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## Effectiveness of hybrid robotic rehabilitation system on upper limb recovery of people with central injuries: a systematic review with meta-analysis

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To enable PROSPERO to focus on COVID-19 registrations during the 2020 pandemic, this registration record was automatically published exactly as submitted. The PROSPERO team has not checked eligibility.

### Citation

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### Review question

What is the effectiveness of hybrid robotic rehabilitation system on upper limb recovery of people with central injuries?

### Searches

The search for relevant studies will be conducted on PEDro (Physiotherapy Evidence Database), EMBASE (Excerpta Medica Database), MEDLINE (Medical Literature Analysis and Retrieval System Online), CINAHL (Cumulative Index to Nursing and Allied Health Literature), COCHRANE (Cochrane Collaboration), AMED (Allied and Complementary Medicine Database), IEEE Xplore (Institute of Electrical and Electronics Engineering) and Compendex (Compendex Engineering Index) without language or date restrictions. In addition, a hand searching will be conducted in the reference lists of all eligible studies and previous reviews in this area. Search terms will be related to: "Hybrid Orthosis", "Upper limb" and "Randomized controlled trial" (see Appendix 1 on the Addenda for detailed search strategy).

This review will be included prospective randomized or quasi-randomized controlled studies including inpatients and outpatients from any primary, secondary or tertiary care setting and community. Studies will be eligible if they included participants with limited upper limb functioning caused by any health condition, regardless of age or sex (inpatients or outpatients from any clinical/hospital care settings including primary, secondary or tertiary services and community individuals). The intervention of interest will be HRRS, which is defined as the systems that rehabilitate or compensate motor functions through the combined action of muscle activation with FES and mechanical/electromechanical forces supplied to joints. Studies will be eligible if HRRS is compared with minimal intervention or other intervention (OI). We defined minimal intervention as when the control group received no intervention, received sham or placebo intervention, or was on a waiting list. We considered any other active intervention that was not hybrid orthosis, such as conventional therapy and physical therapy. The outcome of interest in this review will be related to body function and structure, activity and participation according to the International Classification of Functioning, Disability and Health

### Types of study to be included

This review will be included prospective randomized or quasi-randomized controlled studies

### Condition or domain being studied

People with upper limb disorders caused by central lesions

### Participants/population

Studies will be eligible if they included participants with limited upper limb functioning caused by central lesions, regardless of age or sex (inpatients or outpatients from any clinical/hospital care settings including

primary, secondary or tertiary services and community individuals).

### Intervention(s), exposure(s)

The intervention of interest will be HRRS, which is defined as the systems that rehabilitate or compensate motor functions through the combined action of muscle activation with FES and mechanical/electromechanical forces supplied to joints (DEL-AMA et al., 2012)

### Comparator(s)/control

Studies will be eligible if HRRS is compared with minimal intervention or other intervention (OI). We defined minimal intervention as when the control group received no intervention, received sham or placebo intervention, or was on a waiting list. We considered any other active intervention that was not hybrid orthosis, such as conventional therapy and physical therapy.

### Main outcome(s)

The outcome of interest in this review will be related to body function and structure, activity and participation according to the International Classification of Functioning, Disability and Health (SIVAN et al., 2011; WORLD HEALTH ORGANIZATION, 2001). Body function and structure are considered as the physiological functions of body systems, including psychological function and body structure, i.e., anatomical parts of the body, such as organs, limbs and their components. Activity refers to execution of a task or action by an individual and participation is involvement of an individual in a life situation (WORLD HEALTH ORGANIZATION, 2001).

#### \* Measures of effect

Sample size, mean and standard deviation (SD) for each group will be extract at short, medium- and long-term follow-ups when available. Pooled effects will be estimated using standardized mean differences (SMDs) with 95% confidence intervals (CI).

### Additional outcome(s)

Subgroup qualitative analysis will be conducted to investigate the impact of methodological quality issues on the pooled effects. Studies with PEDro scores of at least 6 out of 10 were considered high quality in this review.

#### \* Measures of effect

Sample size, mean and standard deviation (SD) for each group will be extract at short, medium- and long-term follow-ups when available. Pooled effects will be estimated using standardized mean differences (SMDs) with 95% confidence intervals (CI).

### Data extraction (selection and coding)

Data will be extracted by a reviewer (FMRMF) and double checked by a second reviewer (GPR). Disagreements will be resolved by consensus. Authors will be contacted to clarify eventual doubts. Data extracted at baseline will include: number of participants; mean age; percentages of males and females; cause of the upper limb disorder and its duration, evaluated joints; type of HRRS; comparison groups; frequency and total duration of intervention; and outcome measures. The outcome data extracted will included the sample size, mean and standard deviation (SD) for each group at the short-, medium- and long-term follow-ups, when available. When multiple time points are available within the same follow-up period, the time point closer to the end of the intervention will be considered.

### Risk of bias (quality) assessment

The GRADE (Grading of Recommendations Assessment, Development and Evaluation) system will be used to summarize the overall quality of evidence for each outcome (BALSHEM et al., 2011) . We will rate evidence from the high-quality level and downgrade it one point if one of the following prespecified criteria is present: low methodological quality (average PEDro score < 6); inconsistency of estimates among pooled studies ( $I^2 > 50\%$ ) or when its assessment is not possible (no pooling); indirectness of participants (over 50% of the studies do not describe inclusion criteria); and imprecision (pooling < 400 participants for each outcome) (HIGGINS; GREEN; COCHRANE COLLABORATION, 2008).

### Strategy for data synthesis

Data for each outcome will be pooled when there is sufficient homogeneity among studies. Heterogeneity

among studies will be assessed using  $I^2$  statistics =  $[(Q - df)/Q] \times 100\%$ , where  $Q$  is the  $\chi^2$  statistic and  $df$  is its degrees of freedom (HIGGINS et al., 2003; HIGGINS; THOMPSON, 2002). Low heterogeneity is defined as  $I^2 \leq 50\%$ , and moderate to high heterogeneity is defined as  $I^2 > 50\%$  (HIGGINS; GREEN; COCHRANE COLLABORATION, 2008). Pooled effects will be estimated using standardized mean differences (SMDs) with 95% confidence intervals (CI). A fixed-effects model will be used to conduct the meta-analysis when  $I^2 \leq 50\%$ , and a random-effects model otherwise. To judge the clinical relevance of HRRS, the effect size will be assessed using Cohen's  $d$  coefficient according to the following parameters: 0.2 as small effect, 0.5 as medium effect, and 0.8 as large effect (COHEN, 1988). A funnel plot will be used to investigate publication bias when at least 10 studies are pooled (HIGGINS; GREEN; COCHRANE COLLABORATION, 2008). The meta-analysis will be performed using the software Comprehensive Meta-Analysis, version 3.3.070.

### Analysis of subgroups or subsets

Central tendency and variability measures will be extracted for short-, medium- and long-term effects. Standard deviation (SD) will be imputed using 95% CI.

### Contact details for further information

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Universidade Federal de Minas Gerais

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### Type and method of review

Intervention, Systematic review

### Anticipated or actual start date

04 January 2021

### Anticipated completion date

04 January 2022

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### Conflicts of interest

### Language

English

### Country

Brazil, France

### Stage of review

Review Ongoing

### Subject index terms status

Subject indexing assigned by CRD

### Subject index terms

MeSH headings have not been applied to this record

### Date of registration in PROSPERO

21 January 2021

### Date of first submission

21 December 2020

### Stage of review at time of this submission

| Stage                                                           | Started | Completed |
|-----------------------------------------------------------------|---------|-----------|
| Preliminary searches                                            | Yes     | No        |
| Piloting of the study selection process                         | No      | No        |
| Formal screening of search results against eligibility criteria | No      | No        |
| Data extraction                                                 | No      | No        |
| Risk of bias (quality) assessment                               | No      | No        |
| Data analysis                                                   | No      | No        |

*The record owner confirms that the information they have supplied for this submission is accurate and complete and they understand that deliberate provision of inaccurate information or omission of data may be construed as scientific misconduct.*

*The record owner confirms that they will update the status of the review when it is completed and will add publication details in due course.*

### Versions

21 January 2021  
21 January 2021  
21 January 2021  
02 February 2021

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PROSPERO

This information has been provided by the named contact for this review. CRD has accepted this information in good faith and registered the review in PROSPERO. The registrant confirms that the information supplied for this submission is accurate and complete. CRD bears no responsibility or liability for the content of this registration record, any associated files or external websites.