The Prevalence of Depersonalization-Derealization Disorder:

A Systematic Review

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Abstract

Aims: Depersonalization-Derealization disorder (DDD) is a psychiatric condition

characterized by persistent feelings of detachment from one's self and of unreality

about the outside world. This review aims to examine the prevalence of DDD amongst

different populations.

Methods: A systematic review protocol was developed before literature searching.

Original articles were drawn from three electronic databases and included only studies

where prevalence rates of DDD were assessed by standardized diagnostic tools. A

narrative synthesis was conducted.

Results: 23 papers were identified and categorized into three groups of participants:

general population, mixed in/outpatient samples, and patients with specific disorders.

The prevalence rates ranged from 0-1.9% amongst the general population, 5-20% in

outpatients and 17.5-41.9% in inpatients. In studies of patients with specific disorders,

prevalence rates varied: 1.8-5.9% (substance abuse), 3.3-20.2% (anxiety), 3.7-20.4%

(other dissociative disorders), 16.3% (schizophrenia), 17% (borderline personality

disorder), ~50% (depression). The highest rates were found in people who experienced

interpersonal abuse (25-53.8%).

Conclusions: The prevalence rate of DDD is around 1% in the general population,

consistent with previous findings. DDD is more prevalent amongst adolescents and

young adults as well as in patients with mental disorders. There is also a possible

relationship between interpersonal abuse and DDD, which merits further research.

Key words: Dissociative disorders; epidemiology; systematic review

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Introduction

Depersonalization (DP) and derealization (DR) are symptoms characterized by, respectively, feelings of unreality and detachment from one's self and one's surroundings (American Psychological Association, 2013). Depersonalization and derealization (DP/DR) symptoms can occur as transient experiences in otherwise healthy individuals at times of stress or physical exhaustion and have been used as terms to describe the phenomenon of "burnout" (Maslach & Jackson, 1981). On the other hand,

Depersonalization-Derealization Disorder (DDD) occurs when this symptom cluster is persistent and distressing and is paired with functional impairment, rendering it a clinical diagnosis alongside other mental disorders or as a primary condition (American Psychological Association, 2013).

There is limited literature regarding the prevalence rate of DDD. A recent review focusing on the epidemiology of DP/DR both as a symptom and as a disorder was conducted by Hunter et al. (2004), which included relevant papers published between 1966 to October 2002. According to this review, the prevalence rate of DDD was 1-2% in community samples when using interviews as the diagnostic tool, while transient symptoms of DP/DR were more prevalent in the general population with lifetime rates of 26-74%.

Prevalence rates likely vary due to inconsistent definitions of DP/DR as a symptom or as a clinically significant disorder, paired with the use of a range of diagnostic tools. In order to evaluate as many studies as possible that contained some data on the prevalence of DP/DR, Hunter et al. (2004) did not set strict exclusion criteria for the quality of the studies under review. As a result, the previous review may have included some studies of lower quality, potentially influencing the strength of the evidence. Therefore, this review

aims to update the previous work, adopting a more systematic approach following PRISMA guidelines.

Two clinical interviews are commonly used when making a DDD diagnosis in clinical practice: the Structured Clinical Interview for DSM-IV Dissociative Disorders (SCID-D; Steinberg, 1994) and the Dissociative Disorders Interview Schedule (DDIS; Ross et al., 1989). The SCID-D is a semi-structured interview (Steinberg, 1994) and the DDIS is a clinician-administered structured interview (Ross et al., 1989). Both are used to identify dissociative disorders according to the DSM-IV (Ross et al., 1989; Steinberg, 1994). In order to capture all potentially useful data, studies using either one of these interviews were included, as well as studies that incorporated a standardized scale with a clinical cut-off score. The Cambridge Depersonalization Scale (CDS; Sierra & Berrios, 2000) and the depersonalization subscale of the Dissociative Experiences Scale (DES; Bernstein & Putnam, 1986) are two of the most frequently used standardized diagnostic scales of DDD that include a clinical cut-off score. Simeon et al. (1998) also suggest that the taxon version of the DES, consisting of 8 items regarding pathological dissociation in the DES (Waller et al., 1996), could be more useful than the mean DES score when detecting DDD (Simeon et al., 1998).

Our systematic review aims to describe the prevalence rates of DDD in a range of populations, updating previous findings. To address this, we have reviewed quantitative studies published since October 2002 that provide relevant information about the prevalence rates of DDD and conducted a narrative synthesis to explore the findings from the selected studies.

Method

A systematic review protocol in PROSPERO format was developed and the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines were followed. The review protocol is shown in *Appendix*.

Search strategy

The search was limited to studies published in English after October 2002 but geographical locations were not limited. Three electronic databases were initially searched and screened in March 2020: PsycINFO, MEDLINE and Web of Science. Grey literature was also screened in Google Search. The results from these sources were combined with duplications removed. The search process was conducted by two independent reviewers to minimise error and repeated in February 2021 to include any eligible papers published since March 2020. The original search history is shown in *Appendix*.

Search terms

When defining the search terms, two main concepts were identified: 1, "depersonalization/ derealization", and 2, "prevalence". For each concept, the Boolean operator "OR" was used to group all of the search terms. Both text words (free-text searches) and relevant subject terms (MeSH terms) were used as search terms and the results were combined within each concept. Concept 1 (depersonalization, derealization, depersonalization disorder, derealization disorder) and concept 2 (epidemiolog*, prevalen*, occurrence, frequency) were combined using the Boolean operator "AND". To cover all potential literature, we used truncation and searched both UK and US spellings.

Inclusion/Exclusion criteria

We only included quantitative studies that provided or allowed for the calculation of prevalence rates. Reviews were excluded although reference lists were manually screened to retrieve any other relevant studies.

Studies were excluded if: (a) they focused only on the relationship between DDD and its risk factors; (b) they only provided prevalence rates of other mental disorders or DP/DR symptoms; (c) they provided prevalence rates of DDD without use of a diagnostic interview or a standardized measure and clinical cut-off scores; (d) they focused on burnout or burnout syndrome rather than DDD.

The standardized clinical interviews for depersonalization include the SCID-D (Steinberg, 1994) and the DDIS (Ross et al., 1989). Clinically significant DDD can also be indicated by a cut-off score of ≥ 70 in the CDS (Sierra & Berrios, 2000) or by a sub-scale score of ≥ 30 in the Derealization/Depersonalization sub-scale of the DES (Bernstein & Putnam, 1986). Studies using the CDS-2, with a cut-off score of 3(Michal, Zwerenz, et al., 2010; Sierra & Berrios, 2000), the CDS-9, with a cut-off score of 19 (Michal et al., 2004; Sierra & Berrios, 2000), or the taxon version of the DES, with a cut-off score of 13 (Bernstein & Putnam, 1986) were also eligible for inclusion. Any translated versions of the above interviews and scales that were validated and provided a cut-off score were also accepted. Studies using other scales were excluded.

Procedure

Two independent reviewers conducted study selection, quality assessment and data extraction process. In each phase, any discrepancies between the two reviewers were resolved through discussion with a third reviewer.

After removing duplicates in the initial database searches, the two independent

reviewers screened titles and abstracts of the literature against inclusion criteria and then retrieved full texts of potential studies to assess their eligibility. Reasons for exclusion were recorded.

Then, two reviewers assessed the quality of eligible studies independently, using an adapted version of the Quality assessment checklist for prevalence studies (Nguyen et al., 2016). The adapted checklist consists of nine items that assess the risk of bias in nine domains with one summary item indicating the overall risk of bias (Nguyen et al., 2016). It was selected as it provides more specific criteria regarding the level of risk of bias than the original checklist (Hoy et al., 2012). A score of zero in each item indicated low risk and 1 indicated high risk. The overall risk of bias was indicated by the total score (Low risk: 0-3; Moderate risk: 4-6; High risk: 7-9). Before merging the results from the reviewers, the interrater reliability was calculated by an intraclass correlation coefficient.

Only papers at low or moderate risk of bias were included in the data extraction process, and a standardized form adapted from the Cochrane Data collection form template was used (Higgins, 2008). A narrative synthesis was conducted to explore the findings from the included studies.

Results

Study selection

In total, 1,786 papers were identified in the initial search, with 1,151 remaining after removing duplicates. The PRISMA flow diagram for a summary of the selection process is shown in *Fig. 1*. 67 potential papers were identified and the full texts were assessed for eligibility. In the study by Baker et al. (2003), all participants were DDD patients recruited from a specialist clinic, so this paper was excluded in our review. Additionally, three eligible

papers ,(Foote et al., 2008; Michal, Wiltink, Till, Wild, Münzel, et al., 2010; Tschan et al., 2013) were excluded as the same samples were used in three other included papers (Foote et al., 2006; Michal et al., 2009; Michal, Wiltink, Till, Wild, Blettner, et al., 2010). When screening full texts, we excluded one paper (Duffy, 2000) due to the discrepancy between the publication year of this paper presented in the database (2002) and presented in the paper (2000). When repeating the search process in February 2021, we found one eligible paper (Schlax et al., 2020) published since March 2020. Thus, 23 papers were identified and included in the following analysis.

Study characteristics

The range of publication dates was from 2006 to 2020. There was a good international distribution of the studies including Turkey (n=5), Germany (n=4) and the United States (n=3). Other studies were from Canada (n=1), Israel (n=1), Mexico (n=1), Northern Ireland (n=1), Puerto Rico (n=1), Serbia (n=1), Spain (n=2), Switzerland (n=1), and the United Kingdom (n=1). There was also one transcultural study (Sierra et al., 2006).

Most studies used structured or semi-structured interviews to obtain a diagnosis of DDD, such as the DDIS (n=4) or the SCID-D (n=8). Ten papers used the CDS (n=6), the CDS-2 (n=3) or the CDS-9 (n=1). Only one study used the Derealization/Depersonalization sub-scale of the DES.

Sample characteristics

Sample sizes ranged from 20 to 13,182. In most of the studies, the mean age of participants ranged from 30 to 50 years (n=17) and the proportion of female participants was above 50% (n=18). Two studies did not report mean age of the sample (Gonzalez-Torres et al., 2010; Mueller-Pfeiffer et al., 2012) and two studies did not give the percentage of females (Gonzalez-Torres et al., 2010; Michal, Wiltink, Till, Wild, Blettner, et al., 2010).

Nine papers assessed the prevalence rate of DDD amongst the general population. Five studies were conducted amongst patients with mixed or non-specified disorders, with participants being outpatients (n=2), inpatients (n=1) and mixed or unspecified patients (n=2). 12 papers evaluated the prevalence of DDD amongst patients with specific disorders or conditions, including anxiety disorders (n=4), dissociative spectrum disorders (n=2), substance use disorders (n=2), interpersonal abuse (n=2), borderline personality disorder (n=1), schizophrenia or schizophrenia spectrum disorders (n=1) and depression (n=1). However, there are three studies that involved two or three specific groups (Aponte-Soto et al., 2019; Gonzalez-Torres et al., 2010; Somer et al., 2015).

Quality assessment

Most of the studies were at low risk (n=15) and the remaining papers showed moderate risk (n=8). We therefore included all 23 papers in our analysis. The intraclass correlation coefficient was 0.947, indicating excellent inter-rater reliability.

Results: General population

Table 1 presents the study characteristics and results of nine studies conducted amongst the general population.

The prevalence rates of DDD were similar across five studies (Gonzalez-Torres et al., 2010; Johnson et al., 2006; Michal et al., 2009; Michal, Wiltink, Till, Wild, Blettner, et al., 2010; Schlax et al., 2020), ranging from 0.76% (Schlax et al., 2020) to 1.9% (Michal et al., 2009), even with sample sizes varying from 172 (Gonzalez-Torres et al., 2010) to 13,182 (Schlax et al., 2020). There were a few outliers. Aponte-Soto et al. (2019) found that the prevalence rate was 0% among a sample of 40 adults. Beyond this, the prevalence rate was 9.7% amongst a community adult sample in Israel (Somer et al., 2015), inconsistent with the

general trend of ~1% rates. Additionally, in two studies conducted in adolescents (Michal et al., 2015) and undergraduate students (Myers & Llera, 2020), the prevalence rates were 11.9% and 11%, respectively. It should be noticed that there are potential overlaps between the participants in study by Michal, Wiltink, Till, Wild, Blettner, et al. (2010) and study by Schlax et al. (2020), as they both investigated participants from the Gutenberg Health Study.

Results: Patients with non-specific or mixed disorders

Table 2 presents the study characteristics and results of five studies conducted among patients with unspecified or mixed disorders.

Outpatients

Two studies reported DDD prevalence rates among adult outpatients with unspecified or mixed disorders, varying from 5% (Foote et al., 2006) to 20% (Dorahy et al., 2006). The difference between the prevalence rates could be due to the small sample size of 20 patients in the Dorahy et al. (2006) study and, although these were outpatients, they were described as complex in presentation.

Inpatients

In a transcultural study (Sierra et al., 2006), participants were psychiatric inpatients from three countries: United Kingdom (n=31), Spain (n=68), and Colombia (n=41), assessed using the CDS. Reported prevalence rates were 41.9%, 35.8% and 17.5%, respectively.

Mixed or unspecified patients

Similar prevalence rates of DDD were found in mixed or unspecified patients: 6% in Mexico (Garcia et al., 2006) and 4.4% in Switzerland (Mueller-Pfeiffer et al., 2012).

Results: Patients with specific disorders

See *Table 3*. According to the nature of the disorders or conditions, similar studies were grouped together.

Anxiety disorders

Four studies provided DDD prevalence amongst patients with anxiety disorders: patients with panic disorder (Mendoza et al., 2011; Ural et al., 2015), patients with obsessive-compulsive disorder (Belli et al., 2012) and patients with social anxiety disorder (Belli et al., 2017).

Three studies were conducted in Turkey using the SCID-D (Belli et al., 2012; Belli et al., 2017; Ural et al., 2015) and one was conducted in Spain using the CDS (Mendoza et al., 2011). The prevalence rates ranged from 3.3% (Ural et al., 2015) to 20.2% (Mendoza et al., 2011). In those Turkish studies, the prevalence rates were 3.3% in panic disorder (Ural et al., 2015), 6.3% in social anxiety disorder (Belli et al., 2017), and 10.3% in obsessive-compulsive disorder (Belli et al., 2012). We included patients diagnosed with DDD as well as other dissociative disorders (i.e. dissociative amnesia + depersonalization) when calculating the prevalence in the two studies (Belli et al., 2017; Belli et al., 2012). Ural et al. (2015) reported the number of patients with DDD only, hence the prevalence rate was likely to be underestimated. However, the proportions of patients only diagnosed with DDD were relatively consistent across those three studies: 3.3% (Ural et al., 2015), 5.3% (Belli et al., 2017), and 3.84% (Belli et al., 2012). For panic disorders specifically, the prevalence rate is much higher in the Spanish population (20.2%; Mendoza et al., 2011) than the Turkish population (3.3%; Ural et al., 2015).

Dissociative spectrum disorders

There were two studies conducted among patients with dissociative spectrum disorders. However, there was a large discrepancy in the prevalence of DDD between these two populations, at 3.7% in patients with conversion disorder (Yayla et al., 2015) and 20.4% in patients with non-epileptic attack disorder (Mitchell et al., 2012).

Substance use disorders

Two studies explored the prevalence of DDD in patients with substance use disorders. The prevalence rates were 1.8% in the Turkish sample with alcohol dependency (Evren et al., 2005), and 5.9% in the Israeli sample with opiate use disorder (Somer et al., 2015). Most of the patients in these two studies were males, with only 5.9% females in both studies.

Interpersonal abuse

The prevalence of DDD assessed by the DDIS was 53.8% in women who had experienced domestic violence (Somer et al., 2015). Aponte-Soto et al. (2019) found a lower prevalence rate of 25% using the CDS among a sample of 40 adults with a history of interpersonal abuse.

Other specific disorders

Three studies examined the prevalence of DDD in other specific disorders. Gonzalez-Torres et al. (2010) found prevalence rates of DDD to be 16.3% in Spanish inpatients with schizophrenia or schizophrenia spectrum disorders with 1.4% of their first-degree relatives also meeting diagnostic criteria for DDD. Another study in Serbia (Žikić et al., 2009) found the prevalence of DDD was 47.6% in people with depression, and a study of 21 Canadian outpatients with borderline personality disorder (Korzekwa et al., 2009) detected a prevalence rate of 19%, assessed by the SCID-D.

Results: prevalence assessment tools

We examined prevalence rates of DDD in each study according to the type of assessment tools used (self-reported scales versus diagnostic interviews). See *Table 4*

Discussion

A systematic review based on the prespecified protocol in PROSPERO format and PRISMA guidelines was conducted to examine the prevalence rates of DDD. 23 studies dating from October 2002 to February 2021 were identified. The studies were mainly conducted amongst three types of populations (although some incorporated more than one type) including the general population, patients with non-specific or mixed disorders, and patients with specific disorders. The prevalence rate amongst the general population was relatively consistent across five of nine studies, with an estimate of around 1% (0.76-1.9%; Aponte-Soto et al., 2019; Schlax et al., 2020). The findings amongst patients with specific disorders or unspecified disorders were mixed, however, it is clear that DDD is more prevalent in patients with mental health conditions, as compared to the general population. For the studies conducted amongst the patients with unspecified or mixed disorders, the prevalence rates ranged from 4.4% (Mueller-Pfeiffer et al., 2012) to 41.9% (Sierra et al., 2006). In those patients with specific disorders, prevalence rates varied from 1.8% (Evren et al., 2005) to 53.8% (Somer et al., 2015).

Compared with the review by Hunter et al. (2004), this review only included studies focusing on the prevalence rates of clinical DDD. One strength of the current study is that a review protocol was developed prior to beginning the literature search, and papers were screened based on the specific inclusion and exclusion criteria. Additionally, only high or moderate quality papers using standardized assessment tools or scales with high reliability and validity were included.

Prevalence assessment tools and prevalence rates

Around half of the studies used structured clinical interviews (e.g. DDIS, SCID-D) to diagnose the participants. It is worth noting that the overall prevalence rates of DDD were higher in studies diagnosing participants by self-report scales, which tends to be the case across disorders. In this case, using structured clinical interviews to diagnose participants likely provided more reliable results.

General population

Our findings show that DDD is more prevalent in the younger population, with a prevalence of around 11%. Consistent with these findings, existing evidence indicates that dissociative symptoms are more prevalent in adolescents (Carlson & Putnam, 1993). High levels of anxiety in the mid-teens (Abe & Suzuki, 1986) could be a potential factor when explaining the higher prevalence of DDD in the younger population. Existing literature also suggests that there is a significant association between anxiety and depersonalization in the general population (Trueman, 1984), and across a range of emotions, anxiety was found to be the strongest predictor of depersonalization (Simeon et al., 2003).

When comparing these findings to the results by Hunter et al. (2004), there is a general consensus between the prevalence rates of around 1% of DDD amongst community samples, suggesting that the prevalence rate of DDD is consistent over time in the general population.

Patients with mixed disorders

The prevalence of DDD in patients ranges from 5-20% in outpatient samples and 17.5%-41.9% in inpatient samples, suggesting that DDD is more common in patients with more severe mental health conditions.

Patients with specific disorders

Anxiety disorders

In patients with panic disorders, although we observed a discrepancy in the prevalence rates of DDD across the studies, the prevalence rate of 20% in Spanish patients with panic disorder (Mendoza et al., 2011) is similar to the previous findings (Hunter et al., 2004), which was 24% (Segui et al., 2000).

Dissociative spectrum disorders

The wide range of prevalence rate of DDD in patients with other dissociative spectrum disorders may be accounted for by the variations in diagnostic measures used, for example, Yalya et al. (2015) used the SCID-D while Mitchell et al. (2012) used the self-reported scale. In this case, the high prevalence of 20% in the study by Mitchell et al. (2012) could be caused by patient overestimation of their symptoms.

The types of dissociation found in conversion disorder could be another possible factor for the low prevalence of DDD in patients with conversion disorder (Yayla et al., 2015). According to Holmes et al. (2005), detachment (disconnection from the world and/or a sense of self) and compartmentalisation (where normally integrated cognitive or physical functions are disconnected) are two qualitatively different categories of dissociation. In this case, conversion disorder, which is characterized by compartmentalisation, differs from DDD, characterized by detachment.

Interpersonal abuse

There is consistent evidence that DDD is prevalent in those who have experienced interpersonal abuse (25-53.8%; Aponte-Soto et al., 2019; Somer et al., 2015). As the interpersonal abuse experience is unlikely to be the consequence of the DDD, one can

speculate that interpersonal abuse is a risk factor for DDD. Along these lines, Simeon et al. (2001) found that childhood interpersonal abuse is a strong predictor for the diagnosis of DDD.

Other specific disorders

The prevalence of DDD in individuals with schizophrenia spectrum disorders is consistent with the rates found in the review by Hunter et al. (2004): 6.9% and 11.1% when using clinical diagnostic measures. In patients with depression, the finding of DDD in 47.6% of the population (Žikić et al., 2009) is similar to previous results indicating a prevalence of 60% among 25 depressed inpatients using a self-report questionnaire (Hunter et al., 2004; Noyes & Kletti, 1977). However, Zikic et al. (2009) assessed DDD prevalence using the CDS and some of these items overlap with depressive symptoms, which may lead to overestimation of DDD.

Limitations & Future directions

This review has some limitations. Firstly, we did not measure publication bias in our review. As negative findings, (i.e. studies that did not detect participants diagnosed with DDD) are less likely to be published, the prevalence of DDD could be overestimated. Secondly, considering the high heterogeneity of included studies (i.e. the variability of the study population), we were unable to conduct a meta-analysis. The small number of included studies is another limitation. For instance, we only found one paper involving patients with depression, thus the results could be unrepresentative. Beyond this, although we have assessed the quality of the included papers, the potential selection bias and response bias still existed. Specifically, according to the quality assessment results, almost half of the included studies were at risk of response bias. In addition, all of the included studies were published in English and there was a lack of studies from Australia, Southeast Asia, Central America, and

African countries. Therefore, future research could focus on the prevalence of DDD in those settings and explore whether different cultural backgrounds have an impact. It should be noted that our review only provided descriptive epidemiological information about the frequency of DDD. Therefore, although we establish an association between DDD and potential risk factors, we still cannot infer any causal relationships between them. Cross sectional studies, case control studies and cohort studies are needed in the future to examine the relationship between interpersonal abuse and DDD.

Conclusion

This review summarises the results of epidemiological studies assessing the prevalence of DDD, providing an update to our knowledge of the prevalence of DDD amongst different populations. The included studies were from a range of countries, allowing for a broader understanding of the prevalence rate of DDD across the globe. Overall, results indicate that the prevalence rate of DDD ranges from 1-2% in the general population, remaining consistent with the previous review (Hunter et al., 2004). We also find a trend that DDD is more prevalent amongst adolescents and young adults than the general adult population. Although the prevalence rates amongst patients with unspecified or specific disorders varies, it remains consistent that DDD is more prevalent in patients with mental health conditions than in the general population, suggesting that patients who already have other diagnoses are more vulnerable to DDD.

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Conflicts of interest

No potential competing interest was reported by the authors.

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Appendix

Appendix 1: Systematic review protocol in PROSPERO format

Review Title

The prevalence of depersonalization and derealization (DP/DR) disorders: a systematic review

Review question

What are the prevalence rates of DP/DR disorders in different population (e.g. non-clinical population, patients with mixed or unspecified disorders, and patients within specific disorders)?

Searches

Databases

The following databases will be searched and reviewed:

MEDLINE

PsycINFO

Web of science.

These databases will be combined and searched. Grey literature will also be searched in Google Search to ensure that other relevant literature not retrieved in electronic databases can be covered.

Before the final analysis, the search process will be repeated to include new published literature and avoid any potential mistakes in first search.

Search terms

Concept 1: depersonalization OR derealization OR depersonalisation OR derealisation OR

Depersonalization/Derealization Disorder OR Depersonalisation disorder OR Depersonalization

Disorder OR derealization disorder OR derealisation disorder

Concept 2: epidemiolog* OR prevalen* OR occurrence OR frequency

Both UK and US spellings will be searched. Within each concept, the Boolean operator "OR" will be used to group all the search terms within the corresponding concept, and the Boolean operator "AND" will be used to combine two concepts.

Limitation

All searches will be limited to humans and English language. The results will be limited to studies published after October 2002. The deduplication process will be conducted after finishing the search.

Types of study to be included

We will only include quantitative studies that provide prevalence rates of DP/DR disorders or provide information to calculate prevalence rates. Reviews will be excluded but the reference lists of reviews will be manually checked to retrieve relevant studies.

Condition or domain being studied

Inclusion: Eligible studies must focus on the prevalence rate of DP/DR disorders, which is defined by structured clinical interviews using DSM-IV, DSM-V or ICD-10 criteria, such as the Structured Clinical Interview for DSM-IV Dissociative Disorders (SCID-D; Steinberg, 1994) and the Dissociative Disorders Interview Schedule (DDIS; Ross et al., 1989). Additionally, studies that used Cambridge Depersonalisation Scale (CDS; Sierra &Berrios, 2000) and taxon version of Dissociation Experiences Scale (DES; Bernstein and Putnam, 1986) and provided clinical cut-off scores (70 for CDS and 13 for taxon DES) will also be included. Studies using the 2-item version CDS (CDS-2; Sierra and Berrios, 2000; Michal et al., 2010) with a cut-off of 3 and the 9-item version CDS with a cut-off of 19 (CDS-9; Sierra and Berrios, 2000; Michal et al., 2004) are also accepted. The outcome of eligible studies should include the prevalence rate of DP/DR disorders.

Exclusion: Studies that focus on the relationship between DP/DR disorders and risk factors will be excluded. Studies that only provide prevalence rates of other mental disorders will be excluded. Studies that only give prevalence of symptoms of DP/DR without a standardized

measures and clinical cut-off will be excluded. Studies that focus on burnout or burnout syndrome

will also be excluded.

Participants/population

Inclusion: Both clinical and non-clinical population will be included.

Exclusion: N/A

Intervention(s)

N/A

Comparator(s)/control

N/A

Context

Geographical locations are not limited.

Main outcome(s)

The main outcome is the prevalence rates of DP/DR disorders. Studies that do not provide

prevalence or the information that can be used to calculate prevalence will be excluded. For studies

that only provide information to calculate prevalence rates, prevalence will be calculated by

dividing the number of people identified as having DP/DR disorders by the total sample size.

Secondary outcome(s)

None.

Data extraction (selection and coding)

The search results will be deduplicated, downloaded and imported to Endnote 7 for storing

and screening. Two independent reviewers will screen the titles and abstracts of the identified papers

30

against the pre-specified criteria. Reviewers will record the titles of the excluded studies and reasons of excluding them. Then, these reviewers will retrieve the full texts of the remained studies and assess their eligibility using the criteria. The disagreements about inclusion will be resolved by discussion with a third reviewer.

Data extraction will be conducted by the reviewers using a standardized form, which is adapted from Cochrane Data collection form template (Higgins et al., 2019). Extracted information included authors, year of publication, country, study population, sample size, mean age (SD) or age range of the study sample, prevalence assessment methods and cut-off score, and relevant results of the study (prevalence rates of DDD). Any discrepancies between two reviewers will be resolved through discussion with a third reviewer.

Risk of bias (quality) assessment

To assess the quality of prevalence studies, Quality assessment checklist for prevalence studies (adapted from Hoy et al.) will be used. The checklist includes nine domains (items 1 to 9) and the scores of these items will be combined to generate an overall score to provide an overall assessment of the study quality (item 10). In each domain, reviewer will provide information about whether the risk of bias is low or high. A score of zero in each item indicates low risk in corresponding domain, while a score of one indicates high risk. A low overall risk will be indicated by an overall score between zero to three, while a high overall risk will be indicated by an overall score between seven to nine. An overall score between four to six indicates a moderate overall risk of the study.

An intraclass correlation coefficient will be calculated to indicate the inter-rater reliability of risk of bias assessment results. Any disagreement between reviewers will be resolved by a discussion with a third reviewer. The final result of risk of bias will be presented in an independent table.

Strategy for data synthesis

We will conduct a narrative analysis to provide findings from selected studies. The study population characteristics (e.g. age, sample size, gender), settings, prevalence assessment methods and prevalence rates will be synthesized.

Analysis of subgroups or subsets

Subgroup analysis will be conducted to explore the prevalence rates among non-clinical population, clinical population with unspecified or mixed disorders and clinical population within specific disorders.

Type and method of review

Systematic review

Anticipated or actual start date

15/03/2020

Anticipated completion date

01/08/2020

Funding sources/sponsors

University College London, Division of Psychiatry

Conflicts of interest

None known

Language

English

Country

England

Stage of review

Review completed

Appendix 2: Original search history

Database: APA PsycInfo <1806 to March Week 4 2020>	
Search Strategy:	

1 (depersonalization or derealization or depersonalisation or derealisation or Depersonalisation disorder or Depersonalization Disorder or derealization disorder).mp. [mp=title, abstract, heading word, table of contents, key concepts,

original title, tests & measures, mesh] (4368)

2 exp Depersonalization/ or exp "Depersonalization/Derealization Disorder"/ (933)

3 1 or 2 (4368)

4 (epidemiolog* or prevalen* or occurrence or frequency).mp. [mp=title, abstract, heading word, table of contents, key concepts, original title, tests & measures, mesh] (409186)

5 3 and 4 (596)

6 limit 5 to (human and english language and yr="2002 -Current") (348)

Database: Ovid MEDLINE(R) <1946 to March Week 3 2020> Search Strategy:

1 (depersonalization or derealization or depersonalisation or derealisation or Depersonalisation disorder or Depersonalization Disorder or derealization disorder or

derealisation disorder).mp. [mp=title, abstract, original title, name of substance word, subject heading word, floating sub-heading word, keyword heading word, organism supplementary concept word, protocol supplementary concept word, rare disease supplementary concept word, unique identifier, synonyms] (3020)

word, unique identifier, synonyms] (3020)

2 Depersonalization/ (1547)

3 1 or 2 (3020)

- 4 (epidemiolog* or prevalen* or occurrence or frequency).mp. [mp=title, abstract, original title, name of substance word, subject heading word, floating sub-heading word, keyword heading word, organism supplementary concept word, protocol supplementary concept word, rare disease supplementary concept word, unique identifier, synonyms] (2912240)
- 5 3 and 4 (971)
- 6 limit 5 to (english language and humans and yr="2002 -Current") (664)

Web of Science core collection

		Search History			
Se t	Results	Save History / Create AlertOpen Saved History	Edit Sets	Combine Sets AND OR Combine	Delete Sets Select All Delete

774 #2 AND #1 Edit 3 *Indexes=SCI-EXPANDED, SSCI,* A&HCI, CPCI-S, CPCI-SSH, BKCI-S, BKCI-SSH, ESCI, CCR-EXPANDED, IC *Timespan*=2002-2020 # **3,769** TS= (depersonalization OR derealization **Edit** 2 OR depersonalisation OR derealisation OR Depersonalization/Derealization Disorder OR Depersonalisation disorder OR Depersonalization Disorder OR derealization disorder OR derealisation disorder) *Indexes=SCI-EXPANDED, SSCI,* A&HCI, CPCI-S, CPCI-SSH, BKCI-S, BKCI-SSH, ESCI, CCR-EXPANDED, IC *Timespan=All years* 3,974,51 TS =(epidemiolog* OR prevalen* OR **Edit** occurrence OR frequency) *Indexes=SCI-EXPANDED, SSCI,* A&HCI, CPCI-S, CPCI-SSH, BKCI-S, BKCI-SSH, ESCI, CCR-EXPANDED, IC Timespan=All years

Appendix 3: Quality assessment tool (Hoy et al., 2012; Nguyen et al., 2016)

Jan.	r of publication:		
i ea	rol publication:		
Stud	ly title:		
Risk	c of bias items	Risk of bias levels	Points scored
1.	Was the study's target population a close representation of the national population in relation to relevant variables, e.g. age, sex, occupation?	Yes (LOW RISK): The study's target population was a close representation of the national population.	0
	1	No (HIGH RISK): The study's target population was clearly NOT representative of the national population.	1
2.	Was the sampling frame a true or close representation of the target	Yes (LOW RISK): The sampling frame was a true or close representation of the target population.	0
	population?	No (HIGH RISK): The sampling frame was NOT a true or close representation of the target population.	1
3.	Was some form of random selection used to select the sample, OR, was a census undertaken?	Yes (LOW RISK): A census was undertaken, OR, some form of random selection was used to select the sample (e.g. simple random sampling, stratified random sampling, cluster sampling, systematic sampling).	0
		No (HIGH RISK): A census was NOT undertaken, AND some form of random selection was NOT used to select the sample.	1
4.	Was the likelihood of non-response bias minimal?	Yes (LOW RISK): The response rate for the study was ≥75%, OR, an analysis was performed that showed no significant difference in relevant demographic characteristics between responders and non-responders	0
		No (HIGH RISK): The response rate was <75%, and if any analysis comparing responders and non-responders was done, it showed a significant difference in relevant demographic characteristics between responders and non-responders	1
5.	Were data collected directly from the	Yes (LOW RISK): All data were collected directly from the subjects.	0
	subjects (as opposed to a proxy)?	No (HIGH RISK): In some instances, data were collected from a proxy.	1
ó.	Was an acceptable case definition used in the study?	Yes (LOW RISK): An acceptable case definition was used.	0
7.	Was the study instrument that	No (HIGH RISK): An acceptable case definition was NOT used Yes (LOW RISK): The study instrument had been shown to have	0
	measured the parameter of interest (e.g. prevalence of low back pain)	reliability and validity (if this was necessary), e.g. test-re- test, piloting, validation in a previous study, etc.	Ü
	shown to have reliability and validity (if necessary)?	No (HIGH RISK): The study instrument had NOT been shown to have reliability or validity (if this was necessary).	1
8.	Was the same mode of data collection used for all subjects?	Yes (LOW RISK): The same mode of data collection was used for all subjects.	0
	omen a construction of • of the construction	No (HIGH RISK): The same mode of data collection was NOT used for all subjects.	1
9.	Were the numerator(s) and denominato r(s) for the parameter of interest appropriate	Yes (LOW RISK): The paper presented appropriate numerator(s) AND denominator(s) for the parameter of interest (e.g. the prevalence of low back pain).	0
		No (HIGH RISK): The paper did present numerator(s) AND denominator(s) for the parameter of interest but one or more of these were inappropriate.	1
10.	Summary on the overall risk of study	LOW RISK	0-3
	bias	MODERATE RISK	4-6
		HIGH RISK	7-9

Appendix 4: Results of quality assessment

Table 1. Results of quality assessment (risk of bias) of 22 included papers

Authors	Year of publication	1. Was the study's target population a close representation of the national population in relation to relevant variables, e.g. age, sex, occupation?	2. Was the sampling frame a true or close representa tion of the target population ?	3. Was some form of random selection used to select the sample, OR, was a census undertaken?	4. Was the likelihood of non-response bias minimal?	5. Were data collected directly from the subjects (as opposed to a proxy)?	6. Was an acceptable case definition used in the study?	7. Was the study instrument that measured the parameter of interest (e.g. prevalence of low back pain) shown to have reliability and validity (if necessary)?	8. Was the same mode of data collection used for all subjects?	9. Were the numerator (s) and denominator(s) for the parameter of interest appropriate?	Summary score on the overall risk of study bias
Ural et al.	2015	1	0	0	0	0	1	0	0	0	2
Mendoza et al.	2011	1	0	1	1	0	0	0	0	0	3
Gonzalez-Torres et al.	2010	1	1	1	1	0	0	0	0	1	5
Sierra et al.	2006	1	0	1	1	0	0	0	0	0	3
Zikic et al.	2009	1	1	0	0	0	1	0	0	0	3
Garcia et al.	2006	1	1	1	0	0	1	0	0	0	4
Korzekwa et al.	2009	1	0	0	0	0	1	0	0	0	2
Johnson et al.	2006	1	0	0	1	0	1	0	0	1	4
Evren et al.	2007	1	0	0	0	0	1	0	0	0	2
Somer et al.	2015	1	0	1	1	0	1	0	0	1	5
Mitchell et al.	2012	1	0	0	0	0	1	0	0	1	3
Belli et al.	2012	1	0	0	0	0	1	0	0	0	2

Michal et al.	2011	1	0	0	1	0	1	0	0	0	3
Dorahy et al.	2006	1	1	1	0	0	1	0	0	0	4
Mueller-Pfeiffer et al.	2012	1	0	0	0	0	0	0	0	0	1
Belli et al.	2016	1	0	0	0	0	1	0	0	0	2
Michal et al.	2014	1	0	0	0	0	0	0	0	0	1
Foote et al.	2006	1	0	0	0	0	1	0	0	0	2
Michal et al.	2009	0	0	0	1	0	0	0	0	0	1
Yalya et al.	2015	1	0	1	1	0	1	0	0	0	4
Aponte-Soto et al.	2019	1	0	1	1	0	0	0	0	1	4
Myers & Llera	2020	1	1	1	1	0	1	0	0	0	5
Schlax et al.	2020	1	0	0	0	0	0	0	0	0	1

^a Low risk of bias: overall score of 0-3; Moderate risk of bias: overall score of 4-6; High risk of bias: overall score of 7-9

Table 1. Study characteristics of studies involving the general population

Authors & Year of publication	Country	Population	Sample size	Mean age (SD) /age range	% female (n)	Prevalence assessment method	Cut-off score	Results from original papers	Notes	DP/DR prevalence (%) among general population
Myers & Llera (2020)	United States	College psychology undergraduates	N=198	M = 19.72; $SD = 2.32$	72% (1% gender variant/non- conforming, 27% male)	CDS	70	n=22 participants (11%) reported experiencing clinical levels of dissociation.		11%
Michal et al. (2014)	Germany	Pupils aged 12–18 years	N = 3,809	- No DP: M = 15.6; SD= 1.7. - DP: M = 15.8; SD = 1.6	51.7% (n=1971) - No DP: 51.1 % (n=1716) - DP: 56.4% (n=255)	CDS-2	3 (≥3)	n=452 (11.9%) pupils aged 12-18 years scored 3 or higher in CDS-2.		11.9%
Michal et al. (2010)	Germany	Randomly selected community sample aged 35-74 enrolled in Gutenberg Health Study from April 2007 to October 2008	N= 4,912; **5,000 participants were enrolled but only 4912 completed the survey	- CDS-2 < 3 (n = 4859): M = 55.4; SD = 10.9. - CDS-2 >/= 3 (n = 41): M = 53.3; SD = 10.0	Not given	CDS-2	≥3	n=41 (0.8%) partiticpants scored on or higher than 3 in CDS-2, indicating a clinically significant DDD		0.8%
Schlax et al. (2020)	Germany	German general population aged 35-74 enrolled in Gutenberg Health Study from April 2007 to April 2012	N= 13 182; **15 010 participants were enrolled but only 13 182 included in the analysis	M= 54.8 ; SD= 10.9	49.5% (n=6,526)	CDS-2	≥3	n=100 (0.76%) partiicpants scored on or higher than 3 in CDS-2		0.76%
Michal et al. (2009)	Germany	German general population	N=1,287	Total sample: M = 48.9; SD = 18.3; range=14-90 DP-C (clinical DP): M = 50.4; SD =17.7 DP-I (impairment caused by DP): M = 55.2; SD = 17.8	54.5% - DP-C (clinical DP): 64% - DP-I (impairment caused by DP): 61.6%	CDS-9	19(≥19)	n=25 (1.9%) participants scored in the range of clinically significant DDD/		1.9%

Table1. (Continued)

Authors & Year of publication	Country	Population	Sample size	Mean age (SD) /age range	% female (n)	Prevalence assessment method	Cut-off score	Results from original papers	Notes	DP/DR prevalence (%) among general population
Gonzalez- Torres et al. (2010)	Spain	(1) Patients with schizophrenia or schizophrenia spectrum disorders (2) First-degree healthy relatives with no psychiatric history (3) healthy controls from the general population	N=392: - patients: n=147 - first-degree relatives: n=73 - control subjects:n=172	Overall: Median = 30.0, interquartile range = 24-42 - Patients: Median = 32, interquartile range = 25-40 - Relatives: Median = 35.5, interquartile range = 28-43.3 - Controls: Median = 27, interquartile range = 23-40.3	Not given	CDS (Spanish version)	71	(1) Among 141 patients, n=24 (17%) had DDD according to the cut-off point of 71. (2) Among 71 relatives, n=1 (1%) had DDD according to the cut-off point of 71. (3) Among 172 controls, n=2 (3%) had DDD according to the cut-off point of 71.	prevalence rates are listed as	1.16%
Aponte-Soto et al. (2019)	Puerto Rico	Hispanic adults with and without a history of interpersonal abuse: (1) participants with a history of interpersonal abuse (HIA); (2) control community group (CCG) without HIA	N=80 - HIA: n=40 - CCG: n=40	M = 31.00; SD = 9.9; range = 21-65	65% (n=52) - HIA: 65% (n=26) - CCG: 65% (n=26)	CDS (Spanish version)	70 (>70)	(1) 25% of HIA group scored > 70 on CDS.(2) No participants scored > 70 on CDS in the CCG group.	**Original paper didn't report specific number of participants diagnosed with DDD	0%
Somer et al. (2015)	Israel	 (1) Opiate use disorder patients (2) Arab women subjected to domestic violence (3) Non-clinical controls from a community sample (graduate students and their friends and family members) 	N = 261 - Arab women: n = 80; - OUD patients: n = 68; - Nonclinical participants: n = 103	 Arab women: M = 33.29; OUD patients: M = 32.44; nonclinical: M = 34.80 	59% (n=154) - Arab women group: 100% (n=80) - OUD patient group: 5.9% (n=4) - nonclinical Israeli group: 68% (n=70)	DDIS	Meets DDIS criteria for DDD diagnosis	(1) n = 43 (53.8%) Arab women subjected to violence had DDD. (2) n=4 (5.9%) opiate use disorder patients had DDD. (3) n=10 (9.7%) non-clinical controls had DDD		9.7%
Johnson et al. (2006)	United States	Adults in the community	N=658	M = 33.1; $SD = 2.9$	53.0% (n=349)	SCID-D	Meets DSM-IV criteria for DDD diagnosis	n=5 (0.8%) participants had DDD, including 3 females and 2 males.		0.8%

^a We recalculated the prevalence rates of DDD for all included papers. When the original papers had calculation errors in prevalence rates, we reported the correct prevalence rates in the results. We included patients with mixed DDD (i.e. dissociative amnesia+ Depersonalisation/ Derealisation) when calculating the prevalence rates. The median and interquartile range of age were reported if the original paper didn't provide mean age (SD) or age range of the sample.

^bCDS= Cambridge Depersonalisation Scale; CDS-2= 2-item version Cambridge Depersonalisation Scale; CDS-9= 9-item version Cambridge Depersonalisation Scale; DDIS= Dissociative Disorders Interview Schedule; SCID-D= Structured Clinical Interview for DSM-IV Dissociative Disorders

Table 2. Study characteristics of studies involving patients with mixed or unspecified disorders

Patient type	Authors & Year of publication	Country	population	Sample size	Mean age (SD) /age range	% female (n)	Prevalence assessment method	Cut-off score	Results from original papers	Notes	Prevalence among patients (%)
Outpatients	Dorahy et al. (2006)	Northern Ireland	adult psychiatric patients aged 18-65	N=20	M = 40.7; SD = 8.8; range = 26-57	95% (19)	DDIS	Meets DDIS criteria for DDD diagnosis	(1) n=2 (10%) patients diagnosed with depersonalisation disorder, (2) n=2 (10%) patients with depersonalisation disorder and dissociative amnesia	**new prevalenc e= (2+2)/20= 20%	20%
	Foote et al. (2006)	United States	inner-city outpatient psychiatric population aged 18-65	N=82 ** 231 eligible participants in original sample, but only 82 were interviewed with the DDIS	M = 37.4; SD = 11.4 (for entire sample of 231)	64% (147 of 231)	DDIS	Meets DDIS criteria for DDD diagnosis	Among 82 patients who were interviewed met the criteria for a DSM-IV dissociative disorder diagnosis, 4 (5%) patients had a diagnosis of depersonalisation disorder		5%
Inpatients	Sierra et al. (2006)	United Kingdom, Spain, and Colombia	psychiatric inpatients aged 18-65	N=140; - n=31 from United Kingdom - n=68 from Spain - n=41 from Colombia	UK: M = 38.52; SD=13 Spain: M = 35.13; SD=9 Colombia: M = 33.78, SD=13	47.9% (67) - UK: 45.2% (14) - Spain: 44.1% (30) - Colombia: 56.1% (23)	CDS (English version for UK group and Spanish version for Spanish and Colombian groups)	71 (>70)	(1)13 (41.9%) patients from UK scored above 70 in CDS (2)24 (35.8%) patients from Spain scored above 70 in CDS (3)7 (17.5%) patients from Colombia scored above 70 in CDS		41.9% in UK inpatients; 35.8% in Spain inpatients; 17.5% in Colombia inpatients
Mixed or unspecified patients											
Not mentioned	Garcia et al. (2006)	Mexico	Mexician psychiatric patients receiving treatment aged 18-63	N=100	M = 32.4; SD = 12.5; range = 18-63	63% (63)	DDIS	Meets DDIS criteria for DDD diagnosis	n=6 (6%) patients diagnosed with depersonalisation disorder		6%
Outpatients and day care patients	Mueller- Pfeiffer et al. (2012)	Switzerla nd	psychiatric outpatients and day care patients	N = 160	Median = 32.0	67.3% (107)	SCID-D	Meets DSM-IV criteria for DDD diagnosis	n = 7 (out of a possible 160) individuals diagnosed with depersonalisation disorder (4.4%), and all are female		4.4%

^a We recalculated the prevalence rates of DDD for all included papers. When the original papers had calculation errors in prevalence rates, we reported the correct prevalence rates in the results. We included patients with mixed DDD (i.e. dissociative amnesia+ Depersonalisation) Derealisation) when calculating the prevalence rates. The median and interquartile range of age were reported if the original paper didn't provide mean age (SD) or age range of the sample.

b CDS= Cambridge Depersonalisation Scale; DDIS= Dissociative Disorders Interview Schedule; SCID-D= Structured Clinical Interview for DSM-IV Dissociative Disorders

Table 3. Study characteristics of the studies involving patients with specific disorders

Specific Disorders	Authors & Year of publication	Country	Population	Patient type	Sample size	Mean age (SD) /age range	Gender (% female)	Prevalence assessment method	Cut-off score	Results from original papers	Notes	prevalence (%)
Anxiety Disorders												
Panic disorder	Mendoza et al. (2011)	Spain	Adult patients with panic disorders (with and without agoraphobia)	Outpatients	N=104	M = 37.5; SD = 8.8	71.1% (75)	CDS (Spanish version)	70 (>=70)	$n=21\ (20.2\%)\ had\ a\ CDS\ score\ higher\ than\ 69$ during the panic attack, including 2 males and 19 females		20.2%
Panic disorder	Ural et al. (2015)	Turkey	Psychotropic drug-naïve adult patients with panic disorder	Outpatients	N = 92	M = 31.98; SD = 7.32; range = 18-52	63.0% (58)	SCID-D (Turkish version)	Meets DSM-IV criteria for DDD diagnosis	n = 3 (3.3%) diagnosed with depersonalisation disorder according to the SCID-D.		3.3%
Social anxiety disorder	Belli et al. (2017)	Turkey	Psychotropic drug-naïve patients with social anxiety disorder (SAD)	Outpatients	N=94 - low Dissociation Questionnaire score (< 2.5): n= 56 - high Dissociation Questionnaire score (>2.5): n = 38	Not given	55.32% (52)	SCID-D (Turkish version)	Meets DSM-IV criteria for DDD diagnosis	(1) n = 30 (31.91%) of the 94 patients were found to have dissociative disorder comorbidity (2) n = 5 (5.3%) patients had depersonalisation disorder; (3) One patient (1.0%) had dissociative amnesia and depersonalisation,	**new prevale nce= (5+1)/9 4=6.3%	6.3%
Obsessive- compulsive disorder	Belli et al. (2012)	Turkey	Patients with obsessive- compulsive disorder (OCD)	Outpatients	N=78	M = 31.22; SD = 8.83; range = 18-54	76.9% (60)	SCID-D (Turkish version)	Meets DSM-IV criteria for DDD diagnosis	(1) n=3 patients (3.84%) were diagnosed as having with dissociative depersonalisation disorder (2) 3 patients (3.84%) were diagnosed as having dissociative amnesia disorder + dissociative depersonalisation disorder; 1 patient (1.28%) was diagnosed as having dissociative depersonalisation disorder + dissociative identity disorder, and 1 patient (1.28%) was diagnosed as having dissociative amnesia disorder + dissociative depersonalisation disorder + dissociative identity disorder	**new prevale nce= (3+3+1 +1)/78= 10.26%	10.26%

Table 3. (Continued)

Specific Disorders	Authors & Year of publication	Country	Population	Patient type	Sample size	Mean age (SD) /age range	Gender (% female)	Prevalence assessment method	Cut-off score	Results from original papers	Notes	prevalence (%)
Dissociative spectrum disorders												
Conversion disorder	Yalya et al. (2015)	Turkey	Patients with conversion disorder (CD) aged 18-65	Outpatients	N=54: -people diagnosed with dissociative disorders (DD+): n = 20 - people without dissociative disorders (DD-): n = 34	- DD+: M = 28.05; SD = 7.54; - DD-: M = 29.21; SD = 8.05	90.7% (49)	SCID-D	Meets DSM-IV criteria for DDD diagnosis	n=2 (3.7%) participants had DDD	**in original paper, the prevalence was 3.7% in abstract but 1.08% in result part. Correct prevalence= 2/54=3.7%	3.7%
Non-epileptic attack disorder	Mitchell et al. (2012)	United Kingdom	Adult patients with non-epileptic attack disorder (NEAD)	Not given	N = 50: - NEAD only: n = 39 - dual diagnosis: n = 11	Total sample: M = 42.0; SD = 14.5 NEAD only: M = 41.6; SD = 15.1 - Dual diagnosis: M = 43.4; SD = 12.4	70% (35) - NEAD only :69.2% (27) - Dual diagnosis: 72.7% (8)	Derealisatio n/Deperson alisation subscale of DES		20.4% of the sample reported pathological levels of depersonalisation and derealisation symptoms (sub-score≥30).	**didn't report specific number	20.4%
Substance use disorders												
Alcohol dependence	Evren et al. (2007)	Turkey	Inpatients with alcohol dependency	Inpatients	N=111	M = 43.6; SD = 9.9; range = 18-68	5.4% (6)	SCID-D (Turkish version)	Meets DSM-IV criteria for DDD diagnosis	(1) n=1 (0.9%) patient diagnosed with a depersonalisation disorder. (2) 1 patient had amnesia and derealisation	**new prevalence= (1+1)/111 =1.8%	1.8%
Opiate use disorder	Somer et al. (2015)	us (2 to (3 fr (g th) patients with opiate se disorder) Arab women subjected domestic violence) Non-clinical controls om a community sample raduate students and eir friends and family embers)	Inpatients	N = 261 - Arab women: n = 80; - OUD patients: n = 68; - nonclinical: n = 103	- Arab women: M = 33.29; - OUD patients: M = 32.44; - Nonclinical: M = 34.80	59% (154) - Arab women group: 100% (80) - OUD patient group: 5.9% (4) - Nonclinical Israeli group: 68% (70)	DDIS	Meets DDIS criteria for DDD diagnosis	(1) n = 43 (53.8%) Arab women subjected to violence had DDD (2) n=4 (5.9%) opiate use disorder patients had DDD (3) n=10 (9.7%) non-clinical controls had DDD		5.9%

Table 3. (Continued)

Specific Disorders	Authors & Year of publication	Country	Population	Patient type	Sample size	Mean age (SD) /age range	Gender (% female)	Prevalence assessment method	Cut-off score	Results from original papers	Notes	prevalence (%)
Interpersonal abuse	•									,		,
Domestic violence	Somer et al. (2015)	Israel	(1) Opiate use disorder patients (2) Arab women subjected to domestic violence (3) Non-clinical controls from a community sample (graduate students and their friends and family members)	Inpatients	N = 261 - Arab women: n = 80; - OUD patients: n = 68; - nonclinical: n = 103	- Arab women: M = 33.29; - OUD patients: M = 32.44; - Nonclinical: M = 34.80	59% (154) - Arab women group: 100% (80) - OUD patient group: 5.9% (4) - nonclinical Israeli group: 68% (70)	DDIS	Meets DDIS criteria for DDD diagnosis	(1)n = 43 (53.8%) Arab women subjected to violence had DDD (2) n=4 (5.9%) opiate use disorder patients had DDD (3) n=10 (9.7%) non-clinical controls had DDD		53.8%
Interpersonal abuse	Aponte-Soto et al. (2019)	Puerto Rico	Hispanic adults with and without a history of interpersonal abuse - participants with a history of interpersonal abuse (HIA); - control community group (CCG) without HIA	participants from mental health clinics		M = 31.00; SD = 9.9; range = 21-65	65% (52) - HIA: 65% (26) - CCG: 65% (26)	CDS (Spanish version)	70 (>70)	(1)25% of HIA group scored > 70 on CDS (2) No participants scored >70 on CDS in the CCG group	**didn't report specific number	25%

Table 3. (Continued)

Specific Disorders	Authors & Year of publication	Country	Population	Patient type	Sample size	Mean age (SD) /age range	Gender (% female)	Prevalence assessment method	Cut-off score	Results from original papers	Notes	prevalence (%)
Other specific disorders		,		•					·			
Depression	Zikic et al. (2009)	Serbia	Patients suffering from unipolar depression without psychotic features aged 18- 65	inpatients and outpatients	N=84	DP group: M = 44.5; low DP group (scored lower than 70): M = 45.7	76.2% (64)	CDS	70(≥70)	n=40 (47.6%) subjects scored higher or equal to 70, including 8 males (20%) and 32 females (80%)		47.6%
Schizophrenia or schizophrenia spectrum disorders	Gonzalez- Torres et al. (2010)	Spain	(1) Patients with schizophrenia or schizophrenia spectrum disorders (2) First-degree healthy relatives with no psychiatric history (3) Healthy controls from the general population	inpatients	N=392: - n=147 patients; - n=73 first-degree relatives; - n=172 control subjects	Overall: Median = 30.0, interquartile range = 24-42 - Patients: Median = 32, interquartile range = 25-40 - Relatives: Median = 35.5, interquartile range = 28-43.3 - Controls: Median = 27, interquartile range = 23-40.3	Not given	CDS (Spanish version)	71	(1) Among 141 patients, n=24 (17%) had DDD according to the cut-off point of 71. (2) Among 71 relatives, n=1 (1%) had DDD according to the cut-off point of 71. (3) Among 172 controls, n=2 (3%) had DDD according to the cut-off point of 71.	** errors in calculation in original paper. Correct prevalence were: (1) patients: 24/147=16. 3% (3) controls: 2/172 = 1.16%.	16.3% in patients with schizophrenia or schizophrenia spectrum disorders; 1% in relatives of patients;
Borderline personality disorder	Korzekwa et al. (2009)	Canada	Adult patients with borderline personality disorder (BPD)	outpatients	N=21 **54 eligible patients (BPD) but only 21 patients completed the SCID-D-R	M = 38; SD = 8 * for N=21	76%	SCID-D	Meets DSM-IV criteria for DDD diagnosis	n=4 (19%) patients with BPD had DDD according to SCID-D-R		19%

^a We recalculated the prevalence rates of DDD for all included papers. When the original papers had calculation errors in prevalence rates, we reported the correct prevalence rates in the results. We included patients with mixed DDD (i.e. dissociative amnesia+ Depersonalisation/ Derealisation) when calculating the prevalence rates. The median and interquartile range of age were reported if the original paper didn't provide mean age (SD) or age range of the sample.

bCDS= Cambridge Depersonalisation Scale; DES= Dissociation Experiences Scale; DDIS= Dissociative Disorders Interview Schedule; SCID-D= Structured Clinical Interview for DSM-IV Dissociative Disorders.

Table 4. The specific prevalence assessment tools and the results (prevalence rates) in studies using self-reported scales and diagnostic interviews

reported scales and diagnostic interviews			
	lence assessment tools	Authors (year of publication)	Prevalence (%)
Studies using self-reported scales			
	Cambridge Depersonalisation Scale	Sierra et al. (2006)	41.9% in UK inpatients; 35.8% in Spain inpatients; 17.5% in Colombia inpatients
		Zikic et al. (2009)	47.6%
		Myers & Llera (2020)	11%
		Aponte-Soto et al. (2019)	25% in participants with a history of interpersonal abuse; 0% in control group
	Cambridge Depersonalisation Scale (Spanish version)	Gonzalez-Torres et al. (2010)	16.3% in patients with schizophrenia or schizophrenia spectrum disorders; 1% in relatives of patients; 1.16% in healthy control
		Mendoza et al. (2011)	20.2%
	2-item Cambridge Depersonalisation Scale (CDS-2)	Michal et al. (2010)	0.8%
		Michal et al. (2014)	11.9%
		Schlax et al. (2020)	0.76%
	9-item Cambridge Depersonalisation Scale (CDS-9)	Michal et al. (2009)	1.9%
	Derealisation/Depersonalisati on subscale of Dissociation Experiences Scale (DES)	Mitchell et al. (2012)	20.4%
Studies using diagnostic interviews			
	Dissociative Disorders Interview Schedule (DDIS)	Garcia et al. (2006)	6%
		Dorahy et al. (2006)	10%
		Foote et al. (2006)	5%
		Somer et al. (2015)	53.8% in Arab women 5.9% in OUD patients 9.7% in the nonclinical group
	Structured Clinical Interview for DSM-IV Dissociative Disorders (SCID-D)	Mueller-Pfeiffer et al. (2012)	4.4%
		Korzekwa et al. (2009)	19%
		Belli et al. (2012)	3.84%
		Johnson et al. (2006)	0.8%
		Yalya et al. (2015)	3.7%
		Ural et al. (2015)	3.3%
	SCID-D (Turkish version)	Evren et al. (2007)	0.9%
		Belli et al. (2017)	5.3%

Fig 1. PRISMA flow diagram

(n = 23)