Prevalence of childhood mental disorders in high-income countries: a systematic review and meta-analysis to inform policymaking

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ABSTRACT

Question Mental disorders typically start in childhood and persist, causing high individual and collective burdens. To inform policymaking to address children’s mental health in high-income countries we aimed to identify updated data on disorder prevalence.

Methods We identified epidemiological studies reporting mental disorder prevalence in representative samples of children aged 18 years or younger—including a range of disorders and ages and assessing impairment (searching January 1990 through February 2021). We extracted associated service-use data where studies assessed this. We conducted meta-analyses using a random effects logistic model (using R metafor package).

Findings Fourteen studies in 11 countries met inclusion criteria, published from 2003 to 2020 with a pooled sample of 61545 children aged 4–18 years, including eight reporting service use. (All data were collected pre-COVID-19.) Overall prevalence of any childhood mental disorder was 12.7% (95% CI 10.1% to 15.9%; I²=99.1%). Significant heterogeneity persisted to diagnostic measurement and study location. Anxiety (5.2%), attention-deficit/hyperactivity (3.7%), oppositional defiant (3.3%), substance use (2.3%), conduct (1.3%) and depressive (1.3%) disorders were the most common. Among children with mental disorders, only 44.2% (95% CI 37.6% to 50.9%) received any services for these conditions.

Conclusions An estimated one in eight children have mental disorders at any given time, causing symptoms and impairment, therefore requiring treatment. Yet even in high-income countries, most children with mental disorders are not receiving services for these conditions. We discuss the implications, particularly the need to substantially increase public investments in effective interventions. We also discuss the policy urgency, given the emerging increases in childhood mental health problems since the onset of the COVID-19 pandemic (PROSPERO CRD42020157262).

BACKGROUND

Mental disorders typically start in childhood and adolescence (hereafter childhood) and significantly interfere with well-being and development.¹ ² Now the leading cause of childhood disability globally,³ these disorders often persist into adulthood, adding greatly to the burden for individuals.³⁴ The collective social and economic burdens are also high due to associated healthcare, education, child protection and justice system costs—and to the cost of lost human potential.⁵–⁹ Service shortfalls have compounded the burdens, even in high-income countries such as the UK, USA and Canada.¹⁰ This is despite growing health expenditures in these countries over the years.¹¹–¹³ These shortfalls also persist despite longstanding recognition that all children have the right to adequate health, social and educational services—including mental health services¹⁴—and despite considerable research evidence describing effective interventions for preventing and treating childhood mental disorders.¹⁵–¹⁷

To address children’s mental health needs, policymakers require pooled prevalence data from multiple high-quality epidemiological studies using current or recent diagnostic standards.¹⁸ ¹⁹ Studies using rigorous diagnostic measures in large representative/probabilistic samples and examining a range of mental disorders across multiple age groups provide accurate and comprehensive population estimates—and therefore the most accurate estimates of community burden. At the same time, data on associated patterns of mental health service use can also assist policymakers in identifying and remediating gaps.

Systematic review and meta-analysis methods are an optimal way of synthesising large bodies of research evidence to inform policymaking.²⁰ ²¹ However, recent reviews on the prevalence of childhood mental disorders have had policy-relevant limitations including: accepting less rigorous studies,²²–²⁴ reporting on a relatively limited number of disorders,²⁵; and/or omitting service use data.²²–²⁶ As well, to our knowledge no reviews have included studies applying the latest diagnostic standards from the American Psychiatric Association’s (APA’s) Diagnostic and Statistical Manual of Mental Disorders (DSM) or the World Health Organization’s (WHO’s) International Classification of Diseases (ICD).²⁷ ²⁸

Objective To inform policymaking intended to address children’s mental health in high-income countries, we therefore conducted a systematic review and meta-analysis to identify comprehensive and updated data on the prevalence of childhood mental disorders and, where possible, associated service use. To ensure robust data, we aimed to identify recent high-quality epidemiological studies. We focused on high-income countries.
because most low- and middle-income countries have yet to mobilise children’s mental health services on a large scale—due to insufficient financial and human resources.\textsuperscript{39,40}

**METHODS**

**Search strategy**

We followed the Meta-analysis of Observational Studies in Epidemiology (MOOSE)\textsuperscript{31} and Preferred Reporting Items for Systematic Review and Meta-Analyses (PRISMA)\textsuperscript{32} guidelines for this systematic review and meta-analysis. (Online supplemental appendix A provides MOOSE and PRISMA checklists.) We registered this review with PROSPERO (number CRD42020157262; see www.crd.york.ac.uk/PROSPERO/; no amendments added).

To identify relevant studies, we searched EMBASE, MEDLINE and PsycINFO databases using the following terms: mental disorders; child; adolescent; epidemiology; prevalence; incidence; health survey; survey; population; community; representative; stratified; probability. (We did not conduct separate searches for service use because we aimed to extract these data from the prevalence studies that met our eligibility criteria.) Search dates were January 1990 through February 2021. The year 1990 was chosen to capture the WHO’s ICD 10th and later editions (1990–2020)\textsuperscript{28} and the APA’s DSM fourth and later editions (1994–2013).\textsuperscript{13} We limited our searches to prevalence studies conducted with children aged 18 years or younger and published in peer reviewed journals. We did not include language limiters and used Google Translate to assess studies published in languages other than English. We also hand-searched relevant systematic reviews retrieved through our database searches to identify additional publications (eg, Erskine \textit{et al}\textsuperscript{23} and Polanczyk \textit{et al}\textsuperscript{25}). For applicability to policymaking in high-income countries, we excluded studies conducted in low- and middle-income countries (according to World Bank classifications).\textsuperscript{34} (Online supplemental appendix B gives search strategy details.)

**Study selection**

After title screening, two authors independently assessed all abstracts. Relevant studies were retrieved and independently assessed, again by two authors, identifying those that met all inclusion criteria, which were established a priori. To ensure inclusion of rigorous studies, we applied quality indicators requiring representative/probabilistic sampling and reliable and valid child diagnostic measures, including assessment of impairment; we excluded studies that used only clinical records for diagnoses. We required studies to provide overall prevalence for any disorder since this figure is particularly relevant for policymakers. For comprehensiveness, we also required studies to report on three or more individual disorders or disorder groups. We excluded studies that only reported lifetime prevalence because of potential recall bias and because such data provide less accurate estimates of current service needs.\textsuperscript{35} Next, we identified all supplemental publications making use of data from the accepted original studies, searching Web of Science by study names and/or titles and/or authors. Throughout the process, any disagreements were resolved by consensus involving two or more authors. (Online supplemental appendix C provides study inclusion criteria; figure 1 shows the search process; Online supplemental appendix D provides included studies.)

**Data extraction and quality assessment**

We extracted data on the prevalence of any mental disorder and on the prevalence of individual disorders or disorder groups where possible. We also extracted data on potential moderators of prevalence including: overall study design; study location (continent); sampling area (national vs regional); data collection years; sampling frame; child age and sex; diagnostic standard; diagnostic measure; informants; diagnostic algorithm for reporting/combining data from different informants; and time-frames for assessing symptoms and impairment.\textsuperscript{25}

To determine the prevalence of mental health-related service use, we extracted data from all accepted studies that assessed this variable among children who had mental disorders—including any service contacts for mental health needs regardless of type, setting or provider. One author extracted all salient data; a second author then independently verified this work, with input from other authors as needed. We also contacted study authors when we required additional information.

Beyond applying our inclusion criteria, we also assessed study quality using an adapted risk-of-bias tool developed by Hoy \textit{et al} for prevalence studies.\textsuperscript{36} This tool assessed selection, non-response and measurement biases, as well as biases related to analysis. Scores of 0–3 were deemed low risk of bias, 4–6 moderate, and 7–10 high. (Online supplemental appendix E gives the quality assessment tool and results.)

**Data synthesis**

We conducted a meta-analysis using a random effects logistic model to calculate the pooled overall prevalence estimate for any mental disorder and its 95% confidence interval (95% CI). We performed the same analysis to calculate pooled prevalence estimates for individual disorders and service use among children with mental disorders when relevant data were available from at least two studies. For our meta-analyses, where possible we used prevalence data that study authors reported regardless of diagnostic algorithms used. These algorithms included: using clinical judgement; applying an ‘OR’ rule whereby disorders were counted if diagnostic thresholds were met by any informant (child OR parent); or relying on a single informant. Thirteen studies provided overall estimates for the whole sample, or a weighted average, that we used in our analyses. For one study that reported separate prevalence estimates by informant, with no weighted average for the whole sample, we selected estimates based on the most appropriate informant (for example, parent report for conduct disorder, child report for anxiety), then calculated a weighted average.\textsuperscript{37,38}

Regarding impairment, while precise definitions were not provided for all accepted studies, most reported definitions aligned with moderate-to-severe levels. Therefore, for the one study that reported separate estimates for mild, moderate and severe/‘serious’ impairment, we reported data that aligned with the other studies—in this case, ‘serious’ impairment which was defined as moderate impairment in most areas or severe impairment in at least one area.\textsuperscript{39} Heterogeneity was assessed using I\textsuperscript{2}. We conducted univariate meta-regression analyses to identify potential sources of heterogeneity for the pooled prevalence estimates for any mental disorder. We also performed a random effects meta-analysis to calculate pooled prevalence of mental health-related service use among children with mental disorders across studies. All statistical analyses were conducted using the metafor package for R.\textsuperscript{38}

**FINDINGS**

**Study characteristics**

We identified 159 studies for full-text assessment. Fourteen studies, described in 249 articles, met all inclusion criteria. Published between 2003 and 2020, these studies were conducted...
in 11 countries: the USA (four studies) and Australia, Canada, Chile, Denmark, Great Britain, Israel, Lithuania, Norway, South Korea and Taiwan (one study each). Sample sizes ranged from 957 to 10,438; child ages ranged from 4 to 18 years; 51.2% of participants were girls. Eleven of 14 studies used DSM-IV or DSM-IV-TR diagnostic standards, while two used ICD-10 and one used DSM-5. Most studies reported diagnoses using assessments of symptoms and impairment over one-year periods. (*Table 1* describes the characteristics of the 14 included studies; online supplemental appendix F lists excluded studies.)

Included studies determined diagnoses by interviewing children, parents and/or teachers using measures that included: Child and Adolescent Psychiatric Assessment; Composite International Diagnostic Interview; Development and Well-Being Assessment (DAWBA); Diagnostic Interview Schedule for Children-IV; Kiddie Schedule for Affective Disorders and Schizophrenia-Epidemiological; and Mini-International Neuropsychiatric Interview for Children and Adolescents. Among the seven studies that reported their specific criteria for assessing impairment, five required at least one severe or at least two moderate ratings of impairment in school achievement and/or relationships with family, friends or teachers, while two studies required moderate impairment in one of these domains.

Studies also varied in their use of informants for determining diagnoses. For four studies, diagnoses were established using computer algorithms based on symptoms reported by either parents or children. For the five studies using DAWBA, symptom responses from all informants were considered, then diagnoses were made using clinical judgement. For the remaining studies, prevalence estimates were informed by a single source: one using child report only; two using parent report only; and two using child and parent reports with estimates provided separately based on informant. All studies were deemed low risk regarding selection, non-response, measurement and analysis biases. (See online supplemental appendix E)

**Pooled prevalence of any mental disorder**

The pooled overall prevalence of any mental disorder was 12.7% (95% CI 10.1% to 15.9%; $I^2=99.1%$), based on 14 studies with a pooled sample size of 61,545 children. The following

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*Figure 1* Search process. DSM, Diagnostic and Statistical Manual of Mental Disorders; ICD, International Classification of Diseases.
<table>
<thead>
<tr>
<th>Study</th>
<th>Study location</th>
<th>Data collection year(s)</th>
<th>Sampling frame</th>
<th>Sample size*</th>
<th>Ages (years)</th>
<th>Diagnostic standard</th>
<th>Diagnostic measure†</th>
<th>Informant(s)</th>
<th>Timeframe‡ (months)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Canino 2004</td>
<td>Puerto Rico (USA)</td>
<td>1999–2000</td>
<td>Households</td>
<td>1897</td>
<td>4–17</td>
<td>DSM-IV</td>
<td>DISC-IV</td>
<td>Child or parent</td>
<td>12</td>
</tr>
<tr>
<td>Costello 2003</td>
<td>North Carolina (USA)</td>
<td>1993–2000</td>
<td>Schools</td>
<td>1420</td>
<td>9–16</td>
<td>DSM-IV</td>
<td>CAPA</td>
<td>Child or parent</td>
<td>3</td>
</tr>
<tr>
<td>Kessler 2012</td>
<td>USA</td>
<td>2001–2004</td>
<td>Households + schools</td>
<td>6483</td>
<td>13–17</td>
<td>DSM-IV</td>
<td>CIDI</td>
<td>Child or parent</td>
<td>12</td>
</tr>
<tr>
<td>Park 2015</td>
<td>Seoul (South Korea)</td>
<td>2005–2006</td>
<td>Schools</td>
<td>1645</td>
<td>6–12</td>
<td>DSM-IV</td>
<td>DISC-IV</td>
<td>Parent</td>
<td>12</td>
</tr>
</tbody>
</table>

†Diagnoses determined using clinical judgement combining information from multiple informants. / Diagnoses determined relying on a single informant based either on disorders and/or age ranges; figure 2 gives details on data used in meta-analysis. (Online supplemental appendix D lists references s1 to s15). ‘or’ Diagnoses determined applying ‘OR’ rule whereby disorders were counted if diagnostic thresholds met by any informant (either child or parent).

*51.2% of participants were girls.
†Fully structured measures included CAPA, CIDI, DISC-IV and MINI-KID; semi-structured measures included DAWBA and K-SADS-E.
‡Duration over which symptoms/impairment were assessed; when a range of timeframes were provided, 12 month prevalence was chosen.
§While 11- to 17-year-olds were informants for one disorder, only parent reported estimates were used in meta-analyses due to insufficient data for calculating overall rate for 11- to 17-year-olds.

CAPA, Child and Adolescent Psychiatric Assessment; CIDI, Composite International Diagnostic Interview; DAWBA, Development and Well-Being Assessment; DISC-IV, Diagnostic Interview Schedule for Children; DSM, Diagnostic and Statistical Manual of Mental Disorders; ICD, International Statistical Classification of Diseases and Related Health Problems; K-SADS-E, Kiddie Schedule for Affective Disorders and Schizophrenia-Epidemiological; MINI-KID, Mini-International Neuropsychiatric Interview for Children and Adolescents.
covariates were identified as potential sources of heterogeneity: study location (continent); overall study design; diagnostic standard; diagnostic measure; informants; diagnostic algorithm for reporting/combining data from different informants; and timeframes for assessing symptoms and impairment. Covariates with no moderating effects on prevalence included: sampling area (national vs regional); sampling frame; data collection years; and child age and sex. (Figure 2 provides the forest plot for any disorder; online supplemental appendix G provides details on our meta-regression findings.)

Pooled prevalence of individual disorders and disorder groups

Pooled prevalence was calculated for individual disorders and/or disorder groups when such data were reported in two or more studies. The most common conditions were: any anxiety disorder, pooled prevalence 5.2% (95% CI 3.2% to 8.2%); attention-deficit/hyperactivity disorder (ADHD) 3.7% (95% CI 2.3% to 5.7%); oppositional defiant disorder 3.3% (95% CI 2.4% to 4.6%); any substance use disorder 2.3% (95% CI 2.1% to 2.6%); conduct disorder 1.3% (95% CI 0.8% to 2.3%); major depressive disorder 1.3% (95% CI 0.6% to 2.9%); and alcohol use disorder 1.2% (95% CI 1.0% to 1.4%). Prevalence rates for other conditions were: cannabis use disorder 0.6% (95% CI 0.4% to 1.0%); autism spectrum disorder 0.4% (95% CI 0.2% to 0.8%); obsessive-compulsive disorder 0.3% (95% CI 0.1% to 0.6%); bipolar disorder 0.3% (95% CI 0.1% to 1.1%); eating disorders 0.2% (95% CI 0.1% to 0.5%); post-traumatic stress disorder 0.1% (95% CI 0.02% to 0.5%); and schizophrenia 0.1% (95% CI 0.1% to 0.3%). Ten of 14 studies reported on concurrent disorders. Among children with mental disorders, 26.5% (weighted average; range 14.0–37.0%) had two or more disorders. (Table 2 presents pooled prevalence estimates for these disorders and/or disorder groups, including number of studies and pooled sample sizes; online supplemental appendix H presents forest plots for individual disorders and disorder groups.)

Pooled prevalence for service use

For children with mental disorders, we extracted data on any services accessed for mental health concerns, provided in eight studies.39–42 s2, s8, s13, s15 These studies varied considerably regarding measurement of services. Some included only mental healthcare, such as counselling provided by psychologists or psychiatric medications prescribed by physicians,42 s2, s13 while others included a much wider range, such as complementary medicine and youth justice services.39–41 s8, s15 Studies also varied regarding assessment of timeframes for service use. While most measured past-year use,40–42 s2, s13, s15 three-month39 and six-month use42 were each assessed in a single study. Across these studies children were also defined as being recipients of services regardless of the intensity or duration of care provided. Informants also differed across the studies—involving children34, s15 or parents40–42, or both.39, 41, s8
Our meta-analysis found that only 44.2% children with mental disorders received any services for their mental health concerns (95% CI 37.6% to 50.9%; \( I^2 = 95.5\% \)).

### DISCUSSION

To inform policymaking to address children’s mental health in high-income countries, we conducted a systematic review and meta-analysis to determine the prevalence of childhood mental disorders and, where possible, associated service use. Fourteen high-quality studies from 11 high-income countries met the inclusion criteria, with a pooled sample of 61,545 children aged 4 to 18 years; eight studies also examined service use. The overall prevalence of any childhood mental disorder was 12.7%. The most common conditions were: any anxiety disorder (5.2%); ADHD 3.7%; oppositional defiant disorder 3.3%; any substance use disorder 2.3%; conduct disorder 1.3%; and major depressive disorder 1.3%. Among those with disorders, 26.5% had two or more disorders concurrently. Concerningly, only 44.2% of children with mental disorders received any services for these conditions.

These findings add to the extant literature in several ways that are salient for informing policymaking. To provide accurate population estimates, we have reported on only the most rigorous studies. To ensure comprehensiveness, we have included new studies not captured in previous reviews and have provided estimates for 12 of the most common disorders or disorder groups across a range of child ages. To inform service planning, we have also depicted the proportion of children in the general population who meet diagnostic disorder thresholds—including having both symptoms and impairment—and who therefore require mental health treatment services. As well, we have provided salient new data on service use by children with mental disorders.

We believe that our review can enable policymakers to better understand the mental health needs of children in high-income countries. In particular, policymakers can use our prevalence figures as benchmarks—calculating the numbers needing treatment at any given time within a given population or jurisdiction, then comparing the numbers in need with the numbers actually receiving mental health services. Service use does not always equate with need as defined by disorder prevalence. For example, families may seek services for children experiencing distress who do not meet diagnostic criteria. Consequently, prevalence data may underestimate service needs. However, prevalence data still provide critical population-level information for service planning. Importantly, our data cover children who are not receiving services—thereby providing a crucial supplement to administrative data on mental health service use. Beyond this, our data depict levels of increased children’s mental health needs that have emerged during the pandemic and that are predicted to continue.
We have depicted a high prevalence of childhood mental disorders; and tracking population outcomes through national surveillance programmes. As well, given that only an estimated 44.2% of children with mental disorders are receiving any services for these conditions, many countries will need to substantially increase children’s mental health budgets—and protect these budgets.

Given severe service shortfalls, it is also crucial to consider quality assurance and efficiency. Quality assurance involves ensuring that effective—and only effective—interventions are offered, particularly given considerable high-quality research evidence on both prevention and treatment in the children’s mental health field. Quality assurance can also involve tracking child outcomes in general and clinical populations in order to evaluate the impact of policy efforts and to alter course and redeploy resources as needed. Efficiency, meanwhile, can involve building in prevention programming to avert avoidable individual burdens and to in turn reduce the burdens on treatment systems. While prevention programming has often been under-appreciated in the children’s mental health field, economic evaluation data suggest that preventing even just one case of a severe problem, such as childhood conduct disorder, may save an estimated $5.6 million (2021 USD equivalency) in lifetime costs—through averted healthcare, special education, child protection and justice system costs. In addition, adopting innovative service strategies and establishing tiers of service aligned with the type and intensity of need can help extend the reach of practitioners while also reaching more children—another form of efficiency. Beyond this, it is important to understand barriers to accessing services and how these may be addressed—to ensure that all children can access the mental health services they need when and where they need them.

**CONCLUSIONS**

We have depicted a high prevalence of childhood mental disorders coupled with unacceptable service shortfalls in high-income
countries—to a degree that violates children’s rights.14 In essence, we have illuminated an invisible crisis in children’s mental health. In contrast, robust services are in place for child physical health problems such as cancer, diabetes and infectious diseases in most of these countries.11,12 Typical health expenditure levels indicate that high-income countries can also afford to do better.11–13 We therefore believe that our findings constitute a call to public action, taking the steps outlined above. This is particularly urgent given documented increases in children’s mental health needs since the onset of the COVID-19 pandemic—needs which are predicted to continue.14,15 The need for public action therefore could not be more urgent. Such action is essential if all children are to flourish. This collective flourishing will in turn benefit all—when fewer children needlessly experience mental disorders and associated disability, and when more children go on to thrive, contribute and meet their full potential.

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Competing interests KG and CW participated in the Ontario Child Health Study [reference 8] as Co-Principal Investigator and Co-Investigator, respectively. They therefore did not participate in decisions about inclusion, data collection or interpretation regarding this study. KG was also an author on one other included study [reference 10] and did not participate in decisions about inclusion, data collection or interpretation regarding this study. These two authors have no other conflicts to declare. All other authors declare no conflicts.

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Provenance and peer review Not commissioned; externally peer reviewed.

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Systematic review


