Assessing the impact of mental health difficulties on young people’s daily lives: protocol for a scoping umbrella review of measurement instruments

Karolin Rose Krause†,1,2 Sophie Chung,3 Terri Rodak,4 Kristin Cleverley†,5,6 Nancy J Butcher†,7,8 Peter Szatmari†,8

ABSTRACT

Introduction An important consideration for determining the severity of mental health symptoms is their impact on youth’s daily lives. Those wishing to assess ‘life impact’ face several challenges: First, various measurement instruments are available, including of global functioning, health-related quality of life and well-being. Existing reviews have tended to focus on one of these domains; consequently, a comprehensive overview is lacking. Second, the extent to which such instruments truly capture distinct concepts is unclear. Third, many available scales confound symptoms and their impact, thus undermining much needed analyses of associations between the two.

Methods and analysis A scoping umbrella review will examine existing reviews of life impact measures for use with children and youth aged 6–24 years in the context of mental health and well-being research. We will systematically search six bibliographic databases (MEDLINE, Embase, APA PsycINFO, CINAHL, Web of Science, and the COSMIN database of systematic reviews of outcome measurement instruments), and conduct systematic record screening, data extraction and charting based on methodological guidance by the Joanna Briggs Institute. Data synthesis will involve the tabulation of scale characteristics, feasibility and measurement properties, and the use of summary statistics to synthesise how these instruments operationalise life impact. The protocol was registered prospectively with the Open Science Framework (osf.io/ers48).

Ethics and dissemination This study will provide a comprehensive road map for researchers and clinicians seeking to assess life impact in youth mental health, providing guidance in navigating available measurement options. We will seek to publish the findings in a leading peer-reviewed journal in the field. Formal research ethics approval will not be required.

INTRODUCTION

A key consideration for determining the severity of mental health difficulties is the extent to which these difficulties impact on a young person’s daily life. The Diagnostic and Statistical Manual of Mental Disorders-5† determines ‘clinical significance’ in relation to two criteria: individuals must display specific symptoms, and those symptoms must cause considerable distress or impairment in daily life.2 Impaired daily functioning has been shown to influence help-seeking and health providers’ decisions about the type of care an individual should receive.3 4 Assessing life impact can also help contextualise changes in symptom severity scores when assessing treatment efficacy and effectiveness.5–10 From a public health perspective, consideration of life impact has moved common mental health conditions like depression to the fore of public health agendas, by showcasing that the associated burden of disease is comparable to that of cardiovascular or respiratory diseases.2

In child and youth mental health (hereafter we will use the terms ‘youth mental health’ and ‘young people’/‘youth’ for brevity to refer to those aged 6–24 years, in line with definitions of middle childhood, adolescence and young adulthood by the National Institute of Child Health and Human Development
(Paediatric) Terminology and the United Nations), life impact has typically been approached through the lens of functional impairment. Functioning describes a young person’s ability ‘to adapt to varying demands of home, school, peer group or neighbourhood’ in line with age-specific expectations and cultural norms (p. 1060). On a continuum of functioning, impairment marks one end of the spectrum, while high levels of adaptation and competency (eg, thriving, flourishing) mark the other end.

The Children’s Global Assessment Scale (CGAS) is a commonly used single-item measure that provides an overall rating of a young person’s functioning, based on clinician report. Other instruments take a more fine-grained approach by assessing functioning in specific areas of life. For example, the Social Adjustment Scale generates separate subscale scores for social functioning with friends, family, at school and in dating contexts. In addition, measures of symptom-specific or condition-specific impairment, focus on the extent to which psychopathological ‘symptoms interfere with and reduce adequate performance of important and desired aspects of a child’s life’ (p. 455). For example, the Strengths and Difficulties Questionnaire (Impact Supplement) and diagnostic interviews like the Kiddie Schedule for Affective Disorders and Schizophrenia enquire about functional impairment caused by psychopathology symptoms indicated during earlier parts of the respective assessments.

In physical health contexts and some population-based research, the impact of a particular health condition or of a person’s overall health status on their daily life is often conceptualised as health-related quality of life (HRQoL). Quality of life, has been described as ‘the overall positivity with which individuals view their state and circumstances’ (p. 455), and is thought to span physical, mental and social well-being. HRQoL refers more specifically to quality of life in a health or medical context. Relevant instruments include, for example, the brief EuroQol 5D-youth that is commonly used in economic evaluations; the 52-item KIDSCREEN that was developed for the measurement of HRQoL in the general paediatric population, or the PROMIS item bank for paediatric global health, designed to assess overall perceptions of health in youth with chronic health conditions.

Well-being is another domain that researchers may consider when assessing the life impact of mental health conditions. While a consensus definition is lacking, this domain has been described as ‘a combination of positive emotions, engagement, meaningful relationships and a sense of accomplishment, or as flourishing in aspects of feeling and functioning, thus reflecting the positive aspects of mental health’ (p. 771). For example, the Warwick-Edinburgh Mental Well-Being Scale is a self-report instrument validated in adolescents that exclusively assesses positive aspects of mental health.

The conceptual domains of functioning, HRQoL and well-being have different theoretical roots, yet it has been suggested that these terms are often used interchangeably. All three might be considered as avenues for assessing the life impact of mental health difficulties in children and youth. For example, a recently developed core outcome set for child and youth anxiety and depression recommends assessing functioning via three measures: the CGAS as a measure of clinician-rated global functioning; a self-report scale of condition-specific impairment; and the KIDSCREEN as a HRQoL measure. More generally, it is not clear whether scales purported to assess life impact via these domains are truly conceptually distinct, or whether they merely focus on different ends of the functioning continuum. A systematic review of measurement instruments that examines degrees of overlap and complementarity is lacking.

Researchers wishing to assess the life impact of mental health difficulties in young people further face the challenge of selecting the most appropriate instrument. A recent scoping review identified 14 different measures of global functioning, three measures of condition-specific impairment, and 14 measures of HRQoL, across 257 treatment studies for child and youth anxiety, depression, obsessive–compulsive disorder and post-traumatic stress disorder. Several reviews provide overviews of available instruments, their measurement properties and feasibility characteristics, but these have tended to be domain-specific (eg, focusing only on HRQoL); consequently, a comprehensive overview of life impact measures is lacking. On the other hand, broader reviews of mental health assessment tools have not typically been exhaustive in their coverage of life impact measures, and have not tended to examine methodological questions specific to life impact assessment.

A third challenge to the measurement of life impact in mental health is that many available instruments conflate items that assess symptom severity with items assessing the life impact of such symptoms. For example, the CGAS’s description of ‘superior functioning’ includes ‘no symptoms’ as a criterion. Similarly, the Health of the Nation Outcome Rating Scale is a 13-item measure that includes 7 symptom-focused items alongside five functional items (covering school functioning, self-care and relationships with peers and at home). The conflation of symptom severity and life impact items in a single scale hinders analyses of cross-sectional and longitudinal associations between the two domains.

Finally, an important fourth challenge is that many available instruments have been developed in Western high-income countries, and may not have cross-cultural validity or measurement invariance in lower-income or middle-income contexts or in specific cultural communities. As functioning is defined in relation to age-specific and culture-specific expectations and norms, life impact measures that are not culturally sensitive and appropriate may yield misleading data. Even in the contexts where measurement instruments were originally developed, youth may not always have been involved in their creation, which may weaken their content validity.
Objectives and research questions
This scoping umbrella review will examine how functioning, impairment, HRQoL and well-being have been conceptualised and measured in the youth mental health literature. It will seek to provide an overview of the design characteristics, feasibility aspects and measurement properties of available instruments, by considering existing individual reviews as primary studies. We seek to answer the following research questions:

RQ 1. What child-reported, parent-reported and clinician-reported measurement scales are available for assessing life impact in children and youth aged 6–24 years in the context of mental health and well-being research?

RQ 2. What information is available from existing reviews about the design characteristics (eg, target construct, target age range and use context, intended informant), feasibility (ie, length, cost and accessibility, language version availability) and measurement properties (ie, validity, reliability, responsiveness) of these instruments?

RQ 3. What populations and use contexts were these instruments originally designed for, according to their initial development study? Which cultural contexts were the instruments validated in?

RQ 4. According to an instrument’s original development study, were young people consulted as part of the measure development process?

RQ 5. Do measures of functioning, HRQoL and well-being appear to capture distinct conceptual domains, as opposed to assessing the same domain at different ends of the functioning continuum, based on subscale and item content?

RQ 6. To what extent do available measures of life impact conflate the measurement of psychopathology symptoms with the measurement of life impact?

METHODS AND ANALYSIS
Study design
The proposed study is a scoping umbrella review. An umbrella review considers existing review articles as its principal source of evidence and aims to compare, contrast or synthesise their findings.43 While systematic reviews (including umbrella reviews) typically seek to answer clearly defined questions (eg, ‘Which measure of global functioning provides the highest degree of validity and reliability?’), scoping reviews often seek to answer broader questions about the state of the evidence or about predominant methodological approaches in a given area.44 A scoping umbrella review is an appropriate approach for this study because several existing reviews can be synthesised to provide a comprehensive mapping of available instruments and their properties. We will follow the methodological guidance provided by the Joanna Briggs Institute (JBI) for the conduct of umbrella reviews, scoping reviews and reviews of measurement properties.45–47

Protocol
This review protocol complies with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) for Protocols reporting guidelines (online supplemental appendix 1).46 The final review will follow the PRISMA for Scoping Reviews.44 The protocol was registered prospectively with the Open Science Framework (OSF) on 26 May 2021.49 On registration and submission of the protocol, title and abstract screening was complete, but full text screening had not begun. Any important amendments to this protocol will be documented on the OSF registration page.

Inclusion criteria
This scoping umbrella review will consider systematic reviews, scoping reviews, rapid reviews and narrative reviews that seek to provide an overview of available measurement scales to assess functioning, impairment, HRQoL or well-being. These may be reviews that systematically identify a range of measurement instruments, or reviews that synthesise the available literature for a single instrument. Narrative reviews, rapid reviews and scoping reviews will be included in addition to systematic reviews because this scoping umbrella review aims to map the landscape of available life impact measures as comprehensively as possible, rather than to identify the most systematic or robust evidence pertaining to these instruments. Inclusion criteria for reviews are defined to match the PICO components for systematic reviews of measurement properties (ie, Population, Instruments, Construct, Outcome) in line with JBI guidelines.47 The PICO criteria are summarised in table 1.

Population (P)
Review articles must have an explicit focus on measurement in middle childhood (defined here as starting at age 6 in line with proposed age group standards11), adolescence, and/or young adulthood (defined here as ending at age 24, in line with the United Nations’ definition of ‘youth’12). Studies with a majority focus on adults will not be considered, unless they include a separate appraisal of tools for a relevant paediatric age group. We will exclude reviews focused on early childhood (ie, ages 0–5 years), where tailored assessment approaches are likely needed.

We will include reviews that examine the measurement of life impact in populations with a primary mental health or substance use concern, or in the context of assessing mental health and well-being in the general population or in non-specific health contexts. We will exclude reviews that focus on youth with physical health conditions. Instruments identified in such reviews may place a considerable focus on physical body functions that may be less relevant in a mental health context. We will also exclude reviews focused on youth with intellectual disabilities, neurological conditions (eg, epilepsy, cerebral palsy) or autism spectrum disorder (ASD). These profiles may require specialised assessments of life impact, and the conceptual separation of symptoms from functioning may

be particularly complicated (eg, with social functioning constituting a symptom of ASD). A separate review may be warranted to cover life impact measures for children and youth with these conditions.

Instruments (I)
We will consider scales designed for completion by clinicians, external raters, parents or carers, and young people. These may be assessment or outcome measures but must focus on an eligible domain of life impact (ie, see below) rather than symptom severity or psychopathology. We will exclude reviews focused on diagnostic tools or on the assessment of specific mental health conditions, unless the article’s abstract explicitly states that measures of an eligible life impact construct were considered alongside symptom severity measures. We will further exclude performance tests, cognitive tests, language assessments, biometric tests, school-based functional behavioural assessments, and population-level composite indices of well-being or HRQoL.

Constructs (C)
We will consider instruments designed to assess life impact through the measurement of global functioning, social functioning/adaptation, general or condition-specific impairment, HRQoL, well-being (including flourishing), and life satisfaction. Constructs that are not eligible include symptoms of psychopathology, language ability, cognitive ability, executive functioning and motor functioning. Instruments that cover any of these constructs at an item level as part of measuring a broader eligible construct (eg, HRQoL) may be included.

Outcomes (O)
We will include articles that state an intent to review, appraise or map relevant measurement instruments and that provide a structured discussion or a tabulated overview of the instruments identified.

**Table 1** PICO statement for scoping umbrella review

<table>
<thead>
<tr>
<th>P (Population)</th>
<th>I (Instruments)</th>
<th>C (Constructs)</th>
<th>O (Outcomes)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Included</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Children and adolescents aged 6–24 years, with a primary mental health condition/concern, subject to mental health assessment in general population, or in the context of assessing life impact in health contexts broadly speaking.</td>
<td>Youth, parent, clinician or external rater report; initial assessment or outcome measure</td>
<td>Global functioning</td>
<td>Construct domain</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Target age group</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Reporter</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Target use context</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Length</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Accessibility and cost</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Measurement properties</td>
</tr>
<tr>
<td>Excluded</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ages 0–5 or 24+</td>
<td>Performance test; biometric assessment</td>
<td>Global functioning</td>
<td>Target age group</td>
</tr>
<tr>
<td>Children and youth with intellectual disabilities or where mental health is a secondary concern to a primary physical condition; pure physical health context</td>
<td>Social functioning</td>
<td>Reporter</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Cognitive ability</td>
<td>Functional impairment</td>
<td>Target use context</td>
</tr>
<tr>
<td></td>
<td>Executive functioning</td>
<td>HRQoL</td>
<td>Length</td>
</tr>
<tr>
<td></td>
<td>Symptom severity</td>
<td>Well-being</td>
<td>Accessibility and cost</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Flourishing</td>
<td>Measurement properties</td>
</tr>
</tbody>
</table>

HRQoL, health-related quality of life.

**Publication type**
Reviews must have been published from January 1990 onwards. We will limit the language of publication to English to accommodate languages spoken within the review team; to ensure that all records can be screened by at least two review team members; and because item content will be an important aspect to consider and must therefore be accessible to the majority of the review team.

We will include review articles published in peer-reviewed journals, assessment handbooks (if accessible online) and conference proceedings (including workshop summaries and conference papers, but not including conference abstracts). We will further include reviews that were published as grey literature (eg, as reports on organisational websites) and otherwise meet the inclusion criteria.

**Search strategy**
The development of the search strategy is led by a health science librarian (TR) in collaboration with other members of the review team (KKR, PS). The search strategy combines search terms describing the population (eg, “child*” OR “youth” AND “depress*” OR “anxiety disorder*” OR “externalizing problem*”) and domains of interest (eg, “function*” OR “HRQOL”) with search terms limiting the results to reviews (eg, “systematic review” OR “scoping review”) of measurement instruments or outcome measurement approaches (eg, “psychometr*” OR “measurement instrument*”). Our tailored search syntax is informed by existing search filters for measurement instruments that were developed by the University of Oxford’s Patient-Reported Outcome Measurement Group and by the COnsensus-based Standards for the selection of health Measurement INstruments (COSMIN) initiative. Pilot searches informed the final search strategy (see online supplemental appendix 2).

The final search will be performed by the review team’s health science librarian (TR) in Medline, Embase, APA PsycINFO, Cumulative Index to Nursing & Allied Health, CINAHL, Cochrane Library, and the COnsensus-based Standards for the selection of health Measurement INstruments (COSMIN) initiative.
Health Literature (CINAHL) and Web of Science, and by a member of the review team (KRK) in the COSMIN database of systematic reviews of outcome measurement instruments. Retrieved records will be deduplicated using Covidence systematic review software. We will ask a group of subject matter experts to review the list of articles identified through the automatic search, and to suggest additional reviews that may have been missed. We will also conduct a targeted grey literature search via specific databases and websites identified as relevant by the team’s health science librarian (TR). In addition, we will handsearch the reference lists of included reviews to identify and retrieve the original development papers associated with eligible instruments, as well as copies of the instruments themselves, as available. We will consider supplemental searches if key information about a measure’s design characteristics is not available from the identified reviews or the instruments’ original development studies. Due to resource constraints, we will not conduct supplemental searches for a measure’s feasibility characteristics or measurement properties, and will base our reporting for these aspects on the information available from existing reviews.

We will review clearinghouses of measurement instruments for any additional scales that were missed by the included reviews, and will also make a note of any additional instruments identified while screening for eligibility. These additional instruments will not be subject to a systematic appraisal, but will be listed in the final report.

**Study selection**

Eligibility will be assessed via a two-stage screening process. For the title and abstract screening, 20% of all identified records will be screened independently and in duplicate by two reviewers (KRK and SC). A kappa coefficient exceeding 0.7 will indicate substantial inter-rater agreement. We will discuss any discrepancies and agree a final inclusion or exclusion rating. A single reviewer (KRK) will then screen the remaining titles and abstracts. All records retained for full-text screening will be checked for eligibility independently and in duplicate by two raters. Disagreements will be discussed and decisions about inclusion will be made with the help of a third reviewer as needed. Articles that do not meet inclusion criteria will be coded for exclusion in the Covidence software environment with the first exclusion criterion that becomes apparent. Eligible review articles will progress to data charting.

**Data extraction and charting**

Data will be extracted and charted using tailored adaptations of the JBI data extraction templates for systematic reviews of measurement properties and for umbrella reviews. The adapted matrices will be piloted to ensure an appropriate level of detail is charted. Data extraction will be conducted by one review author, and spot checks for comprehensiveness and accuracy will be conducted on at least 20% of the included reviews by a second reviewer.

Any disagreements will be resolved through discussion. Based on the extent of disagreement identified, the two reviewers will consider extending the spot checks to a larger subset of studies.

The information to be charted is shown in table 2. We will refer to the original development studies as needed, to extract whether or not youth or families were involved in measure development. We will also extract in which contexts an instrument has been validated. Where possible, we may review each measure’s item content to indicate whether items cover symptoms of psychopathology as well as life impact, and to examine the extent of overlap between measures purported to assess different life impact domains. Depending on the number and accessibility of the instruments identified, we may seek to undertake a systematic item-level mapping of content.

**Risk of bias assessment**

Scoping-type reviews do not seek to generate critically appraised and summative responses to specific research questions, but instead aim to map the available evidence on a given topic. Therefore, risk of bias assessments are not typically conducted as part of scoping reviews, and are not planned for this scoping umbrella review.

**Strategy for data synthesis**

We will synthesise the findings of existing reviews in relation to instrument design characteristics, feasibility characteristics, and measurement properties by applying the five-step data synthesis process recommended by Miles and Huberman and Whittmore and Knaff. This process consists of (1) data reduction; (2) data display; (3) data comparison; (4) conclusion drawing and (5) verification. During verification, we will review the original development studies associated with each instrument to ensure that the information about key instrument characteristics compiled during the scoping umbrella review is accurate.

We will present the characteristics of the included review articles, as well as the characteristics of the

### Table 2 Overview of data to extract and chart

<table>
<thead>
<tr>
<th>Information category</th>
<th>Detailed information to extract</th>
</tr>
</thead>
<tbody>
<tr>
<td>Publication identifiers</td>
<td>Journal, year, first author</td>
</tr>
<tr>
<td>Review characteristics</td>
<td>Type of publication, type of review, objective of the review, population and setting considered, number and names of databases searched, date range of search, language/geographical restrictions, number of studies included</td>
</tr>
<tr>
<td>Instrument design characteristics</td>
<td>Instrument name, domain measured, number of items, number and names of subscales, target age group, target population group (clinical vs non-clinical), target use context (screening, diagnosis, outcome measurement), reporter(s), response scale, recall period, involvement of youth in instrument development, cultural context of development and validation studies</td>
</tr>
<tr>
<td>Feasibility characteristics</td>
<td>Length, cost and accessibility, available language versions</td>
</tr>
<tr>
<td>Measurement properties</td>
<td>Summary findings relating to validity, reliability, responsiveness</td>
</tr>
</tbody>
</table>

identified measurement instruments in tabular format. We will report high-level quantitative summary statistics (ie, counts or frequencies) to describe the reviews and instruments identified (eg, number of instruments overall; number of instruments per life impact domain; number of instruments by type of reporter). We will further generate summary statistics and visualisations to report on the domains and subdomains of life impact covered by the identified instruments, and the extent to which these instruments appear to confound items measuring symptoms of psychopathology with items measuring life impact, based on an examination of item or subscale content. We will also specifically indicate whether a measure was validated in a population with a mental health concern, or whether it was validated exclusively in community samples.

**Patient and public involvement**

The Centre for Addiction and Mental Health implements a Youth Engagement Initiative that brings the voices of youth aged 14–29 years with lived experience of mental health challenges into research and service design.60–62 We will collaborate with a designated youth research partner in conducting specific aspects of this review that require the qualitative interpretation of measure content. In addition, we will present draft review findings to a virtual focus group including between four and eight youth advisors to solicit their feedback, and incorporate this into our interpretation and contextualisation of the review findings prior to finalising the study manuscript.

**Ethics and dissemination**

**Ethics**

Formal approval by a research ethics board will not be required, as the proposed project is a scoping umbrella review of existing data. We will consult youth as research partners rather than research subjects and will not collect or report any individual-level participant data.

**Dissemination**

We will seek to publish the findings from this scoping umbrella review in a leading peer-reviewed journal in the field of child and adolescent mental health. We will also seek to disseminate findings at national and international conferences, and will consider submitting the final review to the COSMIN database of systematic reviews of measurement properties. We will equally consider additional channels of dissemination, such as blog posts or podcasts.

**DISCUSSION**

Historically, outcome measurement in youth mental health research has focused on symptom severity.53–45 Yet, many common symptom scales are not immediately interpretable with regard to how a score change translates into real-world changes in a young person’s life. Assessing life impact in a structured way through use of suitable measurement scales can provide important complementary information to data collected via diagnostic tools and symptom severity measures.5–7 66 67 Within an empathetic, person-centred framework of care, it is important that clinicians pay attention to youth and family members’ unstructured, narrative accounts of how a mental health condition affects daily life, and that clinicians consider these narratives when making care decisions together with service users.68 In addition, however, the administration of suitable structured measurement instruments can help ensure that life impact is assessed systematically and reliably, so that comparisons can be made between individuals and over time. Similarly, the use of structured assessment tools can enable the systematic consideration of different perspectives (eg, when combining youth-rated, carer-rated and clinician-rated instruments).69 And the inclusion of self-reported datapoints that convey the youth’s perspective directly without mediation by the clinician. Two recent initiatives have highlighted functioning as a core outcome to track when evaluating clinical care for paediatric anxiety and depression,7 and when measuring youth mental health outcomes in population surveys.70 Yet, difficulties have been reported with identifying a gold-standard measure.9

This scoping umbrella review does not aim to comprehensively identify all life impact measures available worldwide, but will focus on instruments that have been reviewed in English language publications. This review also does not seek to yield an authoritative summary of which instruments provide the best measurement properties. This would require an in-depth assessment of the methodological quality of the psychometric evidence underpinning each instrument in line with COSMIN guidelines,71 which in turn would constitute a study in its own right for each instrument identified.72 Instead, this review will examine a range of design, feasibility and measurement properties to facilitate the preselection of candidates for future in-depth psychometric appraisals. It further aims to identify gaps with regards to age groups or use contexts covered, and examine the degree of conceptual overlap between instruments designated to assess different outcome domains (eg, functioning vs HRQoL). As such, it aims to take stock of current measurement practice, to inform discussions about suitable ways forward and to provide a road map to researchers and clinicians seeking to appraise which tool or combination of tools may be appropriate for a given population and context.

**Author affiliations**

1Cundill Centre for Child and Youth Depression, Centre for Addiction and Mental Health, Toronto, Ontario, Canada
2Research Department for Clinical, Educational and Health Psychology, University College London, London, UK
3Independent Health Researcher, London, UK
4CAMH Library, Department of Education, Centre for Addiction and Mental Health, Toronto, Ontario, Canada
5Lawrence Bloomberg Faculty of Nursing and Department of Psychiatry, University of Toronto, Toronto, Ontario, Canada
6The Margaret and Wallace McCain Centre for Child, Youth & Family Mental Health, Centre for Addiction and Mental Health, Toronto, Ontario, Canada
REFERENCES


58 Miles MB, Huberman AM. *Qualitative data analysis: an expanded sourcebook*. sage, 1994.
## PRISMA–P 2015 Checklist

This checklist has been adapted for use with protocol submissions to *Systematic Reviews* from Table 3 in Moher D et al: Preferred reporting items for systematic review and meta-analysis protocols (PRISMA–P) 2015 statement. *Systematic Reviews* 2015 4:1

**Administrative Information**

<table>
<thead>
<tr>
<th>Section/topic</th>
<th>#</th>
<th>Checklist item</th>
<th>Information reported</th>
<th>Line number(s)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Title</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Identification</td>
<td>1a</td>
<td>Identify the report as a protocol of a systematic scoping review</td>
<td>☒</td>
<td>p. 1</td>
</tr>
<tr>
<td>Update</td>
<td>1b</td>
<td>If the protocol is for an update of a previous systematic review, identify as such</td>
<td>☑</td>
<td>NA</td>
</tr>
<tr>
<td>Registration</td>
<td>2</td>
<td>If registered, provide the name of the registry (e.g., PROSPERO) and registration number in the Abstract</td>
<td>☒</td>
<td>p. 1</td>
</tr>
<tr>
<td><strong>Authors</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Contact</td>
<td>3a</td>
<td>Provide name, institutional affiliation, and e-mail address of all protocol authors; provide physical mailing address of corresponding author</td>
<td>☒</td>
<td>pp. 1,6,7</td>
</tr>
<tr>
<td>Contributions</td>
<td>3b</td>
<td>Describe contributions of protocol authors and identify the guarantor of the review.</td>
<td>☒</td>
<td>p. 7</td>
</tr>
<tr>
<td>Amendments</td>
<td>4</td>
<td>If the protocol represents an amendment of a previously completed or published protocol, identify as such and list changes; otherwise, state plan for documenting important protocol amendments</td>
<td>☒</td>
<td>p. 3</td>
</tr>
<tr>
<td><strong>Support</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sources</td>
<td>5a</td>
<td>Indicate sources of financial or other support for the review</td>
<td>☒</td>
<td>p. 7</td>
</tr>
<tr>
<td>Sponsor</td>
<td>5b</td>
<td>Provide name for the review funder and/or sponsor</td>
<td>☒</td>
<td>p. 7</td>
</tr>
<tr>
<td>Role of sponsor/funder</td>
<td>5c</td>
<td>Describe roles of funder(s), sponsor(s), and/or institution(s), if any, in developing the protocol</td>
<td>☒</td>
<td>p. 7</td>
</tr>
<tr>
<td><strong>Introduction</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rationale</td>
<td>6</td>
<td>Describe the rationale for the review in the context of what is already known</td>
<td>☒</td>
<td>pp.1-2</td>
</tr>
<tr>
<td>Objectives</td>
<td>7</td>
<td>Provide an explicit statement of the question(s) the review will address with reference to participants, interventions, comparators, and outcomes (PICO)</td>
<td>☒</td>
<td>p. 3</td>
</tr>
<tr>
<td>Section/topic</td>
<td>Checklist item</td>
<td>Information reported</td>
<td>Line number(s)</td>
<td></td>
</tr>
<tr>
<td>-------------------------</td>
<td>---------------------------------------------------------------------------------</td>
<td>----------------------</td>
<td>----------------</td>
<td></td>
</tr>
<tr>
<td><strong>METHODS</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Eligibility criteria</td>
<td>Specify the study characteristics (e.g., PICO, study design, setting, time frame) and report characteristics (e.g., years considered, language, publication status) to be used as criteria for eligibility for the review</td>
<td>☒</td>
<td>☐</td>
<td>pp. 3-4</td>
</tr>
<tr>
<td>Information sources</td>
<td>Describe all intended information sources (e.g., electronic databases, contact with study authors, trial registers, or other grey literature sources) with planned dates of coverage</td>
<td>☒</td>
<td>☐</td>
<td>pp. 4-5</td>
</tr>
<tr>
<td>Search strategy</td>
<td>Present draft of search strategy to be used for at least one electronic database, including planned limits, such that it could be repeated</td>
<td>☒</td>
<td>☐</td>
<td>Appendix 2</td>
</tr>
<tr>
<td><strong>STUDY RECORDS</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Data management</td>
<td>Describe the mechanism(s) that will be used to manage records and data throughout the review</td>
<td>☒</td>
<td>☐</td>
<td>pp. 5-6</td>
</tr>
<tr>
<td>Selection process</td>
<td>State the process that will be used for selecting studies (e.g., two independent reviewers) through each phase of the review (i.e., screening, eligibility, and inclusion in meta-analysis)</td>
<td>☒</td>
<td>☐</td>
<td>p. 5-6</td>
</tr>
<tr>
<td>Data collection process</td>
<td>Describe planned method of extracting data from reports (e.g., piloting forms, done independently, in duplicate), any processes for obtaining and confirming data from investigators</td>
<td>☒</td>
<td>☐</td>
<td>pp. 5</td>
</tr>
<tr>
<td>Data items</td>
<td>List and define all variables for which data will be sought (e.g., PICO items, funding sources), any pre-planned data assumptions and simplifications</td>
<td>☒</td>
<td>☐</td>
<td>p. 5, Table 2</td>
</tr>
<tr>
<td>Outcomes and prioritization</td>
<td>List and define all outcomes for which data will be sought, including prioritization of main and additional outcomes, with rationale</td>
<td>☒</td>
<td>☐</td>
<td>p. 2, Table 2</td>
</tr>
<tr>
<td>Risk of bias in individual studies</td>
<td>Describe anticipated methods for assessing risk of bias of individual studies, including whether this will be done at the outcome or study level, or both; state how this information will be used in data synthesis</td>
<td>☒</td>
<td>☐</td>
<td>p. 5</td>
</tr>
<tr>
<td><strong>DATA</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Synthesis</td>
<td>Describe criteria under which study data will be quantitatively synthesized</td>
<td>☒</td>
<td>☐</td>
<td>pp. 5-6</td>
</tr>
<tr>
<td>15b</td>
<td>If data are appropriate for quantitative synthesis, describe planned summary measures, methods of handling data, and methods of combining data from studies, including any planned exploration of consistency (e.g., $I^2$, Kendall’s tau)</td>
<td>☒</td>
<td>☐</td>
<td>pp. 5-6</td>
</tr>
<tr>
<td>15c</td>
<td>Describe any proposed additional analyses (e.g., sensitivity or subgroup analyses, meta-regression)</td>
<td>☒</td>
<td>☐</td>
<td>pp. 5-6</td>
</tr>
<tr>
<td>15d</td>
<td>If quantitative synthesis is not appropriate, describe the type of summary planned</td>
<td>☒</td>
<td>☐</td>
<td>pp. 5-6</td>
</tr>
<tr>
<td>Section/topic</td>
<td>#</td>
<td>Checklist item</td>
<td>Information reported</td>
<td>Line number(s)</td>
</tr>
<tr>
<td>-------------------------------</td>
<td>----</td>
<td>-------------------------------------------------------------------------------</td>
<td>-----------------------</td>
<td>-------------------------</td>
</tr>
<tr>
<td><strong>Meta-bias(es)</strong></td>
<td>16</td>
<td>Specify any planned assessment of meta-bias(es) (e.g., publication bias across studies, selective reporting within studies)</td>
<td>No</td>
<td>NA; Scoping review</td>
</tr>
<tr>
<td><strong>Confidence in cumulative evidence</strong></td>
<td>17</td>
<td>Describe how the strength of the body of evidence will be assessed (e.g., GRADE)</td>
<td>No</td>
<td>NA; Scoping review</td>
</tr>
</tbody>
</table>

BMJ Publishing Group Limited (BMJ) disclaims all liability and responsibility arising from any reliance on this supplemental material which has been supplied by the author(s).
Assessing the Impact of Mental Health Difficulties on Young People’s Daily Lives:
Protocol for a Scoping Umbrella Review of Measurement Instruments

Search strategy for Ovid MEDLINE(R)

1 mental health/
2 social adjustment/
3 mental disorders/ or exp anxiety disorders/ or exp "bipolar and related disorders"/ or exp "disruptive, impulse control, and conduct disorders"/ or exp dissociative disorders/ or exp "feeding and eating disorders"/ or exp mood disorders/ or exp "attention deficit and disruptive behavior disorders"/ or exp child behavior disorders/ or exp schizophrenia, childhood/ or exp neurotic disorders/ or exp personality disorders/ or exp "schizophrenia spectrum and other psychotic disorders"/ or exp somatoform disorders/ or exp substance-related disorders/ or exp "trauma and stressor related disorders"/
4 (mood disorder* or affective disorder* or personality disorder* or depressive disorder* or major depression or anxiety disorder* or obsessive compulsive or bipolar or schizo* or "substance use disorder*" or addiction* or conduct disorder* or behavio?r* disorder or stress disorder* or PTSD or CPTSD or complex trauma or developmental trauma).ti,ab,kf.
5 ((mental* or psychiatrist* or internalizing or externalizing or psychosocial or psycho-social) adj3 (diagnos* or disorder* or ill or illness* or problem* or challenge* or issue* or difficult*)).ti,ab,kf.
6 (mental health or wellbeing or well-being).ti,kf.
7 or/1-6
8 (child* or pediatric* or paediatric* or youngster* or youth* or adolescen* or teen* or young adult* or young person* or young people* or juvenile* or student* or underage* or under-age* or emerging adult* or transition* age*).ti,kf,hw,jn.
9 Psychometrics/
10 "Outcome Assessment (Health Care)"/
11 Psychiatric Status Rating Scales/
12 "Surveys and Questionnaires"/
13 self report/
14 Mass Screening/
15 psychometr*.ti,ab,kf.
16 (clinimetr* or clinometr*).ti,ab,kf.
17 ((outcome* or rating or screening) adj2 (assessment* or scale*)).ti,kf.
18 measure*.ti,kf.
19 questionnaire*.ti,kf.
20 instrument*.ti,kf.
21 (("quality of life" or HRQL or HRQoL or QL or QoL or health profile* or health status* or global health or disabilit* or disabl* or function* or wellbeing or well being or flourish* or life satisfaction or impair* or interference or life impact* or adjust* or adapt* or standardi?ed or personali?ed or nomothetic or idiographic or self-report* or patient report* or child report* or youth report* or parent report* or clinician report* or mental health) adj3 (assess* or index or
indices or instrument or instruments or measur* or questionnaire* or profile* or scale* or score* or status* or survey* or apprais* or metric* or inventor* or tool* or indicator*).ti,kf.

22 (health index* or health indices).ti,ab,kf,hw.

23 (PROM or PROMS).ti,ab,kf.

24 (HR-PRO or HRPRO).ti,ab,kf.

25 or/9-24

26 exp "Review Literature as Topic"/

27 "systematic review"/

28 meta-analysis/

29 (systematic review or scoping review or literature review or rapid review or narrative review or meta-analysis or metaanalysis).ti,ab,kf,hw,pt.

30 ("review of reviews" or "overview of reviews").ti,ab,kf.

31 ((review* or map*) adj5 (tool* or instrument* or scale* or measur*)).ti,ab,kf.

32 (literature adj3 review*).ti,ab,kf.

33 or/26-32

34 7 and 8 and 25 and 33

35 limit 34 to yr="1990 -Current"

***************************

BMJ Publishing Group Limited (BMJ) disclaims all liability and responsibility arising from any reliance placed on this supplemental material which has been supplied by the author(s)

BMJ Open
doi: 10.1136/bmjopen-2021-054679