







COMPLEX REGIONAL PAIN SYNDROME WHAT IS THE OUTCOME? - A SYSTEMATIC REVIEW OF THE COURSE AND SEVERITY OF **CRPS SYMPTOMS AT 12 MONTHS AND BEYOND**

Selina Johnson 1,2*; Fiona Cowell 3, Sharon Gillespie 3, Andreas Goebel 1,2.

1. Walton Centre NHS Foundation Trust; Liverpool, UK, 2. Pain Research Institute, Faculty of health and life Sciences, University of Liverpool; Liverpool, UK, 3. Liverpool University Hospitals Foundation Trust (LUHFT), UK.

Background :

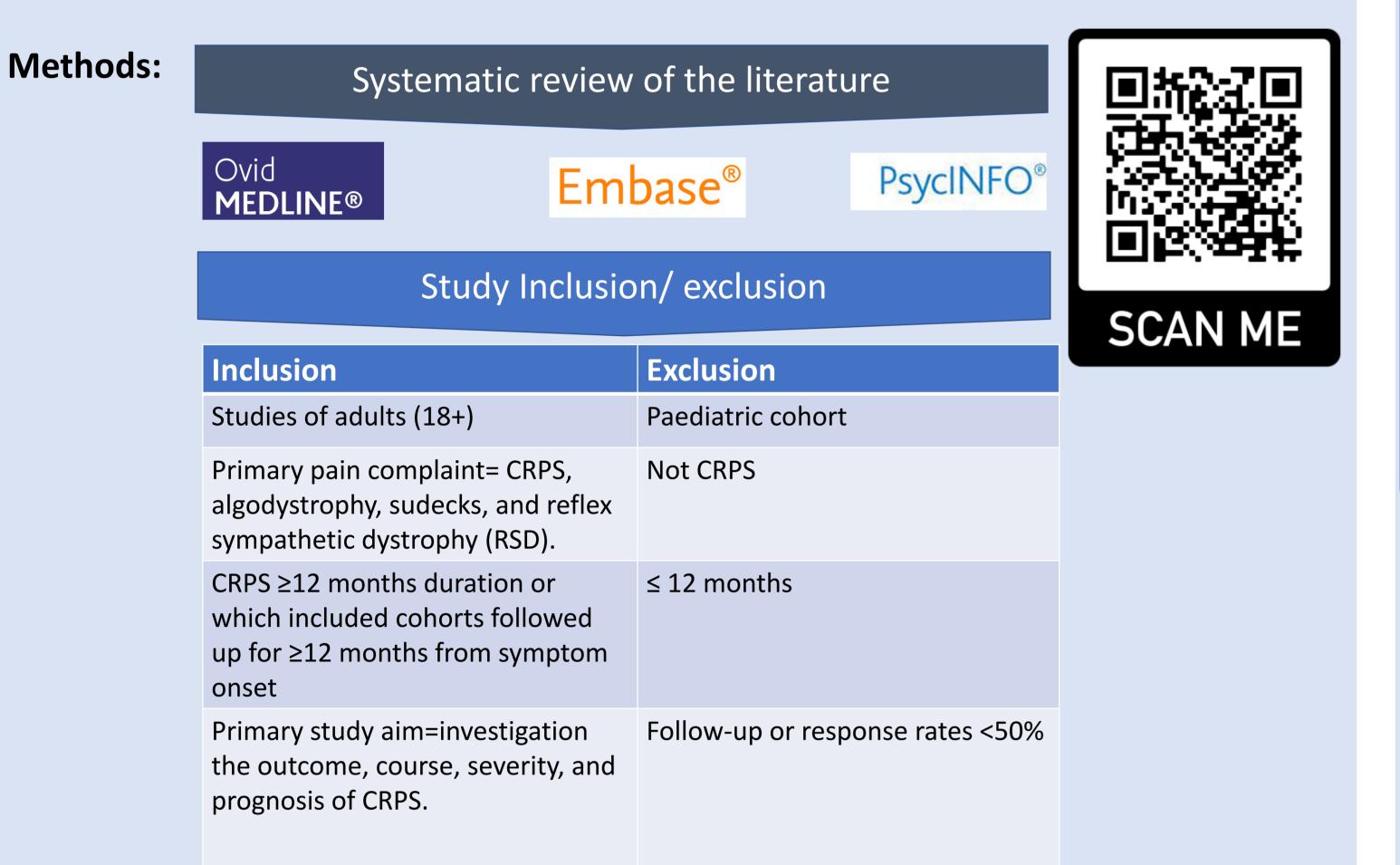
To improve CRPS treatment, it is imperative to understand the *degree, nature, and relative importance* of any ongoing CRPS related problems.

An earlier systematic review comprehensively discussed CRPS symptom recovery in 2012 (1). Information regarding the *functional and occupational impact of CRPS* is needed to understand the individual health economic impact of CRPS, and ensure treatment supports recovery that is meaningful to patients. Therefore, there was good rationale to update the previous review to provide specific information regarding the physical and social/occupational impact of CRPS symptoms.

Results 2:

Overall: Pain and motor dysfunction were found to be the most prevalent ongoing symptoms affecting between 51-89% of all patients at longer term follow up. These features dramatically impact on a person's physical and social abilities. The measurement of physical and social disability was highly variable. Results indicate CRPS is associated with 25% - 66% reduction in grip strength and prevents return to work for 30-40% of cases of at \geq 12 months. The current review provides first time quantitative data on function and work status for CRPS ≥ 12 months and builds on

Aims: This systematic review aimed to summarise the published data concerning the *impact of CRPS symptoms, specifically the physical and occupational impact* of symptoms, at 12 months from symptom onset and beyond.



evidence provided by review.

Motor Function:

51-89% of patients experience weakness, stiffness, and reduced range of movement at \geq 12 months (14 studies)

Prospective studies show 20%-25% reduction in range of movement 25% - 50% reduction in grip strength at \geq 12 months

The impact of such losses will have varying significance dependent on factors including occupation.

Work status:

30-40% of working people Who develop CRPS **DO NOT return to work** (7 studies)

27-35% of those who DO return to work require some form of work role adaptation

Quality assessment

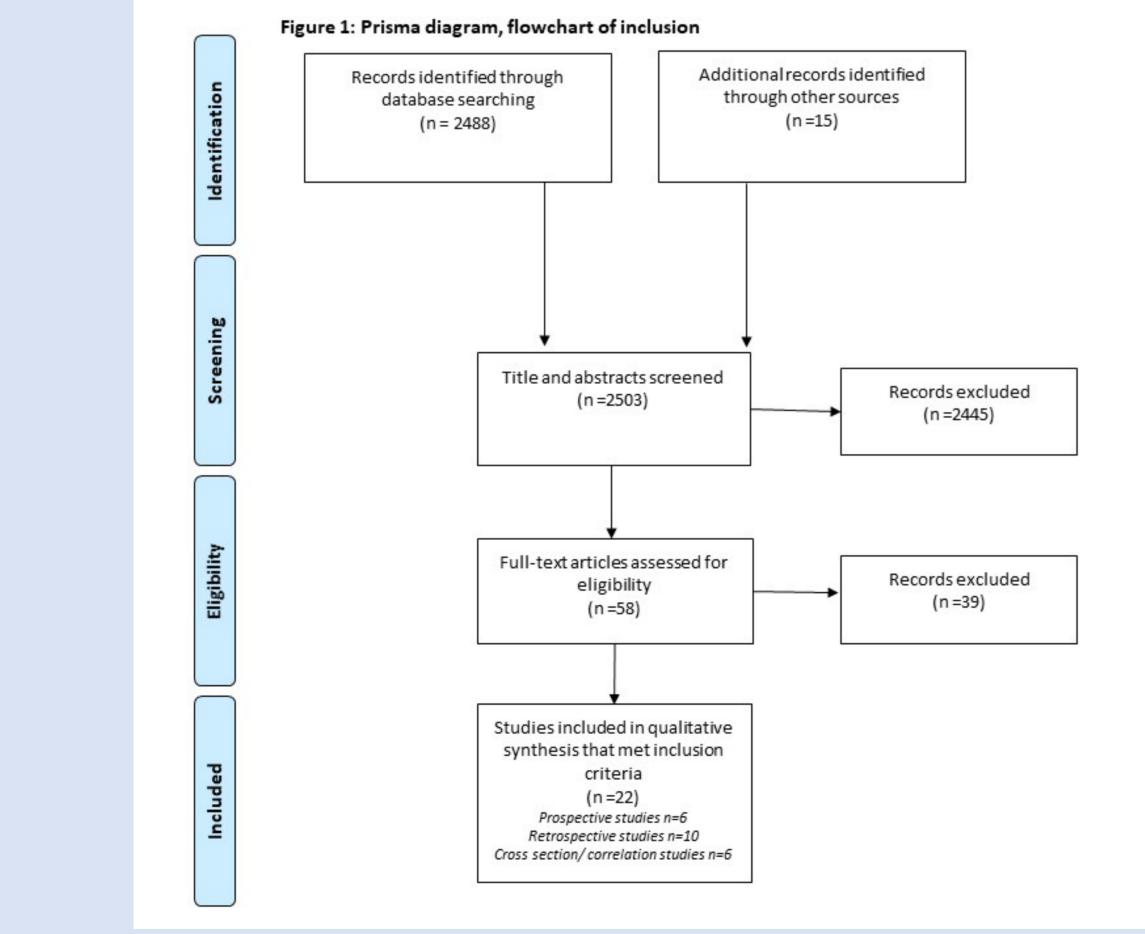
Joanna Briggs Institute (JBI) critical appraisal prevalence study checklist Assessment by 2 authors and discrepancies resolved by a 3rd.

Data extraction

Study type, sample and method, diagnostic criteria, duration of CRPS at baseline, duration of CRPS at follow-up, timing of assessments, and outcome measures (CRPS Signs and symptoms, and physical and social disability).

Data Synthesis data were synthesised according to study type and findings of the relevant outcome measures were reported.

Results 1:



The current health economic impact in terms of occupational recovery is not yet fully understood and what factors return to work.

Physical and social disability:

62%-86 % patients report CRPS affects their Activities of daily life

Heterogeneity in outcomes reflects wide ranging impact of disability.

Conclusions

Results from this review concur with review (1) 10 years ago indicating pain and motor dysfunction are the most persistent disabling features of ongoing CRPS.

We now also report *first time quantitative data specific evidence about losses to* motor function, long-term compromises to work status.

Results indicate despite general improvements in features of CRPS, the ongoing impact of CRPS on hand function and work status is relatively high.

Future research should explore what drives limitation to function and work status and if and how these limitations can be prevented.

REFERENCES:

Bean DJ, Johnson MH, Kydd RR (2014b). Critical Review the Outcome of Complex Regional Pain Syndrome Type 1: A Systematic Review. J Pain, 15(7):677–90. http://dx.doi.org/10.1016/j.jpain.2014.01.500 Author contact details: <u>sjohns@liverpool.ac.uk</u>, twitter @sillyeena