Taylor Hand Function Test (+exp, +exp2) Kocaeli Functional Evaluation Test (+exp2) E1 vs E2 Wolf Motor Function Test (-), Jebsen Taylor Hand Function Test (-) Kocaeli Functional Evaluation Test (-) 			
<u>Fusco et al.</u> (2013) RCT (7) N _{start} =9 N _{end} =9 TPS=Subacute	E1: Dual tDCS E2: Anodal tDCS E3: Cathodal tDCS C: Sham Duration: 15min/d for 2d	 <u>E1/E2/E3 vs C</u> Nine hole peg test (+exp, +exp₂, +exp₃) Grasp force (-) 			
	Cathodal versus dual tDCS stimulation				

Del Felice et al. 2017 RCT crossover (8) N _{Start} =10 N _{End} =10 TPS=Chronic	E: Cathodal Trans Direct Current Stimulation C: Dual tDCS Duration: 20min/d, 5d/wk for 3wk	 Modified Ashworth Scale (+exp) Bhakta Finger Flexion Scale (-) European Stroke Scale (-) Action Research Arm Test (-) Medical Research Council Scale (-) Barthel Index (-)
Anodal tD	CS with strength training compared to sham tD	CS with strength training
Hendy et al. (2014) RCT (7) N _{Start} =10 N _{End} =10 TPS=Chronic	E1: Strength training + anodal tDCS E2: Strength training + sham C: Anodal tDCS Duration: 20min/d, 2d/wk for 5wk	Maximum voluntary dynamic strength for wrist extensors (-)
	Anodal or cathodal tDCS with CIM	лт
Figlewski et al. (2016) RCT (7) N _{Start} =44 N _{End} =44 TPS=Chronic	E: CIMT + Anodal tDCS C: CIMT + Sham tDCS Duration: 6hr/d for 9d	 Wolf Motor Function Test (+exp) Grip Strength (-) Arm Strength (-)
Rocha et al. (2016) RCT (8) N _{Start} =21 N _{End} =21 TPS=Chronic	E1: Anodal tDCS with CIMT E2: Cathodal tDCS with CIMT C: Sham tDCS with CIMT Duration: 1hr/d, 6d/wk for 2wk	E1 vs C • Fugl-Meyer Assessment (+exp) • Motor Acivity Log (-) • Grip Strength (-) <u>E2 vs C</u> • Fugl-Meyer Assessment (-) • Motor Acivity Log (-) • Grip Strength (-)
Cunningham et al. (2015) RCT (6) N _{Start} =12 N _{End} =12 TPS=Chronic	E: anodal tDCS + CIMT C: Sham tDCS + CIMT Duration: 30min/d, 3d/wk for 10wk	 9 Hole Peg Test (-) Motor Activity Log (-) Fugl-Meyer Assessment (-)
	Dual tDCS with cyclic NMES and C	ІМТ
Takebayshi et al. 2017 RCT (7) N _{Start} =20 N _{End} =19 TPS=Chronic	E: Dual tDCS combined with cyclic NMES with CIMT C: CIMT Duration: 2hr (2x/d), 5d/wk for 3wk	 Fugl-Meyer Assessment (+exp) Motor Activity Log (+exp)
Dual or anod	al tDCS with robotics compared to sham tDCS	with robotics or robotics alone
Dehem et al. (2018) RCT-crossover (6) N _{Start} =21 N _{End} =20 TPS=Chronic	E: Dual tDCS with Upper Limb Robotic Assisted Therapy C: Sham tDCS with Upper Limb Robotic Assisted Therapy Duration: 45min/d, 5d/wk for 6wk	 Box and Block Test (+exp) Purdue Pegboard Test (-)
Straudi et al. (2016) RCT (6) N _{Start} =23 N _{End} =23 TPS=Subacute and chronic	E: Robot-assisted therapy + dual tDCS C: Robot-assisted therapy + sham tDCS Duration: 45min/d, 5d/wk for 2wk	 Fugl-Meyer Assessment (-) Box and Block Test (-) Motor Acivity Log (-)
Mazzoleni et al. 2017 RCT (7) N _{Start} =24 N _{End} =24 TPS=Acute	E: Anodal tDCS with Wrist Robot-Assisted Training C: Wrist Robot-Assisted Training Duration: <i>Not Specified</i>	 Fugl-Meyer Assessment (-) Modified Ashworth Scale (-) Motricity Index (-) Box and Block Test (-)
Triccas et al. (2015) RCT (8) N _{start} =23 N _{end} =22 TPS=Subacute	E: Anodal tDCS + robotic ArmeoSpring C: Sham tDCS + robotic ArmeoSpring Duration: 45min/d, 3d/wk for 4wk	 Fugl-Meyer Assessment (-) Action Research Arm Test (-) Motor Activity Log (-) Stroke Impact Scale (-)

Anodal versus cathodal tDCS stimulation with robotics				
Ochi et al. (2013)	E: Anodal tDCS on affected hemisphere + robot	•	Modified Ashworth Scale (+exp)	
RCT (7)	assisted arm training		Fugl-Mever Assessment (-)	
Nstart=18	C: Cathodal tDCS on unaffected hemisphere +		Motor Activity Log (-)	
Nend=16	robot assisted arm training			
TPS=Chronic	Duration: 45min/d. for 5d			
Anoda	al or dual tDCS with brain computer interface-as	ssist	ted motor imagery	
Hong et al. (2017)	E: Brain computer interface -Assisted Motor		Fugl-Meyer Assessment (-)	
PCT (5)	Imagery with Dual tDCS	•		
Noted =19	C: Brain computer interface -Assisted Motor			
	Imagery with Sham tDCS			
TPS-Chronic	Duration: 20min/d. 5d/wk for 2wk			
	Et Anodol tDCS + motor imagon / broin computer	-	Fuel Mover Assessment ()	
$\frac{\text{Ang et al.}}{\text{POT}(c)}$	E. Anodal IDCS + motor imagery brain computer	•	rugi-meyer Assessment (-)	
	C: Sham tDCS I mater imagery brain computer			
NStart = 19 N= -10	c. Sham LDCS + motor imagery brain computer			
TPS-Chronic	Duration: 80min/d. 5d/wk for 4wk			
Chabaiwala at al. 2012				
Snaneiwola et al. 2018		•	rugi-weyer Assessment (+exp)	
RCT (6)	C: Sham tDCS with FES	•	Wolf Motor Function Test Score (+exp)	
NStart =30	Duration: 45min/d, 5d/wk for 4wk	•	Modified Ashworth Scale (-)	
NEnd =30				
IPS=Chronic				
	Anodal tDCS with peripheral nerve sti	imul	ation	
Powell et al. (2016)	E1: Anodal tDCS followed by peripheral nerve	•	Fugl-Meyer Assessment (-)	
RCT (8)	stimulation	•	Stroke Impact Scale (-)	
N _{Start} =11	E2: Peripheral nerve stimulation followed by			
N _{End} =10	tDCS			
TPS=Chronic	Duration: Not Specified			
Sattler et al. (2015)	E: Repetitive peripheral nerve stimulation +	•	Jebsen Hand Function Test (+exp)	
RCT (7)	anodal tDCS	•	Grip Strength (-)	
N _{Start} =20	C: Repetitive peripheral nerve stimulation	•	9 Hole Peg Test (-)	
N _{End} =20	Duration: 20min/d, 5d/wk for 4wk	•	Hand Tapping Test (-)	
TPS=Acute		•	Fugl-Meyer Assessment (-)	
	Dual tDCS with low frequency rTMS and mi	irror	therapy	
D'Agata et al. (2016)	E: Dual tDCS + low frequency (1Hz) rTMS +		Action Research Arm Test (+exp)	
RCT (6)	Mirror Therapy			
N _{Start} =34	C: Sham tDCS + Mirror Therapy			
N _{End} =34	Duration: 1hr/wk, 5d/wk for 2wk			
TPS=Chronic				
	Anodal or cathodal tDCS with virtual	real	ity	
Lee et al. (2014)	E1: cathodal tDCS		<u>E1 vs E2</u>	
RCT (7)	E2: Virtual reality	•	Manual Function Test (+exp)	
Nstart=64	E3: Cathodal tDCS + virtual reality	•	Fugl-Meyer Assessment (+exp)	
NEnd=59	Duration: 90min/d, 3d/wk for 4wk	•	Modified Barthel Index (-)	
TPS=Chronic		•	Manual Muscle Test (-)	
		•	Modified Ashworth Scale (-)	
		•	Box and Block Test (-)	
			<u>E3 vs E2/E1</u>	
		•	Manual Function Test (+exp ₃)	
		•	Fugl-Meyer Assessment (+exp ₃)	
		•	Modified Barthel Index (-)	
		•	Madified Ashurarth Carls ()	
		•	IVIOUIIIEO ASNWORN SCAIE (-)	
		•	BOX and BIOCK LEST (-)	
		+		
$\frac{\text{viana et al.}}{\text{DOT}}$ (2014)	E: VIITUAI reality + anodal tDCS	•	Fugi-ivieyer Assessment (-)	
KUI (9)	C: VIITual reality + snam	•	VVOII IVIOTOF FUNCTION LEST (-)	
	Duration: Thr/d, 3d/wk tor 5WK	•	iviodilied Ashworth Scale (-)	

N _{Start} =20	Grip strength (-)
N _{End} =20	
TPS=Chronic	

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about tDCS

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of anodal tDCS to improve motor function when compared to sham stimulation .	11	Andrade et al. 2017; Marquez et al. 2017; Pavlova et al. 2017; Allman et al. 2016; Ilic et al. 2016; Mortensen et al. 2016; Sik et al. 2015; Hesse et al. 2011; Kim et al. 2010; Boggio et al. 2007; Fregni et al. 2005	
1a	There is conflicting evidence about the effect of cathodal tDCS to improve motor function when compared to sham stimulation or conventional therapy .	9	Maquez et al. 2017; Rabadi et al. 2017; Lee et al. 2015; Fusco et al. 2014; Hesse et al. 2011; Kim et al. 2010; Boggio et al. 2007; Fregni et al. 2005	
1a	Dual tDCS may produce greater improvements in motor function than sham stimulation or conventional therapy.	4	Koh et al. 2017; Sik et al. 2015; Cha et al. 2014; Lindenberg et al. 2010	
1a	Anodal tDCS may not have a difference in efficacy when compared to cathodal tDCS for improving motor function.	3	Hesse et al. 2011; Boggio et al. 2007; Fregni et al. 2005	
1b	Cathodal tDCS may not have a difference in efficacy when compared to dual tDCS for improving motor function.	1	Del Felice et al. 2017	
1a	There is conflicting evidence about the effect of anodal tDCS with CIMT to improve motor function when compared to sham tDCS with CIMT .	3	Figlewski et al. 2016; Rocha et al. 2016; Cunningham et al. 2015	
1b	Cathodal tDCS with CIMT may not have a difference in efficacy when compared to sham tDCS with CIMT for improving motor function.	1	Rocha et al. 2016	
1b	Dual tDCS with cyclic NMES and CIMT may produce greater improvements in motor function than CIMT .	1	Takebayshi et al. 2017	
1b	Dual tDCS with upper limb robotics may not have a difference in efficacy when compared to sham tDCS with upper limb robotics for improving motor function.	1	Straudi et al. 2016	
1a	Anodal tDCS with upper limb robotics may not have a difference in efficacy when compared to sham tDCS with upper limb robotics or upper limb robotics alone for improving motor function.	2	Mazzoleni et al. 2017; Triccas et al. 2015	
1b	Anodal tDCS with upper limb robotics may not have a difference in efficacy when compared to cathodal tDCS with upper limb robotics for improving motor function.	1	Ochi et al. 2013	
1b	Anodal or dual tDCS with brain computer interface- assisted motor imagery interventions may not have a difference in efficacy when compared to sham tDCS with brain computer interface-assisted motor imagery interventions for improving motor function.	2	Hong et al. 2017; Ang et al. 2015	

1b	Dual tDCS with FES may produce greater improvements in motor function than sham tDCS with FES.	1	Shaheiwola et al. 2018
1a	Anodal tDCS with peripheral nerve stimulation may not have a difference in efficacy when compared to peripheral nerve stimulation for improving motor function.	2	Powell et al. 2016; Sattler et al. 2015
1b	Dual tDCS with low frequency rTMS and mirror therapy may produce greater improvements in motor function than sham tDCS with mirror therapy.	1	D'Agata et al. 2016
1a	There is conflicting evidence about the effect of anodal or cathodal tDCS with virtual reality training to improve motor function when compared to virtual reality training with or without sham tDCS.	2	Lee et al. 2014; Viana et al. 2014
1b	Cathodal tDCS may produce greater improvements in motor function than virtual reality training .	1	Lee et al. 2014

STROKE SEVERITY				
LoE	Conclusion Statement	RCTs	References	
1b	There is conflicting evidence about the effect of anodal tDCS to produce greater improvements on measures of stroke severity when compared to sham stimulation .	1	Khedr et al. 2013	
1a	Cathodal tDCS may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improvements on measures of stroke severity.	2	Fusco et al. 2014; Khedr et al. 2013	
1b	Anodal tDCS may not have a difference in efficacy when compared to cathodal tDCS for improvements on measures of stroke severity.	1	Khedr et al. 2013	
1b	Cathodal tDCS may not have a difference in efficacy when compared to dual tDCS for improvements on measures of stroke severity.	1	Del Felice et al. 2017	

DEXTERITY					
LoE	Conclusion Statement	RCTs	References		
1a	There is conflicting evidence about the effect of anodal tDCS to improve dexterity when compared to sham stimulation .	5	Andrade et al. 2017; Pavlova et al. 2017; Kim et al. 2009; Au Yeung et al. 2014; Fusco et al. 2013		
1a	Cathodal tDCS may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improving dexterity.	3	Au Yeung et al. 2014; Fusco et al. 2014; Fusco et al. 2013		
1a	Dual tDCS may produce greater improvements in dexterity than sham stimulation or conventional therapy .	5	Lefebvre et al. 2015; Cha et al. 2014; Lefebvre et al. 2014; Lefebvre et al. 2013; Fusco et al. 2013		
1b	Anodal tDCS with CIMT may not have a difference in efficacy when compared to sham tDCS with CIMT for improving dexterity.	1	Cunningham et al. 2015		
1a	Dual tDCS with upper limb robotics may not have a difference in efficacy when compared to sham tDCS with upper limb robotics for improving dexterity.	2	Dehem et al. 2018; Straudi et al. 2016		
1b	Anodal tDCS with upper limb robotics may not have a difference in efficacy when compared to sham tDCS with upper limb robotics or upper limb robotics alone for improving dexterity.	1	Mazzoleni et al. 2017		
1b	Anodal tDCS with peripheral nerve stimulation may not have a difference in efficacy when compared to peripheral nerve stimulation for improving dexterity.	1	Sattler et al. 2015		
1b	Anodal or cathodal tDCS with virtual reality training may not have a difference in efficacy when compared to virtual reality training with or without sham tDCS for improving dexterity.	1	Lee et al. 2014		
1b	Cathodal tDCS may not have a difference in efficacy when compared to virtual reality training for improving dexterity.	1	Lee et al. 2014		

SPASTICITY				
LoE	Conclusion Statement	RCTs	References	
1b	Anodal tDCS may produce greater improvements in spasticity than sham stimulation .	1	Andrade et al. 2017	
1b	Cathodal tDCS may produce greater improvements in spasticity than sham stimulation or conventional therapy.	1	Wu et al. 2013	
1a	Dual tDCS may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improving spasticity.	2	Koh et al. 2017; Goodwill et al. 2016	
1b	There is conflicting evidence about the effect of cathodal tDCS to improve spasticity when compared to dual tDCS .	1	Del Felice et al. 2017	
1b	Anodal tDCS with upper limb robotics may not have a difference in efficacy when compared to sham tDCS with upper limb robotics or upper limb robotics alone for spasticity.	1	Mazzoleni et al. 2017	
1b	Anodal tDCS with upper limb robotics may produce greater improvements in spasticity than cathodal tDCS with upper limb robotics.	1	Ochi et al. 2013	
1b	Dual tDCS with FES may not have a difference in efficacy when compared to sham tDCS with FES for spasticity.	1	Shaheiwola et al. 2018	
1a	Anodal or cathodal tDCS with virtual reality training may not have a difference in efficacy when compared to virtual reality training with or without sham tDCS for improving spasticity.	2	Lee et al. 2014; Viana et al. 2014	
1b	Cathodal tDCS may not have a difference in efficacy when compared to virtual reality training for improving spasticity.	1	Lee et al. 2014	

ACTIVITIES OF DAILY LIVING				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of anodal tDCS to improve performance of activities of daily living when compared to sham stimulation .	4	Andrade et al. 2017; Mortensen et al. 2016; Khedr et al. 2013; Kim et al. 2010	
1a	There is conflicting evidence about the effect of cathodal tDCS to improve performance of activities of daily living when compared to sham stimulation or conventional therapy .	3	Fusco et al. 2014; Khedr et al. 2013; Kim et al. 2010	
1b	Dual tDCS may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improving performance of activities of daily living.	1	Koh et al. 2017	
1b	Anodal tDCS may not have a difference in efficacy when compared to cathodal tDCS for improving performance of activities of daily living.	1	Khedr et al. 2013	
1b	Cathodal tDCS may not have a difference in efficacy when compared to dual tDCS for improving performance of activities of daily living.	1	Del Felice et al. 2017	
1a	Anodal tDCS with CIMT may not have a difference in efficacy when compared to sham tDCS with CIMT for improving performance of activities of daily living.	2	Rocha et al. 2016; Cunningham et al. 2015	
1b	Cathodal tDCS with CIMT may not have a difference in efficacy when compared to sham tDCS with CIMT for improving performance of activities of daily living.	1	Rocha et al. 2016	
1b	Dual tDCS with upper limb robotics may not have a difference in efficacy when compared to sham tDCS with upper limb robotics for improving performance of activities of daily living.	1	Straudi et al. 2016	
1b	Anodal tDCS with upper limb robotics may not have a difference in efficacy when compared to sham tDCS with upper limb robotics or upper limb robotics alone for improving performance of activities of daily living.	1	Triccas et al. 2015	
1b	Anodal tDCS with upper limb robotics may produce greater improvements in performance of activities of daily living than cathodal tDCS with upper limb robotics.	1	Ochi et al. 2013	
1b	Anodal tDCS with peripheral nerve stimulation may not have a difference in efficacy when compared to peripheral nerve stimulation for improving performance of activities of daily living.	1	Powell et al. 2016	
1b	Anodal or cathodal tDCS with virtual reality training may not have a difference in efficacy when compared to virtual reality training with or without sham tDCS for improving performance of activities of daily living.	1	Lee et al. 2014	

1	b	

Cathodal tDCS may not have a difference in efficacy
when compared to virtual reality training for
improving performance of activities of daily living.Lee et al. 2014

MUSCLE STRENGTH					
LoE	Conclusion Statement	RCTs	References		
1a	Anodal tDCS may not have a difference in efficacy when compared to sham stimulation for improving muscle strength.	9	Andrade et al. 2017; Marquez et al. 2017; llic et al. 2016; Mortensen et al. 2016; Au Yeung et al. 2014; Fusco et al. 2013; Khedr et al. 2013; Stagg et al. 2012; Tanaka et al. 2011		
1a	Cathodal tDCS may not have a difference in efficacy when compared to sham stimulation or conventional therapy for improving muscle strength.	6	Marquez et al. 2017; Au Yeung et al. 2014; Khedr et al. 2013; Fusco et al. 2013; Stagg et al. 2012; Zimmerman et al. 2012		
1a	There is conflicting evidence about the effect of dual tDCS to improve muscle strength when compared to sham stimulation or conventional therapy.	4	Goodwill et al. 2016; Lefebvre et al. 2014; Fusco et al. 2013; Lefebvre et al. 2013		
1a	Anodal tDCS may not have a difference in efficacy when compared to cathodal tDCS for improving muscle strength.	2	Khedr et al. 2013; Stagg et al. 2012		
1b	Cathodal tDCS may not have a difference in efficacy when compared to dual tDCS for improving muscle strength.	1	Del Felice et al. 2017		
1b	Anodal tDCS with strength training may not have a difference in efficacy when compared to sham tDCS with strength training for improving muscle strength.	1	Hendy et al. 2014		
1a	Anodal tDCS with CIMT may not have a difference in efficacy when compared to sham tDCS with CIMT for improving muscle strength.	2	Figlewski et al. 2016; Rocha et al. 2016		
1b	Cathodal tDCS with CIMT may not have a difference in efficacy when compared to sham tDCS with CIMT for improving muscle strength.	1	Rocha et al. 2016		
1b	Anodal tDCS with upper limb robotics may not have a difference in efficacy when compared to sham tDCS with upper limb robotics or upper limb robotics alone for improving muscle strength.	1	Mazzoleni et al. 2017		
1b	Anodal tDCS with peripheral nerve stimulation may not have a difference in efficacy when compared to peripheral nerve stimulation for improving muscle strength.	1	Sattler et al. 2015		
1a	Anodal or cathodal tDCS with virtual reality training may not have a difference in efficacy when compared to virtual reality training with or without sham tDCS for improving muscle strength.	2	Lee et al. 2014; Viana et al. 2014		
1b	Cathodal tDCS may not have a difference in efficacy when compared to virtual reality training for improving muscle strength.	1	Lee et al. 2014		

Key points

The literature is mixed regarding anodal, cathodal, or dual transcranial direct current stimulation, alone or in combination with other therapy approaches, for upper limb rehabilitation following stroke.

Pharmaceuticals Botulinum toxin



Adopted from: http://www.theinvestor.co.kr/view.php?ud=20180104000712

Botulinum toxin exerts a therapeutic effect by reducing overactivity in spastic muscles through blocking the release of acetylcholine at the neuromuscular junction. The benefits of botulinum toxin injections are generally dose-dependent and last approximately 2 to 4 months (Brashear et al. 2002; Francisco et al. 2002; Simpson et al. 1996; Smith et al. 2000). One of the advantages of botulinum toxin is that it is safe to use on small, localized areas or muscles, such as those in the upper extremity. Unlike chemodenervation and neurolytic procedures like phenol or alcohol, botulinum toxin is not associated with skin sensory loss or dysesthesia (Suputtitada & Suwanwela, 2005). Dynamic EMG studies can be helpful in determining which muscles should be injected (Bell & Williams, 2003).

Interventions for 35 RCTs using botulinum toxin included: 19 RCTs looked at botulinum toxin A compared to placebo (Rosales et al. 2018; Elovic et al. 2016; Gracies et al. 2015; Hesse et al. 2012; Marciniak et al. 2012; Wolf et al. 2012; Shaw et al. 2011; Kaji et al. 2010; Shaw et al. 2010; Meythaler et al. 2009; Simpson et al. 2009; Jahangir et al. 2007; Suputtitada and Suwanwela, 2005; Childers et al. 2004; Brashear et al. 2002; Bakheit et al. 2001; Bhakta et al. 2000; Smith et al. 2000; Simpson et al. 1996). Two RCTs looked at botulinum toxin B compared to placebo (Gracies et al. 2014; Brashear et al. 2004). One RCT looked at botulinum toxin A with upper limb rehabilitation compared to botulinum toxin A alone (Devier et al. 2017). Four RCTs looked at OnabotulinumtoxinA compared to letibotulinumtoxinA, NABOTA, Neurnox or tizanidine (Do et al. 2017; Nam et al. 2015; Seo et al. 2015; Simpson et al. 2009). A single RCT looked at high versus low dosage botulinum toxin A (Francisco et al. 2002). A single RCT looked at botulinum toxin A combined with adhesive taping versus botulinum toxin A combined with manual muscle stretching, passive articular mobilization, and palmar splinting (Santamato et al. 2015). Three RCTs looked at ultrasound guided botulinum toxin A injections versus other approaches (Zeuner et al. 2017; Picelli et al. 2014; Santamato et al. 2014). Two RCTs looked at botulinum toxin A combined with NMES (Marvulli et al. 2016; Hesse et al. 1998). A single RCT looked at botulinum toxin A combined with mCIMT compared to botulinum toxin A (Sun et al.

2010). Finally, a single RCT looked at botulinum toxin A combined with task-specific training compared to task-specific training alone (Umar et al. 2018).

The methodological details and results of all 35 RCTs evaluating rTMS for the upper extremity motor rehabilitation are presented in Table 31.

Table 31. RCTs evaluating botulinum toxin injections for upper extremity motor rehabilitation			
Authors (Voor)	Interventions	Outcomo Moasuros	

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Botulinu	m toxin A versus placebo, no injection or	conventional rehabilitation
Rosales et al. (2018) RCT (7) Nstart =42 NEnd =40 TPS=Subacute	E: Abobotulinumtoxin A 500U C: Placebo	 Modified Ashworth Scale (+exp) Upper extremity active motor function (-)
Elovic et al. (2016) RCT (6) N _{Start} =317 N _{End} =299 TPS=Chronic	E: 400U incobotulinumtoxinA C: Placebo	 Ashworth Scale (+exp) Disability Assessment Scale (+exp)
Gracies et al. (2015) RCT (8) NStart=243 NEnd=229 TPS=Chronic	E1: Single 500U AbobotulinumtoxinA E2: Single 1000U AbobotulinumtoxinA C: Placebo	 <u>E1/E2 vs. C</u> Modified Ashworth Scale (+exp, +exp₂) Disability Assessment Scale (-)
<u>Hesse et al</u> . (2012) RCT (7) N _{start} =18 N _{end} =18 TPS=Acute	E: 150U Xeomin C: No injection	 Modified Ashworth Scale score (+exp) Resistance to Passive Movement Scale (+exp) Fugl-Meyer Assessment (-)
Marciniak et al. (2012) RCT (5) N _{Start} =21 N _{End} =19 TPS=Chronic	E: 100-150U of botulinum toxin type A (BTX-A) into the pectoralis major and teres major muscles in the shoulder extensors. C: Placebo	 Modified Ashworth Scale (-) Passive range of motion (+exp) Fugl-Meyer Assessment (+exp) Functional Independence Measure (-) Disability Assessment Scale (+exp)
Wolf et al. (2012) RCT (9) Nstart=25 Nend=22 TPS=Chronic	E: 300U Botox (BTX-A) C: Placebo	Wolf Motor Function test (-)
<u>Shaw et al</u> . (2011) RCT (8) N _{start} =333 N _{end} =329	E: 100-200 U Dysport + 4 weeks therapy C: Therapy only	 Action Research Arm Test (-) Modified Ashworth Scale (+exp) 9-Hole Peg Test (-) Barthel Index (-)
Kaji et al. (2010) RCT (9) Nstart=109 Nend=109 TPS=Chronic	E1: 120 U Botox (BoNTA) C1: Placebo E2: 200 U Botox (BoNTA) C2: Placebo	 <u>E2 vs C2</u> Modified Ashworth Scale (+exp₂) Disability Assessment Scale (+exp₂) <u>E1 vs C1</u> Modified Ashworth Scale (-) Disability Assessment Scale (+exp₁)
<u>Shaw et al.</u> (2010) RCT (6)	E: Botulinum toxin type A (BTX-A, Dysport) injections + upper limb therapy	 Action Research Arm Test (-) Modified Ashworth Scale (+exp)

N _{Start} =333 N _{End} =199 TPS=Subacute <u>Meythaler et al</u> . (2009) RCT (6) N _{start} =21	C: Upper limb therapy E: 100 U Botox (BTX-A) + therapy C: Saline + therapy	 Motricity Index (+exp) Grip Strength (-) 9-Hole Peg Test (-) Barthel Index (-) Motor Activity Log (-) Ashworth Scale (-) Barthel Index (-)
Nend=18 TPS=Chronic		
Simpson et al. (2009) RCT (8) N _{start} =60 N _{end} =41 TPS=Subacute	E1: Up to 500 U of BoNT-Type A E2: Tizanidine C: Placebo	E1 vs C Modified Ashworth Scale (+exp) Disability Assessment Scale (+exp) E2 vs C Modified Ashworth Scale (-) Disability Assessment Scale (-) E1 vs E2 Modified Ashworth Scale (+exp) Disability Assessment Scale (+exp1)
<u>Jahangir et al</u> . (2007) RCT (6) N _{start} =27 N _{end} =27 TPS=Chronic	E: 50 U Botox (BTX-A) C: Placebo	 Modified Ashworth Scale (+exp) Barthel Index (-)
Suputtitada & Suwanwela (2005) RCT (6) N _{start} =45 N _{end} =40 TPS=Chronic	E1: 350U BTX (Dysport) E2: 500U BTX (Dysport) E3: 1000U BTX (Dysport) C: Placebo	 <u>E1/E2/E3 vs C</u> Modified Ashworth Scale (+exp, +exp₂, +exp₃) <u>E2/E3 vs C</u> Action Research Arm Test (+exp₂, +exp₃) <u>E1/E2 vs C</u> Barthel Index (+exp, +exp₂)
Childers et al. (2004) RCT (7) Nstart=91 Nend=91 TPS=Chronic	E1: 90U BTX (type A) E2: 180U BTX (type A) E3: 360U BTX (type A) C: Placebo	 <u>E1/E2/E3 vs C</u> Modified Ashworth Scale (+exp, +exp₂, +exp₃) Functional Independence Measure (-)
Brashear et al. (2002) RCT (7) Nstart=126 Nend=122 TPS=Chronic	E: Botulinum toxin A (50 U) C: Placebo	 Disability Assessment Scale (+exp) Ashworth Scale (+exp)
Bakheit et al. (2001) RCT (8) N _{start} =59 N _{end} =58 TPS=Chronic	E: Total of 1000 IU of BtxA (Dysport) into 5 muscles of the affected arm C: Placebo injections	 Modified Ashworth Scale score (+exp) Active/passive range of motion (-) Barthel Index (-)
Bhakta et al. (2000) RCT (7) Nstart=40 Nend=38 TPS=Chronic	E: Total of 1000 IU Dysport (n=20) C: Placebo (n=20) divided between elbow, wrist, and finger flexors	 Modified Ashworth Scale (+exp) Active range of motion (-)
<u>Smith et al</u> . (2000) RCT (7) N _{start} =25 N _{end} =25 TPS=Chronic	E1: 500 U of botulinum toxin E2: 1000 U of botulinum toxin E3: 1500 U of botulinum toxin C: Placebo	 <u>E1/E2/E3 vs C</u> Modified Ashworth Scale at fingers (+exp_(combined)) Active range of movement (-) Frenchay Arm Test (-)

Cimpoon at al. (4000)	E4. Cingle treatment of 75 H DTV A			
<u>Simpson et al</u> . (1996)	E1: Single treatment of 75 U B1X-A	$\underline{E1/E3VSC}$		
RCT (8)	E2: 150 U BTX-A	Modified Ashworth Scale (+exp ₁ , +exp ₃)		
N _{start} =37	E3: 300 U BTXA	<u>E1/E2/E3 vs C</u>		
Nend=37	C: Placebo	Functional Independence Measure (-)		
TPS-Chronic		Fugl-Meyer Scale (-)		
	Botulinum toxin B versus pl	acebo		
Gracies et al. (2014)	E1: 10000 U Botox (type B)	E1/E2 vs C		
BCT (9)	E2: 15000 LI Botox (type B)	Modified Ashworth Scale (-)		
$N_{\rm e} = 24$	C: Disasha	 Modified Frenchay Scale (-) 		
NStart=24	C. Flacebo			
NEnd=24				
TPS=Chronic				
Brashear et al. (2004)	E: 10000 U of BTX-B	Modified Ashworth scale (-)		
RCT (7)	C: Placebo			
Notor=15				
N -15				
TRend=15				
Botulinum tox	kin A combined with upper limb rehabilita	ation versus botulinum toxin A		
Devier et al. (2017)	E: OnabotulinumtoxinA with upper limb	Fugl-Meyer Assessment (+exp)		
RCT (5)	rehabilitation	Modified Ashworth Scale (-)		
N _{Start} =31	C: OnabotulinumtoxinA	Disability Assessment Scale (-)		
NEnd =29				
TPS=Chronic				
Onabotulinu	⊥ umtoxinA versus letibotulinumtoxinA. NA	BOTA, Neuronox, tizanidine		
	Ful atihatulinumtavinA (Batulav)	Modified Ashwarth Scale ()		
$\frac{D0 \text{ et al. } (2017)}{DCT (9)}$	C. OnebetulinumtovinA	Modified Astriworth Scale (-) Clobal Assessment in Specticity ()		
	C. OnaboluinumloxinA	Giobal Assessment in Spasticity (-)		
NStart =187		Disability Assessment Scale (-)		
N _{End} =169				
TPS=Chronic				
Nam et al. (2015)	E: Botulinum toxin type A (NABOTA) up	Modified Ashworth Scale (-)		
RCT (7)	to 360 U depending on degree of	Disability Assessment Scale (-)		
Nstort=197	spasticity and muscle group			
N _{End} -177	C: Onshotulinum toxin A (Botox) un to			
	260 LI depending on degree of specticity			
TF 3=Subacule	and muscle group			
<u>Seo et al</u> . (2015)	E1: 360 U Neu-BoNT-A (Neuronox)	Modified Ashworth Scale (-)		
RCT (10)	E2: 360 U Botox	Disability Assessment Scale (-)		
N _{Start} =196		•		
N _{End} =170				
TPS=Chronic				
Simpson et al. (2009)	F1: Up to 500 U of BoNT-Type A	F1 vs C		
	E2: Tizanidino	Modified Ashworth Scale (Levo)		
		 Moullieu Ashworth Scale (Texp) Disshility Assessment Scale (Lexp) 		
N _{start} =60	C: Placebo	• Disability Assessment Scale (+exp)		
N _{end} =41				
TPS=Subacute		Modified Ashworth Scale (-)		
		Disability Assessment Scale (-)		
		<u>E1 vs E2</u>		
		Modified Ashworth Scale (+exp)		
		Disability Assessment Scale (+exp)		
	High versus low dosage botuling	um toxin A		
Francisco et al. (2002)	E1: High volume BTX-A (50 units/1 mL	Modified Ashworth Scale: (-)		
RCT (7)	saline: 1.2 mL delivered per muscle)			
N _{start} =13	F2: Low volume BTX-A (100 units/1 ml			
Nord=9	saline: 0.6 mL delivered per muscle)			
Botulinum toxin A combined	with adhesive taning versus betulinum to	vin A combined with manual muscle stratebing		
botuinum toxin A combined with adnesive taping versus botuinum toxin A combined with manual muscle stretching, passive articular mobilization, and palmar splinting				

Santamato et. al (2015) RCT (7) N _{Start} =70 N _{End} =70 TPS=Chronic	E: 50-200 U Botox (type A) + adhesive taping for 10d C: 50-200 U Botox (type A) + manual muscle stretching, passive articular mobilization, and palmar splint	 Modified Ashworth Scale (+exp) Disability Assessment Scale (+exp)
	Ultrasound guided botulinum toxin	A injections
Zeuner et al. (2017) RCT-Crossover (5) Nstart =30 NEnd =23 TPS=Chronic	E: Ultrasound guided Botulinum Toxin A Injections followed by electromyographic (EMG) Guided Botulinum Toxin A Injections (100-400mu) C: EMG Guided Botulinum Toxin A Injections followed by Ultrasound Guided Botulinum Toxin A Injections (100- 400mu)	 Modified Ashworth Scale (-) Barthel Index (-) Disability Assessment Scale (-)
Picelli et al. (2014) RCT (8) N _{Start} =60 N _{End} =60 TPS=Chronic	E1: Botox A Injections (500u) under sonographic guidance E2: Botox A Injection (500u) using electrical stimulation guidance C: Botox A Injection (500u) using manual needle placement	E1 vs C Modified Ashworth Scale (+exp) Tardieu Spasticity angle (+exp) Passive range of motion (+exp) E2 vs C Modified Ashworth Scale (wrist): (+exp ₂) Tardieu Spasticity angle (+exp ₂) Passive range of motion (+exp ₂) E1 vs E2 Modified Ashworth Scale (-) Tardieu Spasticity angle (-) Passive range of motion (-)
Santamato et al. (2014) RCT (4) N _{Start} =30 N _{End} =30 TPS=Chronic	E: BoNT-A injection using ultrasound guidance (dosages determined by investigator) C: BoNT-A using manual needle placement via palpitation and anatomical landmarks (dosages determined by investigator)	Modified Ashworth Scale (+exp)
	Botulinum toxin A combined wi	th NMES
Marvulli et al. (2016) RCT (6) Nstart=36 NEnd=36 TPS=Chronic	E: Botulinum toxin A therapy (118±34 U) + occupational therapy (OT) + functional electrical stimulation C: Botulinum toxin A therapy (116±36 U) + OT Duration: <i>Not Specified</i>	 Modified Ashworth Scale (+exp) Passive range of Motion (+exp) Action Research Arm Test (+exp)
Hesse et al. (1998) RCT (7) N _{start} =24 N _{end} =24 TPS=Chronic	E1: 1000 U Btx A + cyclic NMES E2: 1000 U of Btx A E3: Placebo + cyclic NMES C: Placebo Duration: Daily injections for 3 mo For electrical stimulation: 30 min/d, 2d/ wk for 4 wk	 E1 vs E2 vs E3 vs C Modified Ashworth Scale (-) E1 vs E2/C Reduction in difficulties with cleaning palm (+exp)
	Botulinum toxin A combined with	h mCIMT
Sun et al. (2010) RCT (6) Nstart=32 Nend=32 TPS=Chronic Botulinum to	E: 1,000 U Dysport + mCIMT C: 1,000 U Dysport + conventional rehabilitation Duration : 2hr/d, 3d/wk for 3 mo oxin A combined with task-specific trainin	 Modified Ashworth Scale (+exp) Motor Activity Log (+exp) Action Research Arm Test (+exp)

Umar et al. (2018)	E: Botulinum Toxin A with Task-Specific	•	Fugl-Meyer Assessment (-)
RCT (5)	Training	•	Motor Assessment Scale (-)
N _{Start} =46	C: Task-Specific Training		
N _{End} =41			
TPS=NR			

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at $\alpha{=}0.05$

Conclusions about botulinum toxin

MOTOR FUNCTION					
LoE	Conclusion Statement	RCTs	References		
1a	Botulinum toxin A may not have a difference in efficacy when compared to placebo, no injection or conventional therapy for improving motor function.	8	Rosales et al. 2018; Hesse et al. 2012; Marciniak et al. 2012; Wolf et al. 2012; Shaw et al. 2011; Shaw et al. 2010; Suputitiada and Suwanwela, 2005; Simpson et al. 1996		
2	Botulinum toxin A combined with upper limb rehabilitation may produce greater improvements in motor function than botulinum toxin A alone.	1	Devier et al. 2017		
1b	Botulinum toxin A combined with functional electrical stimulation may produce greater improvements in motor function than botulinum toxin A.	1	Marvulli et al. 2016		
1b	Botulinum toxin A combined with mCIMT may produce greater improvements in motor function than botulinum toxin A.	1	Sun et al. 2010		
2	Botulinum toxin A combined with task-specific training may not have a difference in efficacy when compared to task-specific training alone for improving motor function.	1	Umar et al. 2018		

ACTIVITIES OF DAILY LIVING						
LoE	Conclusion Statement RCTs Reference					
1a	Botulinum toxin A may not have a difference in efficacy when compared to placebo, no injection or conventional therapy for improving performance of activities of daily living.	10	Marcinak et al. 2012; Shaw et al. 2011; Shaw et al. 2010; Meythaler et al. 2009; Jahangir et al. 2007; Suputiada & Suwanwela, 2005; Childers et al. 2004; Bakheit et al. 2001; Smith et al. 2000; Simpson et al. 1996			
1b	Botulinum toxin B may not have a difference in efficacy when compared to placebo for improving performance of activities of daily living.	1	Gracies et al. 2014			
2	Ultrasound guided botulinum toxin A injections may not have a difference in efficacy when compared to electromyography guided botulinum toxin A injections for improving performance of activities of daily living.	1	Zeuner et al. 2017			
1b	Botulinum toxin A combined with mCIMT may produce greater improvements in performance of activities of daily living than botulinum toxin A .	1	Sun et al. 2010			
2	Botulinum toxin A combined with task-specific training may not have a difference in efficacy when compared to task-specific training alone for improving performance of activities of daily living.	1	Umar et al. 2018			

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
1a	Botulinum toxin A may not have a difference in efficacy when compared to placebo, no injection or conventional therapy for improving dexterity.	2	Shaw et al. 2011; Shaw et al. 2010	

RANGE OF MOTION			
LoE	Conclusion Statement	RCTs	References
1a	Botulinum toxin A may not have a difference in efficacy when compared to placebo, no injection or conventional therapy for improving range of motion.	4	Marciniak et al. 2012; Bakheit et al. 2001; Bhakta et al. 2000; Smith et al. 2000
1b	Ultrasound guided botulinum toxin A injections may produce greater improvements in range of motion than manual needle placement injections .	1	Picelli et al. 2014
1b	Electrical stimulation guided botulinum toxin A injections may produce greater improvements in range of motion than manual needle placement injections.	1	Picelli et al. 2014
1b	Ultrasound guided botulinum toxin A injections may not have a difference in efficacy when compared to electrical stimulation guided botulinum toxin A injections for improving range of motion.	1	Picelli et al. 2014
1b	Botulinum toxin A combined with functional electrical stimulation may produce greater improvements in range of motion than botulinum toxin A.	1	Marvulli et al. 2016

MUSCLE STRENGTH					
LoE	Conclusion Statement	RCTs	References		
1b	Botulinum toxin A may produce greater improvements in muscle strength than placebo, no injection or conventional therapy.	1	Shaw et al. 2010		

SPASTICITY			
LoE	Conclusion Statement	RCTs	References
1a	Botulinum toxin A may produce greater improvements in spasticity than placebo, no injection or conventional therapy.	18	Rosales et al. 2018; Elovic et al. 2016; Gracies et al. 2015; Hesse et al. 2012; Marciniak et al. 2012; Shaw et al. 2011; Kaji et al. 2010; Shaw et al. 2010; Weythaler et al. 2009; Simpson et al. 2009; Jahangir et al. 2007; Suputitiada and Suwanwela, 2005; Childers et al. 2004; Brashear et al. 2002; Bakheit et al. 2001; Bhakta et al. 2000; Smith et al. 2000; Simpson et al. 1996
1a	Botulinum toxin B may not have a difference in efficacy when compared to placebo for improving spasticity.	2	Gracies et al. 2014; Brashear et al. 2004
2	Botulinum toxin A combined with upper limb rehabilitation may not have a difference in efficacy when compared to botulinum toxin A alone for improving spasticity.	1	Devier et al. 2017
1b	LetibotulinumtoxinA, NABOTA and neuronox may not have a difference in efficacy when compared to onabotulinumtoxinA for improving spasticity.	3	Do et al. 2017; Nam et al. 2015; Seo et al. 2015
1b	Botulinum toxin A may produce greater improvements in spasticity than tizanidine .	1	Simpson et al. 2009
1b	High volume botulinum toxin A may not have a difference in efficacy when compared to low volume botulinum toxin A for improving spasticity.	1	Francisco et al. 2002
1b	Botulinum toxin A combined with adhesive taping may produce greater improvements in spasticity than botulinum toxin A combined with manual muscle stretching, passive articular mobilization, and palmar splinting.	1	Santamato et al. 2015
2	Ultrasound guided botulinum toxin A injections may not have a difference in efficacy when compared to electromyography guided botulinum toxin A injections for improving spasticity.	1	Zeuner et al. 2017
1b	Ultrasound guided botulinum toxin A injections may produce greater improvements in spasticity than manual needle placement injections.	2	Santamato et al. 2014; Picelli et al. 2014
1b	Electrical stimulation guided botulinum toxin A injections may produce greater improvements in spasticity than manual needle placement injections.	1	Picelli et al. 2014
1b	Ultrasound guided botulinum toxin A injections may not have a difference in efficacy when compared to electrical stimulation guided botulinum toxin A injections for improving spasticity.	1	Picelli et al. 2014
1b	Botulinum toxin A combined with functional electrical stimulation may produce greater improvements in spasticity than botulinum toxin A.	1	Marvulli et al. 2016

1b	Botulinum toxin A combined with cyclic NMES may not have a difference in efficacy when compared to botulinum toxin A, cyclic NMES, or placebo for improving spasticity.	1	Hesse et al. 1998
1b	Botulinum toxin A combined with mCIMT may produce greater improvements in spasticity than botulinum toxin A.	1	Sun et al. 2010

Key points

Botulinum A likely improves spasticity in the upper limb following stroke, but not range of motion or activities of daily living. The effect on general upper limb motor function is conflicting and less clear.

Botulinum toxin A in combination with other types of therapeutic approaches may be beneficial for certain aspects of upper limb function.

Botulinum toxin B has been less well studied to date in comparison to botulinum toxin A.

Steroids



Adopted from: https://en.wikipedia.org/wiki/Corticosteroid

Corticosteroids have been used to treat pain and functional limitations in hemiplegic patients (Dogan et al. 2013). Patients suffering from stroke experience high rates of inflammation and corticosteroids are prescribed to lessen the inflammation (Yasar et al. 2011).

The methodological details and results of a single RCT evaluating intra-articular steroid use for upper extremity motor rehabilitation are presented in Table 32.

Table 32. RCT	intra-articular	steroid use fo	r upper e	xtremity n	notor r	ehabilitation

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Yasar et al. (2011) RCT (9) N _{Start} =26 N _{End} =26 TPS=Subacute	E1: Intra-Articular Steroid Injection E2: Suprascapular Nerve Block Injection Duration: <i>Not Specified</i>	Range of Motion (-)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α=0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about steroids

RANGE OF MOTION					
LoE	Conclusion Statement	RCTs	References		
1b	Intra-articular steroid injections may not have a difference in efficacy when compared to suprascapular nerve block injections for improving range of motion.	1	Yasar et al. 2011		

Key points

Steroid injections may not be beneficial for upper limb rehabilitation following stroke.

Cerebrolysin



Adopted from: http://www.gerovitalshop.eu/it/home/18-cerebrolysin-5ml.html

Cerebrolysin contains low molecular weight neuropeptides and free amino acids which are believed to have neuroprotective properties, inhibit free radical formation, reduce neuroinflammation, and activate calpain apoptosis (Muresanu et al. 2016). The methodological details and results of two RCTs evaluating cerebrolysin for upper extremity motor rehabilitation are presented in Table 33.

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end}	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Time post stroke category		
Chang et al. (2016) RCT (6) Nstart=70 Nend=66 TPS=Acute	E: Cerebrolysin (30mL diluted with 70mL saline) + conventional therapy C: Placebo + conventional therapy Duration: 1x/d for 6wk	 Action Research Arm Test (+exp) National Institute of Health Stroke Scale (+exp) Barthel Index (+exp) Modified Rankin Scale (+exp)
Muresanu et al. (2016) RCT (9) N _{start} =208 N _{end} =196 TPS=Acute	E: Cerebrolysin (30mL diluted with 70mL saline) + physical/occupational therapy C: Placebo + physical/occupational therapy Duration: 1x/d for 3wk	Fugl-Meyer Assessment (+exp)

Table 33. RCTs evaluating cerebrolysin for upper extremity motor rehabilitation

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α=0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about cerebrolysin

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	Cerebrolysin may produce greater improvements in motor function than placebo .	2	Chang et al. 2016; Muresanu et al. 2016	

ACTIVITIES OF DAILY LIVING				
LoE	Conclusion Statement	RCTs	References	
1b	Cerebrolysin may produce greater improvements in range of motion than placebo .	1	Chang et al. 2016	

STROKE SEVERITY				
LoE	Conclusion Statement	RCTs	References	
1b	Cerebrolysin may produce greater improvements in measures of stroke severity than placebo .	1	Chang et al. 2016	

Key points

Cerebrolysin may be beneficial for aspects of upper limb function following stroke.

Levodopa



Adopted from: https://www.maynepharma.com/products/us-products/generic-products/generic-products-catalog/carbidopalevodopa-tablets/

Levodopa has been the hallmark pharmaceutical for the treatment of Parkinson's disease. However, its ability to affect motor movements in Parkison's disease is limited by its narrow therapeutic window, short half-life, and poor bioavailability (Tambassco et al. 2018).

The methodological details and results of two RCTs evaluating levodopa treatment for upper extremity motor rehabilitation in stroke survivors are presented in Table 34.

Table 34. RCTs evaluating levodopa interventions for upper extremity motor rehabilitation

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Rosser et al. (2008) RCT (5) Nstart=18 Nend=18 TPS=Chronic	E: Levodopa (100mg) + Cabidopa (25mg) C: Placebo (125mg) Duration: 1hr physio (3x) + Levodopa (3x)	 Performance in a simple motor task (+exp)
Restemeyer et al. (2007) RCT (9) N _{start} =10 N _{end} =10 TPS=Chronic	E: Levodopa (100mg) C: Placebo (100mg) Duration: 1hr physio (2x) + Levodopa (2x)	 Nine Hole Peg Test (-) Grip strength (-) Action Research Arm Test (-)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at $\alpha{=}0.05$

Conclusions about levodopa

MOTOR FUNCTION			
LoE	Conclusion Statement	RCTs	References
1b	There is conflicting evidence about the effect of Levodopa to improve motor function when compared to placebo .	2	Rosser et al. 2008; Restemeyer et al. 2007

MUSCLE STRENGTH			
LoE	Conclusion Statement	RCTs	References
1b	Levodopa may not have a difference in efficacy when compared to placebo for improving muscle strength.	1	Restemeyer et al. 2007

DEXTERITY			
LoE	Conclusion Statement	RCTs	References
1b	Levodopa may not have a difference in efficacy when compared to placebo for improving dexterity.	1	Restemeyer et al. 2007

Key points

The evidence is mixed regarding Levodopa for upper limb rehabilitation following stroke.

Atorvastatin



Adopted from: https://www.aarp.org/health/drugs-supplements/info-2016/new-guidelines-on-who-should-take-statins-cs.html

HMG-CoA reductase inhibitors (statins) are widely used worldwide due to their antiatherosclerotic, anti-inflammatory, and immunomodulatory properties (Lin et al. 2015). This suggests that statins may have a beneficial role in infection, in fact, statins are found to have beneficial effects on the prevention and treatment of infections in diseases including cerebrovascular accidents (Lin et al. 2015). Statins are also believed to have a neuroprotective effect and are conducive to promoting autophagy in neurological disorders (Lin et al. 2015).

The methodological details and results of a single RCTs evaluating atorvastatin for upper extremity motor rehabilitation are presented in Table 35.

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Zhang et al. (2017) RCT (6) N _{Start} =78 N _{End} =75 TPS=Acute	E: Atorvastatin (20mg) C: Placebo (20mg) Duration: Atorvastatin daily for 6wk	 Modified Rankin Scale (+exp) Barthel Index (+exp) NIHSS (-)

|--|

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at $\alpha{=}0.05$

Conclusions about atorvastatin

ACTIVITIES OF DAILY LIVING			
LoE	Conclusion Statement	RCTs	References
1b	Atorvastatin may produce greater improvements in activities of daily living than placebo.	1	Zhang et al. 2017

STROKE SEVERITY			
LoE	Conclusion Statement	RCTs	References
1b	There is conflicting evidence about the effect of atorvastatin to improve motor function when compared to placebo	1	Zhang et al. 2017

Key points

The evidence is mixed regarding atorvastatin for upper limb rehabilitation following stroke.

Antidepressants



Adopted from: https://www.newportacademy.com/resources/treatment/teens-antidepressants-side-effects-risks-holistic-treatment/

Antidepressants of various kinds are available for medical use, including tricyclics (TCAs), monoamine oxidase inhibitors (MAOIs), selective serotonin reuptake inhibitors (SSRIs), serotonin-noradrenaline reuptake inhibitors (SNRIs, such as venlafaxine, duloxetine and milnacipran), and other agents (mirtazapine, reboxetine, bupropion). SSRIs and SNRIs are two commonly prescribed agents that work by acting to inhibit the reuptake of serotonin and norepinephrine, respectively, from the synaptic cleft (Cipriani et al. 2012). Beyond their ability to improve depression following stroke, antidepressants can be used to enhance upper extremity motor recovery through changes in neurotransmission. There is evidence suggesting that serotoninergic modulation may be involved in motor recovery post stroke. Previous research has suggested that patients who have reacted well to antidepressant treatment may also demonstrate improvements in upper limb motor functioning (Chemerinski et al. 2001). Furthermore, there are reports that single doses of selective serotonin reuptake inhibitors (SSRIs), such as fluoxetine and paroxetine, have resulted in activation of the motor cortices (Dam et al. 1996; Pariente et al. 2001) therefore, manipulation of neurochemicals may influence aspects of function other than psychological distress. Moreover, there is evidence to suggest that noradrenergic reuptake inhibitors (NRIs) increase motor cortex excitability (Plewnia et al. 2002).

The methodological details and results of 7 RCTs evaluating antidepressants for upper extremity motor rehabilitation are presented in Table 36.

Table 36. RCTs evaluating antidepressants interventions for upper extremity motor rehabilitation

Authors (Year)	Interventions	Outcome Measures
Study Design (PEDro Score)	Duration: Session length, frequency	Result (direction of effect)
Sample Size _{start}	per week for total number of weeks	
Sample Sizeend		
Time post stroke category		
<u>Ward et al. (2017)</u>	E: Atomoxetine 40 mg with Task-	 Fugl-Meyer Assessment (+exp)
	Oriented Upper Extremity Training	Action Research Arm Test (-)
NStart = 12	C: Placebo with Task-Oriented Upper	Wolf Motor Function Test (-)
NEnd =9	Extremity Training	
Mehammedianingiad at al. (2014)	E: Lithium corbonata (200mg)	- National Institutos of Haalth Straka
	C: Placebo	National Institutes of Health Stroke Scale (Lexp)
$N_{\text{O}} = -80$	C. Flacebo Duration: Lithium Carbonate 300mg	• Fugl-Mover Assessment (+eyp)
Nstan-00	(2x/d) for 30d	• Tugi-meyer Assessment (+exp)
TPS=Acute		
Chollet et al. (2011)	E: Fluoxetine (20mg)	 Fugl Meyer Assessment (+exp)
RCT (9)	C: Placebo	 National Institutes of Health Stroke
N _{start} =118	Duration: Ingested daily (orally) for 3mo	Scale (-)
N _{end} =113		 Modified Rankin Scale (+exp)
TPS=Chronic		
<u>Zittel et al</u> . (2008)	E: Citalopram (40mg)	 Nine Hole Peg Test (+exp)
RCT (8)	C: Placebo (40mg)	 Hand grip strength (-)
N _{start} =8	Duration: Citalopram (2x)	
N _{end} =8		
TPS=Chronic		-
$\underline{Zittel et al}$. (2007)	E: Reboxetine (6mg)	Lapping speed (+exp)
	C: Placebo (6mg)	Grip strength (+exp)
Nstart=10	Duration: Repoxetine (2x)	
Nend=10 TPS_Chronic		
	Notrintvline + Fluoxetine versus Placeb	0
Mikami et al. (2011)	F1: Nortriptyline (100mg)	F1/F2 vs C
RCT (8)	E2: Fluoxetine (40mg)	 Modified Rankin Scale (+exp1 +exp2)
1 vr follow-up analysis of Robinson et al.	C: Placebo	
2000	Duration: Fluoxetine or Nortriptvline	
N _{start} =104	daily for 12wk	
N _{end} =97		
TPS=Chronic		
Robinson et al. (2000)	E1: Nortriptyline (100mg)	E1 vs E2/C
RCT (8)	E2: Fluoxetine (40mg)	Functional Independence Measure
N _{start} =104	C: Placebo	(+exp1)
N _{end} =97	Duration: Fluoxetine or Nortriptyline	
TPS=Chronic	daily for 12wk	

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp₂ indicates a statistically significant between groups difference at α=0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group - indicates no statistically significant between groups differences at α =0.05
Conclusions about antidepressants

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of antidepressants to improve motor function when compared to placebo treatment .	3	Ward et al. 2017; Mohammadianinejad et al. 2014; Chollet et al. 2011	

MUSCLE STRENGTH				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of antidepressants to improve muscle strength when compared to placebo treatment.	2	Zittel et al. 2008; Zittel et al. 2007	

ACTIVITIES OF DAILY LIVING					
LoE	LoE Conclusion Statement RCTs Reference				
1b	Antidepressants may produce greater improvements in performance of activities of daily living than placebo treatment.	1	Robinson et al. 2000		

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
1a	Antidepressants may produce greater improvements in dexterity than placebo treatment.	2	Zittel et al. 2008; Zittel et al. 2007	

STROKE SEVERITY				
LoE	Conclusion Statement	RCTs	References	
1a	Antidepressants may produce greater improvements in measures of stroke severity than placebo treatment.	3	Mohammadianinejad et al. 2014; Chollet et al. 2011; Mikami et al. 2011	

Key points

Antidepressants may be beneficial for aspects of upper limb function following stroke.

Central nervous system stimulants



Adopted from: https://www.narconon.org/drug-information/amphetamine-health-risks.html

Central nervous system stimulants are drugs that increase cortical excitability, often provided to manage arousal states by enhancing neural transmission. Central nervous system stimulants increase the synaptic concentration and transmission of dopamine, serotonin, and noradrenaline throughout the brain, and neurobehavioral gains ascribed to central nervous system stimulants include enhanced arousal, mental processing speed, and/or motor processing speed (Herrold et al. 2014). Common stimulants used in rehabilitation include amphetamines and methylphenidates. Methylphenidate has been shown to enhance motor recovery after partial cortex ablation in rodents, and to modulate poststroke cerebral reorganization, improving motor function in stroke patients (Wang et al. 2014). Stimulants such as amphetamines have been reported to enhance plasticity through axonal sprouting (Papadopoulos et al. 2009).

The methodological details and results of four RCTs evaluating antidepressants for upper extremity motor rehabilitation are presented in Table 37.

Table 37. RCTs evaluating meridian acupressure and massage therapy interventions for upper extremity motor rehabilitation

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Schuster et al. (2011) RCT (9) N _{start} =16 N _{end} =15 TPS=Chronic	E: Dexamphetamine (10mg) C: Placebo Duration: 20min/d, 2d/wk for 5wk	Chedoke-McMaster Stroke Assessment (+exp)
Tardy et al. (2006) RCT (9) N _{start} =8 N _{end} =8 TPS=Chronic	E: Methylphenidate (20mg) C: Placebo Duration: 2d/wk for 4wk	 Finger tapping scores (+exp) Hand grip strength (-)
Platz et al. (2005a) RCT (9) N _{start} =31 N _{end} =29 TPS=Chronic	E: d-amphetamine (10mg) C: Placebo Duration: 45min/d, 3d/wk for 4wk	• TEMPA (+exp)
	Methylphenidate + tDCS	
Wang et al. (2014) RCT (7) N _{start} =9 N _{end} =9 TPS=Subacute	E1: Dual tDCS + methylphenidate (20mg) E2: Dual tDCS + placebo drug E3: Sham tDCS + methylphenidate C: Sham tDCS + placebo drug Duration: 20min/d, 3d/wk for 4wk	 <u>E1 vs E2/E3</u> Purdue Pegboard Test: (+exp)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α=0.05 in favour of the experimental group

+exp2 indicates a statistically significant between groups difference at α=0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at $\alpha\text{=}0.05$

Conclusions about central nervous stimulants

MOTOR FUNCTION				
LoE Conclusion Statement RCTs Refere				
1a	Dexamphetamine and methylphenidate may produce greater improvements in motor function than placebo treatment.	2	Schuster et al. 2011; Platz 2005a	

MUSCLE STRENGTH				
LoE	Conclusion Statement	RCTs	References	
	Dexamphetamine and methylphenidate may not	1	Tardy et al. 2006	
1b	have a difference in efficacy when compared to			
	placebo treatment for improving muscle strength.			

DEXTERITY				
LoE	Conclusion Statement	RCTs	References	
1b	Dexamphetamine and methylphenidate may produce greater improvements in dexterity than placebo treatment.	1	Tardy et al. 2006	
1b	Methylphenidate combined with dual tDCS may produce greater improvements in dexterity than dual tDCS or methylphenidate.	1	Wang et al. 2014	

Key points

Dexamphetamine or methylphenidate may be beneficial for aspects of upper limb function following stroke.

Methylphenidate combined with dual transcranial direct current stimulation may be beneficial for upper limb rehabilitation following stroke.

Complementary and alternative medicine Acupuncture



Adopted from: https://www.mccaffreyhealth.com/acupuncture-for-chronic-pain/

The use of acupuncture has recently gained attention as an adjunct to stroke rehabilitation in Western countries even though acupuncture has been a primary treatment method in China for about 2000 years (Baldry, 2005). In China, acupuncture is an acceptable, time-efficient, simple, safe and economical form of treatment used to ameliorate motor, sensation, verbal communication and further neurological functions in post-stroke patients," (Wu et al., 2002). According to Rabinstein and Shulman (2003), "Acupuncture is a therapy that involves stimulation of defined anatomic locations on the skin by a variety of techniques, the most common being stimulation with metallic needles that are manipulated either manually or that serve as electrodes conducting electrical currents". There is a range of possible acupuncture mechanisms that may contribute to the health benefits experienced by stroke patients (Park et al. 2006). For example, acupuncture may stimulate the release of neurotransmitters (Han & Terenius, 1982) and have an effect on the deep structure of the brain (Wu et al. 2002). Lo et al. (2005) established acupuncture, when applied for at least 10 minutes, led to long-lasting changes in cortical excitability and plasticity even after the needle stimulus was removed. With respect to stroke rehabilitation, the benefit of acupuncture has been evaluated most frequently for pain relief and recovery from hemiparesis.

18 RCTs for acupuncture were identified. In 11 RCTs Acupuncture was compared to sham or conventional rehabilitation (Chen et al. 2016; Liu et al. 2016; Cui et al. 2014; Bai et al. 2013; Zhuangl et al. 2012; Wayne et al. 2005; Alexander et al. 2004; Sze et al. 2002; Kiendhal et al. 1997; Hu et al. 1993; Naeser et al. 1992). Four RCTs looked at comparisons of different acupuncture techniques (Ni et al. 2013; Fragoso and Ferreira, 2012; Zhao et al. 2009; Gosman-Hedstom et al. 1998). RCTs looked at acupuncture combined with CIMT (Song et al. 2016), TENS (Hopwood et al. 2008). Finally, a single RCT looked at acupuncture compared to cerebroprotein hydrolysate and piracetam (Han et al. 2015).

The methodological details and results of all 18 RCTs evaluating acupuncture for upper extremity motor rehabilitation are presented in Table 38.

Table 38. Summary of RCTS with Examining Acupuncture for upper extremity motor rehabilitation

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
	Acupuncture compared to conventional th	erapy or sham
<u>Chen et al.</u> (2016) RCT (8) N _{Start} =250 N _{End} =250 TPS=Chronic	E: Acupuncture C: Conventional therapy Duration: 45min/d, 6d/wk for 3wk	 National Institute of Health Stroke Scale (+exp) Fugl-Meyer Assessment (+exp)
Liu et al. (2016) RCT (6) N _{Start} =38 N _{End} =31 TPS=Chronic	E: Manual acupuncture + standard care C: Standard care Duration: <i>Not Specified</i>	 National Institute of Health Stroke Scale (-) Fugl-Meyer Assessment (-) Functional Independence Measure (-) Barthel Index (-) Modified Rankin Scale (-)
Cui et al. (2014) RCT (6) Nstart=60 NEnd=60 TPS=NR	E: Yin Yang manipulation C: Conventional needling manipulation Duration: <i>Not Specified</i>	 Elbow spasm (+exp) Clinical Spasticity Index (+exp)
Bai et al. (2013) RCT (9) N _{Star} =120 N _{End} =120 TPS=NR	E1: Acupuncture E2: Physical therapy E3: Acupuncture + physical therapy Duration: <i>Not Specified</i>	E1 vs E2 Fugl-Meyer Assessment (-) Modified Barthel Index (-) E1 vs E3 Fugl-Meyer Assessment (-) Modified Barthel Index (-) E2 vs E3 Fugl-Meyer Assessment (-) Modified Barthel Index (-)
Zhuangl et al. (2012) RCT (7) N _{start} =295 N _{end} =274 TPS=Chronic	E1: Acupuncture E2: Physiotherapy E3: Acupuncture + physiotherapy Duration: 1hr/d, 6d/wk for 4wk	 Fugl-Meyer Assessment (-) Barthel Index (-) Neurologic Defect Scale (-)
Wayne et al. (2005) RCT (9) Nstart=33 Nend=33 TPS=Chronic	E: Acupuncture C: Sham Duration: 45min/d, 2d/wk for 10wk	 Fugl-Meyer Assessment (-) Modified Ashworth scores (-) Arm range of motion (-) Barthel Index (-)
Alexander et al. (2004) RCT (6) N _{start} =32 N _{end} =28 TPS=Acute	E: Acupuncture + Standard Rehabilitation C: Standard Rehabilitation Duration: 30min/d, 2d/wk for 10 wk	 Fugl-Meyer Assessment (-) Functional Independence Measure (-)
<u>Sze et al</u> . (2002) RCT (7) Nstart=106 Nend=106 TPS=Acute	E: Acupuncture + Standard Therapy C: Standard Therapy Duration: 45min/d, 2d/wk for 10wk	 Fugl-Meyer Assessment (-) Barthel Index (-) Functional Independence Measure (-) National Institutes of Health Stroke Scale (-)

Kjendhal et al. (1997) RCT (6) N _{start} =45 N _{end} =41	E: Acupuncture C: Standard Therapy Duration: 30min/d, 3-4d/wk for 6wk	 Motor Assessment Scale (+exp) Sunnaas Index (+exp)
TPS=Subacute Hu et al. (1993)	E: Acupuncture	Scaninavian stroke study Neurological score
RCT (4) N _{start} =30 N _{end} =NR TPS=Acute	C: Supportive Therapy + Conventional Rehabilitation Duration: <i>Not Specified</i>	(+exp) • Barthel Index (-)
Naeser et al. (1992) RCT (6) N _{start} =16 N _{end} =16 TPS=Subacute	E: Acupuncture C: Sham Acupuncture Duration: 1hr/d, 5d/wk for 4wk	 Boston Motor Inventory range of motion (+exp)
	Acupuncture vs acupunctur	e
Ni et al. (2013) RCT (7) N _{Start} =165 N _{End} =165 TPS=NR	E: Standard Acupuncture with Shixuan & Xiaohai acupoints C: Standard Acupuncture only Duration: <i>Not Specified</i>	 Finger grip strength (+exp) Fugl-Meyer Assessment (+exp)
Fragoso & Ferreira (2012) RCT (6) Nstart=32 Nend=32 TPS=Chronic	E1: Acupuncture at Tianquan (PC2) E2: Acupuncture at Quchi (LI11) Duration: 20min/d, 5d/wk for 4wk	 Maximal Isometric Voluntary Contraction during elbow flexion (-)
Zhao et al. (2009) RCT (5) N _{start} =131 N _{end} =120 TPS=Chronic	E: Experimental acupuncture C: Traditional acupuncture Duration: 20min/d, 7d/wk, for 4wk	 Modified Ashworth Scale (+exp) Fugl-Meyer Assessment (+exp) Barthel Index (+exp)
Gosman-Hedstom et al. (1998) RCT (7) N _{start} =104 N _{end} =98 TPS=Acute	E1: Superficial acupuncture E2: Deep acupuncture C: No acupuncture Duration: 1hr/d, 2d/wk for 10 wk	 <u>E1 vs E2 vs C</u> Scaninavian stroke study Neurological score (-) Barthel Index (-) Sunnaas Index (-)
	Acupuncture combined with C	IMT
<u>Song et al.</u> (2016) RCT (5) N _{Start} =30 N _{End} =30 TPS=Acute	E: Scalp cluster acupuncture + constraint- induced movement therapy C: Body acupuncture + traditional rehabilitation Duration: 6hr/d, (needles twisted 2-3x), 6d/wk for 2wk	Fugl-Meyer Assessment (-)
	Acupuncture combined with TEI	NS
Hopwood et al. (2008) RCT (7) N _{start} =105 N _{end} =105 TPS=Acute	E: Acupuncture with TENS C: Acupuncture with sham TENS Duration: 1hr/d, 3d/wk for 4wk	 Barthel Index (-) Motricity Index (-)
Acu	puncture versus cerebroprotein hydrolysa	te and piracetam
Han et al. (2015) RCT (6) N _{Start} =488 N _{End} =488 TPS=NR	E: Meridian sinew row needling combined with dermal needling C: Cerebroprotein hydrolysate (20mL) and piracetam injections (4g) Duration: <i>Not Specified</i>	 Fugl-Meyer Assessment (+exp) Modified Ashworth Scale (+exp)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group +con indicates a statistically significant between groups difference at α =0.05 in favour of the control group - indicates no statistically significant between groups differences at α =0.05

Conclusions about acupuncture

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	Acupuncture may not have a difference in efficacy when compared to conventional therapy or sham for improving motor function.	8	Chen et al. 2016; Liu et al. 2016; Han et al. 2015; Bai et al. 2013; Zhuangl et al. 2012; Wayne et al. 2005; Alexander et al. 2004; Sze et al. 2002	
1b, 2	Standard acupuncture with Shixuan & Xiaohai acupoints and experimental acupuncture may produce greater improvements in motor function than standard or traditional acupuncture.	2	Ni et al. 2013; Zhao et al. 2009	
2	Scalp cluster acupuncture combined with CIMT may not have a difference in efficacy when compared to body acupuncture with traditional rehabilitation for improving motor function.	1	Song et al. 2016	
1b	Meridian sinew row needling combined with dermal needling may produce greater improvements in motor function than cerebroprotein hydrolysate and piracetam.	1	Han et al. 2015	

SPASTICITY

SI ASHOITI					
LoE	Conclusion Statement	RCTs	References		
1a	Acupuncture may produce greater improvements in spasticity than conventional therapy or sham.	3	Cui et al. 2014; Zhao et al. 2009; Wayne et al. 2005		
2	Experimental acupuncture may produce greater improvements in spasticity than traditional acupuncture.	1	Zhao et al. 2009		
1b	Meridian sinew row needling combined with dermal needling may produce greater improvements in spasticity than cerebroprotein hydrolysate and piracetam.	1	Han et al. 2015		

	RANGE OF MOTION				
LoE	Conclusion Statement	RCTs	References		
1a	There is conflicting evidence about the effect of acupuncture to improve range of motion when compared to conventional therapy or sham .	2	Wayne et al. 2009; Naeser et al. 1992		

STROKE SEVERITY				
LoE	Conclusion Statement	RCTs	References	
1a	Acupuncture may not have a difference in efficacy when compared to conventional therapy or sham for improvements on measures of stroke severity.	4	Liu et al. 2016; Zhuangl et al. 2012; Sze et al. 2002; Hu et al. 1993	

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	N

Superficial acupuncture may not have a difference in
efficacy when compared to deep acupuncture for
improvements on measures of stroke severity.Gosman-Hedstrom
et al. 1998

ACTIVITIES OF DAILY LIVING					
LoE	LoE Conclusion Statement RCTs				
1a	Acupuncture may not have a difference in efficacy when compared to conventional therapy or sham for improving performance of activities of daily living.	8	Liu et al. 2016; Bai et al. 2013; Zhuangl et al. 2012; Wayne et al. 2005; Alexander et al. 2004; Sze et al. 2002; Kjendhal et al. 1997; Hu et al.1993		
1b	Acupuncture combined with TENS may not have a difference in efficacy when compared to acupuncture with sham stimulation for improving performance of activities of daily living.	1	Hopwood et al. 2008		
1b	Superficial acupuncture may not have a difference in efficacy when compared to deep acupuncture for improving performance of activities of daily living.	1	Gosman-Hedstom et al. 1998		
2	Experimental acupuncture may produce greater improvements in performance of activities of daily living than traditional acupuncture .	1	Zhao et al. 2009		

MUSCLE STRE	NGTH
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LoE	Conclusion Statement	RCTs	References
1a	Standard acupuncture with Shixuan & Xiaohai acupoints and acupuncture at Tianquan PC2 may produce greater improvements in muscle strength than standard acupuncture only and acupuncture at Quchi LI11.	2	Ni et al. 2013; Fragoso and Ferreira, 2012
1b	Acupuncture combined with TENS may not have a difference in efficacy when compared to acupuncture with sham stimulation for improving muscle strength.	1	Hopwood et al. 2008

Key points

The evidence is mixed regarding acupuncture alone for upper limb rehabilitation following stroke. Acupuncture combined with conventional or other therapy approaches may not be beneficial for upper limb function. Some forms of acupuncture may be more beneficial than others.

Electroacupuncture and transcutaneous electrical acupoint stimulation



Adopted from: https://www.promotionhealthcare.com/electroacupuncture-treatment-pain-injuries/

Electroacupuncture is a variant of acupuncture techniques practiced in traditional Chinese medicine, the difference being that a minute electrical current of similar intensity to that of a bioelectric current produced endogenously in the body is applied to the needles used (Wang et al. 2014). The needle is often placed on meridian points throughout the body (Wang et al. 2014). Similarly, transcutaneous electrical acupoint stimulation (TEAS) stimulates meridian points believed to be associated with a medical condition with electrical impulses given through needles (Zhao et al. 2015). The two techniques have very similar mechanisms of action and their influence on afferent stimulation to the body (Zhao et al. 2015).

11 RCTs were found that evaluated electroacupuncture and transcutaneous electrical acupoint stimulation compared to conventional therapy, sham stimulation, ordinary needling, and strength training (Zhang et al. 2017; Zhao et al. 2015; Au-Yeung et al. 2014; Wang et al. 2014; Wen et al. 2014; Yao et al. 2014; Hsing et al. 2012; Li et al. 2012; Hsieh et al. 2007; Mukherjee et al. 2007; Moon et al. 2003). One RCT looked at electroacupuncture combined with neuronavigation-assisted aspiration compared to neuronavigation-assisted aspiration, electroacupuncture or conventional therapy (Zhang et al. 2017).

The methodological details and results of all 11 RCTs evaluating electroacupuncture and transcutaneous electrical acupoint stimulation for the upper extremity motor rehabilitation are presented in Table 39.

Authors (Year) Study Design (PEDro Score) Sample Size _{start} Sample Size _{end} Time post stroke category	Interventions Duration: Session length, frequency per week for total number of weeks	Outcome Measures Result (direction of effect)
Zhao et al.(2015) RCT (9) N _{start} =60 N _{End} =60 TPS=Chronic	E1: Transcutaneous electrical acupoint stimulation (TEAS) (100Hz) E2: Transcutaneous electrical acupoint stimulation (TEAS) (2Hz) C: Sham stimulation Duration: 0, 2, or 100Hz/d, 5d/wk for 4wk	E1 vs C Modified Ashworth Scale (+exp) Disability Assessment Scale (-) Global Assessment Scale (-) Barthel Index (-) E2 vs C Modified Ashworth Scale (+exp ₂) Disability Assessment Scale (-) Global Assessment Scale (-) Barthel Index (-) E1 v E2 Modified Ashworth Scale (+exp) Disability Assessment Scale (-) Global Assessment Scale (-) Barthel Index (-)
Au-Yeung et al. (2014) RCT (6) N _{Start} =73 N _{End} =60 TPS=Acute	E1: Electroacupoint stimulation E2: Sham stimulation C: Conventional therapy (control) Duration: 20Hz/d, 1h/d, 5d/wk for 4wk	E1 vs. C • Hand grip strength (+exp) • Index grip pinch (+exp) E2 vs C & E1 vs E2 • Hand grip strength (-) • Index grip pinch (-) • Action Research Arm Test (-)
Wang et al. (2014) RCT (6) Nstart=20 NEnd=15 TPS=Chronic	E: Electroacupuncture C: No stimulation with no needle manipulation Duration: 50Hz/d, 20min/d, 2d/wk for 6wk	 Elbow joint muscle tone (+exp) Wrist joint muscle tone (-)
Wen et al. (2014) RCT (7) Nstart=300 NEnd=276 TPS=Acute	E: Electroacupuncture + moxibustion C: Basic therapy Duration: 2 to 15Hz, 5-7d/wk for 4wk	Fugl-Meyer Assessment (-)
<u>Yao et al.</u> (2014) RCT (5) N _{start} =68 N _{End} =65 TPS=Chronic	E: Relaxed needling + electroacupuncture C: Ordinary needling Duration: 5Hz, 30min/d, 3d/wk for 8wk	Fugl-Meyer Assessment (+exp)
Hsing et al. (2012) RCT (7) N _{stan} =62 N _{end} =62 TPS=Subacute	E: Scalp electro-acupuncture C: Sham acupuncture Duration: 2 to 100Hz, 30min/d, 2d/wk for 5wk	 Barthel Index (-) Rankin Scale (-)
Li et al. (2012) RCT (6) N _{start} =120 N _{end} =120 TPS=Acute	E: Electroacupuncture + massage C: Rehabilitation therapy Duration: 25min/d, 5d/wk, 6wk	 Fugl-Meyer Assessment (-) Modified Rankin Scale (+exp)
Hsieh et al. (2007) RCT (8) N _{start} =63 N _{end} =63 TPS=Subacute	E: Electroacupuncture C: No acupuncture Duration: 20min/d, 2d/wk for 4wk	 Functional Independence Measure (-) Fugl-Meyer Assessment (+exp)

Table 39. RCTs evaluating electroacupuncture and transcutaneous electrical acupoint stimulation interventions for upper extremity motor rehabilitation

Mukherjee et al. (2007b) RCT (4) N _{start} =7 N _{end} =7 TPS=Subacute	E: Electroacupuncture + strength training C: Strength training Duration: 2Hz, 40min/d, 2d/wk for 6wk	Modified Ashworth Scale (+exp)
<u>Moon et al</u> . (2003) RCT (5) N _{start} =35 N _{end} =31 TPS=Subacute	E1: Electroacupuncture E2: Moxibustion C: Routine acupuncture Duration: 50Hz, 30min/d, 3d/wk for 3wk	 <u>E1 vs E2/C</u> Modified Ashworth scale (+exp)
Neurona	vigation-assisted aspiration + electroacu	puncture
Zhang et al. (2017) RCT (7) Nstart=240 NEnd=233 TPS=Acute	E1: Neuronavigation-assisted aspiration + electroacupuncture E2: Neuronavigation-assisted aspiration E3: Electroacupuncture C: Conventional therapy Duration: 30min (2x per day) for 8wk	E1 vs E2 • Fugl-Meyer Assessment (+exp) • Modified Ashworth Scale (+exp) • Barthel Index (+exp) E1 vs E3 • Fugl-Meyer Assessment (+exp) • Modified Ashworth Scale (+exp) E1 vs E4 • Fugl-Meyer Assessment (+exp) <u>E3 vs E4</u> • Fugl-Meyer Assessment (+exp3) • Modified Ashworth Scale (+exp3)

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks. +exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group +exp₂ indicates a statistically significant between groups difference at α =0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about electroacupuncture and transcutaneous electrical acupoint stimulation

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of electroacupuncture and transcutaneous electrical acupoint stimulation to improve motor function when compared to conventional therapy, sham stimulation, ordinary needling, and strength training.	6	Zhang et al. 2017; Au- Yeung et al. 2014; Wen et al. 2014; Yao et al. 2014; Li et al. 2012; Hsieh et al. 2007	
1b	Electroacupuncture combined with neuronavigation-assisted aspiration may produce greater improvements in motor function than neuronavigation-assisted aspiration, electroacupuncture and conventional therapy on their own.	1	Zhang et al. 2017	

SPASTICITY				
LoE	Conclusion Statement	RCTs	References	
1a	Electroacupuncture and transcutaneous electrical acupoint stimulation may produce greater improvements in spasticity than conventional therapy, sham stimulation, ordinary needling, and strength training.	5	Zhang et al. 2017; Zhao et al. 2015; Wang et al. 2014; Mukherjee et al. 2007; Moon et al. 2003	
1b	Electroacupuncture combined with neuronavigation-assisted aspiration may produce greater improvements in spasticity than neuronavigation-assisted aspiration, electroacupuncture and conventional therapy on their own.	1	Zhang et al. 2017	

STROKE SEVERITY				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of electroacupuncture and transcutaneous electrical acupoint stimulation to improve scores on measures of stroke severity when compared to conventional therapy, sham stimulation, ordinary needling, and strength training.	2	Hsing et al. 2012; Li et al. 2012	

ACTIVITIES OF DAILY LIVING				
LoE	Conclusion Statement	RCTs	References	
1a	Electroacupuncture and transcutaneous electrical acupoint stimulation may not have a difference in efficacy when compared to conventional therapy, sham stimulation, ordinary needling, and strength training for improving performance of activities of daily living.	3	Zhao et al. 2015; Hsieh et al. 2007; Hsing et al. 2012	
1b	Electroacupuncture combined with neuronavigation-assisted aspiration may produce greater improvements in activities of daily living neuronavigation-assisted aspiration and electroacupuncture on their own.	1	Zhang et al. 2017	

MUSCLE STRENGTH			
LoE	Conclusion Statement	RCTs	References
1b	There is conflicting evidence about the effect of electroacupuncture and transcutaneous electrical acupoint stimulation to improve muscle strength when compared to conventional therapy, sham stimulation, ordinary needling, and strength training.	1	Au-Yeung et al. 2014

Key points

Electroacupuncture with neuronavigation-assisted aspiration may be beneficial for upper limb rehabilitation following stroke, however the evidence is mixed regarding electroacupuncture and transcutaneous electrical acupoint stimulation.

Meridian acupressure and massage therapy



Adopted from: http://physiotherapeutic.ca/servi-physio/111-massage-therapy

Meridian acupressure is a form of treatment whereby finger pressure is applied to meridian points on the body (Yang et al. 2017). There are two types of meridian points: yin and yang (Yang et al. 2017). Yin meridians run from the feet to the torso, and from the torso to the fingertips on the inside of the arms (Cui et al. 2014). On the other hand, yang meridians run from the fingers to the face and from the face to the feet (Cui et al. 2014). Acupressure increases blood (qi) flow to the areas it is applied in (Di et al. 2017).

Massage is the practice of applying structured pressure, tension, motion or vibration — manually or with mechanical aids — to the soft tissues of the body, including: muscles, connective tissue, tendons, ligaments, joints and lymphatic vessels, to achieve a beneficial response (Holland & Pokorny, 2001). As a form of therapy, massage can be applied to parts of the body or successively to the whole body, to heal injury, relieve psychological stress, manage pain, and improve circulation (College of Massage Therapists of Ontario, 2018). The benefits of massage therapy are suggested to be increased blood flow, relief of muscle spasms and release of β -endorphins (Wei et al. 2017). One of the more common forms of massage therapy is the traditional Chinese massage therapy also known as Tui Na (Yang et al. 2017).

The methodological details and results of all 7 RCTs evaluating meridian acupressure and massage therapy for upper extremity motor rehabilitation are presented in Table 40.

Table 40. RCTs evaluating meridian acupressure and massage therapy interventions for upper extremity motor rehabilitation

Authors (Year)	Interventions	Outcome Measures
Study Design (PEDro Score)	Duration: Session length, frequency	Result (direction of effect)
Sample Sizestart	per week for total number of weeks	
Sample Sizeend	· · · · · · · · · · · · · · · · · · ·	
Time post stroke category		
Di et al. 2017	E: Tui Na Therapy	Modified Ashworth Scale (+exp)
RCT (5)	C: Conventional therapy	
Nstart =150	Duration: 30min/d, 5d/wk for 4wk	
N _{End} =150		
TPS=Subacute		
Yang et al. 2017	E: Tui Na	 Modified Ashworth Scale (+exp)
RCT (8)	C: Placebo Tui Na	 Fugl-Meyer Assessment (-)
N _{Start} =90	Duration: 20-25min/d, 5d/wk for 4wk	 Modified Barthel Index (-)
N _{End} =74		
TPS=Subacute		
Yang et al. (2017)	E: Tui Na	 Fugl-Meyer Assessment (-)
RCT (8)	C: Placebo Therapy	 Modified Barthel Index (-)
Nstart=90	Duration: 20-25min/d, 5d/wk for 4wk	
N _{End} =79		
TPS=Subacute		
<u>Cui et al.</u> (2014)	E: Yin Yang manipulation	Elbow spasm (+exp)
RCT (6)	C: Conventional needling manipulation	 Clinical Spasticity Index (+exp)
N _{Start} =60	Duration: Not Specified	
N _{End} =60		
IPS=NR		
Thanakiatpinyo et al. (2014)	E: Thai massage	Modified Ashworth Scale (-)
	C: Physical therapy	Barthel Index (-)
Nstart=50	Duration: 30min/d, 2d/wk for 6wk	
NEnd=45		
$\frac{\text{Yue et al.}}{\text{PCT}(6)}$	E: Acupressure	Barnel Index (+exp) Eucl Meyer Assessment (Leve)
	C: Routine care	• Fugi-meyer Assessment (+exp)
Nstart=70	Duration. 45min/d, 5d/wk, 4wk	
TPS-Chronic		
Kang et al. (2000)	E: Meridian acupressure	
RCT (5)	C: Standard care	 Bassive range of motion (+evp)
Nor (5)	Duration: 10min/d 7d/wk for 2wk	
N _{stal} =56		
TPS=Chronic		
TPS=Chronic		

Abbreviations and table notes: C=control group; D=days; E=experimental group; H=hours; Min=minutes; RCT=randomized controlled trial; TPS=time post stroke category (Acute: less than 30 days, Subacute: more than 1 month but less than 6 months, Chronic: over 6 months); Wk=weeks.

+exp indicates a statistically significant between groups difference at α =0.05 in favour of the experimental group

+exp2 indicates a statistically significant between groups difference at α=0.05 in favour of the second experimental group

+con indicates a statistically significant between groups difference at α =0.05 in favour of the control group

- indicates no statistically significant between groups differences at α =0.05

Conclusions about meridian acupressure and massage therapy

MOTOR FUNCTION				
LoE	Conclusion Statement	RCTs	References	
1a	There is conflicting evidence about the effect of meridian acupressure and massage therapy to improve motor function when compared to conventional therapy or placebo massage therapy .	3	Yang et al. 2017; Yang et al. 2017; Yue et al. 2013	

MUSCLE STRENGTH	
Conclusion Statement	RCTs

LoE	Conclusion Statement	RCTs	References
2	Meridian acupressure and massage therapy may produce greater improvements in muscle strength than conventional therapy.	1	Kang et al. 2009

RANGE OF MOTION				
LoE Conclusion Statement RCTs Reference				
2	Meridian acupressure and massage therapy may produce greater improvements in range of motion than conventional therapy.	1	Kang et al. 2009	

ACTIVITIES OF DAILY LIVING			
LoE	Conclusion Statement	RCTs	References
1a	Meridian acupressure and massage therapy may not have a difference in efficacy when compared to conventional therapy or placebo massage therapy for improving performance of activities of daily living.	4	Yang et al. 2017; Yang et al. 2017; Thanakiatpinyo et al. 2014; Yue et al. 2013

SPASTICITY				
LoE	Conclusion Statement	RCTs	References	
1a	Meridian acupressure and massage therapy may produce greater improvements in spasticity than conventional therapy or placebo massage therapy.	4	Di et al. 2017; Yang et al. 2017; Cui et al. 2014; Thanakiatpinyo et al. 2014	

Key points

Both meridian acupressure and massage therapy may be beneficial for some aspects of upper limb function following stroke.

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