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Prevalence of the Dissociative Subtype of Post-Traumatic Stress Disorder: A Systematic Review and Meta-Analysis

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**Prevalence of the Dissociative Subtype of Post-Traumatic Stress Disorder: A Systematic Review
and Meta-Analysis**

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Abstract

The dissociative subtype of post-traumatic stress disorder (PTSD-DS) was introduced in the fifth edition of the Diagnostic and Statistical Manual of Mental Disorders (DSM-5), and is characterised by symptoms of either depersonalisation or derealisation, in addition to a diagnosis of post-traumatic stress disorder (PTSD). This systematic review and meta-analysis sought to estimate the prevalence of current PTSD-DS, and the extent to which method of assessment, demographic and trauma variables moderate this estimate, across different methods of prevalence estimation. Studies included were identified by searching MEDLINE (EBSCO), PsycInfo, CINAHL, Academic Search Complete, and PTSDpubs, yielding 49 studies that met the inclusion criteria ($N = 8214$ participants). A random effects meta-analysis estimated the prevalence of PTSD-DS as 38.1% (95% CI 31.5–45.0%) across all samples, 45.5% (95% CI 37.7–53.4%) across all diagnosis-based and clinical cut-off samples, 22.8% (95% CI 14.8–32.0%) across all latent class analysis (LCA) and latent profile analysis (LPA) samples, and 48.1% (95% CI 35.0–61.3%) across samples which strictly used the DSM-5 PTSD criteria; all as a proportion of those already with a diagnosis of PTSD. All results were characterised by high levels of heterogeneity, limiting generalisability. Moderator analyses mostly failed to identify sources of heterogeneity. PTSD-DS was more prevalent in children compared to adults, and in diagnosis-based and clinical cut-off samples compared to LCA and LPA samples. Risk of bias was not significantly related to prevalence estimates. The implications of these results are discussed further.

Keywords: Meta-Analysis; Prevalence; Stress Disorders, Post-Traumatic; Systematic Review

Introduction

In the fifth edition of the Diagnostic and Statistical Manual of Mental Disorders (DSM-5; American Psychiatric Association [APA], 2013), post-traumatic stress disorder (PTSD) is classified as a Trauma- and Stressor-Related Disorder. A diagnosis is based on a required number of symptoms across domains of intrusion, avoidance, negative alterations in cognitions and mood, and alterations in arousal and reactivity. Also stipulated in DSM-5 are the criteria required for specifying the dissociative subtype of PTSD (PTSD-DS) where, in addition to first meeting the criteria for PTSD diagnosis, individuals must endorse symptoms of depersonalisation and or derealisation. Depersonalisation involves “persistent or recurrent experiences of feeling detached from, and as if one were an outside observer of, one’s mental processes or body”, whereas derealisation takes the form of “persistent or recurrent experiences of unreality of surroundings” (DSM-5, 2013, pp. 272).

It has been extensively documented that persistent dissociation is linked to post traumatic symptomology (Carlson, Dalenberg, & McDade-Montez, 2012). The subtype model suggests that PTSD and PTSD-DS are distinct from one another (Dalenberg & Carlson, 2012), where PTSD-DS presents with its own epidemiological features (Schivavone, Frewen, McKinnon, & Lanius, 2018). A recent systematic review concluded that there may be an association between PTSD-DS and psychopathological comorbidity and childhood abuse and neglect (Steuwe, Lanius, & Frewen, 2012), adult sexual abuse (Wolf, Miller et al., 2012), and with depression, suicidal thinking, and drug overdoses (Mergler et al., 2017), despite there being a large degree of heterogeneity in the literature concerning risk factors for PTSD-DS (Hansen, Ross, & Armour, 2017). This indicates that PTSD-DS may reflect a more severe form of PTSD (Zoet, Wagenmans, van Minnen, & de Jongh, 2018), although this is not directly assessed in this study.

One criticism of the PTSD-DS diagnosis is that the symptoms of dissociation chosen as necessary criteria to achieve a diagnosis in DSM-5 are too narrow, where it is believed that the current criteria should also include other symptoms of dissociation (Ross, 2021), following evidence that: dissociative amnesia (Wolf et al., 2017), and flashbacks (Dahal, Kumar, & Thapa, 2018; Hyland et al., 2017) are common in individuals with PTSD. Additionally, memory disturbance, disengagement, time loss, and trance (Frewen, Brown, Steuwe, & Lanius, 2015), gaps in awareness, re-experiencing, and sensory misperception (Müllerová, Hansen, Contractor, Elhai, & Armour, 2016; Ross, Baník, Dědová, Mikulášková, & Armour, 2018) are associated with PTSD-DS. However, to some extent, these symptoms are already captured by the existing PTSD criteria.

Several methodologies have been used to determine the prevalence of PTSD-DS, with early studies using taxometric (Waelde, Silvern, & Fairbank, 2005; Waller & Ross, 1997), and signal detection (Ginzburg et al., 2006) analyses. The prevalence of PTSD-DS has also been described in studies where participants were selected primarily due to a specific comorbid difficulty, such as substance abuse disorder and psychosis, using the DSM-5 diagnostic criteria (Gidzgieer et al., 2019; Mergler et al., 2017; van Minnen et al., 2016), and in studies that assessed subsyndromal PTSD (Bennett, Modrowski, Kerig, & Chaplo, 2015; Kerig et al., 2016; Modrowski & Kerig, 2017). Prevalence rates of PTSD-DS have been reported in different ways; some with respect to the total number of participants regardless of whether the sample tested had PTSD, some were only trauma-exposed or from a community sample, whereas other prevalence rates were with respect to those with PTSD. This makes it challenging to make comparisons between studies. Hansen et al.'s (2017) systematic review of latent class and profile analyses (LCA and LPA respectively) indicated the mean prevalence of PTSD-DS as 20.4%. LCA determines hidden groups based on the means of categorical variables, whereas LPA does the same for continuous variables (Oberski, 2016).

Both LCA and LPA are exploratory techniques that determine underlying hidden profiles or groups of individuals from observed data who display similar patterns of symptoms (Muthén, 2004; Oberski, 2016). The ‘best’ number of groups is determined by the most appropriate model fit, and whilst there are many methods for determining the number of classes or profiles, the two most common methods are the Akaike information criterion and Bayesian information criterion (where lower values indicate a better fit). However, the selection of the optimal number of classes or profiles, and the qualitative naming of each group, remains subjective on the part of the researcher which has implications for valid prevalence estimation (Hansen et al., 2017). In addition, Hansen et al. (2017) averaged the prevalence values despite dissociation being defined differently in various studies; some used the DSM-5 criteria stipulating symptoms of either depersonalisation or derealisation, and other studies assessed a wider spectrum of dissociative experiences. Finally, due to methodological constraints, there was no way of breaking down the heterogeneous nature of the population (Hansen et al., 2017).

There is a need to comprehensively systematically review studies to attempt to establish some consensus around how prevalent PTSD-DS is in children and adults. This study aimed to conduct a broad meta-analysis of data from studies investigating current PTSD-DS to reach a reliable estimate of prevalence from studies utilising various methods of prevalence estimation, furthering the systematic review of Hansen et al. (2017). The aim was to provide greater insight into the heterogeneity that is common within participants with PTSD. This might lead to the development of risk factors for this particular subtype and help the structuring of efficacious interventions. This review will be, to the authors’ knowledge, the first of its kind to meta-analyse the prevalence of PTSD-DS in participants with PTSD, assessing moderators that affect PTSD-DS prevalence, and using studies utilising different methods of prevalence estimation. There is disagreement as to what symptoms of dissociation

should be required as necessary criteria to achieve a diagnosis of PTSD-DS, and this review may shed further light on this debate, by comparing the prevalence rates of PTSD-DS when defined by depersonalisation and or derealisation, and when dissociation is defined more broadly.

Method

The protocol for this review was pre-registered on PROSPERO (reference: CRD42021210902) prior to any formal review of searches.

Search Strategy

Relevant studies were identified through a systematic search of the following databases: MEDLINE (EBSCO), PsycInfo, CINAHL, Academic Search Complete, and PTSDpubs. Studies included were those published from 1st January 1980, when the Diagnostic and Statistical Manual of Mental Disorders first defined PTSD according to DSM-III (APA, 1980), and before 14th February 2021 when the searches were conducted.

The following search terms were used for each database, processing study titles and abstracts only: (posttrauma* OR post-trauma* OR "post trauma*" OR PTSD OR PTSS) AND (dissociat* OR depersonali* OR dereali*). Medical Subject Headings (MeSH) terms, and other equivalent key words for other databases, were used for each search term: 'post-traumatic stress disorder', 'post-traumatic stress', 'posttraumatic stress disorder', 'posttraumatic stress' 'post-traumatic stress disorder in children', 'stress disorders, post-traumatic', 'complex PTSD', 'PTSD', 'PTSD (DSM-III)', 'PTSD (DSM-III-R)', 'PTSD (DSM-IV)', 'PTSD (DSM-5)', 'PTSD (ICD-9)', 'PTSD (ICD-10)', 'PTSD (ICD-11)', 'dissociation', and 'depersonalization'.

The reference sections of relevant systematic reviews and meta-analyses were also searched to ensure studies were not missed.

Inclusion and Exclusion Criteria

Studies were included in this review if data were presented on the prevalence of PTSD-DS following a traumatic event. In a bid to take a broad and comprehensive approach, the prevalence of PTSD-DS was defined as the number of participants: who scored above a clinical cut-off on a validated measure or who met DSM diagnostic criteria following a clinical interview or self-report measure, or who were categorised into a distinct class or profile following LCA or LPA. Studies of participants of all ages, any sex, and from either community or clinical samples were included. Studies were excluded: if they were not written in English; if participants were selected primarily due to a specific comorbid disorder; if PTSD was assessed acutely within a month of the index trauma; if exclusively lifetime PTSD or PTSD-DS prevalence was reported; if subsyndromal PTSD was assessed only; if dissociation was triggered via experimental manipulation; or if studies used analyses other than LCA, LPA, diagnostic, or clinical cut-off to determine the prevalence of PTSD-DS. Qualitative methodology, single case studies, reviews and meta-analyses were also excluded.

Screening, Data Extraction, Coding and Synthesis

All studies were screened, and the data extracted by the first author (WW) using a database which indexed the information provided in Table 1. The extracted data for all studies were reviewed by an independent researcher (AO), so as to reduce the likelihood of error (Buscemi, Hartling, Vandermeer, Tjosvold, & Klassen, 2006). Any queries were discussed, and agreement reached between the researchers. Wherever there was continued disagreement, a final decision was made by the senior researcher (RM-S). Where there was missing information, authors were contacted directly.

During data extraction, several rules were followed to ensure consistency between studies. Articles such as Eidhof et al. (2019), Guetta et al. (2019), and Zoet et al. (2018) used multiple measures for the assessment of PTSD, however in these cases the Clinician

Administered PTSD scale (CAPS) was prioritised as it is regarded as the gold standard for assessing PTSD (Weathers et al., 2004). Other studies assessed multiple populations (Hansen, Müllerová, Elklit, and Armour, 2016; Kenny, Helpingstine, Long, & Harrington, 2020; Wolf, Lunney et al., 2012), or used multiple analyses (Choi et al., 2017; 2019; Hansen, Hyland, Armour, & Andersen, 2019), and therefore these were treated separately in this review as individual samples. Care was taken to ensure that no dataset contributed more than one data point in any one meta-analysis (where diagnostic and clinical cut-off samples were prioritised over LCA and LPA samples). Multiple studies investigating the same population were removed, retaining the study with the largest sample size. Many studies (Cloitre, Petkova, Wang, & Lu, 2012; Daniels, Frewen, Theberge, & Lanius, 2016; Swart, Wildschut, Fraijer, Langeland, & Smit, 2020; Tsai, Armour, Southwick, & Pietrzak, 2015) reported means and standard deviations for participant age and sex in aggregated format, rather than for the sample as a whole. For these studies, the means and standard deviations were combined (Altman et al., 2013; Higgins et al., 2012). When absolute frequencies were not reported, these were calculated from the reported percentage prevalence. For the LCA and LPA samples, only those classed as having ‘moderate’ to ‘severe’ symptomology were deemed to meet ‘caseness’ for PTSD and PTSD-DS. The prevalence of PTSD-DS was consistently calculated as a proportion of all participants with PTSD.

Quality Assessment and Risk of Bias

Two authors (WW & AO) assessed the risk-of-bias using a researcher developed tool based on the Assessment Tool for Observational Cohort and Cross-Sectional Studies (National Heart Lung and Blood Institute, 2014), and modified questions from other relevant prevalence and risk factor studies (Hoy et al., 2012; Munn, Moola, Riitano, & Lisy, 2014). The quality assessment checklist (see Supplementary Material) consisted of five items assessing how well the population and index trauma were specified, the rate of participation,

and whether objective and standard criteria were used for the assessment of PTSD and PTSD-DS. Each item used a three-point scale (0-2), and the following categorical system was used to rate the total risk-of-bias score: 0-4 high risk/low quality, 5-6 moderate risk/quality, 7-10 low risk/high quality, following the methodology used by Memarzia, Walker, and Meiser-Stedman (2021). An inter-rater reliability assessment was conducted for all ratings between the two raters (WW & AO) which indicated a good correlation on all items (intraclass correlation = 0.87, 95% CI 0.77–0.93).

Meta-Analytic Method

The meta-analysis was conducted using R (version 4.1.1) which uses the metafor package (version 3.0-2; Viechtbauer, 2010). The extracted prevalence of PTSD-DS, as a proportion of all PTSD cases, was pooled to provide a weighted estimate of the prevalence of PTSD-DS overall (with 95% confidence intervals [CI]).

A random effects model was used given the high degree of variability expected in effect size between samples as it provides a broader and more conservative 95% confidence interval around the estimate of the prevalence.

The estimates of the prevalence underwent an arcsin transformation to ensure that the confidence intervals did not fall below zero for samples where the prevalence estimate was low (Barendregt, Doi, Lee, Norman, & Vos, 2013); results were then back transformed for ease of interpretation.

Cochran's Q test (Cochran, 1954) was used to ascertain if heterogeneity within samples was significant. The I^2 statistic (Higgins & Thompson, 2002) was used to determine the percentage of total variation in sample estimates that is due to between-study heterogeneity.

Moderator analyses of prevalence estimates were conducted to ascertain if sample characteristics impacted the prevalence estimate. These characteristics included: method of

PTSD-DS assessment, which DSM criteria was used, participant age group, occupation, and the type of trauma suffered. These were included as there were multiple samples that allowed for these comparisons to be made. A sensitivity analysis was used to assess the impact of risk-of-bias on the estimated pooled prevalence. This was achieved by repeating the meta-analysis, excluding those samples that constituted a high risk-of-bias. Any differences in the moderator and sensitivity analyses were tested for clinical significance by meta-analytic regression.

A funnel plot was used to assess for publication bias (Higgins et al., 2012), however this is less likely to occur in prevalence studies given there is no assessment of clinical significance, and therefore it is less likely that there is a bias in levels of acceptance to journals (Brewin, Andrews, & Valentine, 2000). The 'trim-and-fill' method was used (Duval & Tweedie, 2000), where any missing null or weaker studies are estimated to improve the symmetry of the sample distribution.

Results

The Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) diagram shows that 337 studies met the eligibility criteria following the initial screen of titles and abstracts (Figure 1). Full text reviews were conducted again, leading to 49 studies being included in the meta-analysis. Four studies were split into two samples due to different characteristics, index traumas or analyses, leaving 53 samples included in this review (Table 1). Around half the samples were treatment-seeking ($k = 23$), and PTSD-focussed (a diagnosis of PTSD was an inclusion criterion; $k = 22$). Nine samples included only female participants, three samples included only males, and the rest were mixed or the sex was not reported. The majority of included samples were adult ($k = 41$); only five exclusively comprised children. Samples mostly originated from high-income countries ($k = 49$).

[Figure 1]

[Table 1]

Risk-of-Bias Assessment

Twelve samples were deemed to be at high risk-of-bias, 16 were moderate risk, whereas 25 were low risk. The proportion of samples rated as low, moderate and high risk across the five quality assessment items can be seen in the Supplementary Material.

Prevalence

The pooled prevalence of PTSD-DS estimates and heterogeneity statistics for all samples can be seen in Table 2. The overall pooled prevalence was 38.1%. For diagnosis-based and clinical cut-off samples the pooled prevalence was 45.5%, while for latent class and profile samples the estimate was 22.8%. Meta-regression analyses indicated that the prevalence of PTSD-DS in the diagnosis-based or clinical cut-off samples was statistically significantly greater than the LCA or LPA samples (see Figure 2 for forest plot). The range of prevalence overall was 0-100%, and the degrees of between sample heterogeneity were extremely high.

[Table 2]

[Figure 2]

Moderator Analyses

All Samples

Moderator analyses were conducted for all samples to assess whether the pooled prevalence estimate of PTSD-DS was associated with demographic, trauma or assessment factors (Table 2). Meta-regression analyses confirmed that the prevalence of PTSD-DS in the child samples was statistically significantly greater than the adult samples, although there were only four child samples for comparison. All other comparisons were non-significant; however, several comparisons were likely underpowered.

Diagnostic and Clinical Cut-off Samples

Further subgroup moderator analyses were conducted separately for the diagnostic and clinical cut-off samples (Table 3), regardless of the dissociation criteria used, given the significant difference in pooled prevalence estimates of PTSD-DS between these samples and those using LCA or LPA. Meta-regression analyses confirmed again that the prevalence of PTSD-DS in the child samples was statistically significantly greater than the adult samples, although there were only four child samples for comparison. All other comparisons were non-significant, however several comparisons were likely underpowered.

[Table 3]

Further moderator analyses were conducted for only those samples utilising DSM-5 criteria for dissociation (depersonalisation and or derealisation; see Supplementary Material). When only samples using DSM-5 diagnostic and clinical cut-off criteria for the assessment of PTSD and PTSD-DS were pooled, the estimated prevalence of PTSD-DS was 48.2%. This provides the most valid estimate of PTSD-DS prevalence according to the DSM-5 criteria. Meta-regression analyses confirmed again that the prevalence of PTSD-DS in the child samples was statistically significantly greater than the adult samples, although there were only four child samples for comparison. All other comparisons were non-significant, however several comparisons were likely underpowered.

Latent Class and Profile Samples

Moderator analyses were conducted separately for the LCA and LPA samples (see Supplementary Material), again given the significant difference in pooled prevalence of PTSD-DS estimates between these samples and those using diagnostic and clinical cut-off methods. Meta-regression analyses confirmed that there were no statistically significant differences, however several comparisons were likely underpowered.

Sensitivity Analyses

When the 28 samples of low and moderate quality were removed, the estimated prevalence of PTSD-DS was not dissimilar to that for all samples (35.7%, 95% CI 24.8–47.3%) with a similar degree of between sample heterogeneity ($k = 25$, $Q(24) = 717.8$, $p < 0.0001$, $I^2 = 98.5\%$). Meta-regression analyses indicated there was not a significant difference between high and low-moderate quality groups ($\beta = 0.0040$ [95% CI -0.1384, 0.1463], $p = 0.96$). Therefore, it can be concluded that there was no support for the quality of the samples affecting the prevalence of PTSD-DS estimates.

Given the differences in prevalence in PTSD-DS between child and adult samples, the child samples were removed to assess whether similar results were achieved as in Table 2. Meta-regression analyses confirmed that the only statistically significant difference existed between the estimated prevalence of PTSD-DS for the diagnosis-based or clinical cut-off samples and LCA or LPA samples ($\beta = -0.2159$ [95% CI -0.3531, -0.0787], $p = 0.002$). All other comparisons were non-significant ($ps = 0.19$ – 0.87), however several comparisons were likely underpowered.

Publication Bias

Visual inspection of the funnel plot (see Supplementary Material) suggests the distribution of samples is asymmetrical, which was confirmed by Egger's test ($p = 0.03$). However, the study of Kenny et al. (2020; sample 29) was very small and should be considered an outlier. When this sample was removed, the Egger's test confirmed the symmetry of the distribution ($p = 0.30$). No null or weaker studies were estimated as missing, indicating little to no publication bias.

Discussion

A comprehensive systematic review and meta-analysis of prevalence data from studies investigating current PTSD-DS utilising various methods of prevalence estimation

was completed. The aim was to generate a reliable estimate for the prevalence of PTSD-DS and to provide greater insight into the heterogeneity that is common within participants with PTSD-DS. The estimated pooled prevalence of PTSD-DS was: 38.1% for all samples, 45.5% for all diagnostic and clinical cut-off samples, and 22.8% for all LCA and LPA samples. The estimated prevalence of PTSD-DS from the LCA and LPA samples was similar to the mean prevalence found in the Hansen et al. (2017) systematic review (20.4%); this is unsurprising given nine of the eleven studies in the Hansen et al. (2017) review were also included in the present study. When only samples strictly using DSM-5 diagnostic and clinical cut-off criteria for the assessment of PTSD and PTSD-DS were pooled, the estimated prevalence of PTSD-DS was 48.1%. The prevalence of PTSD-DS may therefore be significantly greater than previously suggested.

Impact of Diagnostic and Clinical Cut-off Assessment Versus LCA and LPA on Estimated Prevalence of PTSD-DS

The estimated prevalence of PTSD-DS for the diagnostic and clinical cut-off samples was significantly higher than that of the LCA and LPA samples. Use of clinical cut-off measures may overestimate the prevalence of PTSD in adults (Richardson, Frueh, & Acierno, 2010). Moreover, it may be easier to identify individuals with PTSD who show symptoms of depersonalisation or derealisation in a clinical interview or that surpass a clinical cut-off on a dissociation measure, rather than via LCA and LPA methods. On the other hand, latent class and profile analyses may rely on participants reporting multiple significant dissociative symptoms rather than just one symptom to a significant level. Achterhof, Huntjens, Meewisse, and Kiers (2019) questioned the use of LCA and LPA to ascertain the prevalence of Complex PTSD and highlighted that despite the analyses determining distinct profiles, the symptom profile for groups of participants were very close to one another and even

overlapped on occasion. Therefore, it may be questioned whether LCA and LPA reliably and validly estimates subtype prevalence.

Impact of Moderators on Estimated Prevalence of PTSD-DS

There was no significant difference between the estimated prevalence of PTSD-DS when dissociation was assessed by the DSM-5 criteria (presence of either depersonalisation or derealisation) or when defined by a broader spectrum of dissociative symptoms. The aim of the inclusion of the PTSD-DS in DSM-5 was to define a small subgroup of individuals with consistent clinical and epidemiological features (Miller, Wolf, & Keane, 2014; Schiavone et al., 2018), however results from the present study suggest a subtype where the prevalence varies very widely across samples (0-100%) and where the heterogeneity cannot be broken down following moderator analyses. Research literature suggests that the symptomology of PTSD is itself heterogeneous (Elhai, Frueh, Davis, Jacobs, & Hamner, 2003; Galatzer-Levy & Bryant, 2013; Naifeh, Richardson, Del Ben, & Elhai, 2010), where dissociation is one such symptom that can vary.

The estimated prevalence of PTSD-DS was significantly higher for samples of children compared to adults, although there are limited number of samples investigating exclusively children, and the results were dominated by that of Choi et al. (2019; sample 10). There was no one trauma type that best categorised the child samples. Research has shown that dissociation is a common experience for children, that later becomes less prevalent with child development and the transition into adulthood (Brunner, Parzar, Schuld, & Resch, 2000; Coons, 1996; Choi et al., 2017; Shimizu & Sakamoto, 1986). Choi et al. (2019) reported that 53.7% of children with PTSD had the dissociative subtype; a prevalence much higher than in many other adult samples, and the authors cited the prominence of dissociation as a form of coping in response to maltreatment in childhood (Liotti, 2004; Putnam, 1997). Children may be more susceptible to PTSD-DS because they do not have the same capacity

to avoid cues relating to the traumatic event, especially when the trauma was based within the home environment, or with a primary caregiver (Choi et al., 2019). In children, dissociation may offer an alternative method of escape to reduce distress. It might also be considered whether depersonalisation and derealisation are the most appropriate symptoms by which to assess for PTSD-DS in children. The premise of the subtype model is that these dissociative symptoms are rare (Lanius et al., 2014), however it may be that dissociative experiences are more common in youth (Carlson, Yates, & Sroufe, 2009) and may not even be considered as pathological. Further research is required within this area to determine whether children are more at risk from dissociation in the context of PTSD compared to adults, as the lack of power within the samples of children frustrated the moderator analyses.

Other than age group, all other moderator analyses yielded non-significant results indicating no support for any differences between estimated prevalence of PTSD-DS. This is surprising given the extant research on mediators and risk factors in relation to PTSD-DS (Hansen et al., 2017; Schiavone et al., 2018 for review), but these non-significant results are likely to reflect the heterogeneity between these samples and the lack of power in some moderator analyses.

It is important to stress that the pooled prevalence estimates were characterised by a high degree of heterogeneity throughout, and inspection of the forest plot (Figure 2) shows how varied the prevalence of PTSD-DS is across different samples. This is not unexpected given the multiple ways of assessing and conceptualising PTSD-DS, however subsequent sensitivity and moderation analyses failed to reduce the level of heterogeneity. This therefore limits the generalisability of the findings. The consistently high level of heterogeneity may reflect the difficulty in conceptualising and defining a construct such as dissociation in the context of PTSD. Even when only samples adhering to the strict DSM-5 criteria for PTSD-DS were pooled, a high degree of heterogeneity remained.

Clinical Implications and Suggestions for Future Research

This meta-analysis suggests that PTSD-DS is common following trauma exposure, and therefore should be routinely assessed for and formulated. Moreover, the method for determining PTSD-DS was found to have important implications for the estimated prevalence, where samples using diagnostic and clinical cut-off methods reported a higher prevalence than those using LCA and LPA. Future research should also aim to standardise the methodology used to identify and determine PTSD-DS in order to make more valid comparisons between studies.

Additionally, PTSD-DS was found to be more common in children than adults. Clinicians supporting individuals with PTSD should be aware that dissociation is a prevalent and important feature of the overall presentation of PTSD; this may be especially true for children, though this finding was based on only five samples. When the DSM-5 criteria were published it was believed that PTSD-DS cases formed a minority of those with PTSD, however the finding that nearly half of PTSD cases meet the criteria for PTSD-DS suggests that it may be less of a subtype and that dissociation forms a central component to PTSD symptomology. This should be a consideration for how dissociation is specified in future versions of the DSM. Perhaps the conceptualisation of Complex PTSD as defined by the 11th revision of the International Classification of Diseases (World Health Organisation, 2019), where dissociation is stipulated as one of several symptoms seen to be indicative of a more complex form of PTSD, is a more appropriate fit. There is evidence for instance that individuals with Complex PTSD have elevated levels of dissociation (Hyland, Shevlin, Fyvie, Cloitre, & Karatzias, 2019).

Despite the DSM-5 criteria stipulating depersonalisation and derealisation as symptoms required for PTSD-DS, findings of this review suggested that when a wider view of dissociation (i.e., drawing on a broader range of dissociation symptoms) is included in the

criteria, PTSD-DS prevalence does not change significantly. No conclusions can be drawn as to whether it would be more or less appropriate for a narrower (i.e., solely based on depersonalisation and or derealisation) or a broader definition of dissociation, in the context of this subtype, to be used in future versions of diagnostic criteria. However, it does not seem to matter how dissociation is defined when determining the prevalence of PTSD-DS, which raises questions firstly about the strict nature of the DSM criteria when defining this subtype (Ross, 2021), and secondly about the existence of this subtype full stop. Further research is required to establish whether PTSD-DS could be indicative of a distinct form of PTSD that has its own clinical characteristics, and therefore break down the heterogeneity common to populations with the subtype. This would help inform exactly how dissociation should be integrated into future diagnostic criteria of PTSD. Perhaps as Ross (2021) suggests, future diagnostic criteria could stipulate the requirement for the presence of one or more of: depersonalisation, derealisation, dissociative amnesia, and dissociative flashbacks. Non-dissociative PTSD may then form the subtype based on a minority of cases, and dissociative PTSD may form the majority of diagnosed cases.

Limitations

There are several limitations that should be considered for this review. Firstly, whilst many more studies were reviewed in comparison to the most recent systematic review (Hansen et al., 2017), there was still a considerable degree of heterogeneity between samples, reducing the generalisability of the findings. This raises questions around the validity of the underlying diagnostic subtype. Secondly, most studies were conducted in high income countries, and all studies were exclusively written in English, therefore indicating that the results are likely not globally generalisable. Thirdly, some moderator analyses lacked power and further planned moderator analyses were not possible due to a lack of identified studies. Understanding the influence of, for instance, sex, time between index trauma and PTSD

assessment, single- versus multi-event traumas, and individual versus collective trauma could lead to important and interesting findings. Finally, several studies chose to assess PTSD-DS with regard to the most recent trauma that the participant was exposed to, and it is unclear whether other traumas may have taken place, and what impact these may have on the prevalence of PTSD-DS.

Conclusion

This study is the first to meta-analyse data on the prevalence of PTSD-DS. The estimated prevalence of PTSD-DS, with respect to participants diagnosed with PTSD, was 38.1% (95% CI 31.5 – 45.0%) for all samples, 45.5% (95% CI 37.7 – 53.4%) for all diagnosis-based and clinical cut-off samples, 22.8% (95% CI 14.8 – 32.0%) for all LCA and LPA samples, and 48.1% (95% CI 35.0 – 61.3%) for diagnosis-based and clinical cut-off samples which assessed PTSD and PTSD-DS strictly according to the DSM-5 criteria. The prevalence of PTSD-DS was significantly higher for children compared to adults. Factors such as the DSM criteria used for the assessment of both PTSD and dissociation, whether the dissociation assessment was self-report or interview, and participant or trauma characteristics, did not significantly affect the estimated prevalence of PTSD-DS. However, all results were characterised by very high levels of heterogeneity. Further research is required to investigate this construct, and to determine how it should be best conceptualised in future editions of diagnostic criteria.

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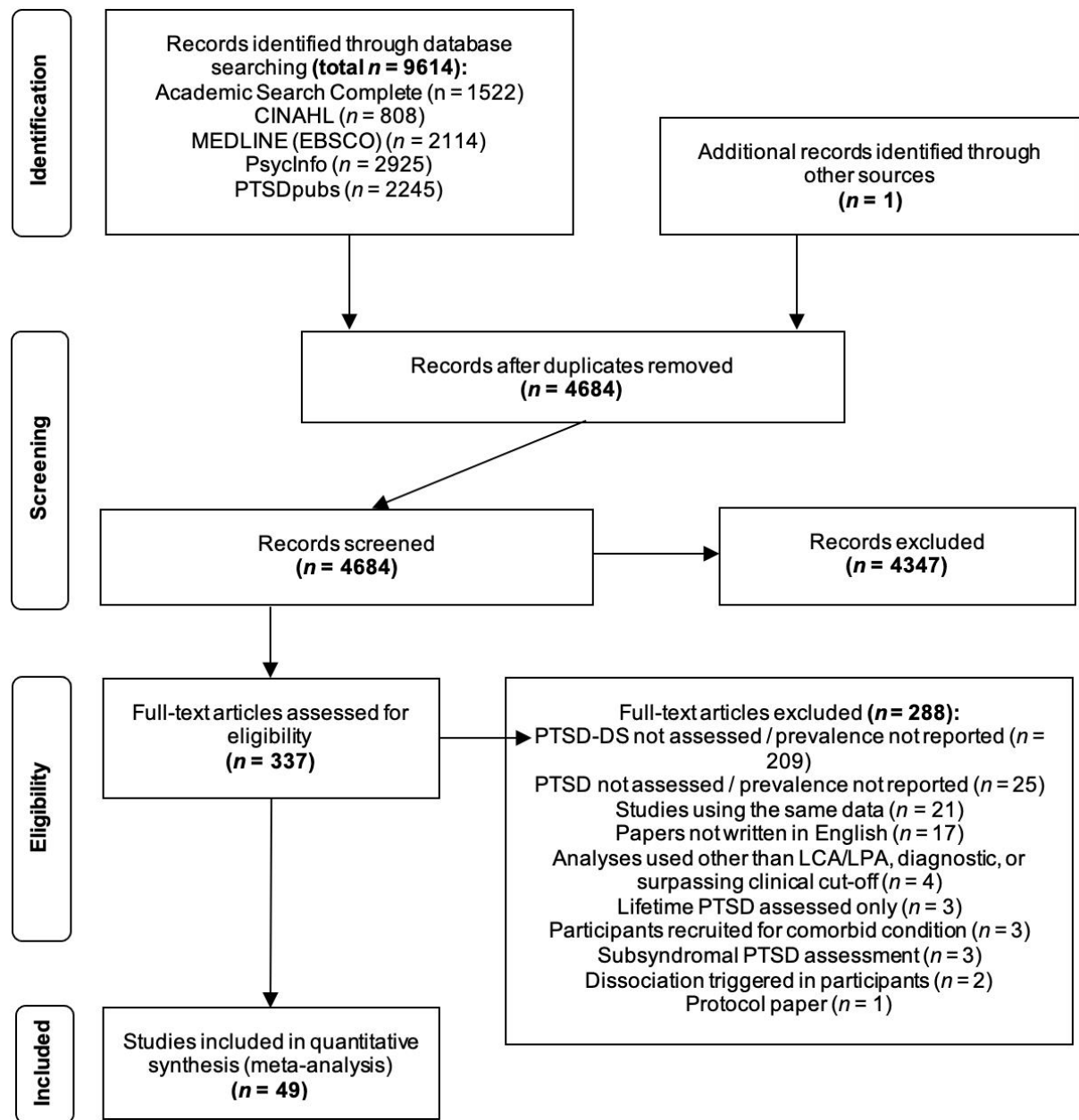


Figure 1

PRISMA diagram showing the process of study identification, screening, and inclusion (n = number of studies)

Table 1*Included sample characteristics*

Sample No.	Sample	Location	Population/trauma type	Proportion female	Age			Method of PTSD assessment	PTSD measure; DSM	PTSD-DS measure; DSM-5/other criteria	N		
					Range	Mean (SD)	Age group				Total	PTSD	PTSD-DS
1	Abu-Rus, Thompson, Naish, Brown, and Dalenberg (2020)	USA	General population (T, P)	46%	NR	37.9 (10.3)	NR	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	345	40	16
2	Acar, Ögülmüş, and Boysan, (2019)	Turkey	Prisoners	3%	18-75	34.5 (9.9)	Adult	Diagnosis	PCL [†] ; DSM-5	DES [†] ; other	399	237	115
3	Armour, Elklit, Lauterbach, and Elhai (2014)	Denmark	Sexual assault and rape (T)	100%	NR	22.4 (9.4)	Both	LPA	HTQ [†] ; DSM-IV	TSC [†] ; other	313	226	41
4	Armour, Karstoft, and Richardson (2014)	Canada	Military veterans (T)	6%	24-93	54.0 (19.0)	Adult	LPA	CAPS; DSM-IV	CAPS; other	432	286	59
5	Blevins, Weathers, and Witte (2014)	USA	Trauma-exposed college students	67%	18-32	20.2 (1.6)	Adult	LCA	PCL-S [†] ; DSM-IV	MDI [†] ; DSM-5	541	206	65
6	Boysan et al. (2017)	Turkey	Psychiatric patients (T, P)	44%	NR	29.0 (9.0)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	90	30	24
7	Briere, Scott, and Weathers (2005)	USA	Trauma-exposed community	48%	NR	45.2 (16.7)	Adult	Diagnosis	DAPS [†] ; DSM-IV	DAPS [†] ; other	372	23	13
8	Burton, Feeny, Connell, and Zoellner (2018)	USA	Chronic PTSD (P)	76%	NR	37.4 (11.3)	Adult	LTA (expanded version of LPA)	PSS-I; DSM-IV	DES-D [†] ; DSM-5	200	129	24
9	Caroppo, Lanzotti, and Janiri (2021)	Italy	Asylum seekers (T)	48%	18-59	25.5 (5.6)	Adult	Diagnosis	SCID-I; DSM-IV	SCID-I; other	180	95	74
10	Choi et al. (2019)	USA	Trauma-exposed adolescents (T)	61%	12-16	14.5 (1.5)	Child	Diagnosis	UCLA PTSD-RI [†] ; DSM-IV	TSCC-A [†] ; DSM-5	3081	734	394
11	Choi et al. (2017)	USA	Trauma-exposed adolescents (T)	61%	12-16	14.5 (1.5)	Child	LCA	UCLA PTSD-RI [†] ; DSM-IV	TSCC-A [†] ; DSM-5	3081	1279	444
12	Cloitre et al. (2012)	USA	Childhood sexual and/or physical abuse (P)	100%	18-65	36.4 (9.4) [§]	Adult	Diagnosis	CAPS; DSM-IV	TSI [†] ; other	104	104	28
13	Criswell, Sherman, and Krippner (2018)	USA	Psychiatric patients (T, P)	73%	20-65	44.0 (NR)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	30	30	13
14	Daniels et al. (2016)	Germany	Trauma-exposed community (P)	61%	23-58	38.0 (11.8) [§]	Adult	Diagnosis & clinical cut-off	CAPS; DSM-IV	CAPS; DSM-5	59	59	15
15	Dorahy et al. (2017)	Northern Ireland	Psychiatric patients (T, P)	32%	19-65 [‡]	40.4 (12.4)	Adult	Diagnosis	Clinical diagnosis; NR	DES [†] ; other	210	65	27
16	Durham, Byllesby, Elhai, and Wang (2020)	USA & Canada	Trauma-exposed community	63%	18-74	36.0 (12.7)	Adult	LPA	PCL [†] ; DSM-5	DES-II [†] ; DSM-5	360	204	51
17	Eidhof et al. (2019)	Netherlands	Trauma-exposed community (T, P)	33%	19-83	48.8 (12.1)	Adult	Diagnosis	CAPS; DSM-5	CAPS [†] ; DSM-5	320	131	31

18	Frewen et al (2015)	Canada	Probable diagnosis of PTSD (T)	71%	NR	33.1 (10.8)	Adult	LPA	PCL [†] ; DSM-5	Dissociation-TRASC item list [†] ; DSM-5	557	311	183
19	Frewen, Zhu, and Lanius (2019)	Canada	Community	52%	NR	36.5 (12.6)	Adult	Diagnosis	PCL [†] ; DSM-5	Dissociation-TRASC item list [†] ; DSM-5	418	98	41
20	Guetta et al. (2019)	USA	Military veterans (P)	16%	21-75	53.8 (11.4)	Adult	LPA	PCL, Trauma Assessment from the NSES; DSM-5	CAPS [‡] ; DSM-5	209	209	31
21	Hansen, Hyland, and Armour (2016)	Denmark	Bank employees following robbery	62%	20-65	42.1 (12.5)	Adult	LCA	HTQ [†] ; DSM-IV	TSC [†] ; DSM-5	371	67	0
22	Hansen et al. (2019)	Denmark	Whiplash injury	62%	18-89	37.5 (13.9)	Adult	Diagnosis	HTQ [†] , TSC [†] ; DSM-5	TSC [†] ; DSM-5	234	21	7
23	Hansen et al. (2019)	Denmark	Whiplash injury	62%	18-89	37.5 (13.9)	Adult	LCA	HTQ [†] , TSC [†] ; DSM-5	TSC [†] ; DSM-5	234	27	0
24	Hansen, Müllerová et al. (2016)	Denmark	Whiplash injury (P)	78%	NR	43.6 (10.4)	Adult	LCA	HTQ [†] , TSC [†] ; DSM-5	TSC [†] ; DSM-5	476	476	178
25	Hansen, Müllerová, et al. (2016)	Denmark	Incest during childhood (T, P)	88%	NR	35.9 (11.0)	Adult	LCA	HTQ [†] , TSC [†] ; DSM-5	TSC [†] ; DSM-5	311	311	139
26	Harricharan et al. (2020)	Canada	Trauma-exposed community (P)	63%	18-60 [*]	39.6 (12.5) [§]	Adult	Diagnosis & clinical cut-off	CAPS; DSM-IV & 5	CAPS; DSM-5	184	133	49
27	Hill et al. (2020)	USA	Trauma-exposed women (T)	100%	18-62	34.1 (13.2)	Adult	Clinical cut-off	PCL [†] ; DSM-5	DSPS [†] ; DSM-5	104	88	73
28	Kenny et al. (2020)	USA	Commercial sexual exploitation (T)	100%	12-18	16.6 (1.2) [‡]	Child	Diagnosis	UCLA PTSD-RI [†] ; DSM-5	UCLA PTSD-RI [†] ; DSM-5	56	15	11
29	Kenny et al. (2020)	USA	At risk of commercial sexual exploitation (T)	100%	12-18	15.3 (1.6) [‡]	Child	Diagnosis	UCLA PTSD-RI [†] ; DSM-5	UCLA PTSD-RI [†] ; DSM-5	40	3	3
30	Kim et al. (2019)	South Korea	Psychiatric patients (T, P)	64%	16-70	38.7 (12.7)	Both	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	249	249	82
31	Lebois et al. (2021)	USA	Interpersonal childhood maltreatment (T, P)	100%	18-61	34.4 (12.2)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	65	65	47
32	Li, Hasset, and Seng (2019)	USA	Pregnant women	100%	NR	NR	NR	Diagnosis	National Women's Study PTSD Module; DSM-IV	DES-T [†] ; other	22	10	4
33	Mulder, Beautrais, Joyce, and Fergusson (1998)	New Zealand	Community	NR	NR	NR	Adult	Diagnosis	SCID; DSM-III	DES [†] ; other	1028	9	3
34	Müllerová et al. (2016)	USA & Canada	Trauma-exposed community	56%	NR	35.2 (11.9)	NR	LPA	PCL [†] ; DSM-5	DSS [†] ; other	309	215	83
35	Naish et al. (2021)	USA	Trauma-exposed community	45%	18-65	40.5 (11.8)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	100	63	31

36	Nejad et al. (2007)	Iran	Military veterans (P)	0%	NR	41.5 (5.1)	Adult	Diagnosis	Clinical diagnosis; DSM-IV	DES [†] ; other	260	130	42
37	Özdemir, Celik, and Oznur (2015)	Turkey	Serving soldiers (P)	0%	NR	30.3 (5.6)	Adult	Diagnosis	SCID-I; DSM-IV	DES [†] ; other	184	84	59
38	Powers et al. (2017)	USA	Trauma-exposed women	100%	18-65 [‡]	39.4 (11.6)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	190	72	2
39	Putnam et al. (1996)	USA & Canada	Psychiatric patients - (T, P)	60%	NR	39.0 (NR)	Adult	Diagnosis	Clinical diagnosis; DSM-III	DES [†] ; other	1566	116	54
40	Richard-Malenfant, Douglass, Higginson, Ray, and Robillard (2019)	Canada	Military veterans (P)	36%	NR	49.3 (9.3)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	14	14	6
41	Ross, Armour, Kerig, Kidwell, and Kilshaw (2020)	USA	Trauma-exposed youth in detention centres	25%	12-19	16.0 (1.3)	Child	Diagnosis	UCLA PTSD-RI [†] ; DSM-5	UCLA PTSD-RI [†] ; DSM-5	448	197	119
42	Ross et al. (2018)	Slovakia	Trauma-exposed university students	83%	NR	22.7 (5.1)	Adult	LPA	PCL [†] ; DSM-5	DSS [†] ; other	689	308	24
43	Sierk, Manthey, Brakemeier, Walter, and Daniels (2021)	Germany	Childhood interpersonal abuse (P)	100%	NR	40.0 (9.8)	Adult	Diagnosis	CAPS; DSM-IV	DES [†] , CDS-30 [†] , CDS-state [†] , CAPS, SCID-D; other	42	42	23
44	Stein et al. (2013)	Global	Community	NR	NR	NR	Adult	Diagnosis	WHO CIDI; DSM-IV	WHO CIDI; DSM-5	25018	747	108
45	Steuwe et al. (2012)	Canada	Trauma-exposed community (T, P)	90%	NR	37.9 (9.4)	NR	Diagnosis	CAPS; DSM-IV	CAPS; DSM-5	134	134	47
46	Swart et al. (2020)	Netherlands	Psychiatric patients (T)	77%	18-68	34.2 (11.9) [§]	Adult	Diagnosis	CAPS; DSM-IV	DES [†] ; DSM-5	150	84	18
47	Tsai et al. (2015)	USA	Military veterans	NR	20-94 [‡]	60.8 (15.2) [§]	Adult	Diagnosis	PCL [†] ; DSM-5 SCID & DIS	CAPS [†] ; DSM-5	1484	64	12
48	van der Kolk et al. (1996)	USA	Psychiatric patients (T)	67%	15+	37.1 (15.0)	Both	Diagnosis	PTSD modules; DSM-III	SIDES; other	395	182	149
49	Verbeck et al. (2015)	USA	Psychiatric patients (T)	49%	18-69	44.0 (10.9)	Adult	Diagnosis	CAPS; DSM-IV	TSI-2 [†] , DES-R [†] ; other	100	47	29
50	Wolf, Lunney et al. (2012)	USA	Military veterans (P)	0%	44-74	50.6 (3.6)	Adult	LPA	CAPS; DSM-IV	CAPS; other	360	360	56
51	Wolf, Lunney et al. (2012)	USA	Military veterans (P)	100%	22-78	44.8 (9.4)	Adult	LPA	CAPS; DSM-IV	TSI [†] ; DSM-5	284	284	85
52	Wolf, Miller et al. (2012)	USA	Military veterans & their partners	36%	21-75 [‡]	51.5 (11.2) [‡]	Adult	LPA	CAPS; DSM-IV	CAPS; other	492	239	30
53	Zoet et al. (2018)	Netherlands	Psychiatric patients (T)	70%	19-63 [‡]	38.2 (10.9) [§]	Adult	Clinical cut-off	CAPS; DSM-IV	CAPS [†] ; DSM-5	168	168	38

Note. SD = standard deviation, T = treatment-seeking inclusion criteria; P = diagnosis of PTSD inclusion criteria; NR = Not Reported; CAPS = Clinician Administered Post-traumatic Stress Disorder Scale; PCL = Post-traumatic Stress Disorder Checklist; DES = Dissociative Experiences Scale; LPA = latent profile analysis; HTQ = Harvard Trauma Questionnaire; TSC = Trauma Symptom Checklist; LCA = latent class analysis; PCL-S = Post-traumatic Stress Disorder Checklist Specific; MDI = Multiscale Dissociation Inventory; DAPS = Detailed Assessment of Posttraumatic Stress; DES-D = depersonalization/derealisation subscale of the DES; PSS = Post-traumatic Stress Disorder Symptom Scale; PSS-I = PTSD Symptom Scale-Interview, SCID-I = Structured Clinical Interview for the DSM-IV Axis I Disorders; UCLA PTSD-RI = University of California at Los Angeles Posttraumatic Stress Disorder Reaction Index; TSCC-A = Trauma Symptom Checklist for Children-Alternate Version; TSI Trauma Symptom Inventory; TRASC = trauma-related altered states of consciousness; NSES = National Stressful Events Survey; DES-T = 8-item taxon version of the Dissociative Experiences Scale; SCID = Structured Clinical Interview for DSM; DSPTS = Dissociative Subtype of PTSD Scale; DSS = Dissociative Symptoms Scale; CDS = Cambridge Depersonalization Scale; SCID-D = Structured Clinical Interview for DSM-IV Dissociative Disorders; WHO CIDI = World Health Organisation Composite International Diagnostic Interview; DIS = Diagnostic Interview Schedule; SIDES = Structured Interview for Disorders of Extreme Stress; DES-R = Dissociative Experiences Scale – Revised

† Measure completed via self-report

‡ Information acquired via correspondence with study author(s)

§ Mean and standard deviation values combined (Altman, Machin, Bryant, & Gardner, 2013; Higgins et al., 2012)

¶ Multiple measures used, however CAPS chosen as the gold standard (Weathers et al., 2004)

Table 2*Pooled prevalence of PTSD-DS as a proportion of PTSD for all samples (k = 51)*

Meta-analysis subgroup	<i>k</i>	<i>n</i>	Pooled Prevalence (%)	95% CI	Q test	<i>I</i> ²
All samples [†]	51	8214	38.1	(31.5, 45.0)	1602.0*	97.4
Method of PTSD-DS Assessment ($\beta = -0.2418$ [95% CI = -0.3780, -0.1056], $p = 0.0005$)						
Diagnosis-based/clinical cut-off	36	4383	45.5	(37.7, 53.4)	923.6*	96.0
LCA/LPA [†]	15	3831	22.8	(14.8, 32.0)	482.5*	97.6
PTSD DSM criteria used ^{†‡} ($\beta = -0.0871$ [95% CI = -0.2328, 0.0586], $p = 0.24$)						
DSM-5	24	3451	42.5	(32.4, 53.0)	624.6*	97.3
DSM-III or DSM-IV	25	4565	34.1	(24.9, 43.9)	936.0*	97.8
Dissociation criteria [†] ($\beta = 0.0342$ [95% CI = -0.1113, 0.1796], $p = 0.65$)						
DSM-5 (Dereal / Depers)	32	5436	36.9	(28.5, 45.8)	895.2*	97.6
Broader dissociation	19	2778	40.2	(29.5, 51.4)	698.3*	97.1
Dissociation measure completion ^{†§} ($\beta = 0.0281$ [95% CI = -0.1189, 0.18], $p = 0.7080$)						
Self-report	31	4997	38.8	(30.6, 47.3)	778.8*	97.2
Interview	19	3175	36.2	(24.8, 48.5)	690.4*	97.9
Age group ^{†¶} ($\beta = 0.3587$ [95% CI = 0.0814, 0.6360], $p = 0.01$)						
Child	4	949	62.9	(39.6, 83.3)	11.4*	82.0
Adult	40	6209	35.0	(27.8, 42.6)	1121.1*	97.3
Occupation [†] ($\beta = -0.1439$ [95% CI = -0.3227, 0.0350], $p = 0.11$)						
Military	9	1670	26.9	(16.2, 39.1)	138.1*	96.3
Civilian	42	6544	40.7	(33.1, 48.5)	1325.7*	97.4
Trauma type [†] ($\beta = 0.1011$ [95% CI = -0.1163, 0.3185], $p = 0.36$)						
Interpersonal	6	763	46.8	(28.3, 65.7)	101.9*	95.9
Other	45	7451	37.0	(29.9, 44.3)	1494.9*	97.5

Note. *k* = number of samples; *n* = number of participants; CI = confidence interval; LCA = latent class

analysis, LPA = latent profile analysis; Dereal = derealisation; Depers = depersonalisation

* $p < 0.0001$, where the degrees of freedom (*df*) = *k* - 1

† Samples 11 and 23 removed to avoid duplication of population samples

‡ Sample 15 removed as no PTSD DSM criteria reported, sample 26 removed as used both DSM-IV and DSM-5 when assessing for PTSD

§ Sample 43 removed as a mix of self-report and interview measures were used

¶ Several samples were removed due to populations formed of both children and adults, or age group not reported

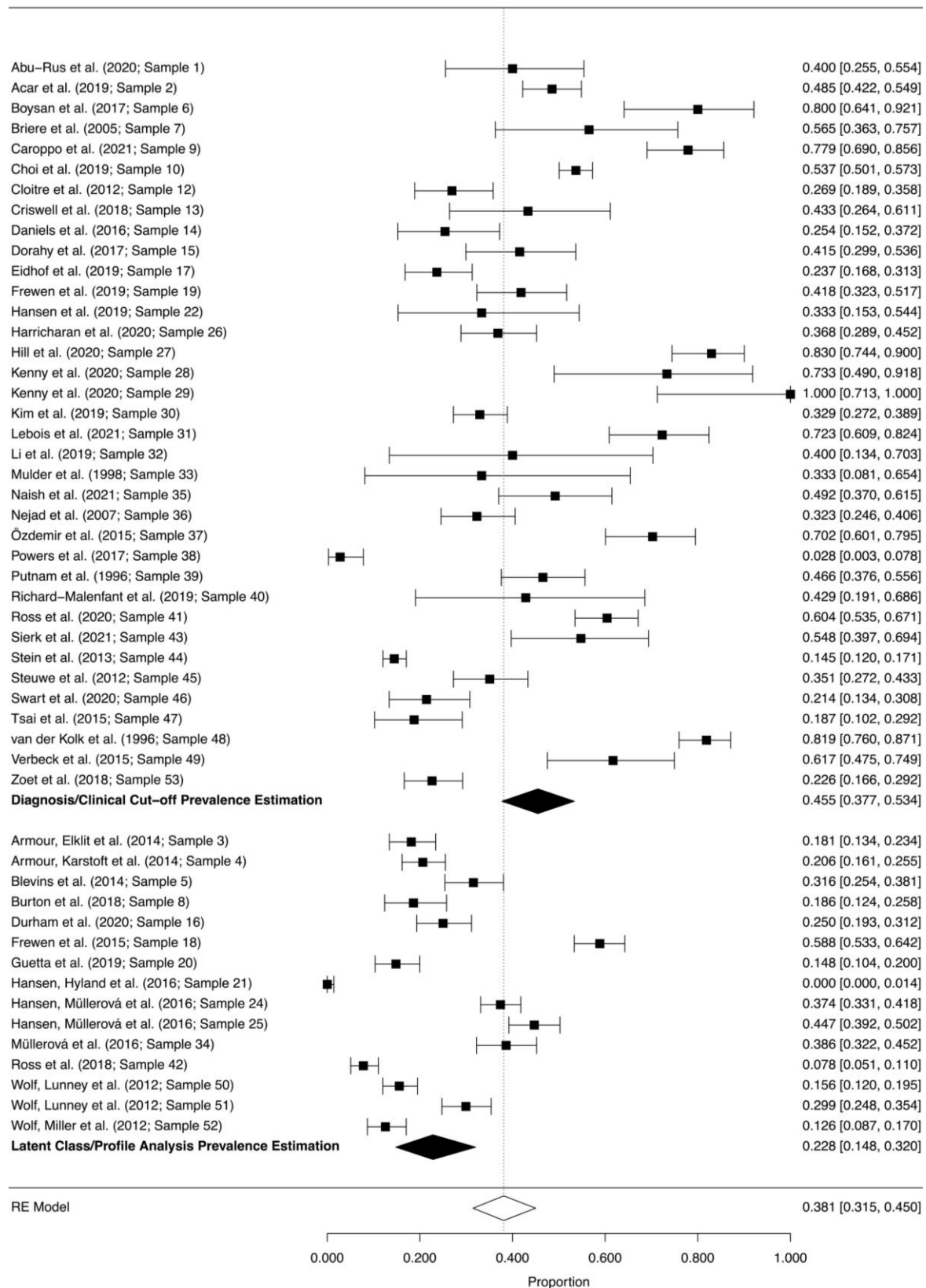


Figure 2

Forest plot of PTSD-DS prevalence estimates grouped by PTSD-DS assessment method (samples 11 and 23 removed to avoid duplication of population samples).

Table 3

Pooled prevalence of PTSD-DS as a proportion of PTSD for all diagnostic and clinical cut-off samples (i.e., excluding LCA and LPA samples; k = 36)

Meta-analysis subgroup	k	n	Pooled Prevalence (%)	95% CI	Q test	I ²
PTSD DSM criteria used [†] ($\beta = -0.0363$ [95% CI = -0.2065, 0.1338], $p = 0.68$)						
DSM-5	17	1417	48.1	(35.0, 61.3)	288.3*	95.7
DSM-III or DSM-IV	17	2768	44.2	(33.6, 55.1)	623.7*	96.5
Dissociation criteria ($\beta = 0.1135$ [95% CI = -0.0471, 0.2740], $p = 0.17$)						
DSM-5 (Dereal / Depers)	23	3239	41.7	(31.5, 52.2)	622.8*	96.9
Broader dissociation	13	1144	52.9	(42.5, 63.3)	173.2*	91.4
Dissociation measure completion [‡] ($\beta = 0.0479$ [95% CI = -0.1171, 0.2130], $p = 0.57$)						
Self-report	20	2260	47.0	(37.8, 56.3)	233.1*	93.9
Interview	15	2081	42.7	(29.2, 56.8)	576.9*	97.3
Age group [§] ($\beta = 0.2794$ [95% CI = 0.0115, 0.5474], $p = 0.04$)						
Child	4	949	62.9	(50.2, 74.7)	11.4**	82.0
Adult	27	2819	42.1	(33.4, 51.2)	616.6*	95.4
Occupation ($\beta = -0.0574$ [95% CI = -0.3115, 0.1968], $p = 0.66$)						
Military	4	292	40.5	(19.1, 63.9)	49.8*	93.2
Civilian	32	4091	46.1	(37.8, 54.6)	873.7*	96.3
Trauma type ($\beta = 0.1184$ [95% CI = -0.1345, 0.3714], $p = 0.36$)						
Interpersonal only	4	226	55.9	(33.4, 77.2)	41.4*	90.5
Other	32	4157	44.2	(35.6, 52.6)	876.4*	96.3

Note. k = number of samples; n = number of participants; CI = confidence interval; Dereal =

derealisation; Depers = depersonalisation

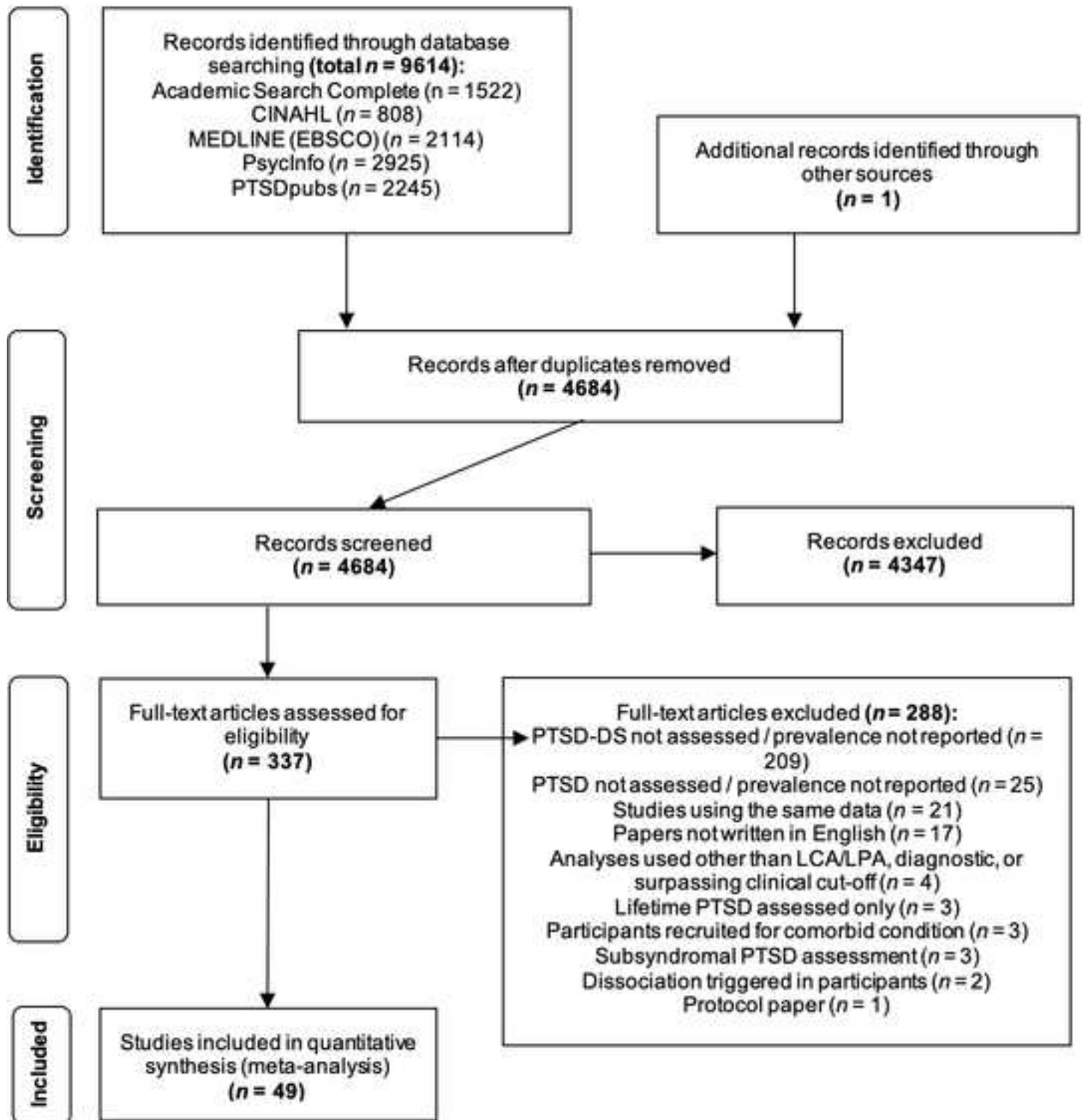
* $p < 0.0001$, where the degrees of freedom (df) = $k - 1$

** $p < 0.01$, where the degrees of freedom (df) = $k - 1$

† Sample 15 removed as no PTSD DSM criteria reported, sample 26 removed as used both DSM-IV and DSM-5 when assessing for PTSD

‡ Sample 43 removed as a mix of self-report and interview measures were used

§ Several samples were removed due to populations formed of both children and adults, or age group not reported



SUPPLEMENTARY INFORMATION

1

Supplementary Information

Supplementary Table 1

Quality Assessment Checklist for Prevalence Meta-Analysis

1	Was the study population and index trauma clearly specified and defined?	
	<i>Descriptive statistics were reported on participant demographics (including age range and mean, gender, ethnicity) and frequency of trauma type/nature within the participant pool reported</i>	2
	<i>Some description statistics provided about the sample but some missing information (e.g. authors did not report frequency of trauma type/nature or provide enough information about demographic variables).</i>	1
	<i>No clear description of sample demographics or index trauma characteristics</i>	0
2	Was the participation rate of eligible persons at least 50%?	
	<i>More than 50% of eligible and approached participants took part</i>	2
	<i>Less than 50% of those approached took part, but there was no significant difference in non-response characteristics (such as age, gender) between those who participated and those who did not</i>	1
	<i>Less than 50% of those approached took part, and differences between those who took part and those who did not were not reported or highlighted significant differences. Or, response was not reported</i>	0
3	Was follow up time for PTSD assessment appropriate and meaningful?	
	<i>An appropriate time frame (>4 weeks) since trauma was reported</i>	2
	<i>No information given regarding time frame since trauma. Or, assessment <4 weeks since trauma</i>	0
4	Were objective, standard criteria used for the assessment of Post-Traumatic Stress Disorder?	
	<i>Diagnostic interview or self-report questionnaire shown to demonstrate good levels of validity and reliability in the assessment of PTSD adhering to DSM criteria for PTSD i.e. cluster-based algorithm</i>	2
	<i>Diagnostic interview or self-report questionnaire shown to demonstrate good levels of validity and reliability in the assessment of PTSD adhering to DSM criteria for PTSD using a cut-off score or grouping analysis such as LPA or LCA</i>	1
	<i>Diagnostic interview or self-report without utilising DSM criteria (e.g. not conforming to cluster-based algorithm or cut-off score or grouping analysis). Or poor validity and reliability.</i>	0
5	Were objective, standard criteria used for the assessment of the Dissociative Subtype of Post-Traumatic Stress Disorder?	
	<i>Diagnostic interview or self-report questionnaire shown to demonstrate good levels of validity and reliability, adhering to DSM-5 criteria for PTSD-DS i.e. based on depersonalisation and derealisation only</i>	2
	<i>Diagnostic interview or self-report questionnaire shown to demonstrate good levels of validity and reliability, however not adhering to DSM-5 criteria for PTSD-DS i.e. based on other domains of dissociation outside of just depersonalisation and derealisation</i>	1

<i>Diagnostic interview or self-report questionnaire shown to demonstrate good levels of validity, however domains of dissociation assessed not reported. Or poor validity and reliability</i>	0
--	---

Note. Where 2 = well addressed, 1 = partially addressed, 0 = poorly addressed/not addressed/not reported

This tool was developed by Mr. William White for a meta-analysis undertaken in partial fulfilment of a Doctorate in Clinical Psychology. The development of this tool was based on the Assessment Tool for Observational Cohort and Cross-Sectional Studies (National Heart Lung and Blood Institute, 2014), combining with modified questions from other prevalence and risk factor studies that would be appropriate for use in this review (Hoy et al., 2012; Munn et al., 2014).

Supplementary Table 2

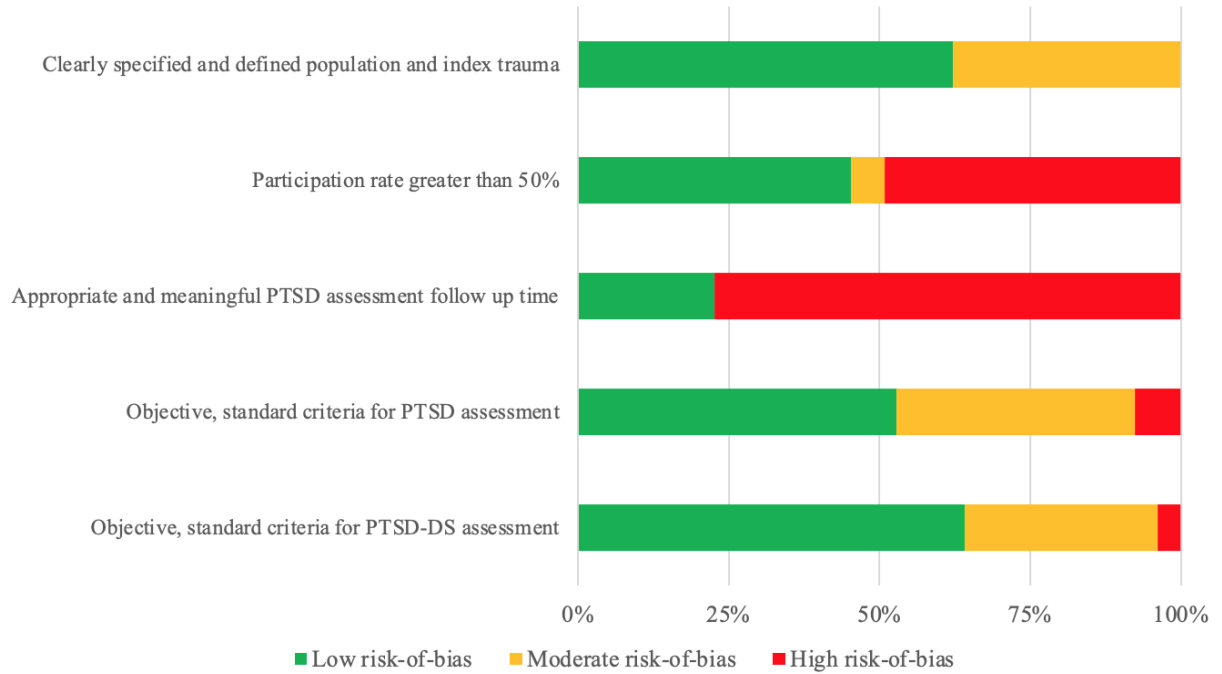
Sample risk-of-bias scores by individual item and total

Sample No.	Author	Item 1	Item 2	Item 3	Item 4	Item 5	Total	Quality
1	Abu-Rus et al. (2020)	1	2	0	2	2	7	High
2	Acar et al. (2019)	1	0	0	2	1	4	Low
3	Armour, Elklit et al. (2014)	2	2	2	1	1	8	High
4	Armour, Karstoft et al. (2014)	2	0	0	1	1	4	Low
5	Blevins et al. (2014)	2	0	0	1	2	5	Medium
6	Boysan et al. (2017)	2	0	2	2	2	8	High
7	Briere et al. (2005)	1	2	0	2	1	6	Medium
8	Burton et al. (2018)	2	0	0	1	2	5	Medium
9	Caroppo et al. (2021)	2	0	0	2	0	4	Low
10	Choi et al. (2019)	2	2	0	2	2	8	High
11	Choi et al. (2017)	2	2	0	1	2	7	High
12	Cloitre et al. (2012)	2	0	2	0	1	5	Medium
13	Criswell et al. (2018)	2	0	2	2	2	8	High
14	Daniels et al. (2016)	1	0	0	1	2	4	Low
15	Dorahy et al. (2017)	1	1	0	0	1	3	Low
16	Durham et al. (2020)	2	2	0	1	2	7	High
17	Eidhof et al. (2019)	2	0	0	2	2	6	Medium
18	Frewen et al. (2015)	1	2	0	1	2	6	Medium
19	Frewen et al. (2019)	1	0	0	2	2	5	Medium
20	Guetta et al. (2019)	1	2	0	1	2	6	Medium
21	Hansen, Hyland et al. (2016)	1	2	2	1	2	8	High
22	Hansen et al. (2019)	2	1	2	2	2	9	High
23	Hansen et al. (2019)	2	1	2	1	2	8	High
24	Hansen, Müllerová et al. (2016)	2	2	0	1	2	7	High
25	Hansen, Müllerová et al. (2016)	2	2	0	1	2	7	High
26	Harricharan et al. (2020)	1	0	0	1	2	4	Low
27	Hill et al. (2020)	1	0	0	1	2	4	Low
28	Kenny et al. (2020)	2	2	0	2	2	8	High
29	Kenny et al. (2020)	2	2	0	2	2	8	High
30	Kim et al. (2019)	2	2	0	2	2	8	High
31	Lebois et al. (2021)	1	2	0	2	2	7	High
32	Li et al. (2019)	2	2	0	2	1	7	High
33	Mulder et al. (1998)	2	2	0	2	1	7	High
34	Müllerová et al. (2016)	2	2	0	1	1	6	Medium
35	Naish et al. (2021)	2	0	2	2	2	8	High
36	Nejad et al. (2007)	2	0	0	0	1	3	Low
37	Özdemir et al. (2015)	2	0	0	2	1	5	Medium
38	Powers et al. (2017)	1	2	2	2	2	9	High
39	Putnam et al. (1996)	1	0	0	0	1	2	Low
40	Richard-Malenfant et al. (2019)	1	0	0	2	2	5	Medium
41	Ross et al. (2020)	2	2	0	2	2	8	High
42	Ross et al. (2018)	2	0	0	1	1	4	Low
43	Sierk et al. (2021)	2	2	2	2	1	9	High
44	Stein et al. (2013)	1	0	2	2	2	7	High
45	Steuwe et al. (2012)	1	0	0	2	2	5	Medium
46	Swart et al. (2020)	2	2	0	2	2	8	High
47	Tsai et al. (2015)	2	0	0	2	2	6	Medium
48	van der Kolk et al. (1996)	1	0	0	2	0	3	Low
49	Verbeck et al. (2015)	2	0	0	2	1	5	Medium
50	Wolf, Lunney et al. (2012)	1	0	0	1	1	3	Low
51	Wolf, Lunney et al. (2012)	1	0	2	1	2	6	Medium
52	Wolf, Miller et al. (2012)	2	2	0	1	1	6	Medium
53	Zoet et al. (2018)	2	2	0	1	2	7	High

Note. 0-4 high risk/low quality, 5-6 moderate risk/quality, 7-10 low risk/high quality

Supplementary Figure 1

Proportion of samples rated as a low, moderate or high risk-of-bias for each quality assessment item



Supplementary Table 3

Pooled prevalence of PTSD-DS as a proportion of PTSD for diagnostic/clinical cut-off samples utilising DSM-5 criteria for dissociation (i.e., excluding LCA and LPA samples and those using broader criteria for dissociation; k = 23)

Meta-analysis subgroup	k	n	Pooled Prevalence (%)	95% CI	Q test	I ²
PTSD DSM criteria used [‡] ($\beta = -0.2041$ [95% CI = -0.4406, 0.0324], $p = 0.09$)						
DSM-5	16	1180	48.2	(34.2, 62.3)	285.9*	95.5
DSM-III or DSM-IV	6	1926	28.3	(17.6, 40.3)	289.7*	96.1
Dissociation measure completion ($\beta = 0.1271$ [95% CI = -0.0882, 0.3423], $p = 0.25$)						
Self-report	10	1435	49.4	(32.0, 66.9)	174.5*	97.1
Interview	13	1804	36.4	(24.3, 49.4)	244.3*	96.3
Age group [‡] ($\beta = 0.3444$ [95% CI = 0.0410, 0.6477], $p = 0.03$)						
Child	4	949	62.9	(50.2, 74.7)	11.4**	82.0
Adult	16	1867	36.7	(24.7, 49.6)	376.3*	96.4

Note. k = number of samples; n = number of participants; CI = confidence interval

* $p < 0.0001$, where the degrees of freedom (df) = $k - 1$

** $p < 0.01$, where the degrees of freedom (df) = $k - 1$

† Sample 26 removed as used both DSM-IV and DSM-5 when assessing for PTSD

‡ Several samples were removed due to populations formed of both children and adults, or age group not reported

Supplementary Table 4

Pooled prevalence of PTSD-DS as a proportion of PTSD for all LCA/LPA samples (i.e., excluding diagnostic and clinical cut-off samples; k = 17)

Meta-analysis subgroup	k	n	Pooled Prevalence (%)	95% CI	Q test	I²
PTSD DSM criteria used ($\beta = -0.0872$ [95% CI = -0.3022, 0.1278], $p = 0.43$)						
DSM-5	8	1750	25.0	(10.9, 42.7)	328.5*	98.6
DSM-III or DSM-IV	9	2850	18.2	(10.1, 28.1)	196.8*	97.4
Dissociation criteria ($\beta = -0.0648$ [95% CI = -0.2912, 0.1616], $p = 0.57$)						
DSM-5 (Dereal / Depers)	11	3503	23.1	(11.3, 37.5)	311.3*	98.8
Broader dissociation	6	1634	18.0	(10.7, 26.7)	83.7*	94.6
Dissociation measure completion ($\beta = 0.0940$ [95% CI = -0.1589, 0.3468], $p = 0.47$)						
Self-report	13	3506	23.1	(12.6, 35.6)	429.9*	98.6
Interview	4	1094	15.9	(12.7, 19.3)	6.7	55.6
Occupation ($\beta = -0.0532$ [95% CI = -0.2918, 0.1853], $p = 0.66$)						
Military	5	1378	18.4	(12.9, 24.6)	32.0*	87.6
Civilian	12	3759	22.5	(11.3, 36.1)	429.7*	98.8

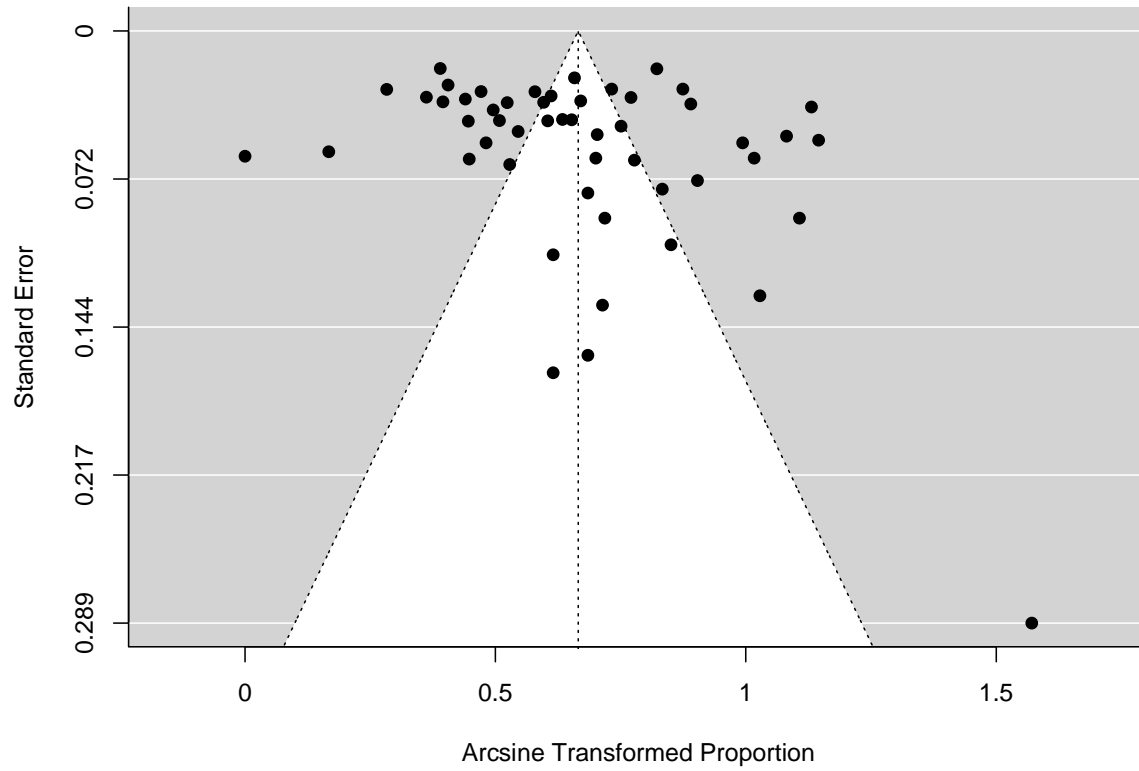
Note. k = number of samples; n = number of participants; CI = confidence interval; Dereal =

derealisation; Depers = depersonalisation

* $p < 0.0001$, where the degrees of freedom (df) = $k - 1$

Supplementary Figure 2

Funnel plot to assessing publication bias





PRISMA 2020 Checklist

Section and Topic	Item #	Checklist item	Location where item is reported
TITLE			
Title	1	Identify the report as a systematic review.	Manuscript title
ABSTRACT			
Abstract	2	See the PRISMA 2020 for Abstracts checklist.	Abstract
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of existing knowledge.	Introduction
Objectives	4	Provide an explicit statement of the objective(s) or question(s) the review addresses.	Introduction
METHODS			
Eligibility criteria	5	Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses.	Inclusion and Exclusion Criteria
Information sources	6	Specify all databases, registers, websites, organisations, reference lists and other sources searched or consulted to identify studies. Specify the date when each source was last searched or consulted.	Search Strategy
Search strategy	7	Present the full search strategies for all databases, registers and websites, including any filters and limits used.	Search Strategy
Selection process	8	Specify the methods used to decide whether a study met the inclusion criteria of the review, including how many reviewers screened each record and each report retrieved, whether they worked independently, and if applicable, details of automation tools used in the process.	Search Strategy & Data Extraction, Coding and Synthesis
Data collection process	9	Specify the methods used to collect data from reports, including how many reviewers collected data from each report, whether they worked independently, any processes for obtaining or confirming data from study investigators, and if applicable, details of automation tools used in the process.	Data Extraction, Coding and Synthesis
Data items	10a	List and define all outcomes for which data were sought. Specify whether all results that were compatible with each outcome domain in each study were sought (e.g. for all measures, time points, analyses), and if not, the methods used to decide which results to collect.	Data Extraction, Coding and Synthesis
	10b	List and define all other variables for which data were sought (e.g. participant and intervention characteristics, funding sources). Describe any assumptions made about any missing or unclear information.	Data Extraction, Coding and Synthesis
Study risk of bias assessment	11	Specify the methods used to assess risk of bias in the included studies, including details of the tool(s) used, how many reviewers assessed each study and whether they worked independently, and if applicable, details of automation tools used in the process.	Quality Assessment and Risk of Bias
Effect measures	12	Specify for each outcome the effect measure(s) (e.g. risk ratio, mean difference) used in the synthesis or presentation of results.	Meta-Analytic Method
Synthesis	13a	Describe the processes used to decide which studies were eligible for each synthesis (e.g. tabulating the study intervention characteristics)	Data



PRISMA 2020 Checklist

Section and Topic	Item #	Checklist item	Location where item is reported
methods		and comparing against the planned groups for each synthesis (item #5)).	Extraction, Coding and Synthesis
	13b	Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data conversions.	Data Extraction, Coding and Synthesis , & Meta-Analytic Method
	13c	Describe any methods used to tabulate or visually display results of individual studies and syntheses.	N/A
	13d	Describe any methods used to synthesize results and provide a rationale for the choice(s). If meta-analysis was performed, describe the model(s), method(s) to identify the presence and extent of statistical heterogeneity, and software package(s) used.	Meta-Analytic Method
	13e	Describe any methods used to explore possible causes of heterogeneity among study results (e.g. subgroup analysis, meta-regression).	Meta-Analytic Method
	13f	Describe any sensitivity analyses conducted to assess robustness of the synthesized results.	Meta-Analytic Method
Reporting bias assessment	14	Describe any methods used to assess risk of bias due to missing results in a synthesis (arising from reporting biases).	Meta-Analytic Method
Certainty assessment	15	Describe any methods used to assess certainty (or confidence) in the body of evidence for an outcome.	Meta-Analytic Method
RESULTS			
Study selection	16a	Describe the results of the search and selection process, from the number of records identified in the search to the number of studies included in the review, ideally using a flow diagram.	Results & Figure 1
	16b	Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded.	Figure 1
Study characteristics	17	Cite each included study and present its characteristics.	Table 1
Risk of bias in studies	18	Present assessments of risk of bias for each included study.	Risk of Bias Assessment & Supplementary information
Results of individual studies	19	For all outcomes, present, for each study: (a) summary statistics for each group (where appropriate) and (b) an effect estimate and its precision (e.g. confidence/credible interval), ideally using structured tables or plots.	Figure 2
Results of syntheses	20a	For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies.	Risk of Bias Assessment, Sensitivity Analysis, & Supplementary information
	20b	Present results of all statistical syntheses conducted. If meta-analysis was done, present for each the summary estimate and its precision (e.g. confidence/credible interval) and measures of statistical heterogeneity. If comparing groups, describe the direction of the effect.	Results, Tables 2-4,



PRISMA 2020 Checklist

Section and Topic	Item #	Checklist item	Location where item is reported
			Figure 2, & Supplementary information
	20c	Present results of all investigations of possible causes of heterogeneity among study results.	Results, Tables 2-4, & Supplementary information
	20d	Present results of all sensitivity analyses conducted to assess the robustness of the synthesized results.	Sensitivity Analysis
Reporting biases	21	Present assessments of risk of bias due to missing results (arising from reporting biases) for each synthesis assessed.	Publication Bias
Certainty of evidence	22	Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed.	Results, Tables 2-4, Supplementary Information
DISCUSSION			
Discussion	23a	Provide a general interpretation of the results in the context of other evidence.	Discussion
	23b	Discuss any limitations of the evidence included in the review.	Discussion
	23c	Discuss any limitations of the review processes used.	Limitations
	23d	Discuss implications of the results for practice, policy, and future research.	Clinical Implications and Suggestions for Future Research
OTHER INFORMATION			
Registration and protocol	24a	Provide registration information for the review, including register name and registration number, or state that the review was not registered.	Method
	24b	Indicate where the review protocol can be accessed, or state that a protocol was not prepared.	Method
	24c	Describe and explain any amendments to information provided at registration or in the protocol.	Prospero protocol
Support	25	Describe sources of financial or non-financial support for the review, and the role of the funders or sponsors in the review.	Funding statement
Competing interests	26	Declare any competing interests of review authors.	Conflict of interest
Availability of data, code and other materials	27	Report which of the following are publicly available and where they can be found: template data collection forms; data extracted from included studies; data used for all analyses; analytic code; any other materials used in the review.	Table 1, analytic code not publicly available



PRISMA 2020 Checklist

From: Page MJ, McKenzie JE, Bossuyt PM, Boutron I, Hoffmann TC, Mulrow CD, et al. The PRISMA 2020 statement: an updated guideline for reporting systematic reviews. *BMJ* 2021;372:n71. doi: 10.1136/bmj.n71

For more information, visit: <http://www.prisma-statement.org/>

**Prevalence of the Dissociative Subtype of Post-Traumatic Stress Disorder: A Systematic Review
and Meta-Analysis**

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Abstract

The dissociative subtype of post-traumatic stress disorder (PTSD-DS) was introduced in the fifth edition of the Diagnostic and Statistical Manual of Mental Disorders (DSM-5), and is characterised by symptoms of either depersonalisation or derealisation, in addition to a diagnosis of post-traumatic stress disorder (PTSD). This systematic review and meta-analysis sought to estimate the prevalence of current PTSD-DS, and the extent to which method of assessment, demographic and trauma variables moderate this estimate, across different methods of prevalence estimation. Studies included were identified by searching MEDLINE (EBSCO), PsycInfo, CINAHL, Academic Search Complete, and PTSDpubs, yielding 49 studies that met the inclusion criteria ($N=8214$ participants). A random effects meta-analysis estimated the prevalence of PTSD-DS as 38.1% (95% CI 31.5–45.0%) across all samples, 45.5% (95% CI 37.7–53.4%) across all diagnosis-based and clinical cut-off samples, 22.8% (95% CI 14.8–32.0%) across all latent class analysis (LCA) and latent profile analysis (LPA) samples, and 48.1% (95% CI 35.0–61.3%) across samples which strictly used the DSM-5 PTSD criteria; all as a proportion of those already with a diagnosis of PTSD. All results were characterised by high levels of heterogeneity, limiting generalisability. Moderator analyses mostly failed to identify sources of heterogeneity. PTSD-DS was more prevalent in children compared to adults, and in diagnosis-based and clinical cut-off samples compared to LCA and LPA samples. Risk of bias was not significantly related to prevalence estimates. ~~This review suggests that a significant proportion of individuals meet criteria for PTSD-DS, however the subtype remains an elusive construct. The implications of these results are discussed further.~~

Keywords: Meta-Analysis; Prevalence; Stress Disorders, Post-Traumatic; Systematic Review

Introduction

In the fifth edition of the Diagnostic and Statistical Manual of Mental Disorders (DSM-5; American Psychiatric Association [APA], 2013), post-traumatic stress disorder (PTSD) is classified as a Trauma- and Stressor-Related Disorder. A diagnosis is based on a required number of symptoms across domains of intrusion, avoidance, negative alterations in cognitions and mood, and alterations in arousal and reactivity. Also stipulated in DSM-5 are the criteria required for specifying the dissociative subtype of PTSD (PTSD-DS) where, in addition to first meeting the criteria for PTSD diagnosis, individuals must endorse symptoms of depersonalisation and/or derealisation. Depersonalisation involves “persistent or recurrent experiences of feeling detached from, and as if one were an outside observer of, one’s mental processes or body”, whereas derealisation takes the form of “persistent or recurrent experiences of unreality of surroundings” (DSM-5, 2013, pp. 272).

It has been extensively documented that persistent dissociation is linked to post traumatic symptomology (Carlson, [Dalenberg, & McDade-Montez-et al.](#), 2012). The subtype model suggests that PTSD and PTSD-DS are distinct from one another (Dalenberg & Carlson, 2012), where PTSD-DS presents with its own epidemiological features (Schiaivone, [Frewen, McKinnon, & Lanius-et al.](#), 2018). A recent systematic review concluded that there may be an association between PTSD-DS and psychopathological comorbidity and childhood abuse and neglect (Steuwe, [Lanius, & Frewen, -et al.](#), 2012), adult sexual abuse (Wolf, Miller et al., 2012), and with depression, suicidal thinking, and drug overdoses (Mergler et al., 2017), despite there being a large degree of heterogeneity in the literature concerning risk factors for PTSD-DS (Hansen, [Ross, & Armour-et al.](#), 2017). This indicates that PTSD-DS may reflect a more severe form of PTSD (Zoet, [Wagenmans, van Minnen, & de Jongh-et al.](#), 2018), although this is not directly assessed in this study.

One criticism of the PTSD-DS diagnosis is that the symptoms of dissociation chosen as necessary criteria to achieve a diagnosis in DSM-5 are too narrow, where it is believed that the current criteria should also include other symptoms of dissociation (Ross, 2021), following evidence that: dissociative amnesia (Wolf et al., 2017), and flashbacks (Dahal, [Kumar, & Thapa-et al.](#), 2018; Hyland et al., 2017) are common in individuals with PTSD. Additionally, memory disturbance, disengagement, time loss, and trance (Frewen, [Brown, Steuwe, & Lanius-et al.](#), 2015), gaps in awareness, re-experiencing, and sensory misperception (Müllerová, [Hansen, Contractor, Elhai, & Armour-et al.](#), 2016; Ross, [Baník, Dědová, Mikulášková, & Armour-et al.](#), 2018) are associated with PTSD-DS. However, to some extent, these symptoms are already captured by the existing PTSD criteria.

Several methodologies have been used to determine the prevalence of PTSD-DS, with early studies using taxometric (Waelde, [Silvern, & Fairbank-et al.](#), 2005; Waller & Ross, 1997), and signal detection (Ginzburg et al., 2006) analyses. The prevalence of PTSD-DS has also been described in studies where participants were selected primarily due to a specific comorbid difficulty, such as substance abuse disorder and psychosis, using the DSM-5 diagnostic criteria (Gidzgie et al., 2019; Mergler et al., 2017; van Minnen et al., 2016), and in studies that assessed subsyndromal PTSD (Bennett, [Modrowski, Kerig, & Chaplo-et al.](#), 2015; Kerig et al., 2016; Modrowski & Kerig, 2017). Prevalence rates of PTSD-DS have been reported in different ways; some with respect to the total number of participants regardless of whether the sample tested had PTSD, some were only trauma-exposed or from a community sample, whereas other prevalence rates were with respect to those with PTSD. This makes it challenging to make comparisons between studies. Hansen et al.'s (2017) systematic review of latent class and profile analyses (LCA and LPA respectively) [indicated the mean prevalence of PTSD-DS as 20.4%.](#), ~~which determines underlying latent profiles or~~

groups of individuals who display similar patterns of symptoms (Muthén, 2004), indicated the mean prevalence of PTSD-DS as 20.4%. LCA determines hidden groups based on the means of categorical variables, whereas LPA does the same for continuous variables (Oberski, 2016). Both LCA and LPA are exploratory techniques that determine underlying hidden profiles or groups of individuals from observed data who display similar patterns of symptoms (Muthén, 2004; Oberski, 2016). The ‘best’ number of groups is determined by the most appropriate model fit, and whilst there are many methods for determining the number of classes or profiles, the two most common methods are the Akaike information criterion and Bayesian information criterion (where lower values indicate a better fit). However, the selection of the optimal number of classes or profiles, and the qualitative naming of each group, remains is subjective on the part of the researcher which has implications for valid prevalence estimation (Hansen et al., 2017). In addition, Hansen et al. (2017) averaged the prevalence values despite dissociation being defined differently in various studies; some used the DSM-5 criteria stipulating symptoms of either depersonalisation or derealisation, and other studies assessed a wider spectrum of dissociative experiences. Finally, due to methodological constraints, there was no way of breaking down the heterogeneous nature of the population (Hansen et al., 2017).

There is a need to comprehensively systematically review studies to attempt to establish some consensus around how prevalent PTSD-DS is in children and adults. This study aimed to conduct a broad meta-analysis of data from studies investigating current PTSD-DS to reach a reliable estimate of prevalence from studies utilising various methods of prevalence estimation, furthering the systematic review of Hansen et al. (2017). The aim was to provide greater insight into the heterogeneity that is common within participants with PTSD. This might lead to the development of risk factors for this particular subtype and help the structuring of efficacious interventions. This review will be, to the authors’ knowledge,

the first of its kind to meta-analyse the prevalence of PTSD-DS in participants with PTSD, assessing moderators that affect PTSD-DS prevalence, and using studies utilising different methods of prevalence estimation. There is disagreement as to what symptoms of dissociation should be required as necessary criteria to achieve a diagnosis of PTSD-DS, and this review may shed further light on this debate, by comparing the prevalence rates of PTSD-DS when defined by depersonalisation and ~~for~~ derealisation, and when dissociation is defined more broadly.

Method

The protocol for this review was pre-registered on PROSPERO (reference: CRD42021210902) prior to any formal review of searches.

Search Strategy

Relevant studies were identified through a systematic search of the following databases: MEDLINE (EBSCO), PsycInfo, CINAHL, Academic Search Complete, and PTSDpubs. Studies included were those published from 1st January 1980, when the Diagnostic and Statistical Manual of Mental Disorders first defined PTSD according to DSM-III (APA, 1980), and before 14th February 2021 when the searches were conducted.

The following search terms were used for each database, processing study titles and abstracts only: (posttrauma* OR post-trauma* OR "post trauma*" OR PTSD OR PTSS) AND (dissociat* OR depersonali* OR dereali*). Medical Subject Headings (MeSH) terms, and other equivalent key words for other databases, were used for each search term: 'post-traumatic stress disorder', 'post-traumatic stress', 'posttraumatic stress disorder', 'posttraumatic stress', 'post-traumatic stress disorder in children', 'stress disorders, post-traumatic', 'complex PTSD', 'PTSD', 'PTSD (DSM-III)', 'PTSD (DSM-III-R)', 'PTSD (DSM-IV)', 'PTSD (DSM-5)', 'PTSD (ICD-9)', 'PTSD (ICD-10)', 'PTSD (ICD-11)', 'dissociation', and 'depersonalization'.

The reference sections of relevant systematic reviews and meta-analyses were also searched to ensure studies were not missed.

Inclusion and Exclusion Criteria

Studies were included in this review if data were presented on the prevalence of PTSD-DS following a traumatic event. In a bid to take a broad and comprehensive approach, the prevalence of PTSD-DS was defined as the number of participants: who scored above a clinical cut-off on a validated measure or who met DSM diagnostic criteria following a clinical interview or self-report measure, or who were categorised into a distinct class or profile following LCA or LPA. Studies of participants of all ages, any sex, and from either community or clinical samples were included. Studies were excluded: if they were not written in English; if participants were selected primarily due to a specific comorbid disorder; if PTSD was assessed acutely within a month of the index trauma; if exclusively lifetime PTSD or PTSD-DS prevalence was reported; if subsyndromal PTSD was assessed only; if dissociation was triggered via experimental manipulation; or if studies used analyses other than LCA, LPA, diagnostic, or clinical cut-off to determine the prevalence of PTSD-DS. Qualitative methodology, single case studies, reviews and meta-analyses were also excluded.

Screening, Data Extraction, Coding and Synthesis

All studies were screened, and the data extracted by the first author (WW) using a database which indexed the information provided in Table 1. The extracted data for all studies were reviewed by an independent researcher (AO), so as to reduce the likelihood of error (Buscemi, [Hartling, Vandermeer, Tjosvold, & Klassen-et-al.](#), 2006). Any queries were discussed, and agreement reached between the researchers. Wherever there was continued disagreement, a final decision was made by the senior researcher (RM-S). Where there was missing information, authors were contacted directly.

During data extraction, several rules were followed to ensure consistency between studies. Articles such as Eidhof et al. (2019), Guetta et al. (2019), and Zoet et al. (2018) used multiple measures for the assessment of PTSD, however in these cases the Clinician Administered PTSD scale (CAPS) was prioritised as it is regarded as the gold standard for assessing PTSD (Weathers et al., 2004). Other studies assessed multiple populations (Hansen, Müllerová, [Elklit, and Armour-et al.](#), 2016; Kenny, [Helpingstine, Long, & Harrington-et al.](#), 2020; Wolf, Lunney et al., 2012), or used multiple analyses (Choi et al., 2017; 2019; Hansen, [Hyland, Armour, & Andersen-et al.](#), 2019), and therefore these were treated separately in this review as individual samples. Care was taken to ensure that no dataset contributed more than one data point in any one meta-analysis (where diagnostic and clinical cut-off samples were prioritised over LCA and LPA samples). Multiple studies investigating the same population were removed, retaining the study with the largest sample size. Many studies (Cloitre, [Petkova, Wang, & Lu-et al.](#), 2012; Daniels, [Frewen, Theberge, & Lanius, -et al.](#), 2016; Swart, [Wildschut, Fraijer, Langeland, & Smit-et al.](#), 2020; Tsai, [Armour, Southwick, & Pietrzak-et al.](#), 2015) reported means and standard deviations for participant age and sex in aggregated format, rather than for the sample as a whole. For these studies, the means and standard deviations were combined (Altman et al., 2013; Higgins et al., 2012). When absolute frequencies were not reported, these were calculated from the reported percentage prevalence. For the LCA and LPA samples, only those classed as having 'moderate' to 'severe' symptomology were deemed to meet 'caseness' for PTSD and PTSD-DS. The prevalence of PTSD-DS was consistently calculated as a proportion of all participants with PTSD.

Quality Assessment and Risk of Bias

Two authors (WW & AO) assessed the risk-of-bias using a researcher developed tool based on the Assessment Tool for Observational Cohort and Cross-Sectional Studies

(National Heart Lung and Blood Institute, 2014), and modified questions from other relevant prevalence and risk factor studies (Hoy et al., 2012; Munn, [Moola, Riitano, & Lisy et al.](#), 2014). The quality assessment checklist (see Supplementary Material) consisted of five items assessing how well the population and index trauma were specified, the rate of participation, and whether objective and standard criteria were used for the assessment of PTSD and PTSD-DS. Each item used a three-point scale (0-2), and the following categorical system was used to rate the total risk-of-bias score: 0-4 high risk/low quality, 5-6 moderate risk/quality, 7-10 low risk/high quality, following the methodology used ~~in~~ [by](#) [Memarzia, Walker, and Meiser-Stedman et al.](#) (2021). An inter-rater reliability assessment was conducted for all ratings between the two raters (WW & AO) which indicated a good correlation on all items (intraclass correlation = 0.87, 95% CI 0.77–0.93).

Meta-Analytic Method

The meta-analysis was conducted using R (version 4.1.1) which uses the metafor package (version 3.0-2; Viechtbauer, 2010). The extracted prevalence of PTSD-DS, as a proportion of all PTSD cases, was pooled to provide a weighted estimate of the prevalence of PTSD-DS overall (with 95% confidence intervals [CI]).

A random effects model was used given the high degree of variability expected in effect size between samples as it provides a broader and more conservative 95% confidence interval around the estimate of the prevalence.

The estimates of the prevalence underwent an arcsin transformation to ensure that the confidence intervals did not fall below zero for samples where the prevalence estimate was low ([Barendregt, Doi, Lee, Norman, & Voset et al.](#), 2013); results were then back transformed for ease of interpretation.

Cochran's Q test (Cochran, 1954) was used to ascertain if heterogeneity within samples was significant. The I^2 statistic (Higgins & Thompson, 2002) was used to determine

the percentage of total variation in sample estimates that is due to between-study heterogeneity.

Moderator analyses of prevalence estimates were conducted to ascertain if sample characteristics impacted the prevalence estimate. These characteristics included: method of PTSD-DS assessment, which DSM criteria was used, participant age group, occupation, and the type of trauma suffered. These were included as there were multiple samples that allowed for these comparisons to be made. A sensitivity analysis was used to assess the impact of risk-of-bias on the estimated pooled prevalence. This was achieved by repeating the meta-analysis, excluding those samples that constituted a high risk-of-bias. Any differences in the moderator and sensitivity analyses were tested for clinical significance by meta-analytic regression.

A funnel plot was used to assess for publication bias (Higgins et al., 2012), however this is less likely to occur in prevalence studies given there is no assessment of clinical significance, and therefore it is less likely that there is a bias in levels of acceptance to journals (Brewin, Andrews, & Valentine *et al.*, 2000). The 'trim-and-fill' method was used (Duval & Tweedie, 2000), where any missing null or weaker studies are estimated to improve the symmetry of the sample distribution.

Results

The Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) diagram shows that 337 studies met the eligibility criteria following the initial screen of titles and abstracts (Figure 1). Full text reviews were conducted again, leading to 49 studies being included in the meta-analysis. Four studies were split into two samples due to different characteristics, index traumas or analyses, leaving 53 samples included in this review (Table 1). Around half the samples were treatment-seeking ($k = 23$), and PTSD-focussed (a diagnosis of PTSD was an inclusion criterion; $k = 22$). Nine samples included

only female participants, three samples included only males, and the rest were mixed of the sex was not reported. The majority of included samples were adult ($k=41$); only five exclusively comprised children. Samples mostly originated from high-income countries ($k=49$).

[Figure 1]

[Table 1]

Risk-of-Bias Assessment

Twelve samples were deemed to be at high risk-of-bias, 16 were moderate risk, whereas 25 were low risk. The proportion of samples rated as low, moderate and high risk across the five quality assessment items can be seen in the Supplementary Material.

Prevalence

The pooled prevalence of PTSD-DS estimates and heterogeneity statistics for all samples can be seen in Table 2. The overall pooled prevalence was 38.1%. For diagnosis-based and clinical cut-off samples the pooled prevalence was 45.5%, while for latent class and profile samples the estimate was 22.8%. Meta-regression analyses indicated that the prevalence of PTSD-DS in the diagnosis-based or clinical cut-off samples was statistically significantly greater than the LCA or LPA samples (see Figure 2 for forest plot). The range of prevalence overall was 0-100%, and the degrees of between sample heterogeneity were extremely high.

[Table 2]

[Figure 2]

Moderator Analyses

All Samples

Moderator analyses were conducted for all samples to assess whether the pooled prevalence estimate of PTSD-DS was associated with demographic, trauma or assessment

factors (Table 2). Meta-regression analyses confirmed that the prevalence of PTSD-DS in the child samples was statistically significantly greater than the adult samples, although there were only four child samples for comparison. All other comparisons were non-significant; however, several comparisons were likely underpowered.

Diagnostic and Clinical Cut-off Samples

Further subgroup moderator analyses were conducted separately for the diagnostic and clinical cut-off samples (Table 3), regardless of the dissociation criteria used, given the significant difference in pooled prevalence estimates of PTSD-DS between these samples and those using LCA or LPA. Meta-regression analyses confirmed again that the prevalence of PTSD-DS in the child samples was statistically significantly greater than the adult samples, although there were only four child samples for comparison. All other comparisons were non-significant, however several comparisons were likely underpowered.

[Table 3]

Further moderator analyses were conducted for only those samples utilising DSM-5 criteria for dissociation (depersonalisation and ~~or~~ derealisation; see Supplementary Material). When only samples using DSM-5 diagnostic and clinical cut-off criteria for the assessment of PTSD and PTSD-DS were pooled, the estimated prevalence of PTSD-DS was 48.2%. This provides the most valid estimate of PTSD-DS prevalence according to the DSM-5 criteria. Meta-regression analyses confirmed again that the prevalence of PTSD-DS in the child samples was statistically significantly greater than the adult samples, although there were only four child samples for comparison. All other comparisons were non-significant, however several comparisons were likely underpowered.

Latent Class and Profile Samples

Moderator analyses were conducted separately for the LCA ~~and~~ LPA samples (see Supplementary Material), again given the significant difference in pooled prevalence of

PTSD-DS estimates between these samples and those using diagnostic and clinical cut-off methods. Meta-regression analyses confirmed that there were no statistically significant differences, however several comparisons were likely underpowered.

Sensitivity Analyses

When the 28 samples of low and moderate quality were removed, the estimated prevalence of PTSD-DS was not dissimilar to that for all samples (35.7%, 95% CI 24.8–47.3%) with a similar degree of between sample heterogeneity ($k=25$, $Q(24)=717.8$, $p<0.0001$, $I^2=98.5\%$). Meta-regression analyses indicated there was not a significant difference between high and low-moderate quality groups ($\beta=-0.0040$ [95% CI -0.1384, 0.1463], $p=0.96$). Therefore, it can be concluded that there was no support for the quality of the samples affecting the prevalence of PTSD-DS estimates.

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Given the differences in prevalence in PTSD-DS between child and adult samples, the child samples were removed to assess whether similar results were achieved as in Table 2. Meta-regression analyses confirmed that the only statistically significant difference existed between the estimated prevalence of PTSD-DS for the diagnosis-based or clinical cut-off samples and LCA or LPA samples ($\beta=-0.2159$ [95% CI -0.3531, -0.0787], $p=0.002$). All other comparisons were non-significant ($p_s=0.19-0.87$), however several comparisons were likely underpowered.

Publication Bias

Visual inspection of the funnel plot (see Supplementary Material) suggests the distribution of samples is asymmetrical, which was confirmed by Egger's test ($p=0.03$). However, the study of Kenny et al. (2020; sample 29) was very small and should be considered an outlier. When this sample was removed, the Egger's test confirmed the symmetry of the distribution ($p=0.30$). No null or weaker studies were estimated as missing, indicating little to no publication bias.

Discussion

A comprehensive systematic review and meta-analysis of prevalence data from studies investigating current PTSD-DS utilising various methods of prevalence estimation was completed. The aim was to generate a reliable estimate for the prevalence of PTSD-DS and to provide greater insight into the heterogeneity that is common within participants with PTSD-DS. The estimated pooled prevalence of PTSD-DS was: 38.1% for all samples, 45.5% for all diagnostic and clinical cut-off samples, and 22.8% for all LCA and LPA samples. The estimated prevalence of PTSD-DS from the LCA and LPA samples was similar to the mean prevalence found in the Hansen et al. (2017) systematic review (20.4%); this is unsurprising given nine of the eleven studies in the Hansen et al. (2017) review were also included in the present study. When only samples strictly using DSM-5 diagnostic and clinical cut-off criteria for the assessment of PTSD and PTSD-DS were pooled, the estimated prevalence of PTSD-DS was 48.1%. The prevalence of PTSD-DS may therefore be significantly greater than previously suggested.

Impact of Diagnostic and Clinical Cut-off Assessment Versus LCA and LPA on Estimated Prevalence of PTSD-DS

The estimated prevalence of PTSD-DS for the diagnostic and clinical cut-off samples was significantly higher than that of the LCA and LPA samples. Use of clinical cut-off measures may overestimate the prevalence of PTSD in adults (Richardson, Frueh, & Acierno et al., 2010). Moreover, it may be easier to identify individuals with PTSD who show symptoms of depersonalisation or derealisation in a clinical interview or that surpass a clinical cut-off on a dissociation measure, rather than via LCA and LPA methods. On the other hand, latent class and profile analyses may rely on participants reporting multiple significant dissociative symptoms rather than just one symptom to a significant level.

Achterhof, Huntjens, Meewisse, and Kiers ~~Achterhof et al.~~ (2019) questioned the use of LCA

and LPA to ascertain the prevalence of Complex PTSD and highlighted that despite the analyses determining distinct profiles, the symptom profile for groups of participants were very close to one another and even overlapped on occasion. Therefore, it may be questioned whether LCA and LPA reliably and validly estimates subtype prevalence.

Impact of Moderators on Estimated Prevalence of PTSD-DS

There was no significant difference between the estimated prevalence of PTSD-DS when dissociation was assessed by the DSM-5 criteria (presence of either depersonalisation or derealisation) or when defined by a broader spectrum of dissociative symptoms. The aim of the inclusion of the PTSD-DS in DSM-5 was to define a small subgroup of individuals with consistent clinical and epidemiological features (Miller, [Wolf, & Keane-et al.](#), 2014; Schiavone et al., 2018), however results from the present study suggest a subtype where the prevalence varies very widely across samples (0-100%) and where the heterogeneity cannot be broken down following moderator analyses. Research literature suggests that the symptomology of PTSD is itself heterogeneous (Elhai, [Frueh, Davis, Jacobs, & Hamner-et al.](#), 2003; Galatzer-Levy & Bryant, 2013; Naifeh, [Richardson, Del Ben, & Elhai-et al.](#), 2010), where dissociation is one such symptom that can vary.

The estimated prevalence of PTSD-DS was significantly higher for samples of children compared to adults, although there are limited number of samples investigating exclusively children, and the results were dominated by that of Choi et al. (2019; sample 10). There was no one trauma type that best categorised the child samples. Research has ~~also~~ shown that dissociation is a common experience for children, that later becomes less prevalent with child development and the transition into adulthood (Brunner, [Parzar, Schuld, & Resch-et al.](#), 2000; Coons, 1996; Choi et al., 2017; Shimizu & Sakamoto, 1986). Choi et al. (2019) reported that 53.7% of children with PTSD had the dissociative subtype; a prevalence much higher than in many other adult samples, and the authors cited the prominence of

dissociation as a form of coping in response to maltreatment in childhood (Liotti, 2004; Putnam, 1997). Children may be more susceptible to PTSD-DS because they do not have the same capacity to avoid cues relating to the traumatic event, especially when the trauma was based within the home environment, or with a primary caregiver (Choi et al., 2019). In children, dissociation may offer an alternative method of escape to reduce distress. It might also be considered whether depersonalisation and derealisation are the most appropriate symptoms by which to assess for PTSD-DS in children. The premise of the subtype model is that these dissociative symptoms are rare (Lanius et al., 2014), however it may be that dissociative experiences are more common in youth (Carlson, Yates, & Sroufe et al., 2009) and may not even be considered as pathological. Further research is required within this area to determine whether children are more at risk from dissociation in the context of PTSD compared to adults, as the lack of power within the samples of children frustrated the moderator analyses.

Other than age group, all other moderator analyses yielded non-significant results indicating no support for any differences between estimated prevalence of PTSD-DS. This is surprising given the extant research on mediators and risk factors in relation to PTSD-DS (Hansen et al., 2017; Schiavone et al., 2018 for review), but these non-significant results are likely to reflect the heterogeneity between these samples and the lack of power in some moderator analyses.

It is important to stress that the pooled prevalence estimates were characterised by a high degree of heterogeneity throughout, and inspection of the forest plot (Figure 2) shows how varied the prevalence of PTSD-DS is across different samples. This is not unexpected given the multiple ways of assessing and conceptualising PTSD-DS, however subsequent sensitivity and moderation analyses failed to reduce the level of heterogeneity. This therefore limits the generalisability of the findings. The consistently high level of heterogeneity may

reflect the difficulty in conceptualising and defining a construct such as dissociation in the context of PTSD. Even when only samples adhering to the strict DSM-5 criteria for PTSD-DS were pooled, a high degree of heterogeneity remained.

Clinical Implications and Suggestions for Future Research

— This meta-analysis suggests that PTSD-DS is common following trauma exposure ~~in children and adults~~, and therefore should be routinely assessed for and formulated. Moreover, the method for determining PTSD-DS was found to have important implications for the estimated prevalence, where samples using diagnostic and clinical cut-off methods reported a higher prevalence than those using LCA and LPA. Future research should also aim to standardise the methodology used to identify and determine PTSD-DS in order to make more valid comparisons between studies.

Additionally, PTSD-DS was found to be more common in children than adults. Clinicians supporting individuals with PTSD should be aware that dissociation is a prevalent and important feature of the overall presentation of PTSD; this may be especially true for children, though this finding was based on only five samples. When the DSM-5 criteria were published it was believed that PTSD-DS cases formed a minority of those with PTSD, however the finding that nearly half of PTSD cases meet the criteria for PTSD-DS suggests that it may be less of a subtype and that dissociation forms a central component to PTSD symptomology. This should be a consideration for how dissociation is specified in future versions of the DSM. Perhaps the conceptualisation of Complex PTSD as defined by the 11th revision of the International Classification of Diseases (World Health Organisation, 2019), where dissociation is stipulated as one of several symptoms seen to be indicative of a more complex form of PTSD, is a more appropriate fit. There is evidence for instance that individuals with Complex PTSD have elevated levels of dissociation (Hyland, Shevlin, Fyvie, Cloitre, & Karatzias, 2019).

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Despite the DSM-5 criteria stipulating depersonalisation and derealisation as symptoms required for PTSD-DS, findings of this review suggested that when a wider view of dissociation (i.e., drawing on a broader range of dissociation symptoms) is included in the criteria, PTSD-DS prevalence does not change significantly. No conclusions can be drawn as to whether it would be more or less appropriate for a narrower (i.e., solely based on depersonalisation and or derealisation) or a broader definition of dissociation, in the context of this subtype, to be used in future versions of diagnostic criteria. However, it does not seem to matter how dissociation is defined when determining the prevalence of PTSD-DS, which raises questions firstly about the strict nature of the DSM criteria when defining this subtype (Ross, 2021), and secondly about the existence of this subtype full stop. This suggests that it does not matter how dissociation is defined when determining a threshold for this subtype.

~~PTSD-DS remains an elusive construct, as evidenced by the considerable heterogeneity found within this study, even when considering specific sub-groups and tightening methodological or diagnostic criteria.~~ Further research is required to establish ~~if there are particular risk factors, and~~ whether PTSD-DS could be indicative of a distinct form of PTSD that has its own clinical characteristics, and therefore break down the heterogeneity common to populations with the subtype. This would help inform exactly how dissociation should be integrated into future diagnostic criteria of PTSD. Perhaps as Ross (2021) suggests, future diagnostic criteria could stipulate the requirement for the presence of one or more of: depersonalisation, derealisation, dissociative amnesia, and dissociative flashbacks. Non-dissociative PTSD may then form the subtype based on a minority of cases, and dissociative PTSD may form the majority of diagnosed cases.

~~Future research should aim to standardise the methodology used to identify and determine PTSD-DS in order to make more valid comparisons between studies.~~

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Limitations

There are several limitations that should be considered for this review. Firstly, whilst many more studies were reviewed in comparison to the most recent systematic review (Hansen et al., 2017), there was still a considerable degree of heterogeneity between samples, reducing the generalisability of the findings. This raises questions around the **reliability** **validity** of the underlying diagnostic subtype. Secondly, most studies were conducted in high income countries, and all studies were exclusively written in English, therefore indicating that the results are likely not globally generalisable. Thirdly, some moderator analyses lacked power and further planned moderator analyses were not possible due to a lack of identified studies. Understanding the influence of, for instance, sex, time between index trauma and PTSD assessment, single- versus multi-event traumas, and individual versus collective trauma could lead to important and interesting findings. Finally, several studies chose to assess PTSD-DS with regard to the most recent trauma that the participant was exposed to, and it is unclear whether other traumas may have taken place, and what impact these may have on the prevalence of PTSD-DS.

Conclusion

This study is the first to meta-analyse data on the prevalence of PTSD-DS. The estimated prevalence of PTSD-DS, with respect to participants diagnosed with PTSD, was 38.1% (95% CI 31.5 – 45.0%) for all samples, 45.5% (95% CI 37.7 – 53.4%) for all diagnosis-based and clinical cut-off samples, 22.8% (95% CI 14.8 – 32.0%) for all LCA and LPA samples, and 48.1% (95% CI 35.0 – 61.3%) for diagnosis-based and clinical cut-off samples which assessed PTSD and PTSD-DS strictly according to the DSM-5 criteria. The prevalence of PTSD-DS was significantly higher for children compared to adults. Factors such as the DSM criteria used for the assessment of both PTSD and dissociation, whether the dissociation assessment was self-report or interview, and participant or trauma

characteristics, did not significantly affect the estimated prevalence of PTSD-DS. However, all results were characterised by very high levels of heterogeneity. Further research is required to investigate this construct, and to determine how it should be best conceptualised in future editions of diagnostic criteria.

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Conflict of Interest: None

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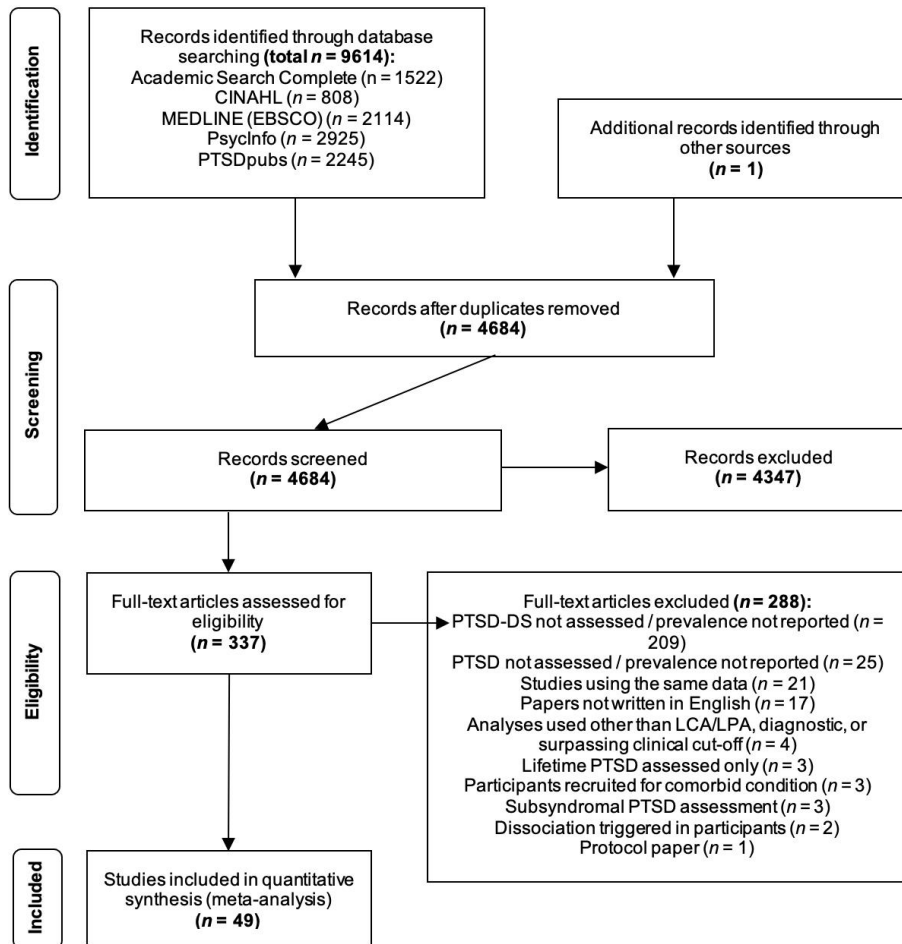
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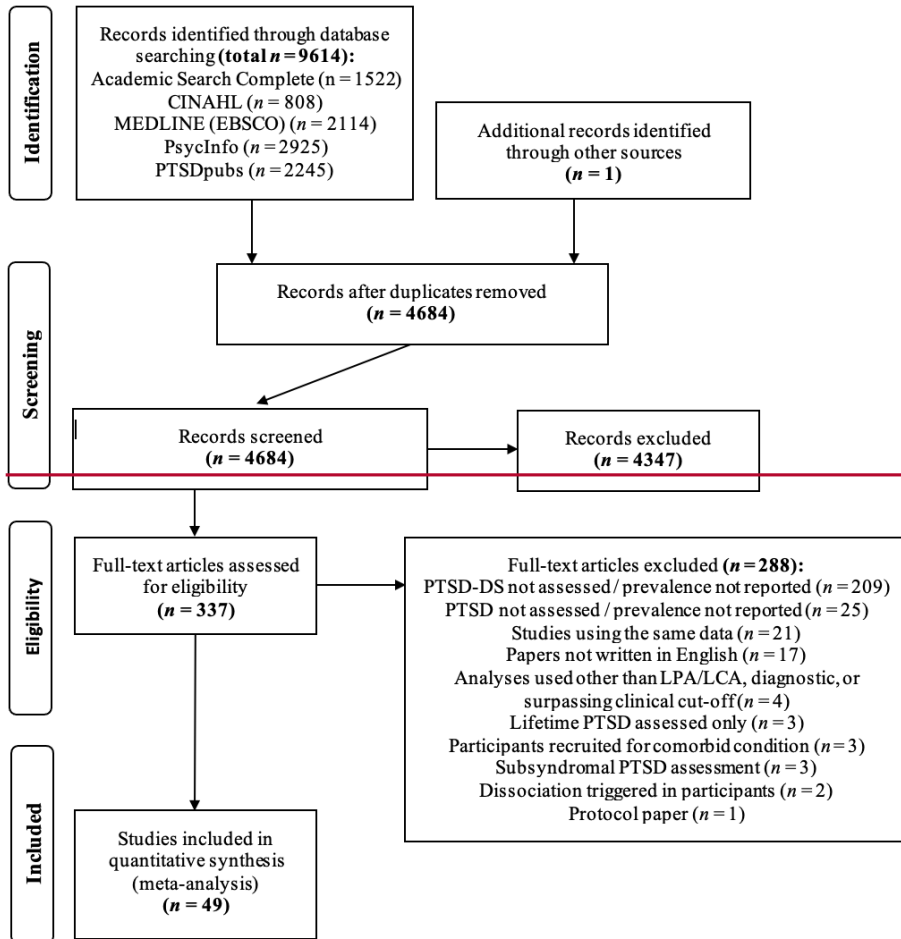


Figure 1

PRISMA diagram showing the process of study identification, screening, and inclusion (n = number of studies)

Table 1*Included sample characteristics*

Sample No.	Sample	Location	Population/trauma type	Proportion female	Age			Method of PTSD assessment	PTSD measure; DSM	PTSD-DS measure; DSM-5/other criteria	N		
					Range	Mean (SD)	Age group				Total	PTSD	PTSD-DS
1	Abu-Rus, Thompson, Naish, Brown, and Dalenberg et al. (2020)	USA	General population (T, P)	46%	NR	37.9 (10.3)	NR	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	345	40	16
2	Acar, Ögülmüş, and Boysan et al. (2019)	Turkey	Prisoners	3%	18-75	34.5 (9.9)	Adult	Diagnosis	PCL [†] ; DSM-5	DES [†] ; other	399	237	115
3	Armour, Elklit, Lauterbach, and Elhai et al. (2014)	Denmark	Sexual assault and rape (T)	100%	NR	22.4 (9.4)	Both	LPA	HTQ [†] ; DSM-IV	TSC [†] ; other	313	226	41
4	Armour, Karstoft, and Richardson et al. (2014)	Canada	Military veterans (T)	6%	24-93	54.0 (19.0)	Adult	LPA	CAPS; DSM-IV	CAPS; other	432	286	59
5	Blevins, Weathers, and Witte et al. (2014)	USA	Trauma-exposed college students	67%	18-32	20.2 (1.6)	Adult	LCA	PCL-S [†] ; DSM-IV	MDI [†] ; DSM-5	541	206	65
6	Boysan et al. (2017)	Turkey	Psychiatric patients (T, P)	44%	NR	29.0 (9.0)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	90	30	24
7	Briere, Scott, and Weathers et al. (2005)	USA	Trauma-exposed community	48%	NR	45.2 (16.7)	Adult	Diagnosis	DAPS [†] ; DSM-IV	DAPS [†] ; other	372	23	13
8	Burton, Feeny, Connell, and Zoellner et al. (2018)	USA	Chronic PTSD (P)	76%	NR	37.4 (11.3)	Adult	LTA (expanded version of LPA)	PSS-I; DSM-IV	DES-D [†] ; DSM-5	200	129	24
9	Caroppo, Lanzotti, and Janiri et al. (2021)	Italy	Asylum seekers (T)	48%	18-59	25.5 (5.6)	Adult	Diagnosis	SCID-I; DSM-IV	SCID-I; other	180	95	74
10	Choi et al. (2019)	USA	Trauma-exposed adolescents (T)	61%	12-16	14.5 (1.5)	Child	Diagnosis	UCLA PTSD-RI [†] ; DSM-IV	TSCC-A [†] ; DSM-5	3081	734	394
11	Choi et al. (2017)	USA	Trauma-exposed adolescents (T)	61%	12-16	14.5 (1.5)	Child	LCA	UCLA PTSD-RI [†] ; DSM-IV	TSCC-A [†] ; DSM-5	3081	1279	444
12	Cloitre et al. et al. (2012)	USA	Childhood sexual and/or physical abuse (P)	100%	18-65	36.4 (9.4) [§]	Adult	Diagnosis	CAPS; DSM-IV	TSI [†] ; other	104	104	28
13	Criswell, Sherman, and Krippner et al. (2018)	USA	Psychiatric patients (T, P)	73%	20-65	44.0 (NR)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	30	30	13
14	Daniels et al. et al. (2016)	Germany	Trauma-exposed community (P)	61%	23-58	38.0 (11.8) [§]	Adult	Diagnosis & clinical cut-off	CAPS; DSM-IV	CAPS; DSM-5	59	59	15
15	Dorahy et al. (2017)	Northern Ireland	Psychiatric patients (T, P)	32%	19-65 [‡]	40.4 (12.4)	Adult	Diagnosis	Clinical diagnosis; NR	DES [†] ; other	210	65	27
16	Durham, Byllesby, Elhai, and Wang et al. (2020)	USA & Canada	Trauma-exposed community	63%	18-74	36.0 (12.7)	Adult	LPA	PCL [†] ; DSM-5	DES-II [†] ; DSM-5	360	204	51

PREVALENCE OF DISSOCIATIVE SUBTYPE OF PTSD

17	Eidhof et al. (2019)	Netherlands	Trauma-exposed community (T, P)	33%	19-83	48.8 (12.1)	Adult	Diagnosis	CAPS; DSM-5	CAPS [†] ; DSM-5	320	131	31
18	Frewen et al. et al. (2015)	Canada	Probable diagnosis of PTSD (T)	71%	NR	33.1 (10.8)	Adult	LPA	PCL [†] ; DSM-5	Dissociation-TRASC item list [†] ; DSM-5	557	311	183
19	Frewen, <u>Zhu, and Lanius-et al.</u> (2019)	Canada	Community	52%	NR	36.5 (12.6)	Adult	Diagnosis	PCL [†] ; DSM-5	Dissociation-TRASC item list [†] ; DSM-5	418	98	41
20	Guetta et al. (2019)	USA	Military veterans (P)	16%	21-75	53.8 (11.4)	Adult	LPA	PCL, Trauma Assessment from the NSES; DSM-5	CAPS [†] ; DSM-5	209	209	31
21	Hansen, Hyland, <u>and Armour-et al.</u> (2016)	Denmark	Bank employees following robbery	62%	20-65	42.1 (12.5)	Adult	LCA	HTQ [†] ; DSM-IV	TSC [†] ; DSM-5	371	67	0
22	Hansen et al. et al. (2019)	Denmark	Whiplash injury	62%	18-89	37.5 (13.9)	Adult	Diagnosis	HTQ [†] , TSC [†] ; DSM-5	TSC [†] ; DSM-5	234	21	7
23	Hansen <u>et al. et al.</u> (2019)	Denmark	Whiplash injury	62%	18-89	37.5 (13.9)	Adult	LCA	HTQ [†] , TSC [†] ; DSM-5	TSC [†] ; DSM-5	234	27	0
24	Hansen, Müllerová et al. et al. (2016)	Denmark	Whiplash injury (P)	78%	NR	43.6 (10.4)	Adult	LCA	HTQ [†] , TSC [†] ; DSM-5	TSC [†] ; DSM-5	476	476	178
25	Hansen, Müllerová <u>et al. et al.</u> (2016)	Denmark	Incest during childhood (T, P)	88%	NR	35.9 (11.0)	Adult	LCA	HTQ [†] , TSC [†] ; DSM-5	TSC [†] ; DSM-5	311	311	139
26	Harricharan et al. (2020)	Canada	Trauma-exposed community (P)	63%	18-60 [‡]	39.6 (12.5) [§]	Adult	Diagnosis & clinical cut-off	CAPS; DSM-IV & 5	CAPS; DSM-5	184	133	49
27	Hill et al. (2020)	USA	Trauma-exposed women (T)	100%	18-62	34.1 (13.2)	Adult	Clinical cut-off	PCL [†] ; DSM-5	DSPS [†] ; DSM-5	104	88	73
28	Kenny et al. et al. (2020)	USA	Commercial sexual exploitation (T)	100%	12-18	16.6 (1.2) [*]	Child	Diagnosis	UCLA PTSD-RI [†] ; DSM-5	UCLA PTSD-RI [†] ; DSM-5	56	15	11
29	Kenny et al. (2020)	USA	At risk of commercial sexual exploitation (T)	100%	12-18	15.3 (1.6) [*]	Child	Diagnosis	UCLA PTSD-RI [†] ; DSM-5	UCLA PTSD-RI [†] ; DSM-5	40	3	3
30	Kim et al. (2019)	South Korea	Psychiatric patients (T, P)	64%	16-70	38.7 (12.7)	Both	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	249	249	82
31	Lebois et al. (2021)	USA	Interpersonal childhood maltreatment (T, P)	100%	18-61	34.4 (12.2)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	65	65	47
32	Li, <u>Hasset, and Seng-et al.</u> (2019)	USA	Pregnant women	100%	NR	NR	NR	Diagnosis	National Women's Study PTSD Module; DSM-IV	DES-T [†] ; other	22	10	4
33	Mulder, <u>Beautrais, Joyce, and Fergusson et al.</u> (1998)	New Zealand	Community	NR	NR	NR	Adult	Diagnosis	SCID; DSM-III	DES [†] ; other	1028	9	3
34	Müllerová <u>et al. et al.</u> (2016)	USA & Canada	Trauma-exposed community	56%	NR	35.2 (11.9)	NR	LPA	PCL [†] ; DSM-5	DSS [†] ; other	309	215	83

PREVALENCE OF DISSOCIATIVE SUBTYPE OF PTSD

35	Naish et al. (2021)	USA	Trauma-exposed community	45%	18-65	40.5 (11.8)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	100	63	31
36	Nejad et al. (2007)	Iran	Military veterans (P)	0%	NR	41.5 (5.1)	Adult	Diagnosis	Clinical diagnosis; DSM-IV	DES [†] ; other	260	130	42
37	Özdemir, Celik, and Oznur-et al. (2015)	Turkey	Serving soldiers (P)	0%	NR	30.3 (5.6)	Adult	Diagnosis	SCID-I; DSM-IV	DES [†] ; other	184	84	59
38	Powers et al. (2017)	USA	Trauma-exposed women	100%	18-65 [‡]	39.4 (11.6)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	190	72	2
39	Putnam et al. (1996)	USA & Canada	Psychiatric patients - (T, P)	60%	NR	39.0 (NR)	Adult	Diagnosis	Clinical diagnosis; DSM-III	DES [†] ; other	1566	116	54
40	Richard-Malenfant, Douglass, Higginson, Ray, and Robillard et al. (2019)	Canada	Military veterans (P)	36%	NR	49.3 (9.3)	Adult	Diagnosis	CAPS; DSM-5	CAPS; DSM-5	14	14	6
41	Ross, Armour, Kerig, Kidwell, and Kilshaw et al. (2020)	USA	Trauma-exposed youth in detention centres	25%	12-19	16.0 (1.3)	Child	Diagnosis	UCLA PTSD-RI [†] ; DSM-5	UCLA PTSD-RI [†] ; DSM-5	448	197	119
42	Ross et al. (2018)	Slovakia	Trauma-exposed university students	83%	NR	22.7 (5.1)	Adult	LPA	PCL [†] ; DSM-5	DSS [†] ; other	689	308	24
43	Sierk, Manthey, Brakemeier, Walter, and Daniels et al. (2021)	Germany	Childhood interpersonal abuse (P)	100%	NR	40.0 (9.8)	Adult	Diagnosis	CAPS; DSM-IV	DES [†] , CDS-30 [†] , CDS-state [†] , CAPS, SCID-D; other	42	42	23
44	Stein et al. (2013)	Global	Community	NR	NR	NR	Adult	Diagnosis	WHO CIDI; DSM-IV	WHO CIDI; DSM-5	25018	747	108
45	Steuwe et al. (2012)	Canada	Trauma-exposed community (T, P)	90%	NR	37.9 (9.4)	NR	Diagnosis	CAPS; DSM-IV	CAPS; DSM-5	134	134	47
46	Swart et al. (2020)	Netherlands	Psychiatric patients (T)	77%	18-68	34.2 (11.9) [§]	Adult	Diagnosis	CAPS; DSM-IV	DES [†] ; DSM-5	150	84	18
47	Tsai et al. (2015)	USA	Military veterans	NR	20-94 [‡]	60.8 (15.2) [§]	Adult	Diagnosis	PCL [†] ; DSM-5 SCID & DIS	CAPS [†] ; DSM-5	1484	64	12
48	van der Kolk et al. (1996)	USA	Psychiatric patients (T)	67%	15+	37.1 (15.0)	Both	Diagnosis	PTSD modules; DSM-III	SIDES; other	395	182	149
49	Verbeck et al. (2015)	USA	Psychiatric patients (T)	49%	18-69	44.0 (10.9)	Adult	Diagnosis	CAPS; DSM-IV	TSI-2 [†] , DES-R [†] ; other	100	47	29
50	Wolf, Lunney et al. (2012)	USA	Military veterans (P)	0%	44-74	50.6 (3.6)	Adult	LPA	CAPS; DSM-IV	CAPS; other	360	360	56
51	Wolf, Lunney et al. (2012)	USA	Military veterans (P)	100%	22-78	44.8 (9.4)	Adult	LPA	CAPS; DSM-IV	TSI [†] ; DSM-5	284	284	85
52	Wolf, Miller et al. (2012)	USA	Military veterans & their partners	36%	21-75 [‡]	51.5 (11.2) [‡]	Adult	LPA	CAPS; DSM-IV	CAPS; other	492	239	30

53	Zoet et al. (2018)	Netherlands	Psychiatric patients (T)	70%	19-63‡	38.2 (10.9)§	Adult	Clinical cut-off	CAPS; DSM- IV	CAPS†; DSM-5	168	168	38
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Note. SD = standard deviation, T = treatment-seeking inclusion criteria; P = diagnosis of PTSD inclusion criteria; NR = Not Reported; CAPS = Clinician Administered Post-traumatic Stress

Disorder Scale; PCL = Post-traumatic Stress Disorder Checklist; DES = Dissociative Experiences Scale; [LPA = latent profile analysis](#); HTQ = Harvard Trauma Questionnaire; TSC = Trauma Symptom Checklist; [LCA = latent class analysis](#); PCL-S = Post-traumatic Stress Disorder Checklist Specific; MDI = Multiscale Dissociation Inventory; DAPS = Detailed Assessment of Posttraumatic Stress; DES-D = depersonalization/derealisation subscale of the DES; PSS = Post-traumatic Stress Disorder Symptom Scale; PSS-I = PTSD Symptom Scale-Interview, SCID-I = Structured Clinical Interview for the DSM-IV Axis I Disorders; UCLA PTSD-RI = University of California at Los Angeles Posttraumatic Stress Disorder Reaction Index; TSCC-A = Trauma Symptom Checklist for Children-Alternate Version; TSI Trauma Symptom Inventory; TRASC = trauma-related altered states of consciousness; NSES = National Stressful Events Survey; DES-T = 8-item taxon version of the Dissociative Experiences Scale; SCID = Structured Clinical Interview for DSM; DSPTS = Dissociative Subtype of PTSD Scale; DSS = Dissociative Symptoms Scale; CDS = Cambridge Depersonalization Scale; SCID-D = Structured Clinical Interview for DSM-IV Dissociative Disorders; WHO CIDI = World Health Organisation Composite International Diagnostic Interview; DIS = Diagnostic Interview Schedule; SIDES = Structured Interview for Disorders of Extreme Stress; DES-R = Dissociative Experiences Scale – Revised

† Measure completed via self-report

‡ Information acquired via correspondence with study author(s)

§ Mean and standard deviation values combined ([Altman, Machin, Bryant, & Gardner, Altman et al., 2013](#); Higgins et al., 2012)

¶ Multiple measures used, however CAPS chosen as the gold standard (Weathers et al., 2004)

Table 2

Pooled prevalence of PTSD-DS as a proportion of PTSD for all samples (k = 51)

Meta-analysis subgroup	k	n	Pooled Prevalence (%)	95% CI	Q test	I ²
All samples [†]	51	8214	38.1	(31.5, 45.0)	1602.0*	97.4
Method of PTSD-DS Assessment ($\beta = -0.2418$ [95% CI = -0.3780, -0.1056], $p = 0.0005$)						
Diagnosis-based/clinical cut-off	36	4383	45.5	(37.7, 53.4)	923.6*	96.0
LCA/LPA [†]	15	3831	22.8	(14.8, 32.0)	482.5*	97.6
PTSD DSM criteria used ^{‡‡} ($\beta = -0.0871$ [95% CI = -0.2328, 0.0586], $p = 0.24$)						
DSM-5	24	3451	42.5	(32.4, 53.0)	624.6*	97.3
DSM-III or DSM-IV	25	4565	34.1	(24.9, 43.9)	936.0*	97.8
Dissociation criteria [†] ($\beta = 0.0342$ [95% CI = -0.1113, 0.1796], $p = 0.65$)						
DSM-5 (Dereal / Depers)	32	5436	36.9	(28.5, 45.8)	895.2*	97.6
Broader dissociation	19	2778	40.2	(29.5, 51.4)	698.3*	97.1
Dissociation measure completion ^{§§} ($\beta = 0.0281$ [95% CI = -0.1189, 0.18], $p = 0.7080$)						
Self-report	31	4997	38.8	(30.6, 47.3)	778.8*	97.2
Interview	19	3175	36.2	(24.8, 48.5)	690.4*	97.9
Age group ^{¶¶} ($\beta = 0.3587$ [95% CI = 0.0814, 0.6360], $p = 0.01$)						
Child	4	949	62.9	(39.6, 83.3)	11.4*	82.0
Adult	40	6209	35.0	(27.8, 42.6)	1121.1*	97.3
Occupation [†] ($\beta = -0.1439$ [95% CI = -0.3227, 0.0350], $p = 0.11$)						
Military	9	1670	26.9	(16.2, 39.1)	138.1*	96.3
Civilian	42	6544	40.7	(33.1, 48.5)	1325.7*	97.4
Trauma type [†] ($\beta = 0.1011$ [95% CI = -0.1163, 0.3185], $p = 0.36$)						
Interpersonal	6	763	46.8	(28.3, 65.7)	101.9*	95.9
Other	45	7451	37.0	(29.9, 44.3)	1494.9*	97.5

Note. k = number of samples; n = number of participants; CI = confidence interval; LCA = latent

class analysis, LPA = latent profile analysis; Dereal = derealisation; Depers = depersonalisation

* $p < 0.0001$, where the degrees of freedom (df) = $k - 1$

[†] Samples 11 and 23 removed to avoid duplication of population samples

^{‡‡} Sample 15 removed as no PTSD DSM criteria reported, sample 26 removed as used both DSM-IV and DSM-5 when assessing for PTSD

^{§§} Sample 43 removed as a mix of self-report and interview measures were used

^{¶¶} Several samples were removed due to populations formed of both children and adults, or age group not reported

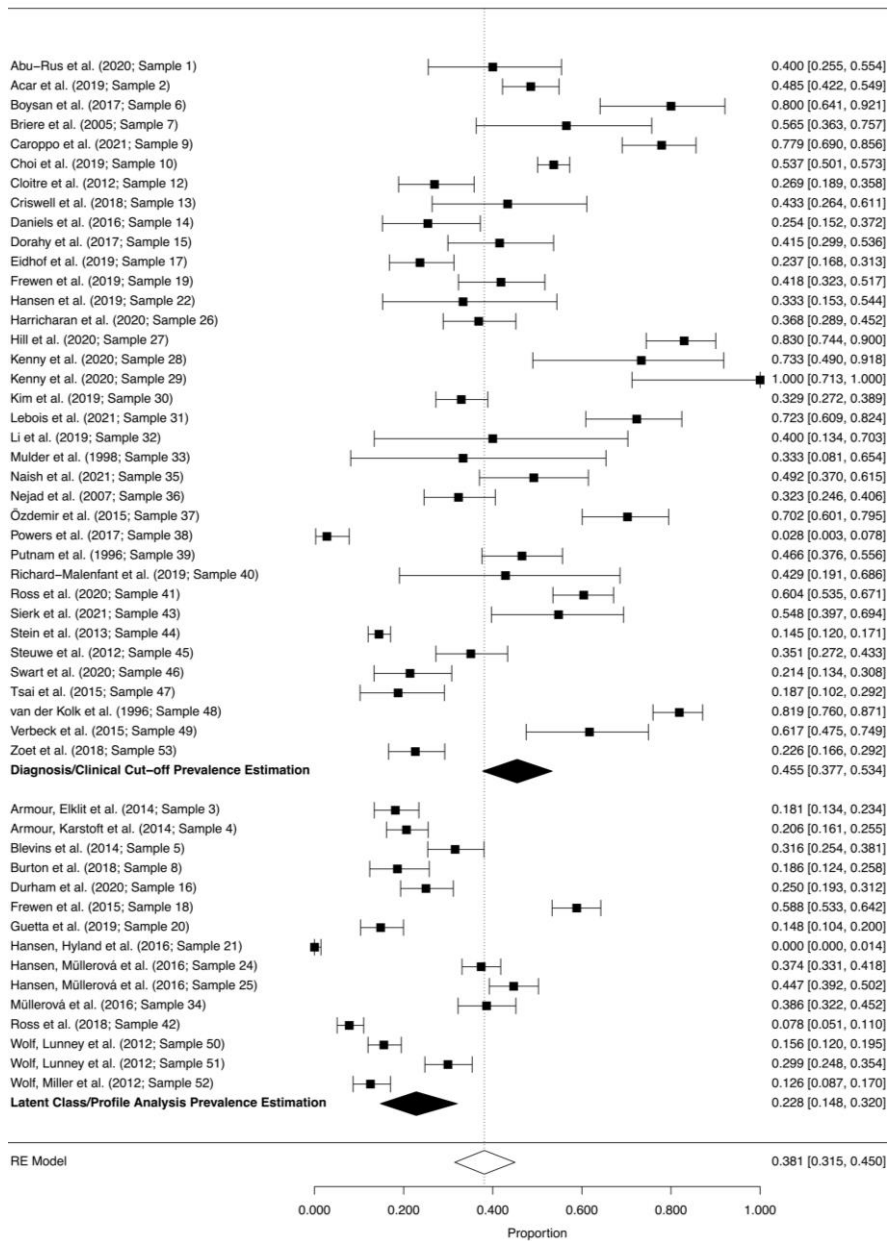


Figure 2
 Forest plot of PTSD-DS prevalence estimates grouped by PTSD-DS assessment method (samples 11 and 23 removed to avoid duplication of population samples).

Table 3

Pooled prevalence of PTSD-DS as a proportion of PTSD for all diagnostic and clinical cut-off samples (i.e., excluding LCA and LPA samples; k = 36)

Meta-analysis subgroup	k	n	Pooled Prevalence (%)	95% CI	Q test	I ²
PTSD DSM criteria used [†] ($\beta = -0.0363$ [95% CI = -0.2065, 0.1338], $p = 0.68$)						
DSM-5	17	1417	48.1	(35.0, 61.3)	288.3*	95.7
DSM-III or DSM-IV	17	2768	44.2	(33.6, 55.1)	623.7*	96.5
Dissociation criteria ($\beta = 0.1135$ [95% CI = -0.0471, 0.2740], $p = 0.17$)						
DSM-5 (Dereal / Depers)	23	3239	41.7	(31.5, 52.2)	622.8*	96.9
Broader dissociation	13	1144	52.9	(42.5, 63.3)	173.2*	91.4
Dissociation measure completion [‡] ($\beta = 0.0479$ [95% CI = -0.1171, 0.2130], $p = 0.57$)						
Self-report	20	2260	47.0	(37.8, 56.3)	233.1*	93.9
Interview	15	2081	42.7	(29.2, 56.8)	576.9*	97.3
Age group [§] ($\beta = 0.2794$ [95% CI = 0.0115, 0.5474], $p = 0.04$)						
Child	4	949	62.9	(50.2, 74.7)	11.4**	82.0
Adult	27	2819	42.1	(33.4, 51.2)	616.6*	95.4
Occupation ($\beta = -0.0574$ [95% CI = -0.3115, 0.1968], $p = 0.66$)						
Military	4	292	40.5	(19.1, 63.9)	49.8*	93.2
Civilian	32	4091	46.1	(37.8, 54.6)	873.7*	96.3
Trauma type ($\beta = 0.1184$ [95% CI = -0.1345, 0.3714], $p = 0.36$)						
Interpersonal only	4	226	55.9	(33.4, 77.2)	41.4*	90.5
Other	32	4157	44.2	(35.6, 52.6)	876.4*	96.3

Note. k = number of samples; n = number of participants; CI = confidence interval; Dereal =

derealisation; Depers = depersonalisation

* $p < 0.0001$, where the degrees of freedom (df) = $k - 1$

** $p < 0.01$, where the degrees of freedom (df) = $k - 1$

† Sample 15 removed as no PTSD DSM criteria reported, sample 26 removed as used both DSM-IV and DSM-5 when assessing for PTSD

‡ Sample 43 removed as a mix of self-report and interview measures were used

§ Several samples were removed due to populations formed of both children and adults, or age group not reported

Supplementary Information

Supplementary Table 1

Quality Assessment Checklist for Prevalence Meta-Analysis

1	Was the study population and index trauma clearly specified and defined?	
	<i>Descriptive statistics were reported on participant demographics (including age range and mean, gender, ethnicity) and frequency of trauma type/nature within the participant pool reported</i>	2
	<i>Some description statistics provided about the sample but some missing information (e.g. authors did not report frequency of trauma type/nature or provide enough information about demographic variables).</i>	1
	<i>No clear description of sample demographics or index trauma characteristics</i>	0
2	Was the participation rate of eligible persons at least 50%?	
	<i>More than 50% of eligible and approached participants took part</i>	2
	<i>Less than 50% of those approached took part, but there was no significant difference in non-response characteristics (such as age, gender) between those who participated and those who did not</i>	1
	<i>Less than 50% of those approached took part, and differences between those who took part and those who did not were not reported or highlighted significant differences. Or, response was not reported</i>	0
3	Was follow up time for PTSD assessment appropriate and meaningful?	
	<i>An appropriate time frame (>4 weeks) since trauma was reported</i>	2
	<i>No information given regarding time frame since trauma. Or, assessment <4 weeks since trauma</i>	0
4	Were objective, standard criteria used for the assessment of Post-Traumatic Stress Disorder?	
	<i>Diagnostic interview or self-report questionnaire shown to demonstrate good levels of validity and reliability in the assessment of PTSD adhering to DSM criteria for PTSD i.e. cluster-based algorithm</i>	2
	<i>Diagnostic interview or self-report questionnaire shown to demonstrate good levels of validity and reliability in the assessment of PTSD adhering to DSM criteria for PTSD using a cut-off score or grouping analysis such as LPA or LCA</i>	1
	<i>Diagnostic interview or self-report without utilising DSM criteria (e.g. not conforming to cluster-based algorithm or cut-off score or grouping analysis). Or poor validity and reliability.</i>	0
5	Were objective, standard criteria used for the assessment of the Dissociative Subtype of Post-Traumatic Stress Disorder?	
	<i>Diagnostic interview or self-report questionnaire shown to demonstrate good levels of validity and reliability, adhering to DSM-5 criteria for PTSD-DS i.e. based on depersonalisation and derealisation only</i>	2
	<i>Diagnostic interview or self-report questionnaire shown to demonstrate good levels of validity and reliability, however not adhering to DSM-5 criteria for PTSD-DS i.e. based on other domains of dissociation outside of just depersonalisation and derealisation</i>	1

<i>Diagnostic interview or self-report questionnaire shown to demonstrate good levels of validity, however domains of dissociation assessed not reported. Or poor validity and reliability</i>	0
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Note. Where 2 = well addressed, 1 = partially addressed, 0 = poorly addressed/not addressed/not reported

This tool was developed by Mr. William White for a meta-analysis undertaken in partial fulfilment of a Doctorate in Clinical Psychology. The development of this tool was based on the Assessment Tool for Observational Cohort and Cross-Sectional Studies (National Heart Lung and Blood Institute, 2014), combining with modified questions from other prevalence and risk factor studies that would be appropriate for use in this review (Hoy et al., 2012; Munn et al., 2014).

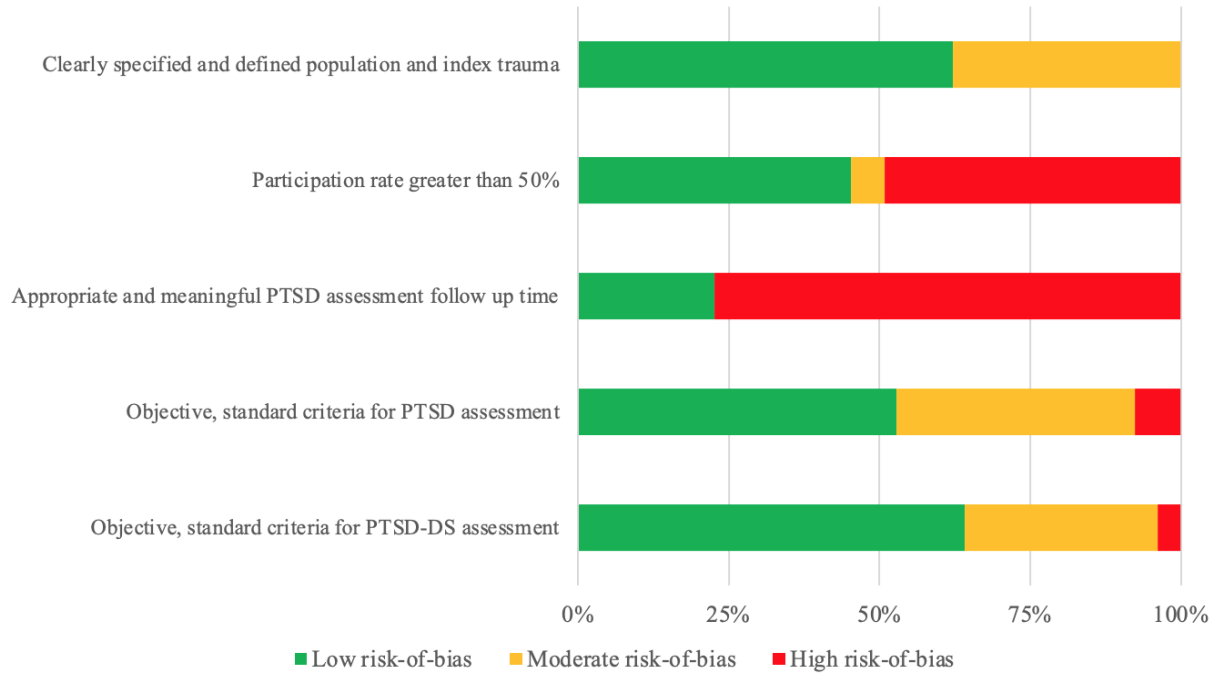
Supplementary Table 2*Sample risk-of-bias scores by individual item and total*

Sample No.	Author	Item 1	Item 2	Item 3	Item 4	Item 5	Total	Quality
1	Abu-Rus et al. (2020)	1	2	0	2	2	7	High
2	Acar et al. (2019)	1	0	0	2	1	4	Low
3	Armour, Elklit et al. (2014)	2	2	2	1	1	8	High
4	Armour, Karstoft et al. (2014)	2	0	0	1	1	4	Low
5	Blevins et al. (2014)	2	0	0	1	2	5	Medium
6	Boysan et al. (2017)	2	0	2	2	2	8	High
7	Briere et al. (2005)	1	2	0	2	1	6	Medium
8	Burton et al. (2018)	2	0	0	1	2	5	Medium
9	Caroppo et al. (2021)	2	0	0	2	0	4	Low
10	Choi et al. (2019)	2	2	0	2	2	8	High
11	Choi et al. (2017)	2	2	0	1	2	7	High
12	Cloitre et al. (2012)	2	0	2	0	1	5	Medium
13	Criswell et al. (2018)	2	0	2	2	2	8	High
14	Daniels et al. (2016)	1	0	0	1	2	4	Low
15	Dorahy et al. (2017)	1	1	0	0	1	3	Low
16	Durham et al. (2020)	2	2	0	1	2	7	High
17	Eidhof et al. (2019)	2	0	0	2	2	6	Medium
18	Frewen et al. (2015)	1	2	0	1	2	6	Medium
19	Frewen et al. (2019)	1	0	0	2	2	5	Medium
20	Guetta et al. (2019)	1	2	0	1	2	6	Medium
21	Hansen, Hyland et al. (2016)	1	2	2	1	2	8	High
22	Hansen et al. (2019)	2	1	2	2	2	9	High
23	Hansen et al. (2019)	2	1	2	1	2	8	High
24	Hansen, Müllerová et al. (2016)	2	2	0	1	2	7	High
25	Hansen, Müllerová et al. (2016)	2	2	0	1	2	7	High
26	Harricharan et al. (2020)	1	0	0	1	2	4	Low
27	Hill et al. (2020)	1	0	0	1	2	4	Low
28	Kenny et al. (2020)	2	2	0	2	2	8	High
29	Kenny et al. (2020)	2	2	0	2	2	8	High
30	Kim et al. (2019)	2	2	0	2	2	8	High
31	Lebois et al. (2021)	1	2	0	2	2	7	High
32	Li et al. (2019)	2	2	0	2	1	7	High
33	Mulder et al. (1998)	2	2	0	2	1	7	High
34	Müllerová et al. (2016)	2	2	0	1	1	6	Medium
35	Naish et al. (2021)	2	0	2	2	2	8	High
36	Nejad et al. (2007)	2	0	0	0	1	3	Low
37	Özdemir et al. (2015)	2	0	0	2	1	5	Medium
38	Powers et al. (2017)	1	2	2	2	2	9	High
39	Putnam et al. (1996)	1	0	0	0	1	2	Low
40	Richard-Malenfant et al. (2019)	1	0	0	2	2	5	Medium
41	Ross et al. (2020)	2	2	0	2	2	8	High
42	Ross et al. (2018)	2	0	0	1	1	4	Low
43	Sierk et al. (2021)	2	2	2	2	1	9	High
44	Stein et al. (2013)	1	0	2	2	2	7	High
45	Steuwe et al. (2012)	1	0	0	2	2	5	Medium
46	Swart et al. (2020)	2	2	0	2	2	8	High
47	Tsai et al. (2015)	2	0	0	2	2	6	Medium
48	van der Kolk et al. (1996)	1	0	0	2	0	3	Low
49	Verbeck et al. (2015)	2	0	0	2	1	5	Medium
50	Wolf, Lunney et al. (2012)	1	0	0	1	1	3	Low
51	Wolf, Lunney et al. (2012)	1	0	2	1	2	6	Medium
52	Wolf, Miller et al. (2012)	2	2	0	1	1	6	Medium
53	Zoet et al. (2018)	2	2	0	1	2	7	High

Note. 0-4 high risk/low quality, 5-6 moderate risk/quality, 7-10 low risk/high quality

Supplementary Figure 1

Proportion of samples rated as a low, moderate or high risk-of-bias for each quality assessment item



Supplementary Table 3

Pooled prevalence of PTSD-DS as a proportion of PTSD for diagnostic/clinical cut-off samples utilising DSM-5 criteria for dissociation (i.e., excluding LCA and LPA samples and those using broader criteria for dissociation; k = 23)

Meta-analysis subgroup	k	n	Pooled Prevalence (%)	95% CI	Q test	I ²
PTSD DSM criteria used [‡] ($\beta = -0.2041$ [95% CI = -0.4406, 0.0324], $p = 0.09$)						
DSM-5	16	1180	48.2	(34.2, 62.3)	285.9*	95.5
DSM-III or DSM-IV	6	1926	28.3	(17.6, 40.3)	289.7*	96.1
Dissociation measure completion ($\beta = 0.1271$ [95% CI = -0.0882, 0.3423], $p = 0.25$)						
Self-report	10	1435	49.4	(32.0, 66.9)	174.5*	97.1
Interview	13	1804	36.4	(24.3, 49.4)	244.3*	96.3
Age group [‡] ($\beta = 0.3444$ [95% CI = 0.0410, 0.6477], $p = 0.03$)						
Child	4	949	62.9	(50.2, 74.7)	11.4**	82.0
Adult	16	1867	36.7	(24.7, 49.6)	376.3*	96.4

Note. k = number of samples; n = number of participants; CI = confidence interval

* $p < 0.0001$, where the degrees of freedom (df) = $k - 1$

** $p < 0.01$, where the degrees of freedom (df) = $k - 1$

† Sample 26 removed as used both DSM-IV and DSM-5 when assessing for PTSD

‡ Several samples were removed due to populations formed of both children and adults, or age group not reported

Supplementary Table 4

Pooled prevalence of PTSD-DS as a proportion of PTSD for all LCA/LPA samples (i.e., excluding diagnostic and clinical cut-off samples; k = 17)

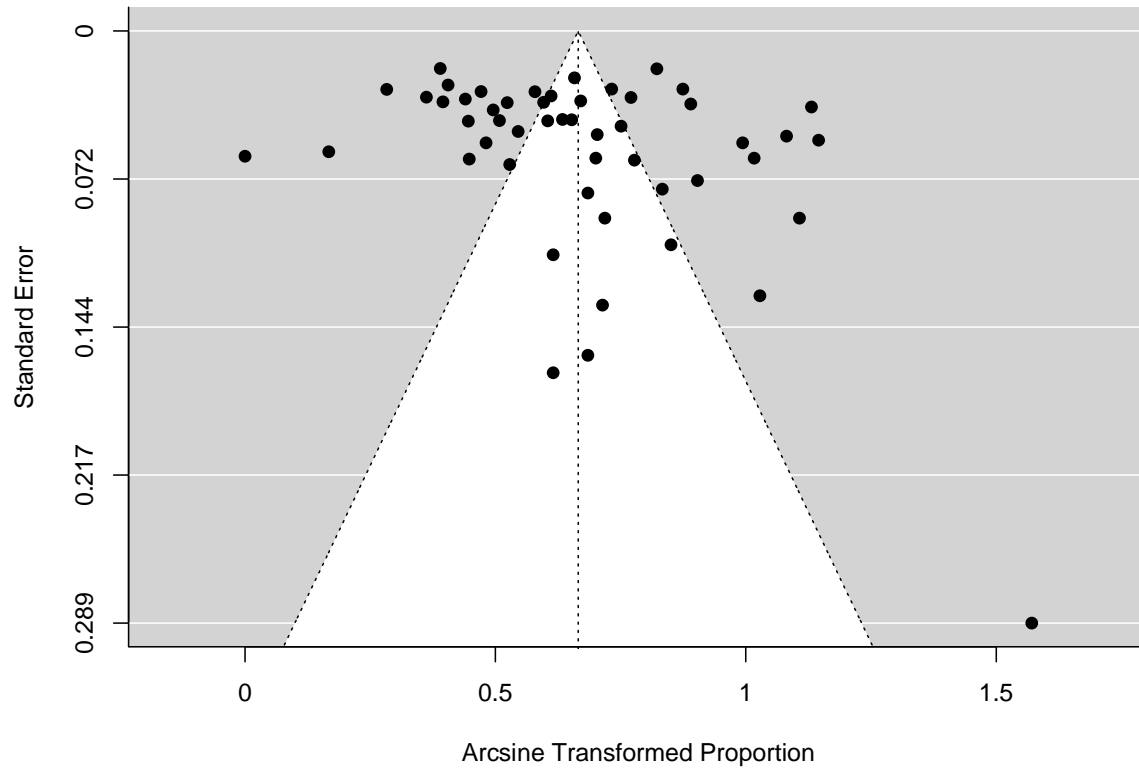
Meta-analysis subgroup	k	n	Pooled Prevalence (%)	95% CI	Q test	I ²
PTSD DSM criteria used ($\beta = -0.0872$ [95% CI = -0.3022, 0.1278], $p = 0.43$)						
DSM-5	8	1750	25.0	(10.9, 42.7)	328.5*	98.6
DSM-III or DSM-IV	9	2850	18.2	(10.1, 28.1)	196.8*	97.4
Dissociation criteria ($\beta = -0.0648$ [95% CI = -0.2912, 0.1616], $p = 0.57$)						
DSM-5 (Dereal / Depers)	11	3503	23.1	(11.3, 37.5)	311.3*	98.8
Broader dissociation	6	1634	18.0	(10.7, 26.7)	83.7*	94.6
Dissociation measure completion ($\beta = 0.0940$ [95% CI = -0.1589, 0.3468], $p = 0.47$)						
Self-report	13	3506	23.1	(12.6, 35.6)	429.9*	98.6
Interview	4	1094	15.9	(12.7, 19.3)	6.7	55.6
Occupation ($\beta = -0.0532$ [95% CI = -0.2918, 0.1853], $p = 0.66$)						
Military	5	1378	18.4	(12.9, 24.6)	32.0*	87.6
Civilian	12	3759	22.5	(11.3, 36.1)	429.7*	98.8

Note. k = number of samples; n = number of participants; CI = confidence interval; Dereal = derealisation; Depers = depersonalisation

* $p < 0.0001$, where the degrees of freedom (df) = $k - 1$

Supplementary Figure 2

Funnel plot to assessing publication bias



Point by Point Responses to Reviewers

Please be advised that all changes can be viewed in “RESPONSE TO REVIEWERS” version of documents.

Comment/feedback	Evidence of where/how covered
<p>A) Abstract (where applicable) must be structured using our standard subheadings only (Background, Methods, Results, Conclusions), not to exceed 250 words (including the 4 subheadings). NB: Review articles only may have an unstructured abstract not exceeding 250 words. Editorials do not require an abstract but may include one an abstract at authors' discretion. Correspondence should be submitted without an abstract.</p>	<p>An unstructured abstract (given the paper is a review) of 248 words, including the keywords, is provided on page 2.</p>
<p>B) Figures, which must be uploaded only as a separate file and not combined with any other element of the submission, should be produced using size 8 point Arial font for the legend. Any wording within a figure should ideally be in Arial - 8 point size is standard, but this may vary depending on space limitations within individual figures. Wherever possible figures for print should be monochrome although colour figures are acceptable for online. You will be asked to pay for unnecessary colour printing. If you wish you may have colour online and black-and-white in print at no charge, in which case you should submit two copies of the figures, identical in every respect other than the colour. Figures should NOT be embedded or included in the main text file.</p>	<p>Both figures 1 and 2 uploaded as separate image files.</p> <p>Text in figure 1 changed to Arial font size 8</p> <p>Figure 2 remains the same, as already Arial font and the appropriate size.</p> <p>Both figure 1 and 2 are monochrome.</p>
<p>C) We are using the APA v6* referencing format, listing all authors up to 7 in number, or the first 6 authors plus the last author, date, article title, journal title in full, volume, page numbers and/or DOI.</p> <p>You need to amend the reference list to our usual style (see instructions for authors and below). A few of them are not quite in the right format for us. I am afraid that if the software will not format them satisfactorily then you will have to edit manually.</p>	<p>In text references, and reference list, edited to conform to APA v6 and Psychological Medicine author guidelines</p>

<p>D) Appendices and supplementary material also should be submitted as a separate file from the main text. These will be published online exactly as they are received, so a clean version without track changes showing, etc, must be submitted. Authors may upload two *clearly labelled* versions of supplementary material, one clean and one showing changes if they feel this appropriate - the tracked version must be listed as 'response to reviewers', the clean one as 'supplementary or appendix'.</p>	<p>Supplementary Information documents provided (one clean, and one with tracked changes)</p>
<p>E) Text files (and tables) should be uploaded in editable form (ie word processor files, not pdf).</p>	<p>All text files with tables uploaded as Microsoft Word files.</p>
<p>F) A clean copy of ALL files will be required prior to final acceptance. If you send a tracked changes copy of any of the files comprising your revised manuscript (useful for editors and reviewers to see where you have made changes), you MUST also send a clean copy. The clean copy should be indicated as the main document/supplementary/table etc etc, with the TRACKED copies as "RESPONSE TO REVIEWERS". You can have more than one file for each designation so it is possible to have both a tracked copy of the manuscript and a letter explaining your changes both indicated as response to reviewers. The tracked copy should NEVER be the main document.</p>	<p>Both a clean copy of the manuscript and Supplementary Information documents have been provided, as well as tracked changes copies (indicated by the file title starting "RESPONSE TO REVIEWERS")</p>
<p>G) You must include in the main document a declaration about any Conflict of Interest. If there is none then please state none.</p>	<p>Conflict of interest statement can be found on page 20 of the main manuscript</p>
<p>H) Point by point responses to reviewers and cover letter to editor (if provided) should be submitted as separate documents and uploaded appropriately as such, not combined. Point by point responses should NEVER be listed as cover letter.</p>	<p>This point by point responses to reviewers document has been uploaded as a separate file, as well as updated cover letter</p>
<p>Reviewer #1: The only thing the authors might consider is explaining latent class and latent profile analyses in a little bit more detail for clinical readers, and the differences between the two, probably on page 4. Also, they could consider adding the full terms for both to the notes at the bottoms of their Tables.</p>	<p>Further explanation of LCA and LPA provided on page 4/5</p> <p>Full terms for LCA and LPA added to the note for each relevant table</p>

<p>Reviewer #1: Why do the authors say on page 17 that PTSD-DS "remains an elusive concept" after many pages of statistics on operationally-defined measures. More elusive than what? Why is it more elusive than PTSD itself? I think this comment should be deleted or else explained better.</p>	<p>This comment has been deleted</p>
<p>Reviewer #1: It's a surprising and counter-intuitive finding that the DP/DR only definition and a broader range of dissociative symptoms yield very similar prevalences - could the authors comment on this a bit more? Does this lead to the conclusion that there is no need to expand the PTSD-DS symptom criteria list? Or should the criteria be broadened because the main consideration is a more accurate and complete clinical description? If the latter, what should be said in the text in future editions of DSM? Some thoughts or suggestions on these questions would be helpful.</p>	<p>Further thoughts and considerations added on pages 17-18</p>
<p>Reviewer #1: Also what do the authors think about having dissociative amnesia in the PTSD symptom criteria but not specified in the PTSD-DS criteria?</p>	<p>The question of whether or not dissociative amnesia (and dissociative flashbacks) should be included in the criteria for PTSD-DS, rather than for PTSD, was not directly focussed on as part of this review. However, evidence suggesting that the current criteria for PTSD-DS is too narrow is already cited on page 4, and the suggestion of Ross (2021) that both dissociative amnesia and flashbacks should be included is additionally commented on page 17-18.</p>
<p>Reviewer #1: Another question is, why is PTSD-DS a subtype? This was based on initial belief ten years back that it accounts for 10%-15% of cases, but if almost 50% of cases meet the criteria, it isn't really a subtype - do the authors have any comments on that?</p>	<p>Again commented on further in "Clinical Implications and Suggestions for Future Research" on page 17.</p>
<p>Reviewer #2: The clinical recommendation that PTSD-DS should be routinely assessed following trauma is important but many clinicians reading this paper would also be interested to consider the symptoms of dissociations that are included in the ICD11 diagnosis of Complex PTSD. Whilst the study focused on DSM5 diagnostic criteria, the discussion of dissociative symptoms in PTSD is lacking</p>	<p>Reference to the ICD11 criteria and dissociation symptoms made in in "Clinical Implications and Suggestions for Future Research" section on page 17, and the Hyland 2020 reference used.</p>

<p>without some reference to symptoms of dissociation in ICD 11 complex PTSD diagnosis and related research (for example Hyland P, Shevlin M, Fyvie C, Cloitre M, Karatzias T. The relationship between ICD-11 PTSD, complex PTSD and dissociative experiences. <i>J Trauma Dissociation</i>. 2020 Jan-Feb;21(1):62-72).</p>	
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Psychological Medicine

Cover letter for resubmission of manuscript for consideration by Psychological Medicine

Ref.: Ms. No. PSM-D-22-00031

Dear Sir or Madam,

We would like to submit our revised manuscript entitled “Prevalence of the Dissociative Subtype of Post-Traumatic Stress Disorder: A Systematic Review and Meta-Analysis” for your consideration, based on the comments and feedback received on 9th May 2022. The protocol for this review was pre-registered on PROSPERO (reference: CRD42021210902) prior to any formal review of searches. This study is the first to meta-analyse the prevalence of the Dissociative Subtype of Post-Traumatic Stress Disorder in both children and adults, following its introduction in the fifth edition of the Diagnostic and Statistical Manual of Mental Disorders. This review raises important issues around the conceptualisation and utility of this subtype.

Please find manuscript and PRISMA 2020 checklist included in this submission.

We look forward to this manuscript being published in Psychological Medicine in due course.

Yours sincerely,

A handwritten signature in black ink, appearing to read 'William White', with a horizontal line underneath.

William White
University of East Anglia