# BMJ Open Effect of adverse perinatal outcomes on postpartum maternal mental health in low-income and middle-income countries: a protocol for systematic review

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### **ABSTRACT**

**Introduction** More than three-fourths of adverse perinatal outcomes (preterm, small for gestational age, low birth weight, congenital anomalies, stillbirth and neonatal death) occur in low-income and middle-income countries. These adverse perinatal outcomes can have both short-term and long-term consequences on maternal mental health. Even though there are few empirical studies on the effect of perinatal loss on maternal mental illness, comprehensive information on the impact of adverse perinatal outcomes in resource-limited settings is scarce. Therefore, we aim to systematically review and synthesise evidence on the effect of adverse perinatal outcomes on maternal mental

Methods and analysis The primary outcome of our review will be postpartum maternal mental illness (anxiety, depression, post-traumatic stress disorder and postpartum psychosis) following adverse perinatal outcomes. All peer-reviewed primary studies published in English will be retrieved from databases: PubMed, MEDLINE, CINAHL Ultimate (EBSCO), PsycINFO, Embase, Scopus and Global Health through the three main searching terms—adverse perinatal outcomes, maternal mental illness and settings, with a variant of subject headings and keywords. We will follow the Joanna Briggs Institute critical appraisal checklist to assess the quality of the studies we are including. The review findings will be reported in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses 2020 statement. Estimate-based meta-analysis will be performed. We will assess heterogeneity between studies using the I<sup>2</sup> statistics and publication bias will be checked using funnel plots and Egger's test. A subgroup analysis will be conducted to explore potential sources of heterogeneity (if available). Finally, the certainty of the evidence will be evaluated using the Grading of Recommendations, Assessment, Development and Evaluation approach.

**Ethics and dissemination** Since this systematic review does not involve human participants, ethical approval is not required. The review will be submitted for publication in a peer-reviewed journal.

PROSPERO registration number CRD42023405980.

### STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ Through linking adverse perinatal outcomes and postpartum maternal mental illness, we can examine the association that have not been examined at low-income and middle-income countries.
- ⇒ Use the Grading of Recommendations, Assessment. Development and Evaluation approach to determine the quality of evidence for each outcome.
- ⇒ We anticipate the possible limitation of language bias; we will include studies published in English.

### INTRODUCTION

Adverse perinatal outcomes include both perinatal morbidity (low birth weight (LBW), small for gestational age (SGA), preterm birth or congenital anomalies) and perinatal mortality (stillbirth and neonatal death). 12 Every year, approximately 2.4 million children die within the first 28 days of life,<sup>3</sup> and there are 2.6 million reported stillbirth<sup>4</sup> worldwide. Perinatal morbidity rates also remain unacceptably high globally with nearly 15 million preterm babies<sup>5</sup> and 20 million babies with LBW each year.<sup>6</sup> Alarmingly, 98% of these cases occur in low-income and middle-income countries.<sup>47</sup> Infants suffering from perinatal morbidity are at increased risk of mortality and developmental problems. For instance, preterm babies are at high risk of hospitalisation and can suffer lifelong effects such as cerebral palsy, mental retardation, visual and hearing impairments, and poor health. 18-11 These adverse perinatal outcomes can have significant impacts on the well-being of their parents.

Evidence suggests that depression during pregnancy can predict preterm birth and LBW, indicating a bidirectional relationship between maternal mental health and adverse perinatal outcomes. 12-14 As a result, adverse perinatal



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outcomes can have short-term and long-term consequences on maternal mental health and well-being. 15 16 Studies have shown that perinatal mortality was associated with an increased risk of depressive and anxiety disorder. 17-19 Similarly, studies showed that mothers of preterm babies or those with SGA experience higher levels of anxiety, depression, post-traumatic stress disorder (PTSD) and postpartum psychosis compared with parents of healthy term-born babies. 15 20-26 This could be due to the fact that preterm babies often require hospitalisation, complex interventions such as ventilation and incubation, and longer stays in neonatal care, which can be stressful for parents.<sup>27</sup> Other research has also found that mothers of LBW babies are more likely to experience depression<sup>21 24 28</sup> and PTSD<sup>29-31</sup> compared with mothers of average birth weight babies. Furthermore, parents of infants born with congenital anomalies are at high risk of developing symptoms of anxiety, 32 33 depression, 34-36 PTSD37 and psychological distress.38

While many systematic review studies have focused on the effect of maternal mental illness on perinatal outcomes, <sup>13</sup> <sup>14</sup> <sup>39–42</sup> only a few studies have examined the effects of adverse perinatal outcomes on maternal mental illness. <sup>16</sup> <sup>17</sup> <sup>24</sup> <sup>28</sup> <sup>37</sup> Nearly 90% of these studies were conducted in high-income countries, 43 with one systematic study assessing maternal mental illness following perinatal loss in a global context. 17 Given the high incidence of adverse perinatal outcomes within low-income and middle-income settings and the lack of research, there is a pressing need to analyse maternal mental illness following adverse perinatal outcomes in low-income and middle-income countries.

To our knowledge, no comprehensive systematic review has been conducted to estimate rates of anxiety, depression, PTSD and postpartum psychosis for mothers following adverse perinatal outcomes in low-income and middle-income countries. Therefore, our aim is to synthesise evidence on effects of adverse perinatal outcomes on postpartum maternal mental illness, including anxiety, depression, PTSD and postpartum psychosis and to identify potential moderators of these associations, such as study quality, type of adverse outcome, economic status of the study country and maternal age. We hypothesise that measures of depression, anxiety, PTSD, postpartum psychosis show a significant increase following adverse perinatal outcomes compared with the normal perinatal outcomes.

# **METHODS**

We will conduct a systematic review of empirical evidence regarding the effect of adverse perinatal outcomes on postpartum maternal mental illness in low-income and middle-income countries. The finding of this review will be reported based on the Preferred Reporting Items for Systematic Reviews and Meta-Analyses 2020 statement. 44

# **Search Strategy**

In this systematic review, the Patient/population-Intervention-Comparison-Outcomes-Type of study-Context approach is used to formulate and address the research questions. 45 The population will consist of adult women aged 18 years and older who have given birth. Exposed group will be postpartum women who experienced at least one of the adverse perinatal outcomes (preterm, SGA, LBW, congenital anomalies, stillbirth and neonatal death). The comparison group will consist of postpartum women who did not experience any adverse perinatal outcome. In this review, the outcomes will be postpartum maternal mental illness which includes anxiety, depression, PTSD and postpartum psychosis. All primary studies that report on the effect of adverse perinatal outcome on maternal mental illness for at least one of the conditions in low-income and middle-income countries (defined by World Bank) will be included.

Studies will be retrieved from seven electronic bibliographic databases, namely PubMed, MEDLINE, CINAHL Ultimate (EBSCO), PsycINFO, Embase, Scopus and Global Health. Google Scholar will also be considered to search for further literatures. Additional relevant studies from the included articles will be assessed and included (citation snowballing) in the final review. To build the search terms, subject headings and keywords related to adverse perinatal outcomes, postpartum maternal mental illness, and low-income and middle-income countries will be used (see sample MEDLINE search strategy on table 1). These terms will be linked using appropriate Boolean Operators such as "AND" and "OR" as per the requirements of the respective databases. 46

### **Inclusion and exclusion**

The review will include all peer-reviewed primary studies published in English from 1 January 2000 to the date of the review (to include the most up to date studies on the topic) that report on postpartum maternal mental illness for at least one adverse perinatal outcome in low-income and middle-income countries. As per the world bank's definition for 2023, countries with a gross national income per capita of US\$1085 or less are categorised as low-income countries, while those with a gross national per capita of US\$1086-US\$13 205) are deemed middle-income countries. 47 Among the identified countries, there are 28 low-income countries and 108 middleincome countries. The review will exclude abstracts, review papers, qualitative studies, theoretical and conceptual articles, conference papers, randomised controlled trials and letters or editorials. Finally, the full text of all relevant studies found to meet the inclusion criteria for the final review will be retained. The detailed inclusion and exclusion criteria are summarised in table 2.

### **Study selection**

Once the search strategy is developed and tested, two authors (SMF and TGH) will retrieve all relevant articles from all the databases and export them to EndNote V.20 to remove duplicates. Then, after removing the duplicates, studies will be exported to Rayyan (a systematic review management software package) for screening.<sup>48</sup>

Terms A Keywords ((			
	Adverse Perinatal outcomes	Common mental illness	Low-income and middle-income countries
	((Adverse or perinatal or pregnancy or gestation or birth or neonatal or obstetric or infant or fetal or congenital or child) ADJ3(outcome* or death or defect or anomal* or complication or event or loss or mortalit* or morbidit* or consequence or sequelae)). Lw.ab. OR (Stillbirth or still-birth or prematurity or preterm or small baby or PTB or SGA or low birthweight or LBW).tw,ab.	((Anxiety or generalized anxiety or depression or depressive or posttraumatic stress) ADJ2(disorder or symptom)).tw,ab. Or ((Postpartum or posthatal or puerperal or PNC or post-delivery or maternity or perinatal) ADJ2 (Anxiety or generalized anxiety or depression or depression or disorder)).tw,ab.	((Developing or undeveloped or low income or low-income* or middle income or middle-income or low resource or middle resource or low-and middle income or low-income or non-industrialised or third world) ADJ3 (countr* or nation* or setting*)). tw,ab. Or (Africa, sub-Sahara or sub-Saharan or Eastern Africa or Middle East Asia or Asia or Latin America or South America). tw,ab. Or (Africa, sub-Saharan or Euchina Faso or Burundi or Central African Republic or Chad or Congo Democratic Republic or Entrea or Ethiopia or Gambia or Guinea or Guinea-Bissau or Korea Democratic People Republic or Liberia or Madagascar or Malawi or Malawi or Mogrer or Rwanda or Sierra Leone or Somalia or Songh Sudan or Syrian Arab Republic or Togo or Uganda or Yemen Republic or Zambia or Angola or Algoria or Angola or Algoria or Angola or Algoria or Angola or Malawi or Republic or Cambodia or Cameroon or Comorox or Congo Republic or Cete d Ivoire or Dijbouti or Egypt Arab Republic or Eswatini or Ghana or Haiti or Honduras or India or Indonesia or Iran Islamic Republic or Kenya or Kiribati or Kyrgyz Republic or Eswatini or Ghana or Nigeria or Pakistan or Papua New Guinea or Micronesia Fed Sts or Mongolia or Morocco or Myanmar or Nepal or Nicaragua or Nigeria or Pakistan or Papua New Guinea or Philippines or Samoa or "Sao Tome and principe" or Senegal or Solomon Islands or Sri Lanka or Tanzania or Tajikistan or Timor-Leste or Tunisia or Uzbekistan or Vanuatu or Vietnam or "West Bank and Gaza" or Zimbabwe or Albania or American samoa or Armenia or Azerbaijan or Belarus or Belize or "Bosnia and Herzegovina" or Botswana or Buzzil or Bulgaria or China or Colombia or Costa Rica or Cuba or Dominica or Jordan or Kazakhstan or Kosovo or Libya or Malaysia or Malaysia or Marshall Islands or Muritius or Maxico or Jordan or Kazakhstan or Rosala or Palau or Paguay or Peru or Tonga or Turkiye or Turkiye or Turkiya or Saint Lucia or "Saint
Subject heading	Q		
	Pregnancy Complications/ or Pregnancy Outcome/ or exp Fetal Death/ or exp Fetal Diseases/ or exp Perinatal Mortality/ or exp Perinatal Death/ or exp Perinatal Death/ or exp Permature Birth/ or Infant, Low Birth Weight/ or exp Congenital Abnormalities/ or Infant, Newborn, Diseases/	exp Depression, Postpartum/ or exp Depression/ or exp anxiety// or exp psychological distress/ or exp Stress Disorders, Traumatic/	Developing Countries/ or exp Resource-Limited Settings/ or exp africa/ or exp Agriaal or exp "africa south of the Sahara"/ or exp Afghanistan/ or exp Abbaniaal or exp Ageriaal or exp American Samoaal or exp Angolad or exp Argentiaal or exp American Samoaal or exp Angolad or exp Argentinaal or exp American Samoaal or exp Angolad or exp Angolad or exp Bulzal or exp Bulgarial or exp Bentin/ or exp Bulzal or exp Cameroon/ or exp Cameroon/ or exp Cameroon/ or exp Cameroon/ or exp Bulzal African Republic/ or exp Cambolad or exp Cameroon/ or exp Dominican Republic or exp Equatorial Bulzal Republic/ or exp Cambolad or exp Cameroon/ or exp Ewatin/ or exp Filipolad or exp Equatorial Guinead or exp Genombia Argential or exp Republic or exp Emandor or exp Genombia Argential or exp India or exp Mauritius/ or exp Mauritius/ or exp Madagascar/ or exp Sendal or

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PICOTC	Definition	Inclusion criteria	Exclusion criteria
Population	Postpartum women	All women aged 18 and older, women who gave birth and women up to 1 year after giving birth	<ul> <li>Women who have experienced substance abuse and intimate partner violence</li> <li>Women who experienced common mental illness during pregnancy (before giving birth)</li> </ul>
Exposure / intervention	Adverse perinatal outcome	Postpartum women who experienced at least one of the adverse perinatal outcomes (preterm, SGA, LBW, congenital anomalies, stillbirth and neonatal death)	Another –
Comparison	With and without adverse perinatal outcome	<ul> <li>Articles reported the maternal mental illness among postpartum women experienced adverse perinatal outcome</li> <li>Articles reported maternal mental illness among postpartum women who did not experience adverse perinatal outcomes</li> </ul>	_
Outcome	Maternal mental illness incurred due to the presence of adverse perinatal outcomes during postpartum period	<ul> <li>Articles quantitatively reported the mental illness among postpartum women experienced adverse perinatal outcomes</li> </ul>	<ul> <li>Qualitatively assessed maternal mental illness</li> </ul>
Type of studies	Types of articles/evidence to be included in the review	▶ All primary observational studies reported on association between adverse perinatal outcomes and postpartum maternal mental illness	<ul> <li>Qualitative studies</li> <li>Editorials and conference papers</li> <li>Review papers</li> <li>RCT studies</li> </ul>
Context/ settings	Low-income and middle-income countries	All low-income and middle-income countries according to the world bank definition of 2023	► Country not low-income and middle- income in 2023, but low income andmiddle income prior to the review
Other criteria	Year	Article published after 1 January 2000	► Article published before 2000
	Language	Published in English	► Studies published in non-English language
	Studies species	Human studies	► Animal studies

Two authors (SMF and TGH) will screen the studies based on the title and abstract. Then the full texts of remaining articles will be retrieved and reviewed by each of two authors independently based on the inclusion and exclusion criteria as presented in table 2. Any disagreement will be discussed and handled by the senior author (AD).

### **Data extraction and analysis**

A consistent Excel form will be used to record extracted data from the studies included for assessment of study quality. The abstracted data will include: (1) The study characteristics: article title, authors, year of publication, country of the study, year of publication, study design (cohort, case–control and cross-sectional); (2) Study population; (3) Participants characteristics: mean/median age, and sample size, baseline characteristics; (4) type of adverse perinatal outcomes (preterm, SGA, LBW, congenital anomalies, stillbirth and neonatal death); (5) Outcomes of the study: maternal mental illness (anxiety, depression, PTSD and postpartum psychosis) and (6) Each primary raw data will be taken to calculate the measurements of association (if any), associated variables with  $\beta$  coefficient or adjusted risk ratio with their CI.

A narrative summary of maternal mental illness will be made for studies that do not report effects by systematically organising the information based on the mental illness classification. Characteristics of the studies will be described in box 1. The postpartum maternal mental

# Box 1 Data abstraction sheet elements to describe the study in the final review

### Data abstraction sheet elements

- 1. Author(s), year.
- 2. Country.
- 3. Study design.
- 4. Study population/participants.
- 5. Sample size.
- 6. Outcome.
- 7. Measurement tool.
- 8. Methods of analysis.
- 9. Exposure.
- 10. Mental illness category.
- 11. Key finding.



illness will include anxiety, depression, PTSD and postpartum psychosis.

In the meta-analysis, we will calculate the OR for binary variables and the standard mean difference for continuous variables with 95% CI. A random-effects model will be selected under the assumption that studies included in the meta- analysis have been carried out with heterogeneous populations. Statistical heterogeneity between studies will be evaluated using the  $I^2$  statistics ( $I^2$  test), which can quantify the heterogeneity ranging from 0% (no heterogeneity) to 100% (the differences between the effect sizes can completely be explained by chance alone). 49 Subgroup and meta-regression analyses will be conducted to explore the potential source of heterogeneity. We will adopt Egger's and Begg's test to check the publication bias and funnel plot asymmetry to assess the effect of publication bias if the number of studies greater than or equal to 10, based on Cochrane Handbook recommendation.<sup>50</sup> In addition, we will perform sensitivity analysis by excluding studies one at a time to ensure the stability of the results. All data will be analysed using the STATA V.16.

### Risk of bias assessment

For the quality appraisal, the Joanna Briggs Institute critical appraisal checklist for observational studies will be used.

# **Quality of evidence assessment**

The quality of evidence for each outcome will be determined using the Grading of Recommendations, Assessment, Development and Evaluation approach.<sup>52</sup> This system helps to evaluate the quality of evidence in the domains of risk of bias, consistency, directness, precision

and publication bias. Accordingly, the quality of the evidence will be classified as high, moderate, low or very low. 53 Interpretation of the grading scores is presented in table 3.

# Patient and public involvement

No patients and public were not involved in the design, or conduct, or reporting, or dissemination plans of our research.

### **ETHICS AND DISSEMINATION**

This systematic review will not directly involve human participants and requires no ethical approval. The findings of this systematic review will be disseminated through peer-reviewed publication and will be presented at international conferences related to this field. In addition, findings will be used by other relevant organisations and stakeholders for decision-making.

# **Strengthens and limitations**

This study will quantify the effect of adverse perinatal outcomes on postpartum maternal mental illness in low-income and middle-income countries, focusing on common conditions such as anxiety, depression and PTSD. This will give a comprehensive picture of the burden of adverse perinatal outcomes on maternal mental illness in low-income and middle-income countries. Consequently, which help to design a system wide integrated strategies to improve maternal mental illness following adverse perinatal outcome in resource-limited settings. By publishing the research protocol, we reinforce the clarity of the strategy and minimise the risk of bias. However, we anticipate some limitations, including

		Criteria for grad	ading	
Quality of evidence	Interpretation of grading score	Type of evidence	Decrease grade if	Increase grade if
High	Further research is very unlikely to change our confidence in the estimate of effect	<ul> <li>▶ Randomised trial=high</li> <li>▶ Observational study=low</li> <li>▶ Any other evidence=low</li> </ul>	► Serious (-1) or very serious (-2) limitations to	➤ Strong evidence of association significant RR of >2 (<0.5) based on consisten
Moderate	Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate		(-1)	evidence from two or more observational studies, with no plausible confounders (+1)  Very strong evidence of
Low	Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate		<ul> <li>Some (-1)         or major (-2)         uncertainty about         directness</li> <li>Imprecision or</li> </ul>	association—RR of >5 (<0.2) based on direct evidence wit no major threats to validity (+2)  ► Evidence of dose response
Very low	Any estimate of effect is very uncertain		sparse data (-1)  High probability of reporting bias (-1)	gradient (+1)  All plausible confounders would have reduced the effect (+1)



the use of English language limiters and heterogeneity of the studies.

### Amendments of the review

Any change to this protocol will be updated on the PROSPERO registration database (registration number: CRD42023405980) and the updated part of the protocol will be published together with the full systematic review.

**Contributors** SMF and TGH conceptualised and designed the protocol, drafted the initial manuscript, and reviewed the manuscript. SMF, TGH and AD defined the concepts and search items, data extraction process and methodological appraisal of the studies. SMF and TGH planned the data extraction and statistical analysis. AD and TGH provided critical insights. All authors have approved and contributed to the final written manuscript.

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Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

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