Overview of Cochrane Systematic Reviews of Rehabilitation Interventions for Persons with Traumatic Brain Injury: A Mapping Synthesis

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Abstract: Background: The World Health Organization has identified an unmet global need for rehabilitation interventions concerning 20 non-communicable diseases, traumatic brain injury included. This overview compiles and synthesizes the quality and quantity of available evidence on the effectiveness of rehabilitation interventions for traumatic brain injury from Cochrane systematic reviews (CSRs). The results will be used to develop the Package of Interventions for Rehabilitation. Methods: All CSRs on TBI tagged in the Cochrane Rehabilitation database published between August 2009 and September 2021 were included. Evidence mapping was implemented to extract study characteristics and evidence from the CSRs. Results: Six CSRs (42 studies; n = 3983) examined the effectiveness of either non-pharmacological or pharmacological interventions after TBI. Among 19 comparisons, 3% were rated as high in quality of evidence, 9% moderate, 54% low, and 34% very low. Non-pharmacological interventions with moderate quality, hospital-based cognitive rehabilitation and cognitive didactic therapy, likely produced minimal to no changes in the return-to-work rate. Anti-epileptic drugs and neuroprotective agents resulted in a minimal difference to the frequency of late seizure episodes in post-traumatic epilepsy. Conclusions: No prominent advances in treatment options were reported in any of the CSRs. The high rate of low and very low quality of evidence makes it difficult to ascertain the effectiveness of several recommended non-pharmacological interventions.

Keywords: brain injuries; traumatic; interventions; treatment outcome; rehabilitation; overview

1. Introduction

The World Health Organization (WHO) has described an unmet global need for the delivery of rehabilitation interventions in health systems, which is amplified in low- and middle-income countries with limited availability of resources [1–3]. The ‘WHO Rehabilitation 2030 Call for Action’ [2] was therefore launched. One of the main actions considered is the development of a Package of Interventions for Rehabilitation (PIR) [3,4]. The PIR aims at promoting favorable outcomes, accessibility, and the integration of multidisciplinary/interdisciplinary rehabilitation services into healthcare systems worldwide [3,4]. The WHO identified 20 major noncommunicable diseases to be investigated to develop the PIR; among these is traumatic brain injury (TBI) [4].

TBI is defined as ‘any alteration in brain function or other evidence of brain pathology caused by an external force’ [5] and it is estimated to affect 69,000 individuals worldwide annually [6]. Alterations in brain function may include any of the following: loss of (or decrease in) consciousness; loss of memory of events immediately preceding or following...
the injury; neurologic deficits (e.g., loss of balance or vision); or altered mental status, such as disorientation or confusion at the time of the injury [5]. TBI can be categorized into three possible diagnostic levels (mild, moderate, or severe), typically after evaluation using the Glasgow Outcome Scale or Glasgow Outcome Scale Extended [7,8] or by assessing structural imaging, loss of consciousness, altered consciousness, or post-traumatic amnesia.

Research has identified falls and road injuries as the two main causes of TBI worldwide [9,10], although causes of TBI have been found to differ across countries, depending on income, geographical region, and political circumstances [9,11]. Other common causes include sports-related concussions, assault, interpersonal violence, and blast injuries [12]. The direct consequences of a single TBI or repetitive insults include many possible long-term sequelae that vary according to age, sex, and the nature of the injury [13,14]. Common secondary pathophysiological conditions include seizures, sleep disorders, neurodegenerative diseases, neuroendocrine dysregulation, and psychiatric issues, each of which may persist throughout the long-term recovery process following moderate-to-severe TBI [15].

Due to these numerous clinical and demographic variables, TBI patients often experience nonlinear recovery trends, and those with moderate and severe cases are reported to show deteriorating Glasgow Outcome Scale Extended scores over time [16]. These unfavorable outcomes can hinder functioning, quality of life, and employment, and may worsen pre-existing conditions [17], further highlighting the chronic health issues associated with TBI as well as the need for complex rehabilitative programs and long-term services to support this group of patients [16].

A major step to the development of the PIR encompasses the “Best Evidence for Rehabilitation” (be4rehab) approach, which is applied to this work. Be4rehab supports the gathering of best evidence on the effectiveness and quality of pharmacological and non-pharmacological rehabilitation interventions for individuals with TBI and the delivery of this overview of Cochrane systematic reviews (CSRs) [4]. Overviews of systematic reviews are a methodological approach proposed by Cochrane to compile and synthesize data from multiple systematic reviews into one single, accessible document. All overviews requested by the WHO are restricted to CSRs to preserve the coherence and quality of the gathered evidence.

Supplemented by evidence mapping to aid in the synthesis of available evidence, this work aims at identifying the broad quality and the quantity of evidence, published in CSRs, on the effectiveness of rehabilitation interventions in person with TBI.

2. Materials and Methods

The WHO PIR adheres to methods designed from the collaborative efforts of the WHO Rehabilitation Programme and Cochrane Rehabilitation, and the directives from the WHO Guidelines Review Committee [4]. We used evidence mapping to synthesize and visualize study characteristics and evidence from CSRs on TBI. The overview was registered in Open Science Framework Registries (https://doi.org/0.17605/OSF.IO/M5XVG) and was reported following the Preferred Reporting Items for Systematic Reviews and Meta-analysis (PRISMA 2020 statement) [18].

2.1. Search Strategy

According to the methodology developed by the Cochrane Rehabilitation [19], CSRs relevant to rehabilitation are continuously tagged to maintain an up-to-date database (https://rehabilitation.cochrane.org/evidence, accessed on 1 September 2019). We initially searched all CSRs related to TBI published between August 2009 and August 2019 and reported the results to the WHO. We subsequently searched the Cochrane Library to August 2021 to preserve the timeliness of evidence. Eligible CSRs included those assessing interventions for persons with TBI provided or prescribed by rehabilitation professionals [19].

We included only tagged CSRs that examined rehabilitation interventions on individuals with TBI, of any age and gender. CSRs focused on persons with acquired brain injury or
non-traumatic brain injury were excluded to ensure that the evidence synthesis is strictly applicable to persons who sustained a TBI.

2.2. Assessment of Methodological Quality of Included Studies

The methodological quality of each CSR was appraised by two assessors using the 16-item A Measurement Tool to Assess Systematic Reviews (AMSTAR) 2 tool. In this updated version, the 16 items are scored on a binary yes or no scale. AMSTAR-2 does not generate an ‘overall score’; a high score may disguise weaknesses in 7 critical items [20]. The assessors adopted a process of ‘considered judgment’, which entails (1) interpreting weaknesses detected by the critical items and (2) reaching a consensus on the methodological quality of each CSR. Disagreements were resolved through discussion with a third assessor.

2.3. Data Extraction and Quality of Evidence Appraisal

The authors referred to the Table of Findings presented in each of the included CSRs; these contain the following data: type of outcome, outcome measure(s), number of primary studies, sample sizes, type of population, intervention, comparator(s), and effect (i.e., no effect, in favor of intervention, or in favor of comparator). Data were collected and entered into an Excel datasheet.

In addition, the quality of evidence for each outcome was extracted using the Grading of Recommendations Assessment, Development, and Evaluation (GRADE) rating system. For CSRs that did not include GRADE ratings, two members of the Cochrane Rehabilitation team independently appraised the quality of evidence for the primary outcomes only using the GRADE approach [21]. Any disagreement was resolved through consensus decision-making involving a third author [22]. The GRADE appraisal approach included two steps: (1) retrieval of the original primary studies included in each CSR; and (2) tabulation of the quality of evidence provided in Summary of Findings tables using GRADEPro software.

2.4. Summarizing the Data with an Evidence Map

Quality of evidence and effect data were transferred into evidence maps developed in Excel. The evidence map integrates the outcome and rehabilitation intervention values for each comparison. The magnitude of the effect (i.e., no effect, in favor of intervention, or in favor of comparator) and the quality of evidence (i.e., very low, low, moderate, or high) were presented laterally and color-coded for each outcome in order to generate a visual aid to facilitate the understanding of the overall judgement of the evidence.

Evidence mapping was employed as a complementary method to collating and appraising evidence from the CSRs, and subsequently used to summarize the results for the overview. The instrument collated outcomes and rehabilitation interventions and resulted in a comprehensive overview of the quality of evidence and effects. Because we did not consider other outcomes and interventions in addition to the those examined in the included CSRs, evidence mapping was not used to identify evidence gaps.

3. Results

The authors identified six tagged CSRs related to TBI: one published in 2013 [23], two in 2015 [24,25], and three in 2017 [26–28] (see Figure 1).

Three CSRs included only participants who sustained a TBI and excluded people with acquired brain injury and non-traumatic injury. Two CSRs included studies with a mixed population only when disaggregated data were reported to ensure that evidence was relevant to TBI. Finally, one CSR reported including studies where the etiology of the TBI is uncertain. The characteristics of the included systematic reviews are reported in Table 1.
Figure 1. Flow chart displaying the tagging process of Cochrane systematic review.

Comprehensively, this mapping review encompasses 42 primary studies, 3983 participants, and 19 comparisons that examined the effectiveness and safety of either non-pharmacological or pharmacological interventions for individuals with TBI. Among non-pharmacological comparisons, four interventions (six outcomes) were categorized as very low quality of evidence, and eight interventions (16 outcomes) were deemed as low quality of evidence. Among the pharmacological comparisons, we found that four interventions (six outcomes) were rated very low and three interventions (three outcomes) were rated low in quality. The AMSTAR 2 assessment tool identified high methodological quality in the six CSRs; even when sources of funding were not reported. Results of the AMSTAR 2 assessment are displayed in Table 2.
Table 1. Characteristics of included systematic reviews.

| Author (Year) | Population | Primary Outcome | Outcome Measure | Intervention | Comparator | Effect | Quality |
|---------------|------------|-----------------|-----------------|--------------|------------|--------|---------|
| Hassett et al., 2017 [26] | People with TBI; any age and sex | Cardiorespiratory fitness | Submaximal incremental cycle ergometer test | Exercise using large muscle | Usual care, a non-exercise intervention, or no intervention | Favor intervention | Low |
| Kumar (2017) [27] | Adults (≥16 years); any sex; any severity | Return to work | Attainment of work within 14 weeks (medium-term) of initiating intervention | Cognitive rehabilitation therapy | No treatment | None | Very low |
| | | Community integration | Return to work status | Cognitive rehabilitation therapy | No treatment | None | Low |
| | | Return to work | Follow-up: 6 months (medium-term) | Cognitive rehabilitation therapy | Conventional therapy | None | Low |
| | | Independence in activities of daily living | Functional independence measure, with 18 items in basic and psychosocial functional activities | Cognitive rehabilitation therapy | Conventional therapy | None | Very low |
| | | Community integration | Community integration questionnaire | Cognitive rehabilitation therapy | Conventional therapy | None | Low |
| | | Return to work | Return to work status | Hospital-based cognitive rehabilitation therapy | Home programme | None | Moderate |
| | | Return to work | Follow-up: 24 months (long-term) | Cognitive didactic therapy | Functional experiential therapy | None | Moderate |
| | | Independence in activities of daily living | Structured interview follow-up: 1 year (medium-term) | Cognitive didactic therapy | Functional experiential therapy | None | Low |
| Synnot (2017) [28] | Children and adults who had skeletal muscle spasticity post injury; Any severity | Spasticity at up to 6 h after treatment | Ashworth Scale, 0-, with a higher score indicating greater spasticity | Intrathecal baclofen 50 µg (injected into the lumbar spine) | Saline placebo | Not reported | Very low |
| | | | | Intrathecal baclofen 50 µg (injected into the lumbar spine) | Saline placebo | Not reported | Very low |
Table 1. Cont.

| Author (Year)               | Population                                      | Primary Outcome                                      | Outcome Measure                                                                 | Intervention                                                                 | Comparator                   | Effect       | Quality  |
|-----------------------------|-------------------------------------------------|------------------------------------------------------|---------------------------------------------------------------------------------|-------------------------------------------------------------------------------|------------------------------|--------------|----------|
| Spasticity at 4–12 weeks   | Modified Ashworth scale, 0–5, at 12 weeks and Tardieu scale, 0–5, at 4 weeks | Botulinum toxin A × 1 dose (500/1000 U) or botulinum toxin A × 1 dose of 200 U + serial casting | Botulinum toxin A × 1 dose (500/1000 U) or botulinum toxin A × 1 dose of 200 U + serial casting | Placebo (±casting)           | Uncertain       | Very low   |
| Adverse events              |                                                 |                                                      |                                                                                 |                                                                                |                              |              |
| Spasticity at up to 6 h after treatment | Modified Ashworth scale, 0–4, with a higher score indicating greater spasticity | Repositioning splints equipped with participant-specific pseudoelastic hinges | Repositioning splints equipped with participant-specific pseudoelastic hinges | Traditional splints with fixed angle braces | Uncertain       | Very low   |
| Adverse events              |                                                 |                                                      |                                                                                 |                                                                                |                              |              |
| Gertler (2015) [24]         | Children and adults with depression after TBI; any severity | Depression                                           | Beck depression inventory-II, Hamilton Rating Scale for Depression, and Hospital Anxiety and Depression Scale; higher score means more depressed | Cognitive behavioral therapy                                                | Wait-list control            | None        | Very low |
| Depression                  | Beck Depression Inventory; higher score means more depressed | Depression                                           | Beck Depression Inventory; higher score means more depressed                    | Cognitive behavioral therapy                                                | Supportive psychotherapy     | None        | Very low |
| Depression                  | Hamilton Rating Scale for Depression; higher score means more depressed | Depression                                           | Repetitive transcranial magnetic stimulation                                     | Repetitive transcranial magnetic stimulation plus tricyclic antidepressant    | Favor control               | Very low   |
| Depression                  | Beck Depression Inventory; higher score means more depression | Depression                                           | Supervised exercises                                                            | Exercise as usual                                                            | None                        | Low         |
Table 1. Cont.

| Author (Year) | Population                                                                 | Primary Outcome | Outcome Measure | Intervention                  | Comparator                        | Effect          | Quality |
|---------------|-----------------------------------------------------------------------------|-----------------|-----------------|-------------------------------|-----------------------------------|-----------------|---------|
| Thompson (2015) [25] | People with TBI who received prophylactic treatment with antiepileptic drugs or neuroprotective agents. Any age; any severity; acute | Early seizures  
Follow-up: 5–7 days | Count of Events | Antiepileptic drugs | Placebo or standard care      | Favor intervention | Low     |
|                |                                                                             | Late seizures   
Follow-up: 3–24 months | Count of Events | Antiepileptic drugs | Placebo or standard care      | None             | Very low |
|                |                                                                             | Early seizure   
Follow-up: 7 days | Count of Events | Neuroprotective agents | Placebo                        | None             | Low     |
|                |                                                                             | Late seizure    
Follow-up: 6 months | Count of Events | Neuroprotective agents | Placebo                        | None             | High    |
|                |                                                                             | Early seizure   
Follow-up: 7 days | Count of Events | Phenytoin         | Other antiepileptic drugs     | None             | Low     |
|                |                                                                             | Late seizure    
Follow-up: 6 months to 2 years | Count of Events | Phenytoin         | Other antiepileptic drugs     | None             | Moderate|
| Wong (2013) [23] | People with TBI. Any age, sex, and severity                                | Post-treatment  
Modified Barthel Index-1 month post-treatment | Barthel index   | Electro-acupuncture plus rehabilitation training | Rehabilitation training | Favor intervention | Low     |
|                |                                                                             | Post-treatment  
Modified Barthel Index-3 months post-treatment | Barthel index   | Electro-acupuncture plus rehabilitation training | Rehabilitation training | Favor control     | Low     |
|                |                                                                             | Post-treatment  
Fugl-Meyer assessment-1 month post-treatment | Fugl-Meyer Assessment | Electro-acupuncture plus rehabilitation training | Rehabilitation training | Favor intervention | Low     |
|                |                                                                             | Post-treatment  
Fugl-Meyer assessment-3 months post-treatment | Fugl-Meyer Assessment | Electro-acupuncture plus rehabilitation training | Rehabilitation training | Favor intervention | Low     |
### Table 1. Cont.

| Author (Year) | Population | Primary Outcome | Outcome Measure | Intervention | Comparator | Effect | Quality |
|---------------|------------|-----------------|-----------------|--------------|------------|--------|---------|
| Post-treatment Glasgow Outcome score | Glasgow Outcome Scale | Needle acupuncture plus conventional medical intervention | Conventional medical intervention | Favor intervention | Low |
| Post-treatment Glasgow Coma score | Glasgow Coma Scale | Needle acupuncture plus conventional medical intervention | Conventional medical intervention | Favor intervention | Low |
| Frequency of normal post-treatment Glasgow Outcome score | Glasgow Outcome Scale | Electro-acupuncture plus conventional medical intervention | Conventional medical intervention | Favor intervention | Low |
| Mortality | | Electro-acupuncture plus conventional medical intervention | Conventional medical intervention | None | Low |
| Frequency of post-treatment Barthel index above 60 | Barthel index | Electro-acupuncture plus hyperbaric oxygen and rehabilitation training | Hyperbaric oxygen and rehabilitation training | Favor intervention | Low |
| Frequency of post-treatment Barthel index above 40 | Barthel index | Electro-acupuncture plus hyperbaric oxygen and rehabilitation training | Hyperbaric oxygen and rehabilitation training | None | Low |

Abbreviation: TBI = traumatic brain injury.
Table 2. AMSTAR 2 Quality Assessment of Cochrane Systematic Reviews.

| Item                                                                 | Hassett 2017 [26] | Kumar 2017 [27] | Synnot 2017 [28] | Gertler 2015 [24] | Thompson 2015 [25] | Wong 2013 [23] |
|---------------------------------------------------------------------|-----------------|-----------------|-----------------|-----------------|-------------------|----------------|
| (1) Did the research questions and inclusion criteria for the review include the components of PICO? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (2) Did the report of the review contain an explicit statement that the review methods were established prior to the conduct of the review and did the report justify any significant deviations from the protocol? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (3) Did the review authors explain their selection of the study designs for inclusion in the review? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (4) Did the review authors use a comprehensive literature search strategy? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (5) Did the review authors perform study selection in duplicate? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (6) Did the review authors perform data extraction in duplicate? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (7) Did the review authors provide a list of excluded studies and justify the exclusions? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (8) Did the review authors describe the included studies in adequate detail? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (9) Did the review authors use a satisfactory technique for assessing the risk of bias (RoB) in individual studies that were included in the review? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (10) Did the review authors report on the sources of funding for the studies included in the review? | N               | N               | N               | N               | N                 | N              |
| (11) If meta-analysis was performed did the review authors use appropriate methods for statistical combination of results? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (12) If meta-analysis was performed, did the review authors assess the potential impact of RoB in individual studies on the results of the meta-analysis or other evidence synthesis? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (13) Did the review authors account for RoB in individual studies when interpreting/discussing the results of the review? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (14) Did the review authors provide a satisfactory explanation for, and discussion of, any heterogeneity observed in the results of the review? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (15) If they performed quantitative synthesis did the review authors carry out an adequate investigation of publication bias (small study bias) and discuss its likely impact on the results of the review? | Y               | Y               | Y               | Y               | Y                 | Y              |
| (16) Did the review authors report any potential sources of conflict of interest, including any funding they received for conducting the review? | Y               | Y               | Y               | Y               | Y                 | Y              |
| Total                                                               | 15              | 15              | 15              | 15              | 15                | 15             |

Abbreviations: Y = Yes, N = No.
The evidence map findings were divided into two categories: (1) non-pharmacological interventions and (2) pharmacological interventions. Table 3 provides an overview of evidence map finding for non-pharmacological interventions for TBI. Table 4 provides an overview of evidence map finding for pharmacological interventions for TBI.

Table 3. Evidence map of non-pharmacological interventions.

| Intervention                          | Comparison                                             | Outcome                          | GRADE  |
|--------------------------------------|--------------------------------------------------------|----------------------------------|--------|
|                                      |                                                        |                                  | H      |
|                                      |                                                        |                                  | M      |
|                                      |                                                        |                                  | L      |
|                                      |                                                        |                                  | VL     |
| Cognitive rehabilitation             | No treatment                                          | Return to work                   | ⊗      |
|                                      |                                                        | Community integration            | ⊗      |
|                                      | Conventional Therapy                                  | Return to work                   | ⊗      |
|                                      |                                                        | Community integration            | ⊗      |
|                                      |                                                        | Activities of daily living       | ⊗      |
| Hospital-based                       | Home-based cognitive rehabilitation                    | Return to work                   | ⊗      |
| Cognitive didactic therapy           | Functional experiential therapy                        | Return to work                   | ⊗      |
|                                      |                                                        | Activities of daily living       | ⊗      |
| Cognitive behavioral therapy         | Supportive psychotherapy                               | Depression                       | ⊗      |
| Supervised exercise                  | Exercise as usual                                      | Depression                       | ⊗      |
| Large muscle group exercise          | Usual care, non-exercise, no intervention              | Cardiorespiratory fitness        | ✓      |
| Repositioning splints                | Traditional splints                                    | Spasticity                       | ?      |
| Electro-acupuncture + Rehabilitation training | Rehabilitation training                              | Modified Barthel Index (1 mo)    | ✓      |
|                                      |                                                        | Modified Barthel Index (3 mo)    | ✗      |
|                                      |                                                        | Fugl-Meyer Assessment (1 mo)     | ✓      |
|                                      |                                                        | Fugl-Meyer Assessment (3 mo)     | ✓      |
| Needle-acupuncture + Conventional medical intervention | Conventional medical intervention                      | Post-Treatment Glasgow Outcome Scale | ✓      |
|                                      |                                                        | Post-Treatment Glasgow Coma Score | ✓      |
| Electro-acupuncture + Conventional medical intervention | Conventional medical intervention                      | Frequency of Normal Glasgow Coma Score | ✓      |
|                                      |                                                        | Mortality                        | ⊗      |
| Electro-acupuncture + Hyperbaric oxygen | Rehabilitation training vs. Hyperbaric oxygen and rehabilitation training | Frequency Barthel > 60           | ✓      |
|                                      |                                                        | Frequency Barthel > 40           | ⊗      |

High = H; M = Moderate; Low = L; VL = Very low; No effect = ⊗; Favor Intervention = ✓; Favor Comparator = ✗; Uncertain = ?.

3.1. Quality of Evidence Mapping for Non-Pharmacological Interventions

3.1.1. Moderate Quality of Evidence

Hospital-based versus home-based cognitive rehabilitation likely has little to no effect on the return-to-work rate for moderate-to-severe TBI (1 study; n = 120) [26]. Similarly, cognitive didactic versus functional experiential therapy likely has little to no effect on the same outcome for moderate-to-severe TBI (1 study, n = 366) [26].
Table 4. Evidence map of pharmacological interventions.

| Intervention                          | Comparison                              | Outcome                        | Grade |
|---------------------------------------|-----------------------------------------|--------------------------------|-------|
| Neuroprotective agents                | Placebo                                 | Early seizure                  | ⊗     |
|                                       |                                         | Late seizure (6 mo)             | ⊗     |
| Antiepileptic drugs                   |                                        | Early seizure                  | ✓     |
|                                       |                                        | Late seizure (3-24 mo)          | ⊗     |
| Phenytoin                             | Antiepileptic drugs                     | Early seizure                  | ⊗     |
|                                       |                                        | Late seizure (6-24 mo)          | ⊗     |
| Repetitive transcranial magnetic      | Repetitive transcranial magnetic        | Depression                     | ⊗     |
| stimulation plus tricyclic            | stimulation plus tricyclic antidepressants |                               |       |
| Baclofen 50 µg                         | Saline placebo                          | Spasticity                     | NR    |
|                                       |                                        | Adverse events                 | NR    |
| Botulinum toxin A × 1 dose (500/1000 U) or botulinum toxin A × 1 dose 200 U+ | Placebo                           | Spasticity                     | ?     |
|                                       |                                        | Adverse events                 | ?     |

Abbreviations: High = H; M = Moderate; Low = L; VL = Very low; No effect = ⊗; Favor Intervention = ✓; Favor Comparator = ⊗; Uncertain = ?; Not reported = NR.

3.1.2. Low Quality of Evidence

Exercise using large muscle groups may have little to no effect on the cardiorespiratory fitness compared to usual care in severe and unspecified TBI severity levels (3 studies, n = 67) [27].

Cognitive rehabilitation may have little or no effect compared to no treatment on community integration in severe TBI (1 study; n = 12) [26], while it may have little to no effect relative to conventional therapy on return to work (1 study; n = 68) [26], and community integration (3 studies; n = 123) [26] in mild-to-severe TBI, respectively.

Electro-acupuncture as an adjunct treatment to rehabilitation training may have a positive effect on sensorimotor impairment (Fugl-Meyer Assessment) at 1 and 3 months, and on disability (Modified Barthel index) at 1 month, but not at 3 months, when the effects favored rehabilitation training alone (unspecified TBI severity; 1 study; n = 150) [23]. When added to conventional medical intervention, electro-acupuncture may make little to no difference to mortality rate, but it may increase the frequency of normal Glasgow Coma Score evaluations in coma patients with severe TBI (1 study, n = 50) [23]. Added to hyperbaric oxygen and rehabilitation training, electro-acupuncture may have an effect on the percentage of patients decreasing to moderate disability (Barthel Index > 60) but there is uncertainty on the effects on reducing its severity (Barthel Index > 40) (unspecified TBI severity; 1 study; n = 122) [23].

3.1.3. Very Low Quality of Evidence

In mild-to-moderate TBI, the true effect of cognitive rehabilitation remains uncertain on return-to-work when compared to no treatment (1 study; n = 50) [26]; on activities of daily living when compared to conventional therapy (unspecified TBI severity; 2 studies, n = 41) [26]; on depression level versus waiting list (3 studies, n = 146) [24] and supportive psychotherapy (1 study; n = 48) [24]. There is also uncertainty on the utility on spasticity (6 h post-treatment) of repositioning splints equipped with participant-specific pseudoelastic hinges versus traditional splints with fixed angle braces for pediatric TBI (unspecified TBI severity; 1 study; n = 25) [28].
3.2. Quality of Evidence Mapping for Pharmacological Interventions

3.2.1. High Quality of Evidence

Neuroprotective agents had little to no effect versus placebo on late seizures 6 months after the start of treatment in moderate-to-severe TBI in participants aged 14 and older (1 study; \(n = 498\)) [25].

3.2.2. Moderate Quality of Evidence

Phenytoin likely resulted in no changes in late seizures 6 to 24 months after the start of the treatment relative to other antiepileptic drugs in moderate-to-severe TBI (2 studies; \(n = 378\)) [25].

3.2.3. Low Quality of Evidence

There may be minimal effect on the frequency of early seizures (7 days) for neuroprotective agents compared to placebo, (moderate-to-severe TBI, 1 study, \(n = 499\)) [25]. Antiepileptic interventions compared with placebo may reduce the frequency of early seizures (moderate-to-severe, 5 studies, \(n = 987\)) [25]. Neuroprotective agents versus other antiepileptic drugs may have minimal effect on adverse events (moderate-to-severe TBI, 2 studies, \(n = 431\)) [25].

3.2.4. Very Low Quality of Evidence

A review comparing baclofen 50 \(\mu\)g versus saline placebo included one study (\(n = 11\)) and examined the effects on spasticity (6 h), and adverse events [28]. The findings could not be extracted since they were not reported in the randomized control trial. The efficacy and safety of the intervention remain thereby unclear.

A review evaluated the efficacy of botulinum toxin A \(\times\) 1 dose (500/1000 U) or botulinum toxin A \(\times\) 1 dose of 200 U + serial casting versus placebo on spasticity (4–12 weeks post treatment), and adverse events (2 studies; \(n = 47\)) [28]. No statistically significant differences were detected between groups and the quality of evidence was rated very low. This hindered the ability to ascertain the true treatment effects of either intervention.

Evaluating 1029 participants and six studies, one CSR examined the difference in effects on late seizure occurrence (3 to 24 months after the start of the treatment) comparing between antiepileptic medications and placebo [25]. No significant differences were found for either outcome. The comparison was judged to provide very low quality of evidence, which indicates that the effects of antiepileptic interventions on these two outcomes remain uncertain.

In a total sample of 67 participants and one study, the reviewers found a significant difference in depression level between the repetitive transcranial magnetic stimulation and repetitive transcranial magnetic stimulation plus antidepressant groups (TBI severity unspecified) [24]. While the treatment effect was in favor of the comparator, repetitive transcranial magnetic stimulation plus tricyclic antidepressants, the true treatment effect remains uncertain due to the very low quality of evidence.

4. Discussion

This overview summarizes evidence on the effects of non-pharmacological and pharmacological interventions for any level of TBI severity, and reports the challenges identified in TBI research that are critical for further developing the integration and augmentation of rehabilitation services.

Amongst the options for non-pharmacological interventions, hospital-based cognitive rehabilitation and cognitive didactic therapy likely produce minimal or no changes in the return-to-work rate (moderate certainty evidence). These findings agree with published reports in the literature on neurocognitive status and the return-to-work rates, ref. [29–31] which maintain that favorable outcomes are facilitated by the inclusion of multidisciplinary/interdisciplinary rehabilitation services, and not by a monotherapy approach, such as cognitive rehabilitation or cognitive training alone [32,33]. Executive functions,
especially sequencing and inhibitory control, are necessary to perform well at work and their status predicts the return-to-work rate following TBI [29]. Ensuring that available cognitive interventions and cognitive strategy training lead to improvements in cognitive functioning and are properly integrated in the rehabilitation management are crucial for increasing return-to-work rates, as well as improving life satisfaction and the wellbeing of individuals with TBI and their families.

The low-certainty of evidence found in acupuncture, splint therapy, and exercise of large muscle groups prevented us from ascertaining the role of these interventions on Glasgow Coma Scale scores, spasticity, and cardiorespiratory fitness, respectively. With respect to acupuncture, the lack of information on the etiology of the TBI from three of the four RCTs prevented us from determining whether the results are equally applicable to acquired brain injury, traumatic brain injury, and non-traumatic brain injury cases. Likewise, there is insufficient quality of evidence to support the roles of cognitive therapeutic approaches as monotherapy in improving community integration, depression, and activities of daily living (very low certainty evidence).

Amongst the pharmacological interventions used to reduce the number and frequency of late-seizure episodes (i.e., 6 months after the start of treatment; high-quality evidence), neuroprotective agents produced little to no difference on the frequency of late-seizures (high-quality evidence) and minimal differences on early seizures (low-quality evidence). The anti-convulsant drug, phenytoin, for example, appeared to have little effect on the number and frequency of late seizures (moderate quality evidence) and little to no effect on early-seizure events (low quality evidence). This finding aligns with current guidelines that support the use of phenytoin to treat early seizures or active seizures, but not late seizures [34].

Our evidence mapping shows that other antiepileptic drugs do not reduce the number and frequency of late seizure events. The literature primarily focuses on early seizures, and data on late seizures after TBI are limited. Discussions of study results typically note that no evidence supports the use of neuroprotective agents and antiepileptic drugs for late seizures, mainly due to the differences observed in studies on pathogenesis of early seizures in post-traumatic epilepsy [34,35]. This feature of post-TBI care warrants further attention since late seizure episodes may impair otherwise positive neurological and rehabilitation outcomes [36].

For the remaining two pharmacological interventions (botulinum toxin A × 1 dose (500/1000 U) or botulinum toxin A × 1 dose of 200 U + serial casting; intrathecal baclofen 50 µg), uncertainty of their effects on spasticity and adverse events remain, as the quality of evidence for these two therapies has been assessed as very low [28].

The absence and/or low quality of evidence for pharmacological interventions to reduce early- and late-seizure frequency, and improve spasticity, may be associated in part with the following situations: (1) research challenges exacerbated by the narrow window for effective intervention; (2) the inability of candidate medications to cross the blood–brain barrier; and (3) possible delays and ethical issues encountered when patients are unable to provide consent [37]. These difficulties are exacerbated among pediatric groups [38], which may explain the limited results for pediatric patients with TBI among the CSRs that analyzed pharmacological interventions.

The low to very-low quality evidence found is in accordance with past reviews that focused on clinical practice guidelines for TBI [39,40], which stressed the persistent paucity of quality evidence and the major gaps between the bench and the bedside in the context of rehabilitation interventions associated with both methodological issues and clinical complexity. The reviewers stated that few published trials examined rehabilitation outcomes, such as cognitive and physical function, with the majority of studies targeting symptom management or reduction [39,40].

For non-pharmacological trials, the primary issues concerned the number of studies and the small sample sizes (cumulative <500 participants), which affected the estimated effect sizes, heterogeneity among the respondents, and the imprecision of the results...
Similar to pharmacological trials, some studies showed a lack of clarity regarding random sequence generation, blinding, and allocation concealment.

Overall, our evidence map shows that no prominent advances were reported in any of the CSRs, confirming the concerns expressed a decade ago by Maas et al. [41], who observed that randomized control trials (RCTs) fail to showcase significant recovery trajectories when assessing the effectiveness of interventions on TBI populations. Other study designs (e.g., observational) could provide additional insights when conducting systematic reviews for patients with TBI.

The landscape displayed by this evidence map places strong emphasis on the need to prioritize and augment rehabilitation research efforts for patients with TBI. Hence, we reiterate four priorities for bolstering the quality of evidence associated with rehabilitation outcomes: (1) revisit the recruitment and consent process and preserve ethical standards; (2) increase efforts and funding to support trials that examine functioning (i.e., cognitive, physical, and emotional); (3) consider multi-site recruitment options to increase participant diversity and sample sizes; (4) clearly identify the etiology of brain injury or offer disaggregate data in studies with mixed brain injury populations; and (5) promote the transparent reporting of adverse events, if applicable.

**Strengths and Limitations**

Evidence maps represent a novel approach that can be employed to detect broader issues, lead to research synthesis, and guide researchers in formulating both future research and studies with a narrower focus [42,43]. Evidence maps have been especially helpful in visualizing research contexts and appreciating how a specific focus fits into the broader research field [44]. In the case presented here, the evidence map aids in understanding how TBI research fits within the context of clinical research and where it stands overall in the field of rehabilitation.

A limitation that requires some discussion pertains to the search strategy. This overview exclusively analyzed systematic reviews published in the Cochrane library, which may have limited the inclusion of other high-quality systematic reviews on TBI. Nevertheless, Cochrane suggests this approach to preserve consistency in the results of the overview since the included works follow the same methodological standard [45].

We acknowledge that the evidence map developed for TBI is unable to address specific questions or nuances regarding the effectiveness of rehabilitation interventions in individuals with TBI.

Despite its limitations, the evidence map we have constructed disseminates evidence from existing literature findings on TBI, draws attention to the current challenges faced by researchers, and can provide an effective tool in guiding future research efforts and policymaking.

**5. Conclusions**

This work clarifies the need to expand research efforts in the context of TBI and clinical rehabilitation research to augment clinical applicability. In general, patients receiving rehabilitation services display a broad range of deficits and needs, which is particularly apparent among patients with TBI. Currently, the efficacy and safety of non-pharmacological and pharmacological interventions that are able to meet the needs of individuals with TBI remain uncertain, jeopardizing the clinical applicability of potentially effective interventions. To address the challenges experienced in clinical rehabilitation research, increasing the number of clinical and non-clinical trials performed that reflect sound methodology remains a priority.
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References

1. Cieza, A.; Causey, K.; Kamenov, K.; Hanson, S.W.; Chatterji, S.; Vos, T. Global estimates of the need for rehabilitation based on the Global Burden of Disease study 2019: A systematic analysis for the Global Burden of Disease Study 2019. Lancet Lond. Engl. 2021, 396, 2006–2017. [CrossRef]

2. Gimigliano, F.; Negrini, S. The World Health Organization “Rehabilitation 2030: A call for action”. Eur. J. Phys. Rehabil. Med. 2017, 53, 155–168. [CrossRef] [PubMed]

3. Negrini, S.; Arienti, C.; Patrini, M.; Kiekens, C.; Rauch, A.; Cieza, A. Cochrane collaborates with the World Health Organization to establish a Package of Rehabilitation Interventions based on the best available evidence. Eur. J. Phys. Rehabil. Med. 2021, 57, 478–480. [CrossRef] [PubMed]

4. Rauch, A.; Negrini, S.; Cieza, A. Toward Strengthening Rehabilitation in Health Systems: Methods Used to Develop a WHO Package of Rehabilitation Interventions. Arch. Phys. Med. Rehabil. 2019, 100, 2205–2211. [CrossRef] [PubMed]

5. Menon, D.K.; Schwab, K.; Wright, D.W.; Maas, A.I. Demographics and Clinical Assessment Working Group of the International and Interagency Initiative toward Common Data Elements for Research on Traumatic Brain Injury and Psychological Health. Position statement. Definition of traumatic brain injury. Arch. Phys. Med. Rehabil. 2010, 91, 1637–1640. [CrossRef]

6. Dewan, M.C.; Rattani, A.; Gupta, S.; Baticulon, R.E.; Hung, Y.C.; Punchak, M.; Agrawal, A.; Adeleye, A.O.; Shrim, M.G.; Rubiano, A.M.; et al. Estimating the global incidence of traumatic brain injury. J. Neurosurg. 2018, 130, 1080–1097. [CrossRef]

7. Kosty, J.A.; Stein, S.C. Measuring outcome after severe TBI. Neurol. Res. 2013, 35, 277–284. [CrossRef]

8. Wilson, J.T.; Pettigrew, L.E.; Teasdale, G.M. Structured interviews for the Glasgow Outcome Scale and the extended Glasgow Outcome Scale: Guidelines for their use. J. Neurotrauma 1998, 15, 573–585. [CrossRef]

9. James, S.L.; Theadom, A.; Ellenbogen, R.G.; Bannick, M.S.; Montjoy-Venning, W.; Lucchesi, L.R.; Abbasi, N.; Abdulkader, R.; Abrah, H.N.; Adsuar, J.C.; et al. Global, regional, and national burden of traumatic brain injury and spinal cord injury, 1990–2016: A systematic analysis for the Global Burden of Disease Study 2016. Lancet Neurol. 2019, 18, 56–67. Available online: https://www.thelancet.com/journals/lanneur/article/PIIS1474-4422(18)30415-0/fulltext (accessed on 30 April 2022). [CrossRef]

10. Brain Injury Facts [Internet]. International Brain Injury Association. Available online: https://www.internationalbrain.org/resources/brain-injury-facts (accessed on 1 May 2022).

11. Iaccarino, C.; Carretta, A.; Nicolosi, F.; Morselli, C. Epidemiology of severe traumatic brain injury. J. Neurosurg. Sci. 2018, 62, 535–541. [CrossRef]

12. Traumatic Brain Injury: Hope through Research! National Institute of Neurological Disorders and Stroke. Available online: https://www.ninds.nih.gov/Disorders/Patient-Caregiver-Education/Hope-Through-Research/Traumatic-Brain-Injury-Hope-Through (accessed on 1 May 2022).

13. Chan, V.; Mollayeva, T.; Ottenbacher, K.J.; Colantonio, A. Clinical profile and comorbidity of traumatic brain injury among younger and older men and women: A brief research notes. BMC Res. Notes 2017, 10, 371. [CrossRef] [PubMed]

14. Najem, D.; Rennie, K.; Ribecco-Lukiewicz, M.; Ly, D.; Haukenfrers, J.; Liu, Q.; Nzau, M.; Fraser, D.D.; Bani-Yaghoub, M. Traumatic brain injury: Classification, models, and markers. Biochem. Cell Biol. 2018, 96, 391–406. [CrossRef] [PubMed]

15. Hammond, F.M.; Corrigan, J.D.; Ketchum, J.M.; Malec, J.F.; Dams-O’Connor, K.; Hart, T.; Novack, T.A.; Bogner, J.; Dahdah, M.N.; Whitenecck, G.G. Prevalence of Medical and Psychiatric Comorbidities following Traumatic Brain Injury. J. Head Trauma Rehabil. 2019, 34, E1–E10. [CrossRef]

16. Forslund, M.V.; Ferrin, P.B.; Rea, C.; Sigurdardottir, S.; Hellstrom, T.; Berntsen, S.A.; Lu, J.; Arango-Lasprilla, J.C.; Andelic, N. Global Outcome Trajectories up to 10 Years After Moderate to Severe Traumatic Brain Injury. Front. Neurol. 2019, 10, 219. [CrossRef]

17. Bramlett, H.M.; Dietrich, W.D. Long-Term Consequences of Traumatic Brain Injury: Current Status of Potential Mechanisms of Injury and Neurological Outcomes. J. Neurotrauma 2015, 32, 1834–1848. [CrossRef]

18. Page, M.J.; McKenzie, J.E.; Bossuyt, P.M.; Boutron, I.; Hoffmann, T.C.; Mulrow, C.D.; Shamseer, L.; Tetzlaff, J.M.; Akl, E.A.; Brennan, S.E.; et al. The PRISMA 2020 statement: An updated guideline for reporting systematic reviews. J. Clin. Epidemiol. 2021, 134, 178–189. [CrossRef]
19. Levack, W.M.M.; Rathore, F.A.; Pollet, J.; Negrini, S. One in 11 Cochrane Reviews Are on Rehabilitation Interventions, According to Pragmatic Inclusion Criteria Developed by Cochrane Rehabilitation. *Arch. Phys. Med. Rehabil.* **2019**, *100*, 1492–1498. [CrossRef]

20. Shea, B.J.; Reeves, B.C.; Wells, G.; Thuku, M.; Hamel, C.; Moran, J.; Moher, D.; Tugwell, P.; Welch, V.; Kristjansson, E.; et al. AMSTAR 2: A critical appraisal tool for systematic reviews that include randomised or non-randomised studies of healthcare interventions, or both. *BMJ* **2017**, *358*, j4008. Available online: https://www.bmj.com/lookup/doi/10.1136/bmj.j4008 (accessed on 10 February 2022). [CrossRef]

21. Guyatt, G.; Oxman, A.D.; Akl, E.A.; Kunz, R.; Vist, G.; Brozek, J.; Norris, S.; Falck-Ytter, Y.; Glasziou, P.; deBeer, H.; et al. GRADE guidelines: 1. Introduction-GRADE evidence profiles and summary of findings tables. *J. Clin. Epidemiol.* **2011**, *64*, 383–394. [CrossRef]

22. Guyatt, G.H.; Oxman, A.D.; Vist, G.E.; Kunz, R.; Falck-Ytter, Y.; Alonso-Coello, P.; Schünemann, H.J. GRADE: An emerging consensus on rating quality of evidence and strength of recommendations. *BMJ* **2008**, *336*, 924–926. Available online: https://www.bmj.com/content/336/7650/924 (accessed on 9 November 2021). [CrossRef]

23. Wong, V.; Cheuk, D.K.; Lee, S.; Chu, V. Acupuncture for acute management and rehabilitation of traumatic brain injury. *Cochrane Database Syst. Rev.* **2013**, *3*, CD007700. Available online: https://www-cochranelibrary-com.ezproxy1.lib.asu.edu/cdsr/doi/10.1002/14651858.CD007700.pub2/full (accessed on 9 November 2021). [CrossRef] [PubMed]

24. Gertler, P.; Tate, R.L.; Cameron, I.D. Non-pharmacological interventions for depression in adults and children with traumatic brain injury. *Cochrane Database Syst. Rev.* **2015**, *12*, CD009871. Available online: https://www-cochranelibrary-com.ezproxy1.lib.asu.edu/cdsr/doi/10.1002/14651858.CD009871.pub2/full (accessed on 9 November 2021). [CrossRef] [PubMed]

25. Thompson, K.; Pohlmann-Eden, B.; Campbell, L.A.; Abel, H. Pharmacological treatments for preventing epilepsy following traumatic head injury. *Cochrane Database Syst. Rev.* **2015**, *8*, CD009900. [CrossRef] [PubMed]

26. Hassett, L.; Moseley, A.M.; Harmer, A.R. Fitness training for cardiorespiratory conditioning after traumatic brain injury. *Cochrane Database Syst. Rev.* **2012**, CD006123. Available online: https://www-cochranelibrary-com.ezproxy1.lib.asu.edu/cdsr/doi/10.1002/14651858.CD006123.pub3/full (accessed on 9 November 2021). [CrossRef] [PubMed]

27. Kumar, K.S.; Samuelkamaleshkumar, S.; Viswanathan, A.; Macaden, A.S. Cognitive rehabilitation for adults with traumatic brain injury to improve occupational outcomes. *Cochrane Database Syst. Rev.* **2017**, *6*, CD007935. Available online: https://www-cochranelibrary-com.ezproxy1.lib.asu.edu/cdsr/doi/10.1002/14651858.CD007935.pub2/full (accessed on 9 November 2021). [CrossRef]

28. Synnot, A.; Chau, M.; Pitt, V.; O’Connor, D.; Gruen, R.L.; Wasiak, J.; Clavisi, O.; Pattuswagen, L.; Phillips, K. Interventions for managing skeletal muscle spasticity following traumatic brain injury. *Cochrane Database Syst. Rev.* **2017**, *11*, CD008929. Available online: https://www-cochranelibrary-com.ezproxy1.lib.asu.edu/cdsr/doi/10.1002/14651858.CD008929.pub2/full (accessed on 9 November 2021). [CrossRef]

29. Wong, A.W.K.; Chen, C.; Baum, M.C.; Heaton, R.K.; Goodman, B.; Heinemann, A.W. Cognitive, Emotional, and Physical Functioning as Predictors of Employment following mild traumatic brain injury: A discriminant analysis. *J. Head Trauma Rehabil.* **2000**, *15*, 1103–1112. [CrossRef]

30. Ownsworth, T.; McKenna, K. Investigation of factors related to employment outcome following traumatic brain injury: A critical review and conceptual model. *Disabil. Rehabil.* **2004**, *26*, 765–783. [CrossRef]

31. Watana, S. Vocational rehabilitation for clients with cognitive and behavioral disorders associated with traumatic brain injury. *Work (Read. Mass.)* **2013**, *45*, 273–277.

32. Bogdanova, Y.; Verfaellie, M. Cognitive sequelae of blast-induced traumatic brain injury: Recovery and rehabilitation. *Neuropsychol. Rev.* **2012**, *22*, 4–20. [CrossRef] [PubMed]

33. Wilson, C.D.; Burks, J.D.; Rodgers, R.B.; Bakare, A.A.; Safavi-Abbasi, S. Early and Late Posttraumatic Epilepsy in the Setting of Traumatic Brain Injury: A Meta-analysis and Review of Antiepileptic Management. *World Neurosurg.* **2013**, *79*, 273–277. [CrossRef] [PubMed]

34. Dang, K.; Gupta, P.K.; Diaz-Arrastia, R. Chapter 14: Epilepsy after Traumatic Brain Injury. In *Translation Research in Traumatic Brain Injury*; CRC Press: Boca Raton, FL, USA, 2016; pp. 299–313. Available online: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6436116/ (accessed on 18 November 2021). [CrossRef]

35. Drake, A.I.; Gray, N.; Yoder, S.; Pramuka, M.; Llewellyn, M. Factors predicting return to work following mild traumatic brain injury: A discriminant analysis. *J. Head Trauma Rehabil.* **2000**, *15*, 1103–1112. [CrossRef]

36. Menon, D.K. Unique challenges in clinical trials in traumatic brain injury. *Crit. Care Med.* **2009**, *37*, S129–S135. [CrossRef]

37. Stanley, R.M.; Johnson, M.D.; Vance, C.; Bajaj, L.; Babcock, L.; Aatabaki, S.; Thomas, D.; Simon, H.K.; Cohen, D.M.; Rubacalva, D.; et al. Challenges Enrolling Children Into Traumatic Brain Injury Trials: An Observational Study. *Acad. Emerg. Med.* **2017**, *24*, 31–39. [CrossRef] [PubMed]

38. Gerber, L.H.; Deshpande, R.; Moosvi, A.; Zafonte, R.; Bushnik, T.; Garfinkel, S.; Cai, C. Narrative review of clinical practice guidelines for treating people with moderate or severe traumatic brain injury. *NeuroRehabilitation* **2021**, *48*, 451–467. [CrossRef] [PubMed]
40. Gerber, L.H.; Bush, H.; Cai, C.; Garfinkel, S.; Chan, L.; Cotner, B.; Wagner, A. Scoping review of clinical rehabilitation research pertaining to traumatic brain injury: 1990–2016. *NeuroRehabilitation* 2019, 44, 207–215. [CrossRef]

41. Maas, A.I.R.; Menon, D.K.; Lingsma, H.F.; Pineda, J.A.; Sandel, M.E.; Manley, G.T. Re-orientation of clinical research in traumatic brain injury: Report of an international workshop on comparative effectiveness research. *J. Neurotrauma* 2012, 29, 32–46. [CrossRef]

42. Bragge, P.; Clavisi, O.; Turner, T.; Tavender, E.; Collie, A.; Gruen, R.L. The Global Evidence Mapping Initiative: Scoping research in broad topic areas. *BMC Med. Res. Methodol.* 2011, 11, 92. [CrossRef]

43. Katz, D.; Williams, A.-L.; Girard, C.; Goodman, J.; Comerford, B.; Behrman, A.; Bracken, M.B. The evidence base for complementary and alternative medicine: Methods of Evidence Mapping with application to CAM. *Altern. Ther. Health Med.* 2003, 9, 22–30.

44. Althuis, M.D.; Weed, D.L. Evidence mapping: Methodologic foundations and application to intervention and observational research on sugar-sweetened beverages and health outcomes. *Am. J. Clin. Nutr.* 2013, 98, 755–768. [CrossRef] [PubMed]

45. Pollock, M.; Fernandes, R.; Becker, L.; Pieper, D.; Hartling, L. Chapter V: Overviews of Reviews. In *Cochrane Handbook for Systematic Reviews of Interventions;* Version 6.2; Higgins, J.P.T., Thomas, J., Chandler, J., Cumpston, M., Li, T., Page, M.J., Welch, V.A., Eds.; Cochrane, 2021. Available online: [www.training.cochrane.org/handbook](http://www.training.cochrane.org/handbook) (accessed on 13 December 2021).