

BMJ Open Association between gestational diabetes mellitus diagnostic criteria and adverse pregnancy outcomes – a systematic review and meta-analysis of adjusted effect sizes from studies using current diagnostic criteria

Elhassan Mahmoud , Abdalla Moustafa Elsayed, Basant Elsayed, Yasmin Elsalakawi, Aswathy Gopinath, Tawanda Chivese

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Qatar University College of Medicine, Doha, Ad Dawhah, Qatar

Correspondence to

Tawanda Chivese;
tchivese@qu.edu.qa

ABSTRACT

Objectives To quantify the association between Gestational Diabetes Mellitus (GDM) and adverse pregnancy outcomes and primarily compare the associations between diagnostic criteria following the International Association of Diabetes and Pregnancy Study Groups (IADPSG) recommendations and non-IADPSG criteria, which use higher blood glucose cut-offs.

Design Systematic review and meta-analysis of observational studies using contemporary GDM diagnostic criteria.

Data sources PubMed, Scopus, Google Scholar, Cochrane Database of Systematic Reviews and the Cumulative Index to Nursing and Allied Health Literature (CINAHL) were searched for articles published between 2010 and 2023. The search was carried out on 15 May 2023.

Eligibility criteria Studies were included if they were observational studies that reported adjusted effect sizes for GDM-related adverse outcomes and compared outcomes between women with and without GDM, used contemporary diagnostic criteria and were conducted after 2010.

Data extraction and synthesis Two reviewers independently extracted data and assessed study quality using the MethodologicAI Standards for Epidemiological Research (MASTER) scale. Bias-adjusted inverse variance heterogeneity meta-analysis models were used to synthesise adjusted effect sizes. The same meta-analytic models were used to synthesise the overall OR and their 95% CIs for comparisons of the criteria which followed the IADPSG recommendations to other criteria, mostly with higher blood glucose cut-offs (non-IADPSG).

Results We included 30 studies involving 642 355 participants. GDM was associated with higher odds of maternal outcomes, namely; caesarean section (adjusted OR (aOR) 1.24, 95% CI 1.01 to 1.51) and pregnancy-induced hypertension (aOR 1.55, 95% CI 1.03 to 2.34). GDM was associated with higher odds of neonatal outcomes, specifically; macrosomia (aOR 1.38, 95% CI 1.13 to 1.69), large for gestational age (aOR 1.42, 95% CI 1.23 to 1.63), preterm birth (aOR 1.41, 95% CI 1.21 to

STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ Utilised contemporary studies with modern Gestational Diabetes Mellitus (GDM) diagnosis criteria, relevant to current debate on screening and diagnosis of GDM.
- ⇒ Included only adjusted effect sizes, minimising the influence of confounding on the relationship between GDM and outcomes.
- ⇒ Limitations include the use of data from observational studies, where confounding factors could not be fully eliminated.
- ⇒ Had a limited number of studies using non-International Association of Diabetes and Pregnancy Study Group criteria, potentially affecting the conclusiveness of the analysis.

1.64), neonatal intensive care unit admission (aOR 1.42, 95% CI 1.12 to 1.78), neonatal hypoglycaemia (aOR 3.08, 95% CI 1.80 to 5.26) and jaundice (aOR 1.47, 95% CI 1.12 to 1.91). Further analyses showed no major differences in adverse pregnancy outcomes between IADPSG and non-IADPSG criteria.

Conclusions GDM is consistently associated with adverse pregnancy, maternal and foetal outcomes, regardless of the diagnostic criteria used. These findings suggest no significant difference in risk between lower and higher blood glucose cut-offs used in GDM diagnosis.

INTRODUCTION

Gestational diabetes mellitus (GDM) is defined as any degree of glucose intolerance with onset or first recognition during pregnancy and it affects 14% of pregnancies globally.^{1 2} After delivery, most women diagnosed with GDM revert to normal glycemic status, however, both the mother and their offspring are at a higher risk of developing type 2 diabetes and cardiovascular disease later



in life.^{3 4} The hyperglycaemia and pregnancy outcomes (HAPO) study showed that there was a linear increase in the risk of adverse pregnancy outcomes with increasing blood glucose, but there are no known cut-offs at which the risk of these outcomes becomes significantly elevated, unlike diabetes outside of pregnancy.^{5 6} Although many guideline bodies have adopted the International Association of Diabetes and Pregnancy Study Groups (IADPSG) recommendations, debate is still ongoing about the appropriate GDM screening strategies, blood glucose cut-offs and timing of GDM testing.⁷⁻⁹ Given the variation of the diagnostic criteria for GDM and screening approaches internationally, the prevalence of GDM varies widely.¹⁰ It is still not clear how the heterogeneity in screening approaches and diagnostic criteria affects the association between GDM and adverse pregnancy outcomes.

There is now abundant evidence that GDM not only causes adverse pregnancy outcomes and future type 2 diabetes and cardiovascular disease, but also has impact on a woman's mental health and is associated with higher costs to the health system.^{3 4 11-14} The landmark HAPO study findings showed that milder levels of hyperglycaemia can adversely affect pregnancy outcomes.⁵ These findings resulted in changes and revisions to many international GDM diagnosis guidelines, based on the recommendations of the IADPSG published in 2010.⁶ The WHO in 2013,¹⁵ the American Diabetes Association (ADA),¹⁶ the Australasian Diabetes in Pregnancy Society (ADIPS)¹⁷ and the Society for Endocrinology, Metabolism and Diabetes of South Africa (SEMDSA)¹⁸ are examples of guideline bodies which adapted their GDM diagnostic guidelines to align with the IADPSG recommendations. The IADPSG recommends universal screening for GDM of all pregnant women without pre-existing diabetes, between 24 and 28 weeks of gestation using a one-step 2 hour 75 g oral glucose tolerance test (OGTT) and to diagnose GDM if a woman has one abnormal test result based on the following cut-offs: fasting plasma glucose (FPG) ≥ 5.1 mmol/L, 1 hour OGTT plasma glucose ≥ 10.0 mmol/L or 2 hour OGTT plasma glucose ≥ 8.5 mmol/L.⁶

Despite the consensus on the adverse effects of hyperglycaemia on pregnancy outcomes, there is still a lack of agreement on GDM screening, testing and diagnosis, evidenced by the existence of more than 30 different GDM diagnostic guidelines in use in many regions and countries worldwide.¹⁹ The differences in these criteria are not only in diagnostic maternal blood glucose levels, but also in the screening approaches, glucose testing methods and timing of GDM screening. Some of the heterogeneity also stems from differences in resource allocation for GDM care, while others arise from uncertainty in the evidence about the appropriate GDM screening and testing approaches. Some notable guideline bodies that have not adopted the IADPSG recommendations are the National Institute for Health and Care Excellence (NICE) which recommends risk factor-based GDM screening and has maintained a higher fasting glucose of ≥ 5.6 mmol/L for GDM diagnosis.²⁰ Another example is the Diabetes

in Pregnancy Study Group India (DIPSI) which recommends testing in a non-fasting state and diagnosis of GDM only if the 2 hour plasma glucose is ≥ 7.8 mmol/L.²¹ The heterogeneity in GDM screening and diagnostic criteria is likely one reason why there is variability in the observed effect magnitudes of the association between GDM and adverse pregnancy outcomes.

Findings on the estimates of the effect of GDM on adverse pregnancy outcomes are still not conclusive. A recent meta-analysis²² evaluated the association between GDM and adverse pregnancy outcomes. However, this meta-analysis included studies based on older diagnostic criteria that are no longer in practice, potentially encompassing cohorts which include overt diabetes and pre-existing diabetes. This limitation may have led to overestimation of the impact of GDM by including undiagnosed pre-existing diabetes in the analysis. Further, some meta-analyses used unadjusted odds ratios (ORs), thereby reported associations that could be confounded.²³ To address these limitations, the current meta-analysis investigated the effect of GDM, diagnosed using contemporary criteria, on adverse pregnancy outcomes, and compared the effect sizes between criteria that conformed to the IADPSG recommendations and non-IADPSG criteria that generally used higher blood glucose cut-offs. By restricting our analysis to studies that report adjusted effect sizes, we aim to minimise the influence of confounders and provide a more accurate estimate of the true association between GDM and adverse pregnancy outcomes under current diagnostic practices.

RESEARCH QUESTIONS

What is the effect of GDM, diagnosed using contemporary criteria, on each adverse pregnancy outcome? Does the effect of GDM on adverse pregnancy outcomes differ between different GDM diagnostic criteria?

METHODS

Study design

A systematic review and meta-analysis of relevant studies was conducted. The study protocol is registered on the International Prospective Register of Systematic Reviews (PROSPERO) (CRD42020155061) and it follows the Preferred Reporting Items for Systematic Reviews and Meta-Analyses protocol extension (PRISMA-P).²⁴

Search strategy for identification of studies

Data sources and electronic searches

PubMed, Scopus, Google Scholar, Cochrane Central Register of Controlled Trials (CENTRAL), the Cumulative Index to Nursing and Allied Health Literature (CINAHL) were searched for articles published between 2010 and 2023. The search was carried out on 15 May 2023. Medical subject headings (MeSH words) and keyword searches for GDM and pregnancy outcomes were used in the search. Supplementary Tables 1-3

contains the search strategy. Further, the reference lists of included papers were also searched. EndNote was used to remove duplicate, and studies were screened for inclusion using the Rayyan systematic review management website (www.rayyan.ai). Two reviewers (EM, AE) independently screened the studies for inclusion within Rayyan. Following the initial screening, four reviewers (EM, AE, BE, YE) evaluated the papers for inclusion using full text, according to the specified inclusion criteria.

Studies inclusion criteria

Inclusion criteria

Studies were included if they were observational cohort, cross-sectional and case-control comparing adverse pregnancy outcomes between women with and those without GDM. The studies were included if they were conducted between 2010, when the IADPSG recommendations were published, to the year 2023 and if they reported adjusted ORs for the association between GDM and adverse pregnancy outcomes. Experimental studies were included only if they compared GDM diagnostic criteria as intervention and comparators.

Exclusion criteria

Studies were excluded if they were conducted prior to 2010, review articles, included animal studies, did not report an effect size or any outcomes related to this study, did not report adjusted effect sizes or included participants with pre-existing diabetes.

Outcomes of interest

Maternal outcomes

Maternal outcomes included caesarean section, pregnancy-induced hypertension (PIH) and pre-eclampsia. Caesarean sections included both elective and emergency. PIH was defined as a systolic blood pressure ≥ 140 mm Hg or diastolic blood pressure ≥ 90 mm Hg diagnosed at ≥ 20 weeks gestation. Pre-eclampsia was defined as hypertension ($\geq 140/90$ mm Hg) and proteinuria.

Foetal outcomes

Foetal outcomes included large-for-gestational-age (LGA), small-for-gestational-age (SGA), macrosomia, preterm birth, shoulder dystocia, neonatal hypoglycaemia, neonatal intensive care unit admission (NICU), jaundice and respiratory distress syndrome (RDS). Macrosomia was defined as birth weight greater than 4000 g. LGA was defined as birth weight above the 90th percentile for gestational age. SGA was defined as birth weight of less than 10th percentile for gestational age. Preterm birth was defined as birth before 37 completed weeks of gestation.

Data extraction and management

For duplicate publications, we only included the article that contains the most information, and all others were excluded. The following data were extracted from the

articles: study characteristics such as the publication year, duration of the study, region, country, study design, sample size, GDM diagnostic criteria used, numbers of participants with the outcomes of interest and the effect size with their corresponding CIs. Data were extracted into a predesigned and piloted Microsoft Office Excel spreadsheet. For each study, two reviewers independently extracted the data and compared thereafter. Disparity in data extracted was resolved via discussion between all the reviewers.

Assessment of risk of bias

The risk of bias and external validity of the included studies was assessed using the Methodological Standards for Epidemiological Research (MASTER) scale.²⁵ Two reviewers independently assessed each study, and differences were resolved by discussion. If no consensus was reached, a third reviewer was consulted to resolve the conflict.

Data synthesis

Study characteristics and other data were narratively described and were presented as tables. Because the included studies were observational, of varying quality, a bias-adjusted inverse variance heterogeneity (quality effects) model was used as to synthesise overall effect sizes for the meta-analysis, with quality weights derived from the MASTER scale. Estimates from the random-effects model were also computed for comparison purposes, since this is the most widely used model in literature. The I^2 statistic and the Cochrane's Q p-values were both used to assess the heterogeneity. Doi plots and funnel plots were used for the assessment of publication bias. To explore the association between GDM diagnostic criteria and the odds of adverse outcomes, further analyses were carried out by comparing IADPSG to non-IADPSG. Non-IADPSG criteria in this study were Carpenter-Coustan (CC) (two studies^{26,27}), 2008 Canadian Diabetes Association (CDA) (one study²⁸), ADA 2014 (one study²⁹), WHO 1999 (one study³⁰) and the ADIPS (one study³¹). The studies using CC criteria employed universal OGTT screening. The cut-offs used in these studies varied. For CC and ADA 2014 criteria, fasting glucose ≥ 5.3 mmol/L, 1-hour ≥ 10.0 mmol/L and 2-hour ≥ 8.6 mmol/L were used ($n=7612$). The WHO 1999 cut-offs included fasting glucose ≥ 7.0 mmol/L or 2-hour glucose ≥ 7.8 mmol/L ($n=42\,656$). The 2008 CDA criteria used fasting glucose ≥ 5.3 mmol/L, 1-hour ≥ 10.6 mmol/L and 2-hour ≥ 8.9 mmol/L ($n=270\,843$). The ADIPS cut-offs used included fasting glucose ≥ 5.5 mmol/L and 2-hour ≥ 8.0 mmol/L ($n=32\,013$). For analysis purposes, the non-IADPSG criteria were grouped together, since they used a higher FPG and are therefore expected to result in stronger associations with adverse pregnancy outcomes. The analysis was carried out using Stata V.17 software.

Patient and public involvement

No patients or members of the public were involved in this study.

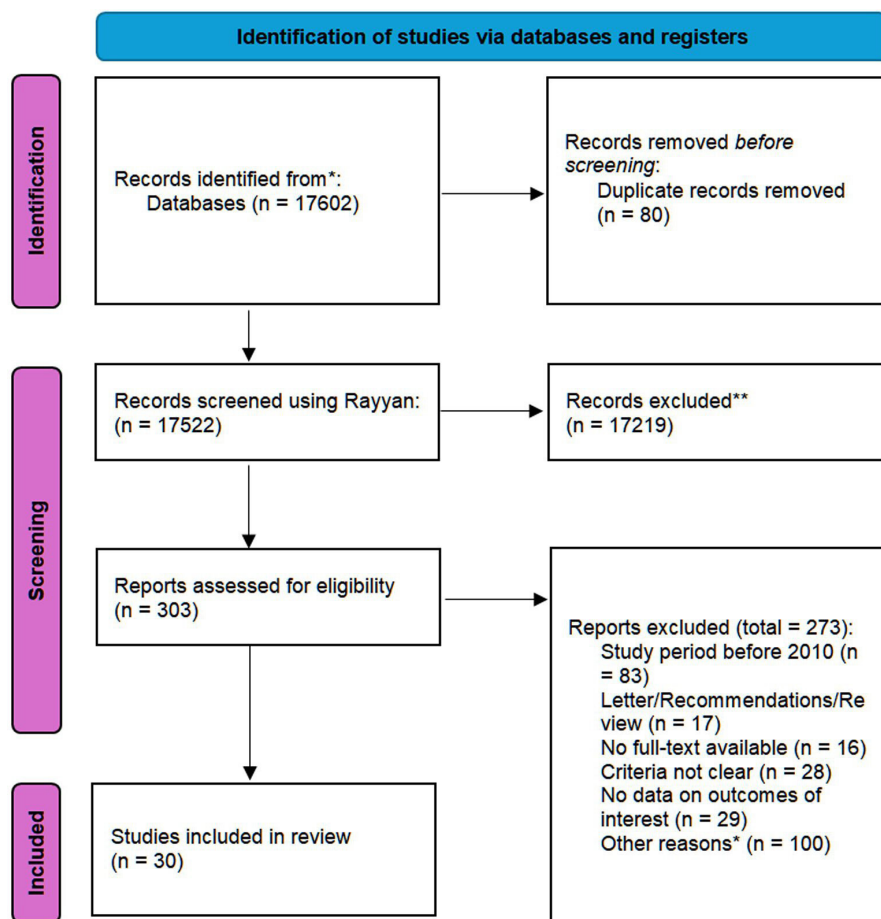


Figure 1 PRISMA flow chart showing the search. *Other reasons—did not exclude pre-existing diabetes, did not report relevant effect sizes (adjusted OR/RR). PRISMA, Preferred Reporting Items for Systematic reviews and Meta-Analyses.

RESULTS

Search results

A total of 17513 records were identified. There were 80 duplicate records that were removed. Figure 1 shows the PRISMA flow chart for the search process. Out of 305 study records selected at the initial title and abstract screening, 273 were excluded as they did not meet the inclusion criteria. The reasons for the exclusions were as follows: studies conducted before 2010 (n=83), letters/recommendations/reviews (n=17), studies where full texts were not available (n=16), studies where the criteria used were not clear (n=28), studies with no outcomes of interest (n=29) and studies excluded for other reasons (n=100). The list of excluded studies and reasons for exclusion are in online supplemental table 4. A total of 30 studies^{26–55} with 642 355 participants were finally included.

Characteristics of included studies

Table 1 shows the characteristics of the included studies. Of the 30 included studies, most (n=17) were from Asia,^{26 27 29 37–39 41–44 47 48 51–55} four were from Europe,^{30 34 36 49} three were from the Middle East,^{32 33 50} two were from Australia,^{31 40} two were from Africa,^{45 46} one was from South America³⁵ and one was from North America²⁸ (table 1). The studies were from these countries: Australia,^{31 40} Brazil,³⁵ Croatia,^{30 36} India,⁴⁸ Iran,²⁶ Saudi

Arabia,^{32 50} Qatar,³³ Italy,^{34 49} Canada,²⁸ Vietnam,^{29 38 53} South Korea,²⁷ China^{37 41–44 47 51 52 54 55} and Ethiopia.^{45 46} All the studies employed either cross-sectional or cohort designs. Four of these studies contained two independent populations that were analysed separately in the meta-analysis. In table 1, these populations are labelled as ‘Author, Year A’ for the first population and ‘Author, Year B’ for the second population. While the total number of studies is 30, the inclusion of these separate populations increased the total number of assessed populations in the meta-analysis to 34. The years of data collection were from 2010 to 2023. All studies have employed universal screening.

Quality of included studies

Overall, most of the studies had relatively high scores in the quality assessment using the MASTER scale⁵⁶ (online supplemental figure 1). Four studies^{33–35 55} scored 28/36, four studies^{29 38 40 48} had a score of 27/36 and four studies^{27 42 49 52} had a score of 26/36. The scores of the remaining studies ranged from 22/36 to 25/36. The main deficiencies were in equal retention, equal ascertainment, equal prognosis and sufficient analysis domains (online supplemental figure 1).

Maternal outcomes

Table 2 shows the results of the overall syntheses for the association between GDM and adverse pregnancy outcomes.

Table 1 Characteristics of included studies

Study	Study duration	Country	Sample size	Region	Study design	Criteria	Screening
Alfadhli <i>et al</i> , 2015 ³²	2011–2014	Saudi Arabia	954	Middle East	Cohort	IADPSG	Universal
Bashir <i>et al</i> , 2020 ³³	2015–2016	Qatar	2221	Middle East	Cohort	IADPSG	Universal
Capula <i>et al</i> , 2013 ³⁴	2010–2012	Italy	2448	Europe	Cohort	IADPSG	Universal
Carvalho <i>et al</i> , 2023 ³⁵	2020–2020	Brazil	1618	South America	Cross-sectional	IADPSG	Universal
Darbandi <i>et al</i> , 2022 ²⁶	2018–2018	Iran	3675	Asia	Cross-sectional	Non-IADPSG (CC)	Universal
Djelmis <i>et al</i> , 2016 ³⁶	2012–2014	Croatia	4646	Europe	Cohort	IADPSG	Universal
Erjavec <i>et al</i> , 2016 ³⁰	2010–2010	Croatia	42 656	Europe	Cross-sectional	Non-IADPSG (WHO-1999)	Universal
Erjavec <i>et al</i> , 2016 ³⁰	2014–2014	Croatia	39 092	Europe	Cross-sectional	IADPSG	Universal
He <i>et al</i> , 2023 ³⁷	2012–2021	China	115 097	Asia	Cohort	IADPSG	Universal
Hiersch <i>et al</i> , 2019 ²⁸	2012–2016	Canada	266 942	North America	Cohort	Non-IADPSG (CDA)	Universal
Hiersch <i>et al</i> , 2019 ²⁸	2012–2016	Canada	3901	North America	Cohort	Non-IADPSG (CDA)	Universal
Hirst <i>et al</i> , 2012 ³⁸	2010–2011	Vietnam	2772	Asia	Cohort	IADPSG	Universal
Kawasaki <i>et al</i> , 2023 ³⁹	2015–2019	Japan	1807	Asia	Cohort	IADPSG	Universal
Kim <i>et al</i> , 2019 ²⁷	2014–2016	Korea	1907	Asia	Cohort	Non-IADPSG (CC)	Universal
Kim <i>et al</i> , 2019 ²⁷	2014–2016	Korea	1969	Asia	Cohort	IADPSG	Universal
Laafira <i>et al</i> , 2016 ⁴⁰	2011–2014	Australia	3105	Australia	Cohort	IADPSG	Universal
Li <i>et al</i> , 2014 ⁴¹	2011–2011	China	54 275	Asia	Cross-sectional	IADPSG	Universal
Lin <i>et al</i> , 2022 ⁴²	2012–2020	China	2151	Asia	Cohort	IADPSG	Universal
Mak <i>et al</i> , 2019 ⁴³	2015–2015	China	1901	Asia	Cohort	IADPSG	Universal
Mei <i>et al</i> , 2021 ⁴⁴	2016–2018	China	333	Asia	Cohort	IADPSG	Universal
Muche <i>et al</i> , 2020 ⁴⁵	2018–2019	Ethiopia	694	Africa	Cohort	IADPSG	Universal
Muche <i>et al</i> , 2020 ⁴⁶	2018–2019	Ethiopia	684	Africa	Cohort	IADPSG	Universal
Nguyen <i>et al</i> , 2020 ²⁹	2015–2016	Vietnam	2030	Asia	Cohort	Non-IADPSG (ADA-2014)	Universal
Pan <i>et al</i> , 2015 ⁴⁷	2010–2012	China	17 808	Asia	Cohort	IADPSG	Universal
Punnose <i>et al</i> , 2022 ⁴⁸	2011–2017	India	2638	Asia	Cohort	IADPSG	Universal
Ronco <i>et al</i> , 2023 ⁴⁹	2010–2020	Italy	2364	Europe	Cohort	IADPSG	Universal
Wahabi <i>et al</i> , 2017 ⁵⁰	2013–2015	Saudi Arabia	9723	Middle East	Cohort	IADPSG	Universal
Wan <i>et al</i> , 2019A ³¹	2010–2013	Australia	3419	Australia	Cohort	Non-IADPSG (ADIPS)	Universal
Wan <i>et al</i> , 2019B ³¹	2010–2013	Australia	28 594	Australia	Cohort	Non-IADPSG (ADIPS)	Universal
Wang <i>et al</i> , 2021 ⁵²	2012–2013	China	8844	Asia	Cohort	IADPSG	Universal
Wang <i>et al</i> , 2023 ⁵¹	2018–2020	China	2031	Asia	Cohort	IADPSG	Universal
Yang <i>et al</i> , 2018 ⁵⁵	2011–2015	China	1232	Asia	Cohort	IADPSG	Universal
Yue <i>et al</i> , 2022 ⁵³	2016–2018	Vietnam	4703	Asia	Cohort	IADPSG	Universal
Zou <i>et al</i> , 2022 ⁵⁴	2016–2018	China	4121	Asia	Cohort	IADPSG	Universal

ADA, American Diabetes Association; ADIPS, Australasian Diabetes in Pregnancy Society; CC, Carpenter-Coustan; CDA, Canadian Diabetes Association; IADPSG, International Association of Diabetes and Pregnancy Study Groups.

A total of 18 studies^{26–28 30–33 35 36 40 42 43 45 48–50 52 53} reported data on total C-sections, with adjusted ORs (aORs) between 0.8^{31 42} and 2.3.³⁶ The overall aOR of total C-section was 1.24 (95% CI 1.01 to 1.51) with high

heterogeneity ($I^2=85.9\%$) (online supplemental figure 2). GDM was associated with a 25% increase in the odds of pre-eclampsia, in overall synthesis (aOR 1.25, 95% CI 1.00 to 1.56, $I^2=31.8\%$, $n=8$ studies^{27 28 31 33–35 38 49}) (online



Table 2 Results of overall syntheses for the association between GDM and each adverse pregnancy outcome

Outcome	Overall aOR (95% CI)	I ² (%)	LFK*	Number of studies
Maternal outcomes				
Total C section	1.24 (1.01, 1.51)	85.9	1.7	18
Pre-eclampsia	1.25 (1.00, 1.56)	31.8	1.6	8
PIH	1.55 (1.03, 2.34)	69.4	-2.8	7
Birth size-related neonatal outcomes				
Macrosomia	1.38 (1.13, 1.69)	75.0	4.2	19
LGA	1.42 (1.23, 1.63)	60.1	2.8	19
SGA	0.91 (0.80, 1.04)	40.1	0.8	14
Shoulder dystocia	1.20 (0.86, 1.66)	0.0	-1.0	4
Other neonatal outcomes				
Preterm birth	1.41 (1.21, 1.64)	62.3	0.0	17
NICU admission	1.42 (1.12, 1.78)	78.7	0.0	14
Neonatal hypoglycaemia	3.08 (1.80, 5.26)	86.3	1.1	7
Jaundice	1.47 (1.12, 1.91)	65.0	-5.0	6
RDS	1.22 (1.01, 1.47)	40.1	2.7	6

*The LFK is a measure of symmetry of publication bias plots and reflects major asymmetry when its absolute value is greater than 2 (or -2). aOR, adjusted OR; GDM, Gestational Diabetes Mellitus; LGA, large-for-gestational-age; NICU, neonatal intensive care unit admission; PIH, pregnancy-induced hypertension; SGA, small-for-gestational-age.

supplemental figure 3). Finally, in overall synthesis of seven studies,^{27 28 31 33–35 38 45 47–49} GDM showed an estimated 55% increase in the odds of PIH (aOR 1.55, 95% CI 1.03 to 2.34, I²=69.4%; online supplemental figure 4). The analyses suggested minor evidence of publication bias for all maternal outcomes, except for PIH which showed major evidence (online supplemental figures 5–7). In further analyses, compared with the IADPSG, non-IADPSG criteria showed similar odds of pre-eclampsia, PIH and total C-section (table 3).

Birth size-related neonatal outcomes

Data from 19 studies were included in the analysis of macrosomia.^{26 27 29 31–33 36 37 40 41 43 46–48 50 52–55} The overall aOR for macrosomia was 1.38 (95% CI 1.13 to 1.69) with moderate heterogeneity (I²=75.0%) (online supplemental figure 8). Overall, GDM was associated with 1.42-fold higher odds of LGA (aOR 1.42, 95% CI 1.23 to 1.63, I²=60.1%, n=19^{27 29 31 33–38 42–44 46–49 52–54}) (online supplemental figure 9). However, the synthesis suggested no significant associations between GDM and the odds of SGA (aOR 0.91, 95% CI 0.80 to 1.04, I²=40.1%, n=14;^{27 31 33 34 38 39 42–44 46 49 52–54} online supplemental figure 10) or shoulder dystocia (aOR 1.20, 95% CI 0.86 to 1.66, I²=0.0%, n=4;^{27 31 32 50} online supplemental figure 11). The analyses suggested evidence of publication bias for macrosomia and LGA, minor evidence for shoulder dystocia and no evidence of publication bias for SGA (online supplemental figures 12–15). In further analyses, compared with the non-IADPSG, the IADPSG criteria showed similar odds of macrosomia, LGA and SGA (table 3).

Other neonatal outcomes

In an analysis of 17 studies,^{26–28 31–34 38 39 43 46–48 50–53} GDM was associated with increased odds of preterm birth (online supplemental figure 16), with an overall aOR of 1.41 (95% CI 1.21 to 1.64) and moderate heterogeneity (I²=62.3%). For NICU admission, data from 14 studies^{27 28 31–35 38 39 42 43 48 50 53} showed that GDM was associated with a 1.42-fold increased odds (aOR 1.42, 95% CI 1.12 to 1.78) with high heterogeneity (I²=78.7%) (online supplemental figure 17). The overall aOR for neonatal hypoglycaemia was 3.08 (95% CI 1.80 to 5.26, I²=86.3%, n=7^{27 28 31–33 38 42}) (online supplemental figure 18). GDM was associated with 1.47-fold higher odds of neonatal jaundice (aOR 1.47, 95% CI 1.12 to 1.91, I²=65.0%, n=6;^{27 28 31–34} online supplemental figure 19). Moreover, GDM was associated with a 1.22-fold increased odds of neonatal RDS (aOR 1.22, 95% CI 1.01 to 1.47, I²=40.1%, n=6;^{28 31–34 42} online supplemental figure 20). The analyses suggested evidence of publication bias for jaundice and RDS, minor evidence for neonatal hypoglycaemia and no evidence of publication bias for preterm birth and NICU admission (online supplemental figures 21–25). Analyses by diagnostic criteria showed that, compared with non-IADPSG, IADPSG criteria showed similar odds of jaundice, RDS, neonatal hypoglycaemia, preterm birth and NICU admission (table 3).

DISCUSSION

In this meta-analysis of 30 studies, we found strong associations between GDM diagnosed using contemporary

Table 3 Results of analyses by criteria for the association between GDM and each adverse pregnancy outcome

Outcome	Criteria	Overall aOR (95% CI)	P for interaction
Maternal outcomes			
Total C-section	IADPSG	1.34 (1.12, 1.60)	0.398
	Non-IADPSG	1.20 (1.02, 1.43)	
Pre-eclampsia	IADPSG	1.08 (0.60, 1.94)	0.565
	Non-IADPSG	1.29 (1.11, 1.49)	
PIH	IADPSG	1.34 (0.82, 2.16)	0.636
	Non-IADPSG	1.57 (0.98, 2.54)	
Birth size-related neonatal outcomes			
Macrosomia	IADPSG	1.42 (1.24, 1.63)	0.577
	Non-IADPSG	1.04 (0.34, 3.13)	
LGA	IADPSG	1.41 (1.20, 1.66)	0.759
	Non-IADPSG	1.48 (1.14, 1.94)	
SGA	IADPSG	0.94 (0.80, 1.10)	0.298
	Non-IADPSG	0.81 (0.65, 1.01)	
Shoulder dystocia	IADPSG	1.36 (0.63, 2.95)	0.761
	Non-IADPSG	1.16 (0.60, 2.26)	
Other neonatal outcomes			
Preterm birth	IADPSG	1.44 (1.21, 1.71)	0.797
	Non-IADPSG	1.39 (1.15, 1.86)	
NICU admission	IADPSG	1.32 (1.11, 1.58)	0.723
	Non-IADPSG	1.41 (1.04, 1.92)	
Neonatal hypoglycaemia	IADPSG	3.09 (1.52, 6.29)	0.956
	Non-IADPSG	3.01 (1.64, 5.51)	
Jaundice	IADPSG	1.54 (1.24, 1.92)	0.816
	Non-IADPSG	1.46 (0.96, 2.22)	
RDS	IADPSG	1.32 (1.01, 1.74)	0.574
	Non-IADPSG	1.19 (0.92, 1.54)	
aOR, adjusted OR; GDM, Gestational Diabetes Mellitus; LGA, large-for-gestational-age; NICU, neonatal intensive care unit admission; PIH, pregnancy-induced hypertension; RDS, respiratory distress syndrome; SGA, small-for-gestational-age.			

aOR, adjusted OR; GDM, Gestational Diabetes Mellitus; LGA, large-for-gestational-age; NICU, neonatal intensive care unit admission; PIH, pregnancy-induced hypertension; RDS, respiratory distress syndrome; SGA, small-for-gestational-age.

criteria and adverse pregnancy outcomes. The highest associations were observed for neonatal hypoglycaemia, PIH, jaundice, NICU admission, macrosomia, LGA and preterm birth. We found no major differences in the effect of GDM between IADPSG-based criteria and criteria that used higher glucose cut-offs.

We found no major differences between IADPSG and non-IADPSG criteria on the effect of GDM on adverse pregnancy, maternal and foetal outcomes. When comparing IADPSG to stricter GDM criteria, this meta-analysis showed that no outcome differed by criteria. Our findings are similar to those of older meta-analyses which have also found that the risk of adverse pregnancy outcomes was not largely different across the different diagnostic criteria.^{23 57–59} A key difference between our synthesis and the older previously publishes studies is that we included contemporary studies, with adjusted effect magnitudes, that were conducted after 2010 when the IADPSG recommendations were published. Our findings

and those of previously published studies raise the question about the benefits of using lower glucose cut-offs for the diagnosis GDM. It has been argued that the use of criteria with lower fasting glucose cut-offs combined with universal screening, like the IADPSG, leads to an increase in GDM prevalence, without a concurrent increase in benefit (ie, reduced pregnancy outcomes and postpartum type 2 diabetes).¹⁰

Our findings have several implications. For healthcare systems, adopting the IADSPG criteria, that is, universal screening and lower glycaemic thresholds compared with targeted screening and generally higher glycaemic diagnostic thresholds, may strain resources, as more women would require screening, monitoring and interventions. This could lead to an increase in healthcare costs,^{60 61} which will lead to an increased burden, especially in settings where resources are already constrained. On the other hand, selective or targeted screening may result in some proportions of women progressing with

undiagnosed hyperglycaemia in pregnancy, and the consequent higher risk of adverse pregnancy outcomes. The NICE, for example, has opted to keep their guidelines which use risk factor-based screening and higher glycaemic thresholds. It is crucial to balance the costs and benefits of adopting either the IADPSG recommendations or selective screening, higher glycaemic threshold approaches such as that used by the NICE. These considerations may be different for different health systems, depending on affordability and healthcare system capacity. For clinicians, these findings highlight the need for careful consideration when diagnosing and managing GDM, as they should be mindful of the potential for overdiagnosis and overtreatment, and they should tailor management strategies based on each patient's individual risk factors, ensuring that interventions are justified and beneficial. For women, the increased likelihood of a GDM diagnosis that comes with universal screening and lower glycaemic thresholds may result in increased anxiety and an increased likelihood of medical interventions, without a clear improvement of outcomes. GDM diagnosis has been associated with a higher occurrence of mental health problems in pregnant women.^{62 63} It is therefore critical to provide women with clear and balanced information along with the implications, and to promote shared decision-making. More research is needed to identify appropriate blood glucose cut-offs where the benefit of GDM diagnosis outweighs the unintended negative consequences.

GDM was associated with around a 25% increase in the odds of pre-eclampsia and total C-section and 56% increase in the odds of PIH. A previous meta-analysis showed a 50% increase in pre-eclampsia and a 40% increase in C-sections in women with than in those without gestational diabetes mellitus.²² The HAPO study found that the occurrence of pre-eclampsia was positively associated with blood glucose level even after adjusting for clinical centre, age, Body Mass Index, height, smoking status, alcohol consumption, family history of diabetes, gestational age at OGTT and urinary tract infection.^{13 64} GDM causes increase in the insulin secretion by the foetal pancreas which itself is an anabolic hormone and leads to increase in the foetal weight. Fetuses with high birth size are usually delivered by caesarean sections as vaginal deliveries carry high risks to both mothers and babies.⁶⁵ The pathophysiology of pre-eclampsia is not well understood, and the association observed in these studies may be bidirectional. Irrespective of direction of association, the findings of this meta-analysis confirm the need to screen and monitor women with GDM for pre-eclampsia and PIH. Notably, pre-eclampsia and PIH are all associated with higher rates of both emergency and elective C-sections, and therefore may partly explain the higher risk of C-section in women with GDM.

The current meta-analysis showed that GDM was associated with higher the odds of neonatal hypoglycaemia, LGA, macrosomia, preterm birth, jaundice, NICU admission, RDS and shoulder dystocia. The higher odds of

birth-size-related complications, LGA, macrosomia and shoulder dystocia, are likely because of maternal hyperglycaemia, which leads to a high glucose intrauterine environment which promotes foetal hyperglycaemia and hyperinsulinemia, which in turn induce excess fat deposition in the fetus.^{66 67} Notably, the highest OR was observed for neonatal hypoglycaemia, with three-fold higher odds for GDM exposed neonates compared with the non-GDM exposed neonates. However, it is important to consider that this risk could be exaggerated due to the possibility of allocation bias for this outcome. Neonates born to mothers with GDM are more likely to be routinely tested for blood glucose levels shortly after birth due to the known risks of hypoglycaemia, whereas neonates of non-GDM pregnancies do not typically undergo such testing unless clinically indicated. This difference in clinical practice likely increases the detection rate of hypoglycaemia in the GDM group, which could lead to an overestimation of the association between GDM and neonatal hypoglycaemia. Previous meta-analyses have generally found that GDM was associated with adverse pregnancy outcomes.^{21 22} However, our findings differ from those of many of these previous meta-analyses in that our aORs, although still suggesting a higher risk of adverse pregnancy outcomes with GDM, are generally lower than those reported by the other meta-analyses.²³ This discrepancy is mostly due to the other meta-analyses having used unadjusted effect sizes. GDM is thought to cause RDS by interfering with the production of surfactant lipids and proteins.⁶⁸ Notably, some previous meta-analyses have reported contrasting findings in terms of the associations observed. Ye *et al*, using a meta-analysis of unadjusted ORs and studies with criteria that are no longer in use, found no association between shoulder dystocia and GDM was not significant.²² Tehrani *et al* used a meta-analysis of unadjusted ORs and reported a 20% decrease in the odds SGA, contrary to our finding.²³

A strength of this study is the use of contemporary studies using contemporary GDM diagnosis criteria, therefore contributing to the current debate about the appropriate screening tests and testing strategy for GDM. We only included adjusted effect sizes, thus minimising the effect of confounding on the relationship between GDM and the outcomes, which is the main limitation of existing meta-analyses. However, this study has some limitations. Since this study uses data from observational studies, the role of confounding cannot be fully eliminated. Our findings require confirmation by experimental randomised controlled trials which compare these criteria. Additionally, most of the included studies were conducted in Asia (54%), and relatively fewer studies from the other regions. This may limit the generalisability of our findings to non-Asian populations. Finally, the small number of studies using non-IADPSG criteria, most of which employed cut-offs relatively close to those recommended by IADPSG, limits the strength of the comparison between IADPSG and non-IADPSG criteria, as the non-IADPSG group may

not fully represent the diversity of diagnostic approaches in use.

Conclusion

GDM showed consistent associations with pregnancy, maternal and foetal outcomes, with no major differences in the effects when different contemporary criteria were used.

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ORCID iD

Elhassan Mahmoud <http://orcid.org/0000-0001-8528-0272>

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Supplementary Materials for “Association between gestational diabetes mellitus diagnostic criteria and adverse pregnancy outcomes– a systematic review and meta-analysis of adjusted effect sizes from studies using current diagnostic criteria”

Supplementary Table 1: PubMed Search Strategy

Population:		
#1	MeSH terms:	Diabetes, Gestational
#2	Text Word:	Gestational Diabetes OR GDM OR Gestational Diabetes Mellitus OR Pregnancy-induced diabetes OR Diabetes in Pregnancy OR Hyperglycaemia in Pregnancy OR Hyperglycemia in Pregnancy OR gestational hyperglycemia OR gestational diabetes* OR gestational glucose OR pregnancy diabetes* OR pregnancy glucose OR maternal hyperglycemia OR maternal diabetes* OR maternal glucose OR HIP OR glucose intolerance in pregnancy
#3	#1 OR #2	
Outcomes		
#4	Text Word:	Fetal outcomes OR Foetal outcomes OR Macrosomia OR Large for Gestational Age OR Perinatal Mortality OR Shoulder Dystocia OR Congenital Malformation OR Miscarriage OR Spontaneous Abortion OR Neonatal Hypoglycaemia OR Neonatal Hypoglycemia OR Hyperbilirubinaemia OR Hyperbilirubinemia OR Birth Asphyxia OR Admission to the Neonatal Intensive Care Unit OR Overweight OR Obesity OR Offspring OR Child OR Childhood OR Children OR spontaneous miscarriage OR pregnancy loss OR instrumental birth OR caesarean section OR C-section OR hypertensi* OR PIH OR pregnancy-induced hypertension OR preeclampsia OR eclampsia OR mortality OR death OR stillbirth OR congenital OR anomaly OR impair* OR disability* OR prematur* OR preter* OR low birth weight OR LBW OR large for gestational age OR LGA OR small for gestational age OR SGA OR neonatal hypo* OR jaundice OR respiratory distress syndrome OR RDS OR NICU OR neonatal intensive care unit
#5	#3 AND #4	
#6	#5 NOT (review OR metaanalysis OR systematic review OR meta-analysis OR literature review)	

Filters

1. 2010-2023

2. Humans

Supplementary Table 2: Cochrane Search Strategy

Population:		
#1	Key Word:	Gestational Diabetes OR GDM OR Gestational Diabetes Mellitus OR Pregnancy-induced diabetes OR Diabetes in Pregnancy OR Hyperglycaemia in Pregnancy OR Hyperglycemia in Pregnancy
#4	#1 OR #2 OR #3	
#5	Pregnancy	
Outcomes		
#6	Key Word:	Fetal Outcomes OR Foetal Outcomes OR Macrosomia OR Large for Gestational Age OR Perinatal Mortality OR Shoulder Dystocia OR Congenital Malformation OR Miscarriage OR Spontaneous Abortion OR Neonatal Hypoglycaemia OR Neonatal Hypoglycemia OR Hyperbilirubinaemia OR Hyperbilirubinemia OR Birth Asphyxia OR Admission to the Neonatal Intensive Care Unit OR Overweight OR Obesity OR Long Term Outcomes in Offsprings OR co-ordinated care OR coordinated integrated care OR co-ordinated integrated care OR multicare OR multiservice OR multiclinic
#10	#4 AND #5 AND #8 AND #9	

Supplementary Table 3: Scopus Search Strategy

Population:		
#1	Key Word:	Gestational Diabetes OR GDM OR Gestational Diabetes Mellitus OR Pregnancy-induced diabetes OR Diabetes in Pregnancy OR Hyperglycaemia in Pregnancy OR Hyperglycemia in Pregnancy
Outcomes:		
#2	Key Word:	Macrosomia OR Mortality OR Shoulder Dystocia OR Congenital OR Malformation OR Miscarriage OR Abortion OR Hypoglycaemia OR Hypoglycemia OR Hyperbilirubinaemia OR Hyperbilirubinemia OR Birth Asphyxia OR Overweight OR Obesity
#3	#1 AND #2	

Supplementary Table 4: Table of excluded studies

Study	Reason for exclusion
Stanescu 2014	Arguments/controversies
Abd Latif 2022	No relevant effect size
Abdelwahab 2023	No full-text article
Abell 2017	Study period before 2010
Absalom 2019	No relevant effect size
Abu 2022	No full-text article
Alberico 2014	Criteria not clear
Ali 2018	No relevant effect size
Al-Shwyiat 2022	No outcomes of interest
Anderberg 2010	Study period before 2010
Asemi 2021	Letter
Au 2016	No relevant effect size
Aulinas 2013	Study period before 2010
Aung 2015	No relevant effect size
Aviram 2016	Study period before 2010
Badakhsh 2016	Criteria not clear
Baharvand 2022	No outcomes of interest
Bahl 2022	No outcomes of interest
Bai 2023	No outcomes of interest
Bartáková 2017	No relevant effect size
Bashir 2018	No relevant effect size
Bashir 2019	No relevant effect size
Bashir 2021	No relevant effect size
Basri2018	No relevant effect size
Bassaw 2012	Criteria not clear
Basu 2012	Study period before 2010
Bauer 2023	No relevant effect size
Bayoumi 2021	Correction
Beetham 2022	No outcomes of interest
Benhalim 2019	No relevant effect size
Benhalima 2013	No relevant effect size
Berggren 2011	No relevant effect size
Berghella 2019	Letter
Bhavadharini 2021	No relevant effect size
Bianchi 2018	No relevant effect size
Black 2010	Study period before 2010
Blickstein 2018	Study period before 2010
Blomberg 2023	No outcomes of interest
Bodmer-Roy 2012	Study period before 2010

Bogdanet 2017	No relevant effect size
Bomba-Opoń 2022	No outcomes of interest
Bordin 2020	Study period before 2010
Boriboonhirunsarn 2023	No outcomes of interest
Brankica 2016	No relevant effect size
Briana 2022	No full-text article
Brown 2022	No relevant effect size
Buffarini 2019	Study period before 2010
Cai 2016	No relevant effect size
Catalano 2012	Study period before 2010
Chen 2021	No outcomes of interest
Chen 2022	No outcomes of interest
Chen 2022	No relevant effect size
Chen 2023	No outcomes of interest
Cheng 2019	Included pre-gestational diabetes
Cheung 2018	No relevant effect size
Chew 2013	Included pre-gestational diabetes
Cho 2016	Study period before 2010
Choi 2022	No relevant effect size
Chung 2022	Criteria not clear
Cosson 2013	No relevant effect size
Cosson 2022	Criteria not clear
Côté-Corriveau 2021	Letter
Dalfrà 2011	Study period before 2010
Davis 2018	Study period before 2010
de Wit 2021	No relevant effect size
Deng 2022	Criteria not clear
Ding 2018	No relevant effect size
Domanski 2018	Study period before 2010
Donovan 2017	No relevant effect size
Duran 2014	No relevant effect size
Ehmann 2019	Criteria not clear
Ekeroma 2015	No relevant effect size
Esakoff 2011	Study period before 2010
Ethridge 2014	No relevant effect size
Feghali 2018	Study period before 2010
Feleke 2022	No outcomes of interest
Feleke 2022	No relevant effect size
Feng 2017	No relevant effect size
Foeller 2015	Study period before 2010
Gao 2022	No outcomes of interest

García-Patterson 2020	No outcomes of interest
Gasim 2012	Study period before 2010
Glover 2016	Study period before 2010
Goedegebure 2018	No relevant effect size
Gojnic 2022	No relevant effect size
Gopalakrishnan 2015	No outcomes of interest
Gorban 2021	No relevant effect size
Gorgal 2012	No relevant effect size
Greenberg 2021	No full-text article
Gregory 2022	No outcomes of interest
Grotenfelt 2019	Study period before 2010
Gu 2019	No outcomes of interest
He 2020	No relevant effect size
Hernandez-Rivas 2013	Study period before 2010
Hildén 2019	No relevant effect size
Hildén 2020	Study period before 2010
Hillier 2022	Study period before 2010
Hosseini 2018	No outcomes of interest
Huang 2016	No full-text article
Huhn 2017	No full-text article
Ikenoue 2014	No relevant effect size
Immanuel 2021	Criteria not clear
Jain 2016	No relevant effect size
Jao 2013	Letter
Jin 2020	No relevant effect size
Kalra 2013	No relevant effect size
Karcaaltincaba 2011	Study period before 2010
Kaul 2022	Criteria not clear
Keikkala 2020	No relevant effect size
Kgosidialwa 2015	Study period before 2010
Kim 2021	No relevant effect size
Kirke 2014	No outcomes of interest
Koivunen 2017	Criteria not clear
Koivunen 2020	Study period before 2010
König 2014	Study period before 2010
Koning 2018	No relevant effect size
Kosman 2016	Included pre-gestational diabetes
Kösüs 2013	No outcomes of interest
Kragelund 2021	Study period before 2010
Kragelund 2021	Study period before 2010
Kumari 2018	No relevant effect size

Kung 2022	Criteria not clear
Kwong 2019	Study period before 2010
Lapolla 2011	No relevant effect size
Lara-Barea 2022	No outcomes of interest
Lee 2018	No relevant effect size
Lee 2020	No relevant effect size
Leybovitz-Haleluya 2018	No relevant effect size
Li 2020	No outcomes of interest
Li 2020	No relevant effect size
Liu 2012	Study period before 2010
Liu 2020	No outcomes of interest
Liu 2020	Criteria not clear
Lloreda-Garcia 2016	Criteria not clear
Lu 2016	Study period before 2010
Lu 2019	Study period before 2010
Lu 2023	Criteria not clear
Lucovnik 2020	No outcomes of interest
Luengmettakul 2015	Criteria not clear
Macaulay 2018	No relevant effect size
Macrì 2018	Study period before 2010
Makwana 2017	No relevant effect size
Maresh 2021	No outcomes of interest
Mayo 2015	Study period before 2010
McIntyre 2018	No relevant effect size
Mdoe 2021	No relevant effect size
Meek 2015	Study period before 2010
Miailhe 2015	No relevant effect size
Miao2017	No relevant effect size
Mikkelsen 2011	Study period before 2010
Minsart 2014	No relevant effect size
Mitanchez 2014	Review
Morikawa 2017	No relevant effect size
Mwanri 2014	No relevant effect size
Myszkowski 2023	No relevant effect size
Nabi 2022	No relevant effect size
Nayak 2013	No relevant effect size
Nelson 2023	Initiative
Nguyen 2016	Study period before 2010
Nicolosi 2020	Study period before 2010
No article	No full-text article
Ogonowski 2015	Study period before 2010

Olerich 2022	Criteria not clear
Oster 2014	Study period before 2010
O'Sullivan 2012	No full-text article
O'Sullivan 2016	Study period before 2010
Ovesen 2015	Study period before 2010
Park 2015	Study period before 2010
Parveen 2022	No relevant effect size
Pavic 2021	Study period before 2010
Perak 2021	No outcomes of interest
Picon 2022	No full-text article
Pintaudi 2018	No relevant effect size
Poulain 2015	Study period before 2010
Pouliot 2019	No relevant effect size
Protsenko 2010	No full-text article
Qadir 2012	No full-text article
Ramanjaneya 2021	Corrigendum
Redman 2021	Criteria not clear
Rehder 2011	No full-text article
Reichelt 2017	Study period before 2010
Reitzle 2023	No relevant effect size
Ritchie 2023	No relevant effect size
Rotem 2022	No relevant effect size
Ryan 2018	No relevant effect size
Ryan 2020	Study period before 2010
Sacks 2015	Study period before 2010
Sagili 2015	No relevant effect size
Sajani 2014	No full-text article
Sarkar 2022	No full-text article
Saxena 2011	Study period before 2010
Schmidt 2022	No full-text article
Schneider 2011	Study period before 2010
Seely 2023	No relevant effect size
Selen 2022	No relevant effect size
Sesmilo 2017	Study period before 2010
Sesmilo 2020	Study period before 2010
Seval 2016	Study period before 2010
Shah 2020	Study period before 2010
Shahbazian 2016	No relevant effect size
Shahbazian 2016	No relevant effect size
Shang 2014	No relevant effect size
Shang 2014	Study period before 2010

Shi 2020	No outcomes of interest
Shindo 2020	Study period before 2010
Shub 2019	Study period before 2010
Siegel 2017	Study period before 2010
Silva 2021	Review
Silveira 2021	Criteria not clear
Singh 2018	Criteria not clear
Sirimarco 2017	No relevant effect size
Sletner 2017	Study period before 2010
Soliman 2018	No relevant effect size
Soliman 2018	No relevant effect size
Somasundaram 2016	Review
Song 2014	Study period before 2010
Song 2019	No outcomes of interest
Song 2020	No relevant effect size
Sperling 2023	No relevant effect size
Srichumchit 2015	Study period before 2010
Stuebe 2015	Study period before 2010
Sudasinghe 2018	No relevant effect size
Sugiyama 2017	Study period before 2010
Sun 2020	No relevant effect size
Sunder 2022	No relevant effect size
Sunjaya 2018	Includes pre-gestational diabetes
Surendran 2019	Review
Sweeting 2016	Study period before 2010
Tan 2017	No relevant effect size
Todi 2020	No outcomes of interest
Tomić 2013	Study period before 2010
Tong 2022	Criteria not clear
Tong 2022	Criteria not clear
Tundidor 2012	Study period before 2010
Uma 2017	No relevant effect size
Vale 2023	Criteria not clear
Valias 2022	No relevant effect size
Vélez 2020	Study period before 2010
Violante-Ortíz 2023	No full-text article
Voldner 210	Study period before 2010
von Katterfeld 2012	Study period before 2010
Wahabi 2013	No relevant effect size
Wahi 2011	Study period before 2010
Wahlberg 2016	Study period before 2010

Wang 2013	Study period before 2010
Wei 2014	Study period before 2010
Wei 2016	Criteria not clear
Wells 2015	Study period before 2010
Wielandt 2015	Study period before 2010
Wilmot 2014	Recommendations
Wilmot 2014	Recommendations
Wilmot 2014	Recommendations
Wilmot 2014	Recommendations
Wolka 2022	Included pre-gestational diabetes
Wong 2012	Study period before 2010
Xaverius 2022	Criteria not clear
Yan 2017	Study period before 2010
Yang 2019	No relevant effect size
Yang 2019	Study period before 2010
Yang 2023	Criteria not clear
Ye 2021	No relevant effect size
Yesildager 2016	Criteria not clear
Yin 2022	Comment
Youngwanichsetha 2014	Criteria not clear
Yu 2014	No relevant effect size
Zahid 2022	Study period before 2010
Zawiejska 2014	Study period before 2010
Zeki 2018	Study period before 2010
Zhang 2018	No relevant effect size
Zhao 2020	Criteria not clear
Zhao 2023	No relevant effect size
Zheng 2022	No relevant effect size
Żurawska-Kliś 2016	No relevant effect size

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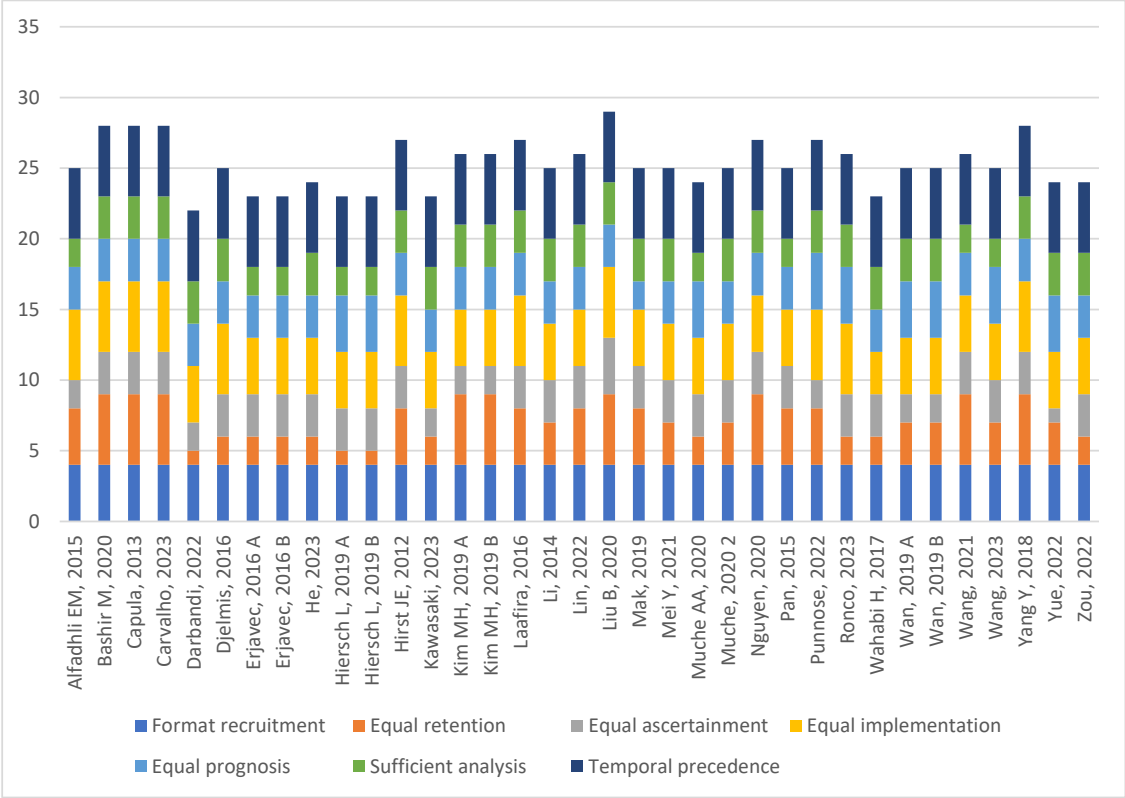
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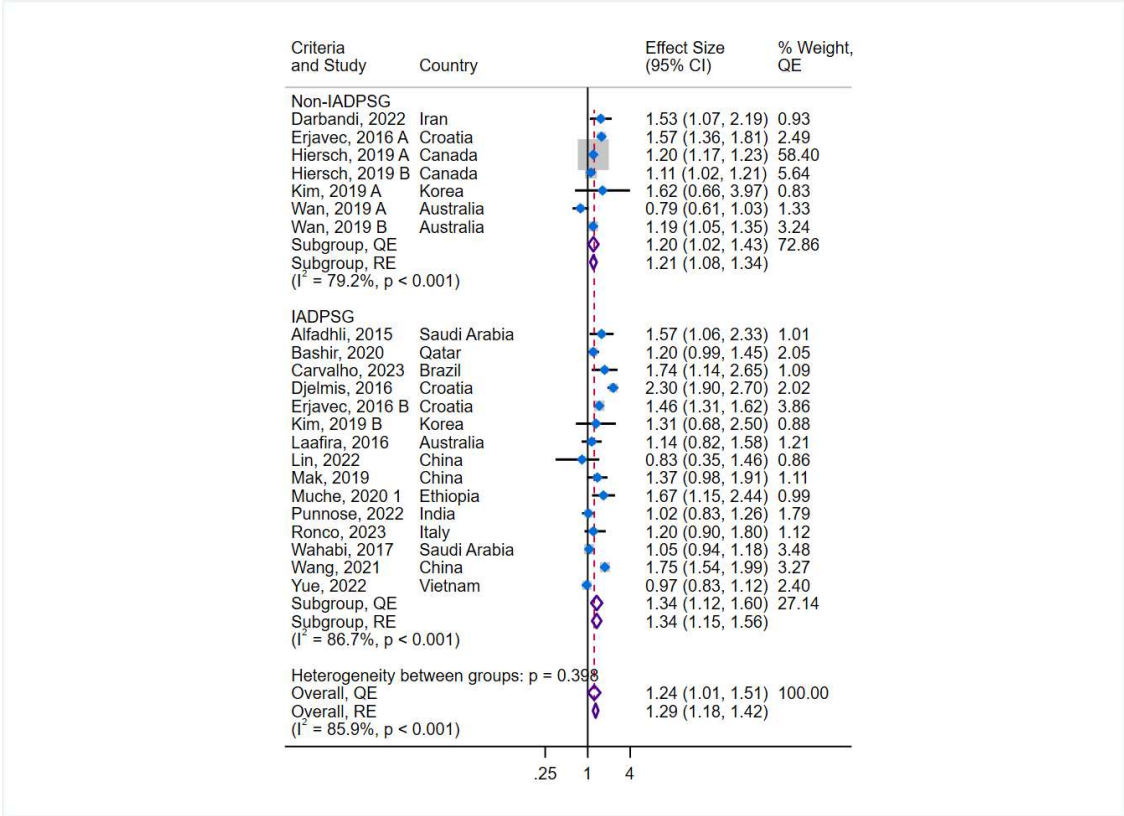
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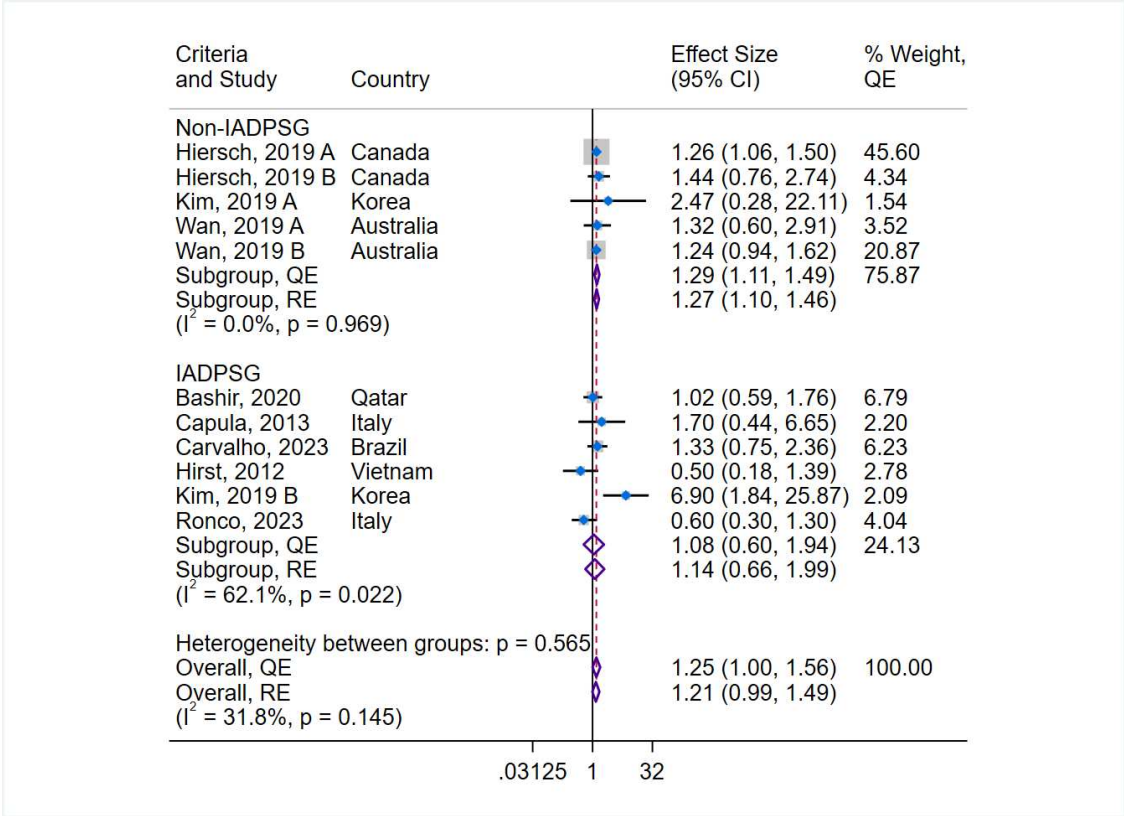
Supplementary Fig. 1: MASTER Scale Assessment



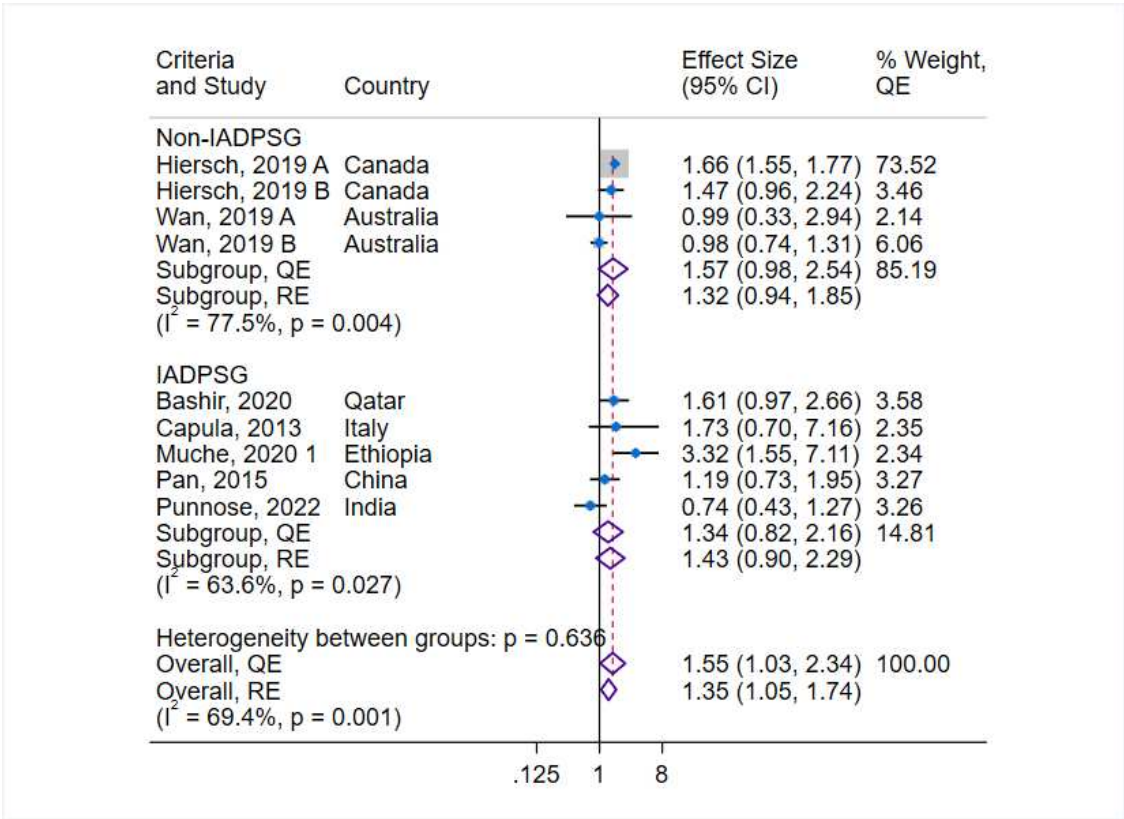
Supplementary Fig. 2: Forest plot for cesarean sections



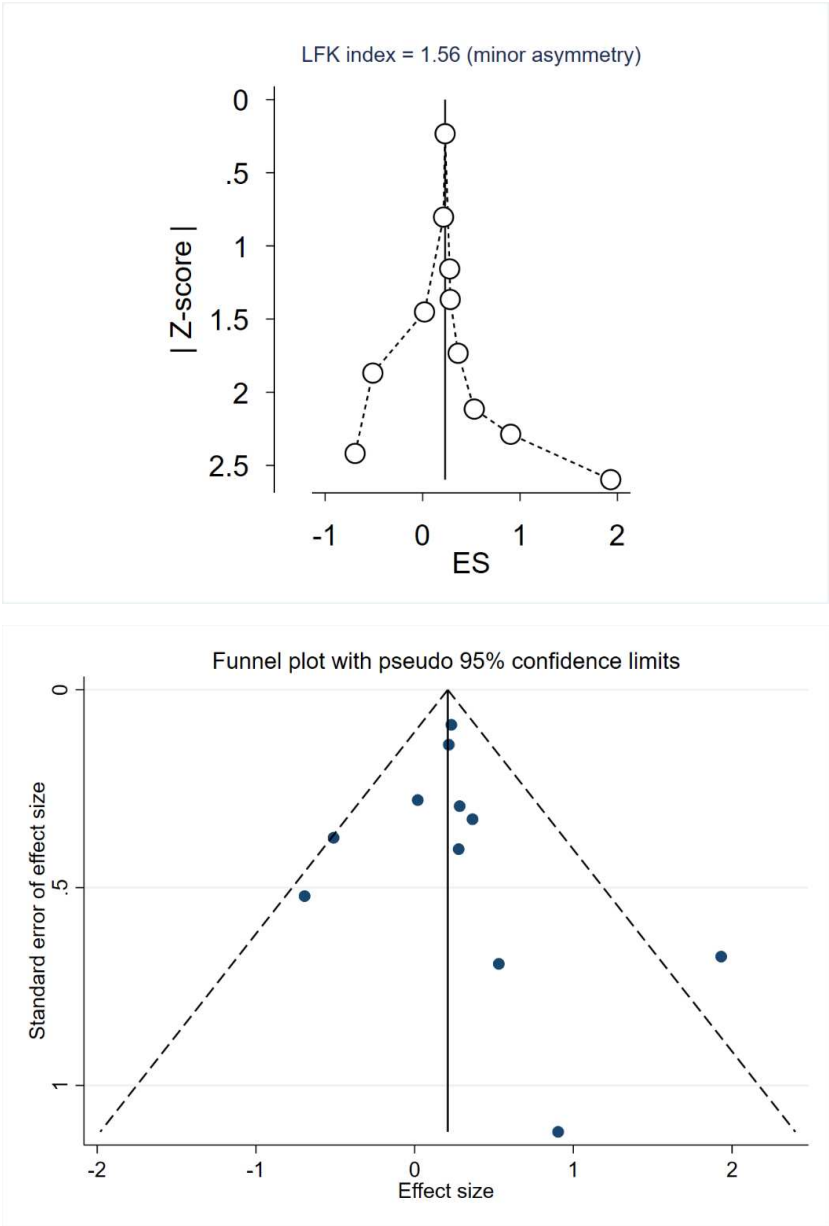
Supplementary Fig. 3: Forest plot for preeclampsia



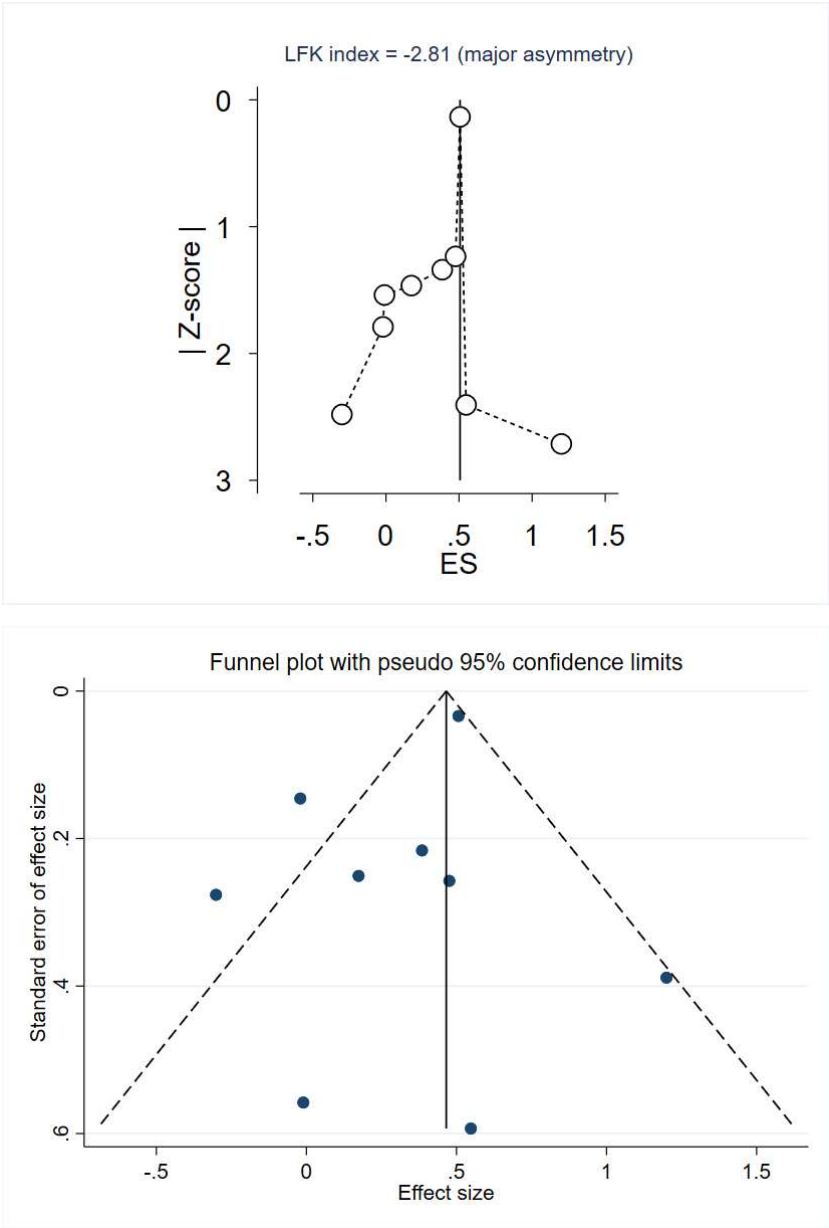
Supplementary Fig. 4: Forest plot for PIH



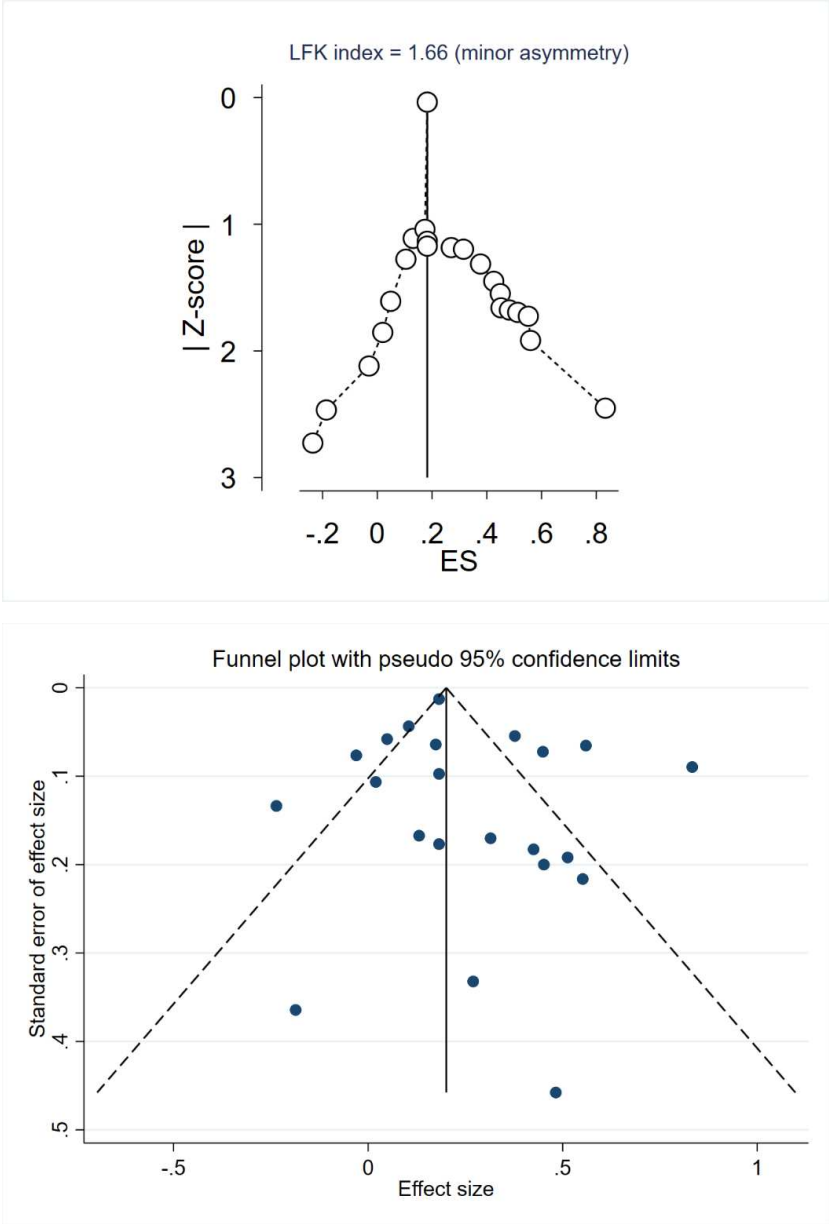
Supplementary Fig. 5: Doi and funnel plots for publication bias assessment for preeclampsia



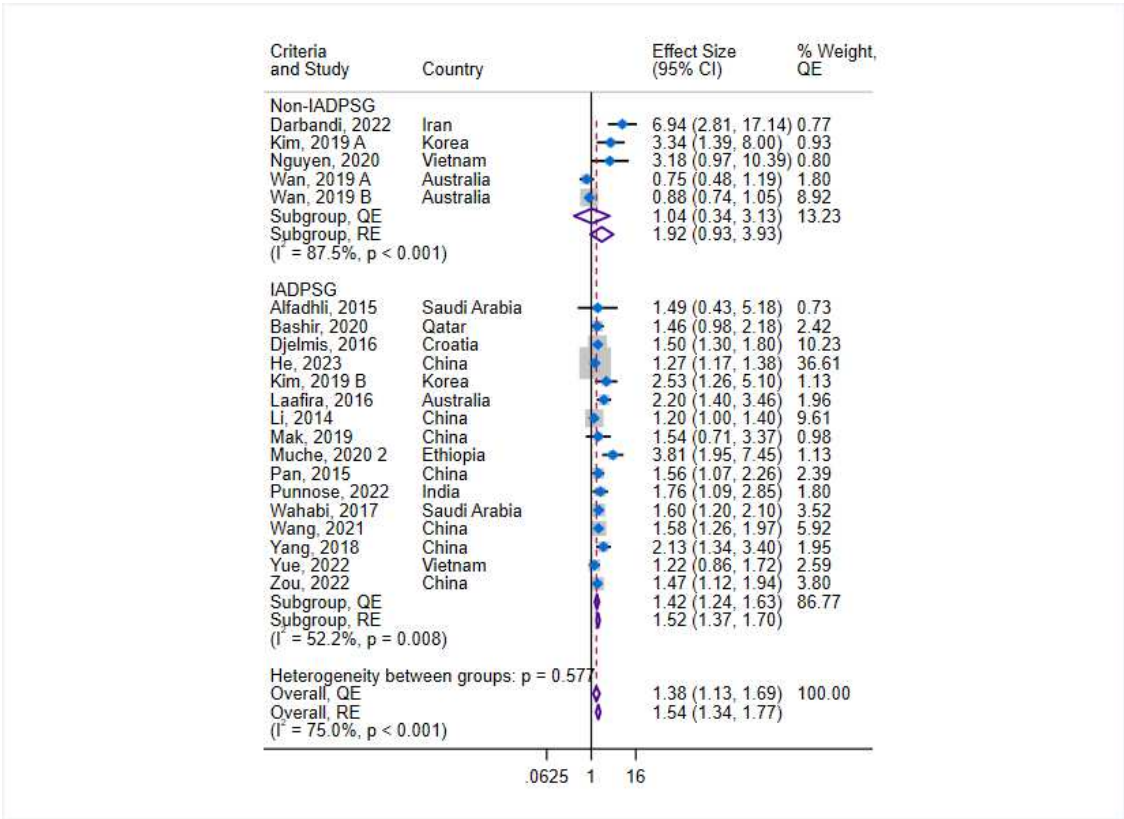
Supplementary Fig. 6: Doi and funnel plots for publication bias assessment for PIH



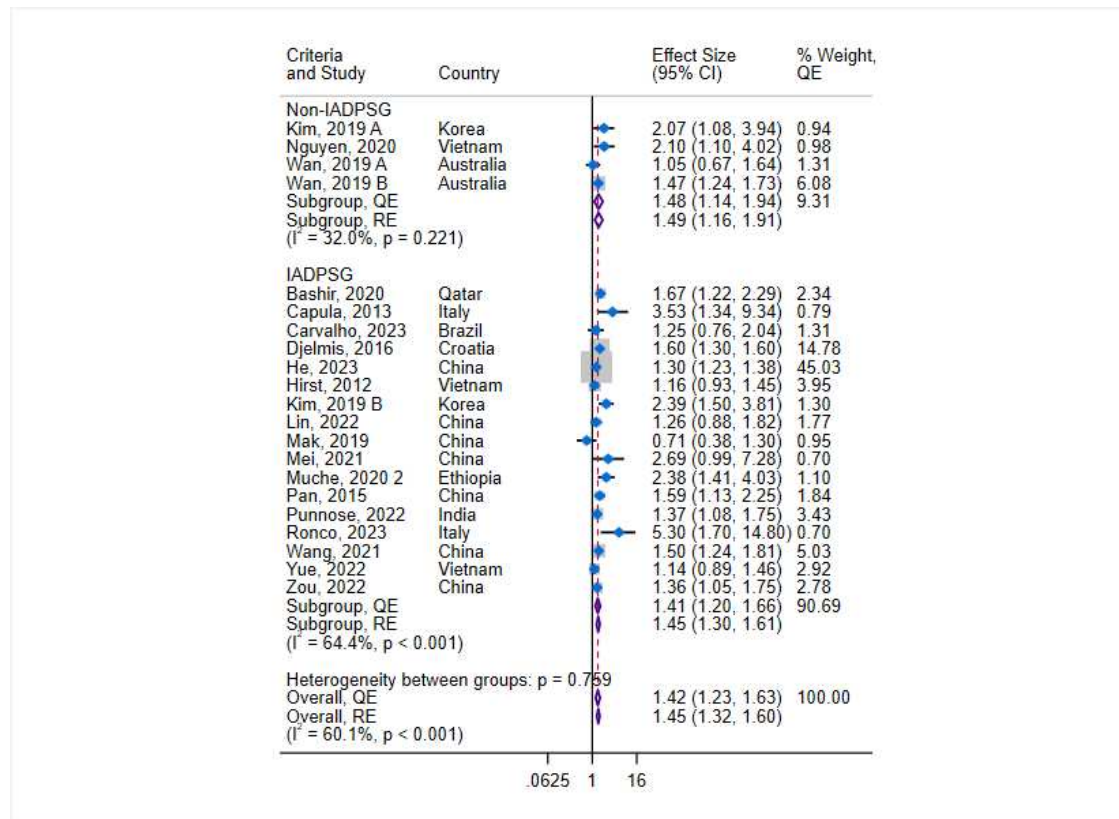
Supplementary Fig. 7: Doi and funnel plots for publication bias assessment for caesarean sections



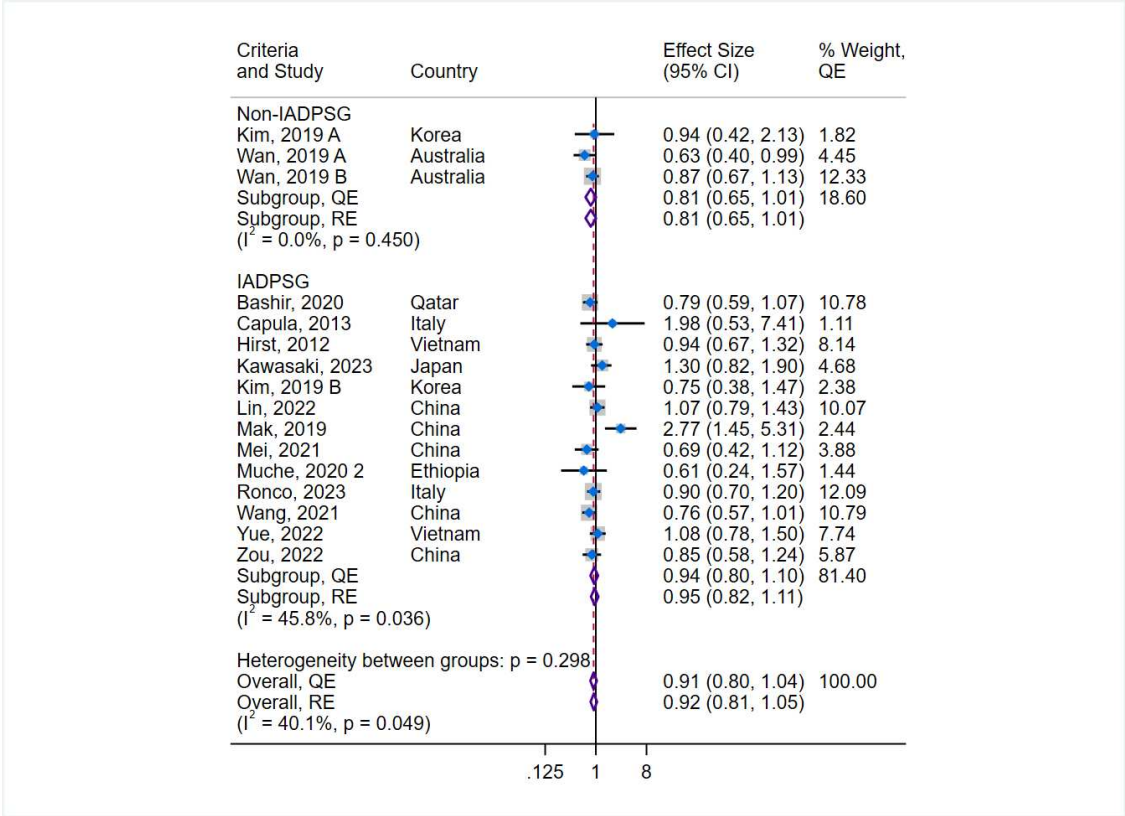
Supplementary Fig. 8: Forest plot for macrosomia



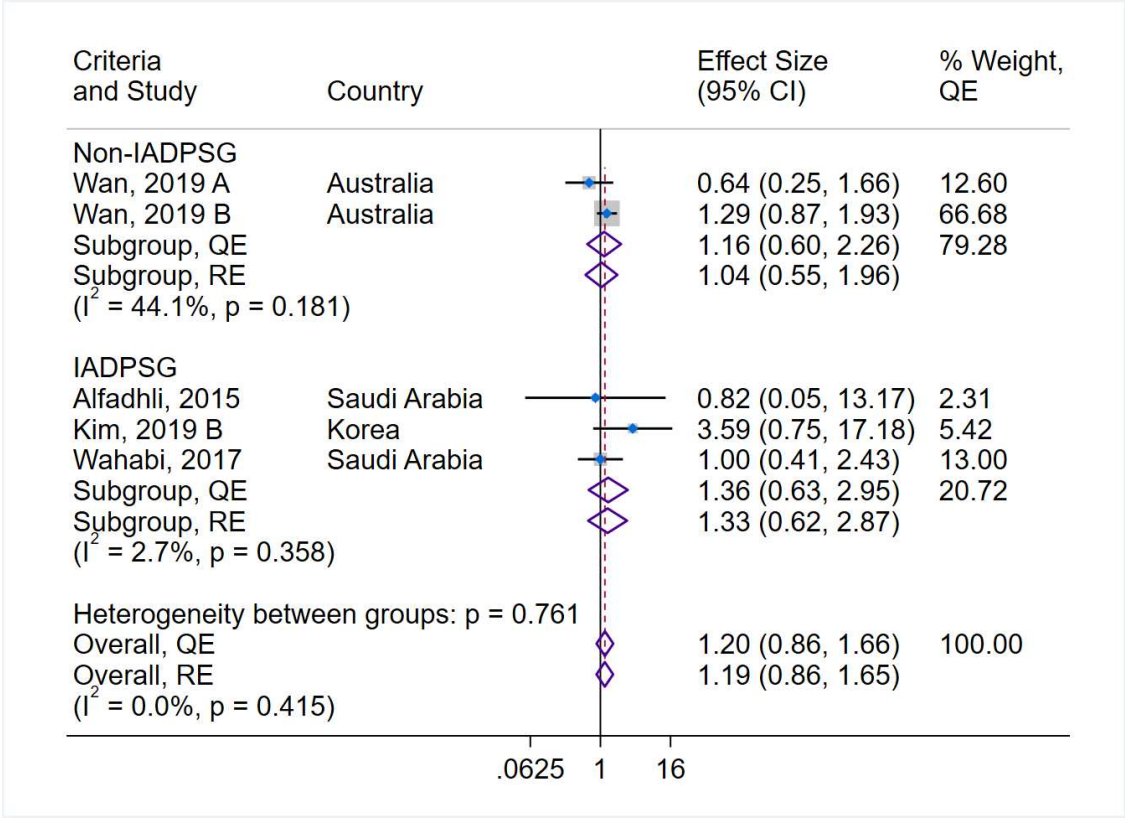
Supplementary Fig. 9: Forest plot for LGA



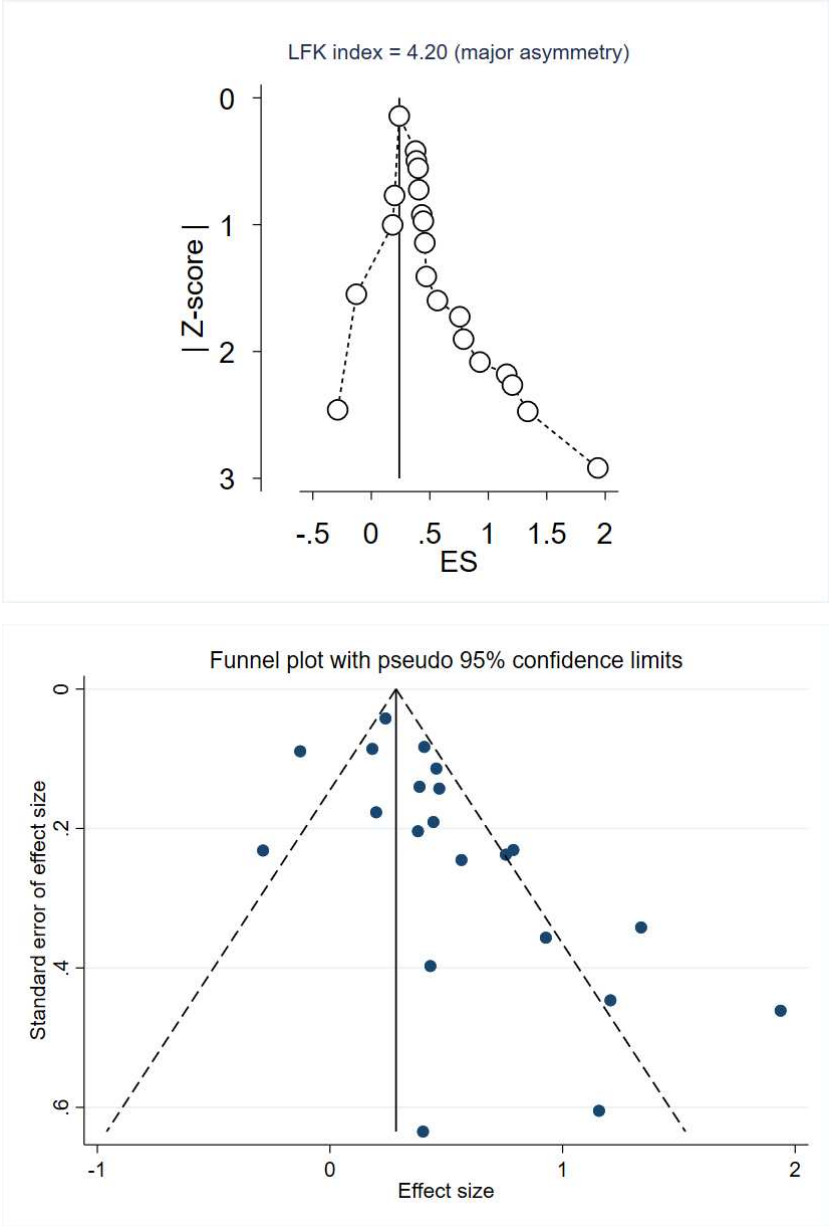
Supplementary Fig. 10: Forest plot for SGA



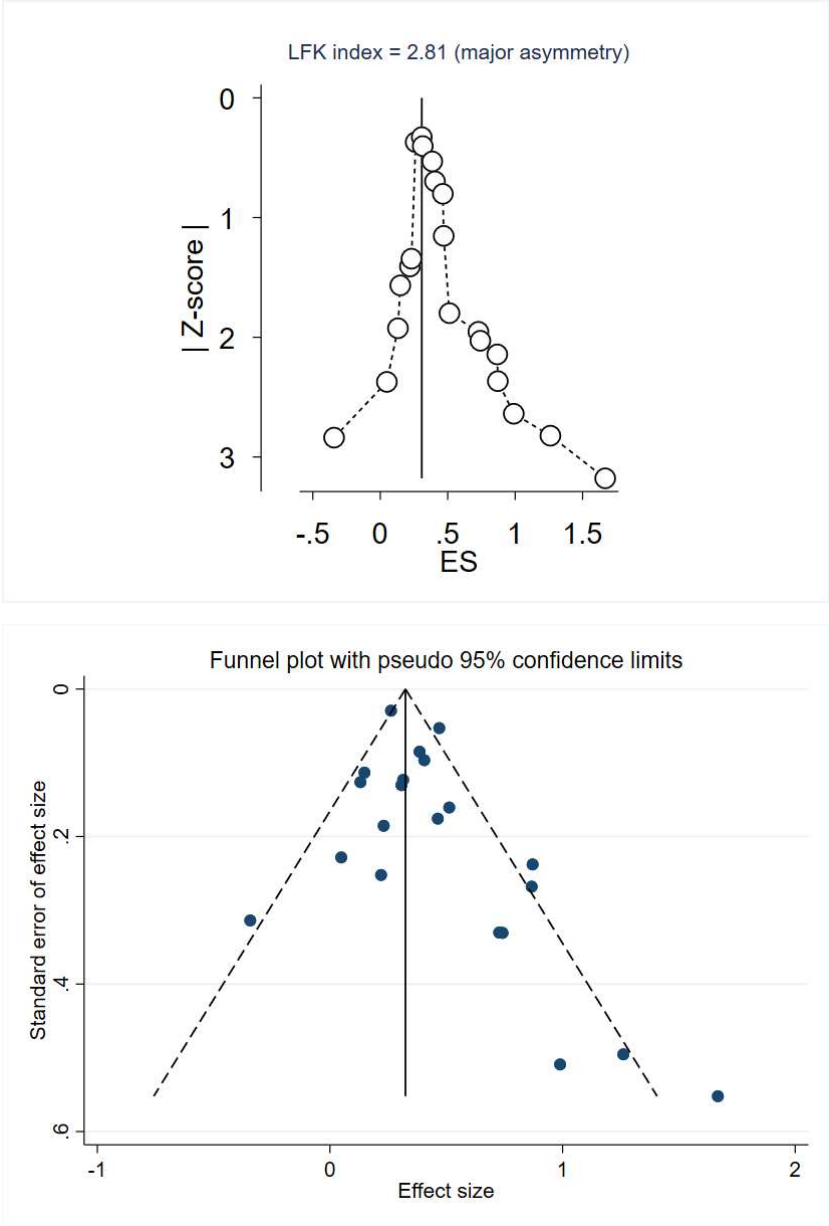
Supplementary Fig. 11: Forest plot for shoulder dystocia



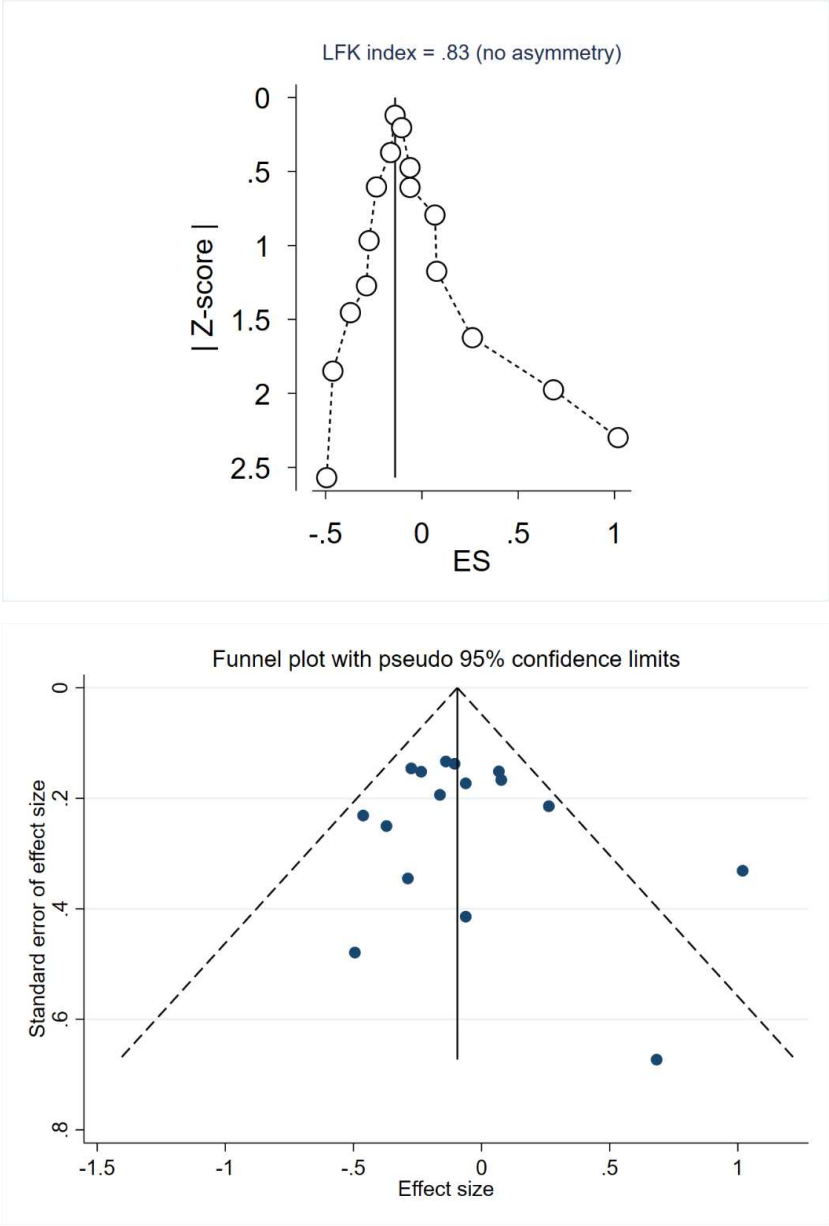
Supplementary Fig. 12: Doi and funnel plots for publication bias assessment for macrosomia



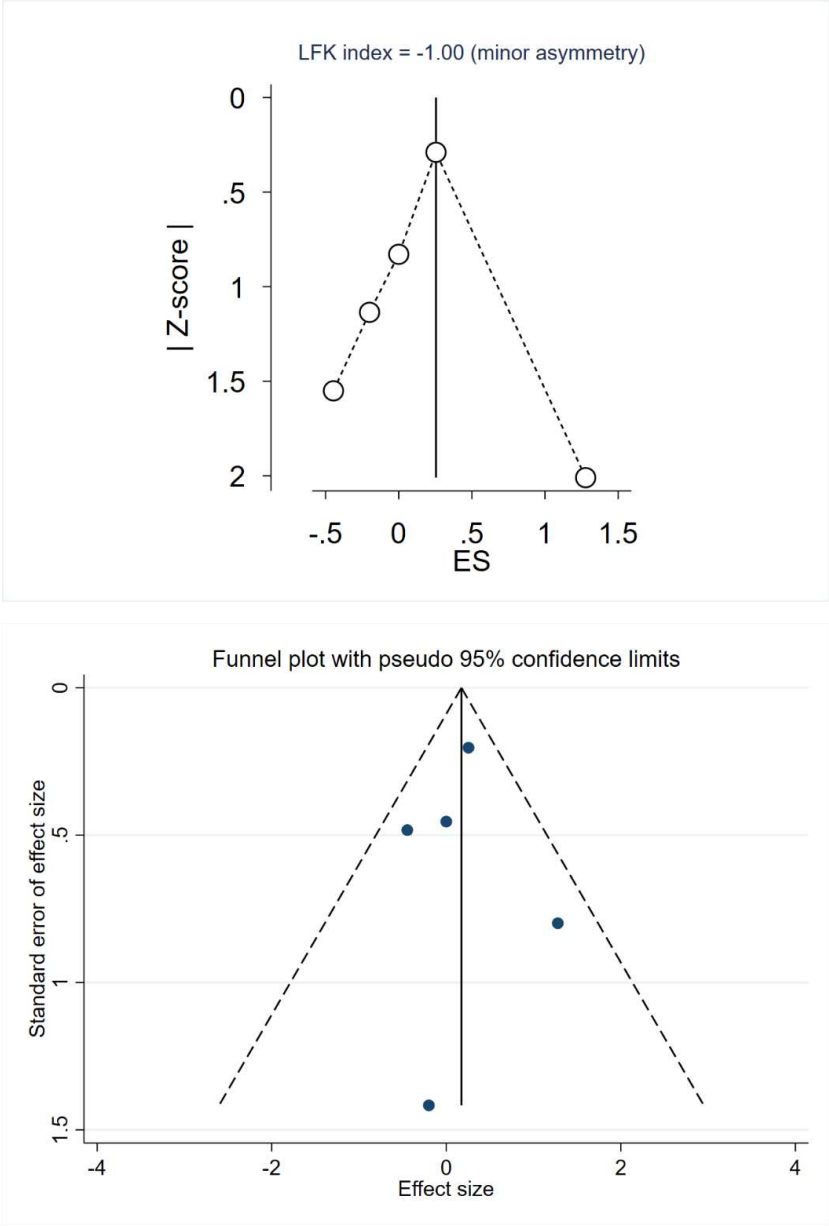
Supplementary Fig. 13: Doi and funnel plots for publication bias assessment for LGA



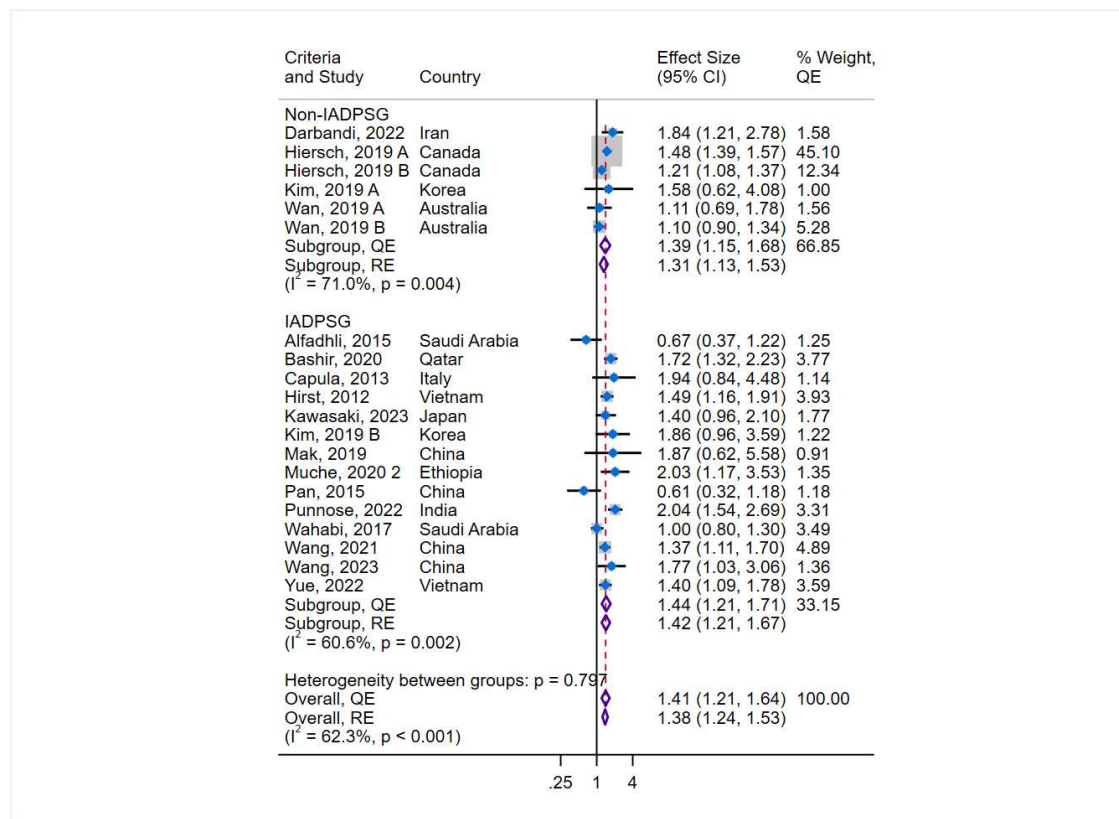
Supplementary Fig. 14: Doi and funnel plots for publication bias assessment for SGA



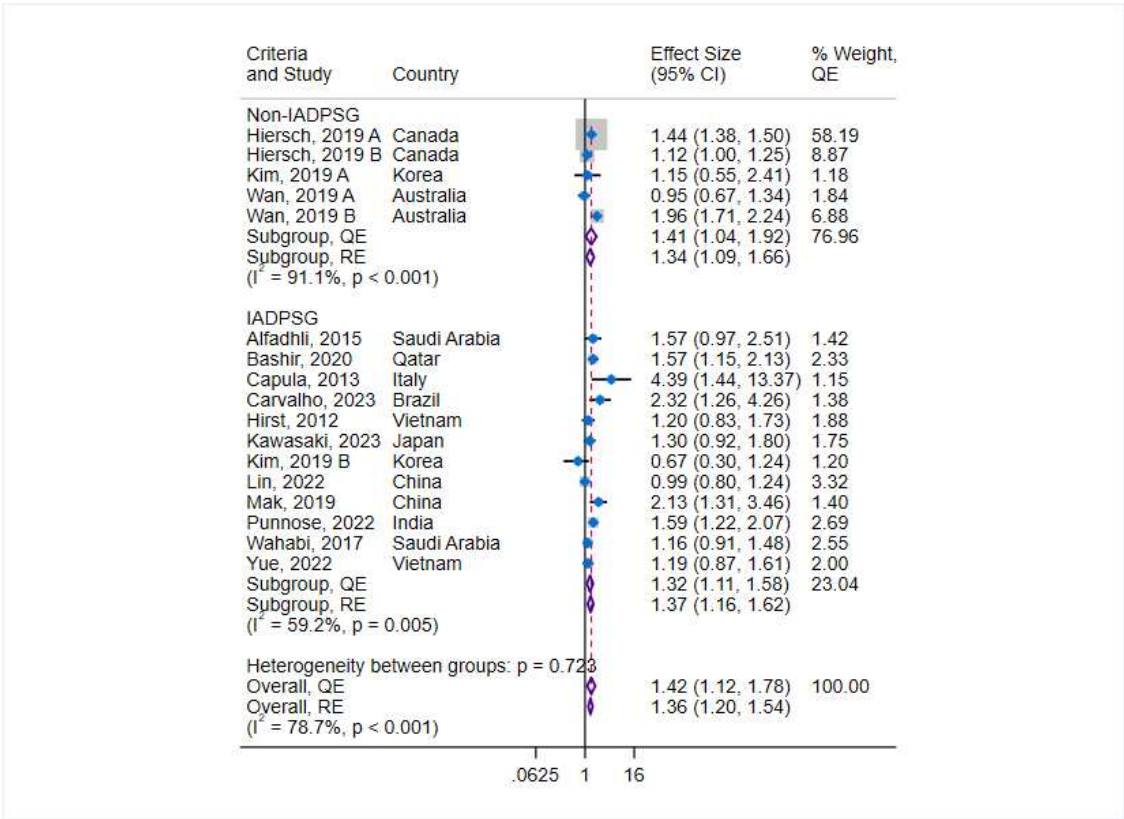
Supplementary Fig. 15: Doi and funnel plots for publication bias assessment for shoulder dystocia



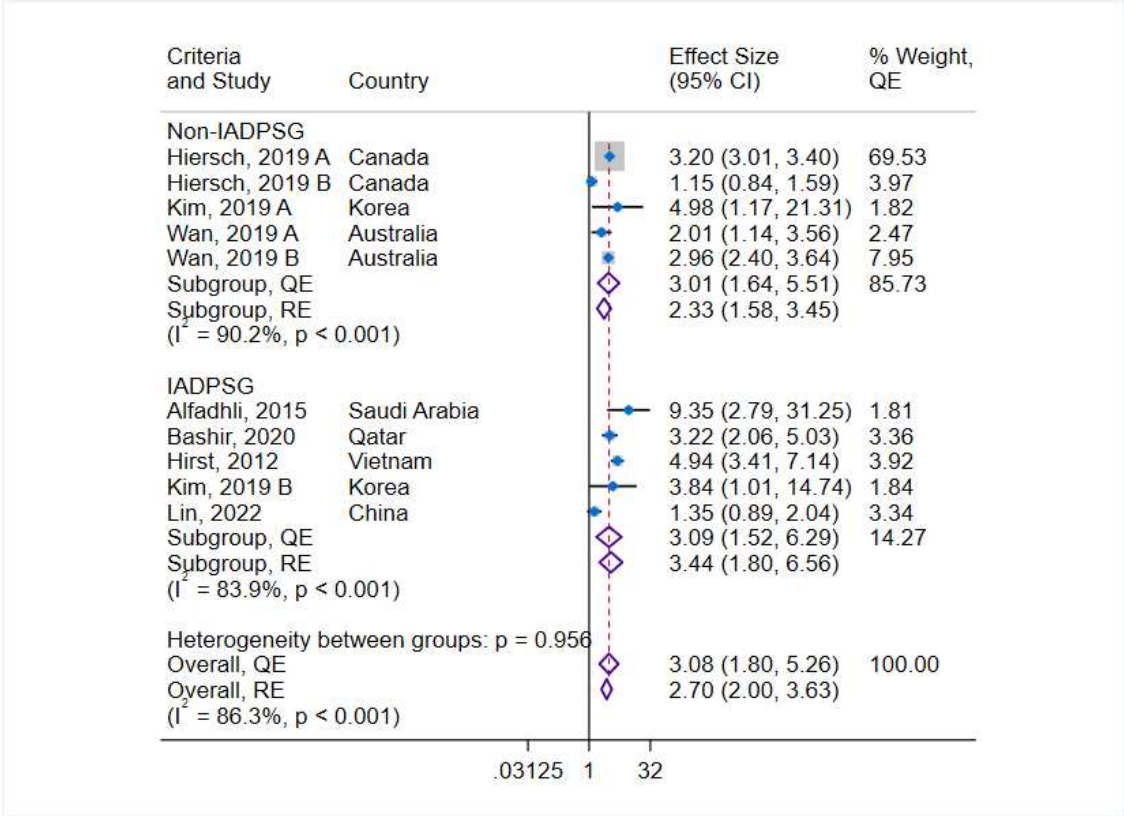
Supplementary Fig. 16: Forest plot for preterm birth



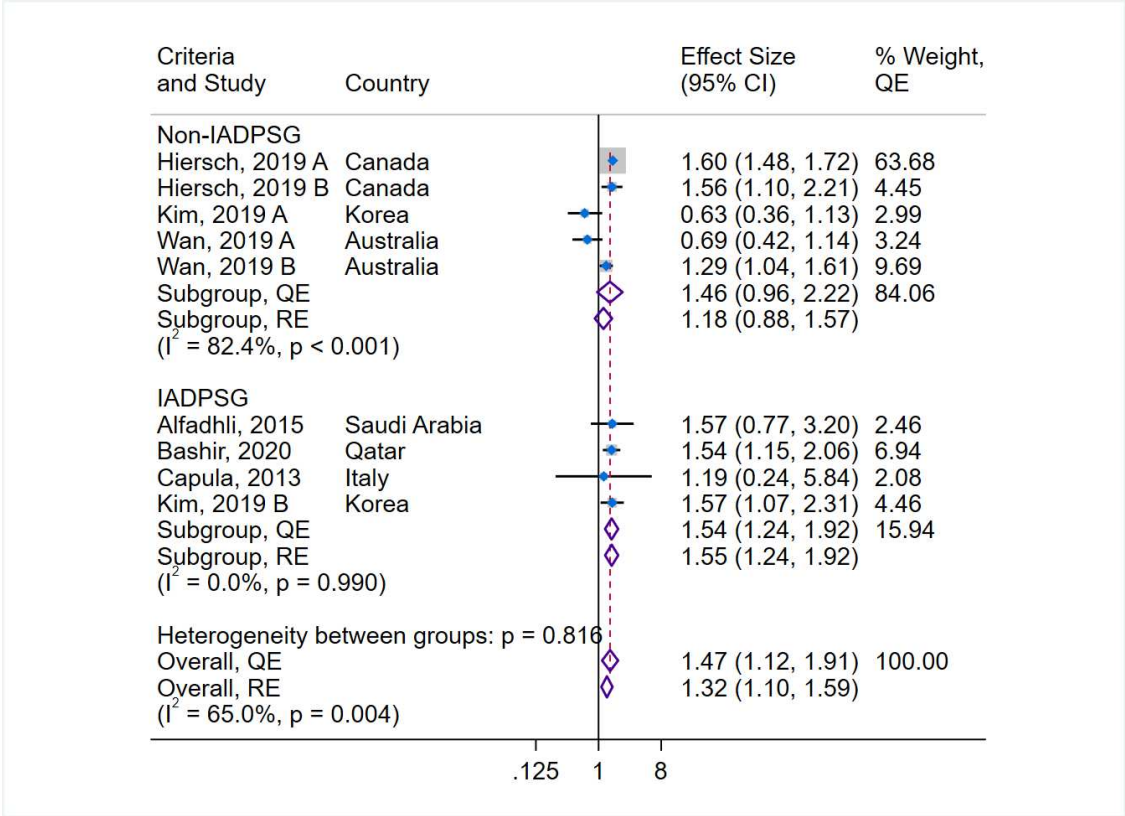
Supplementary Fig. 17: Forest plot for NICU admission



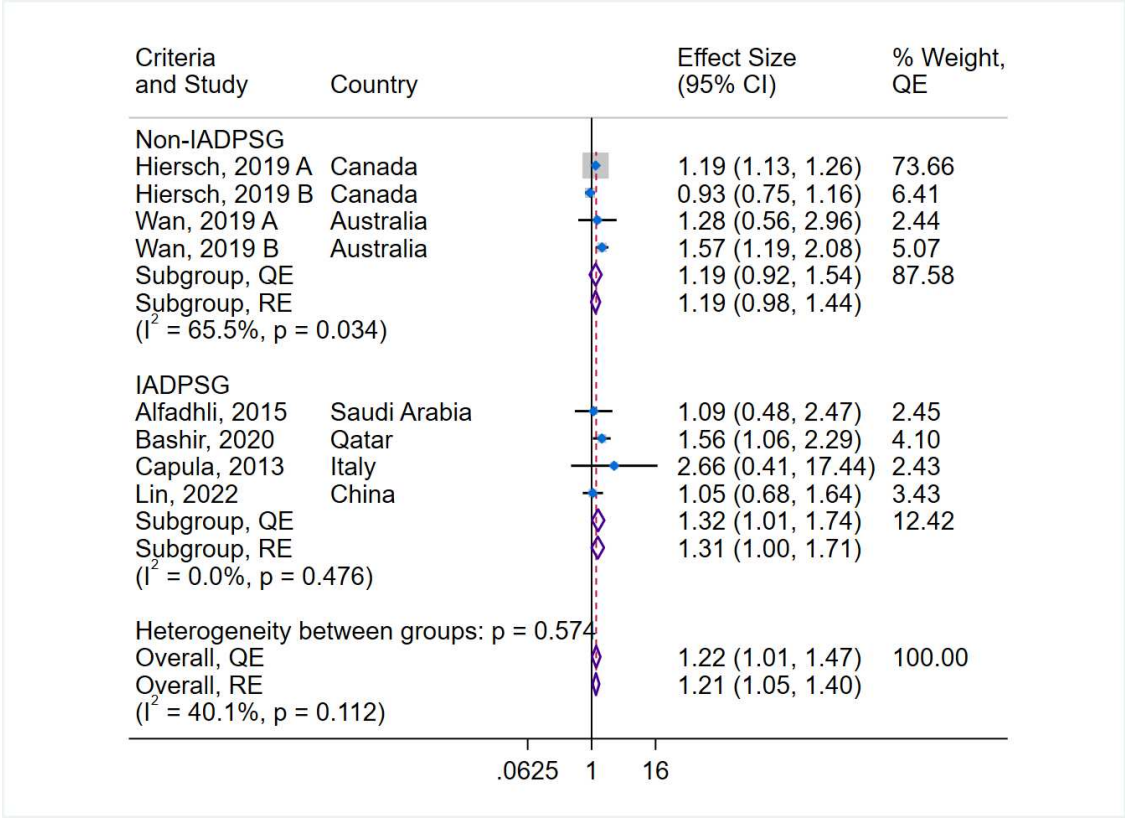
Supplementary Fig. 18: Forest plot for neonatal hypoglycemia



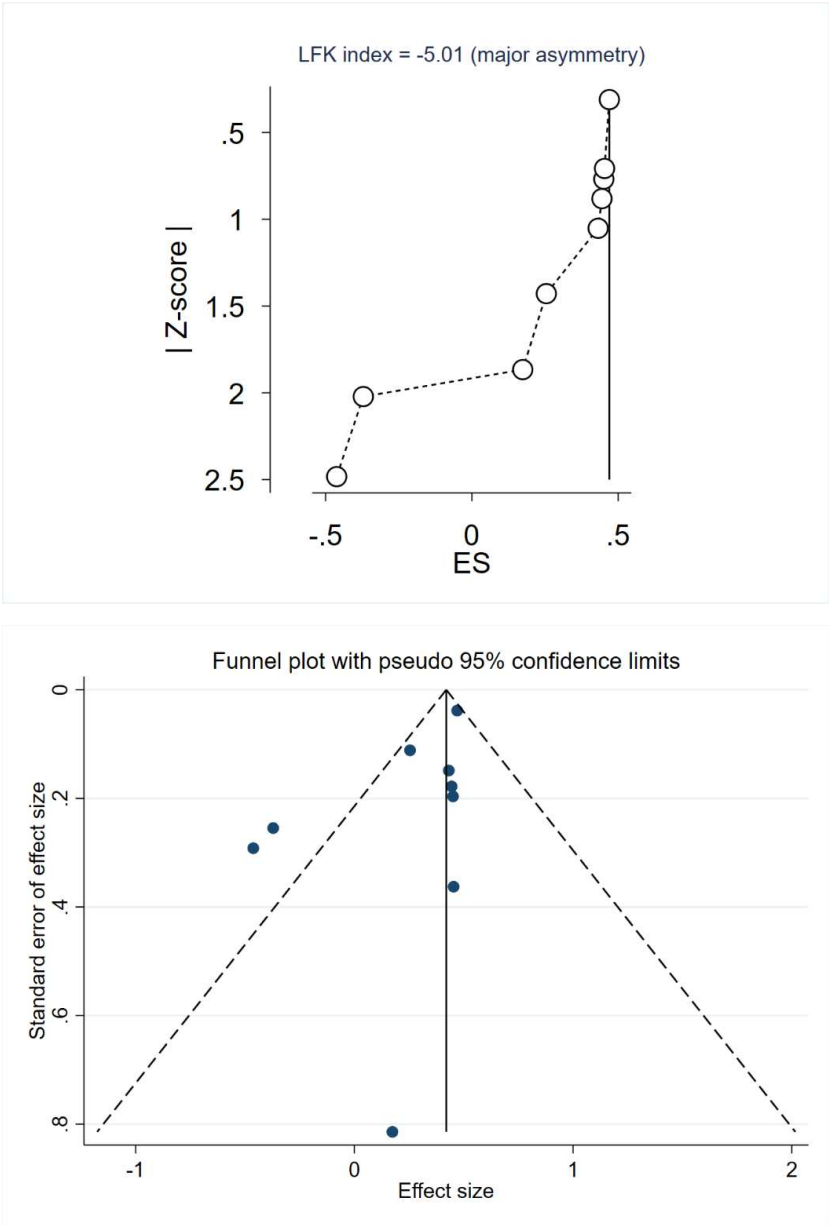
Supplementary Fig. 19: Forest plot for jaundice



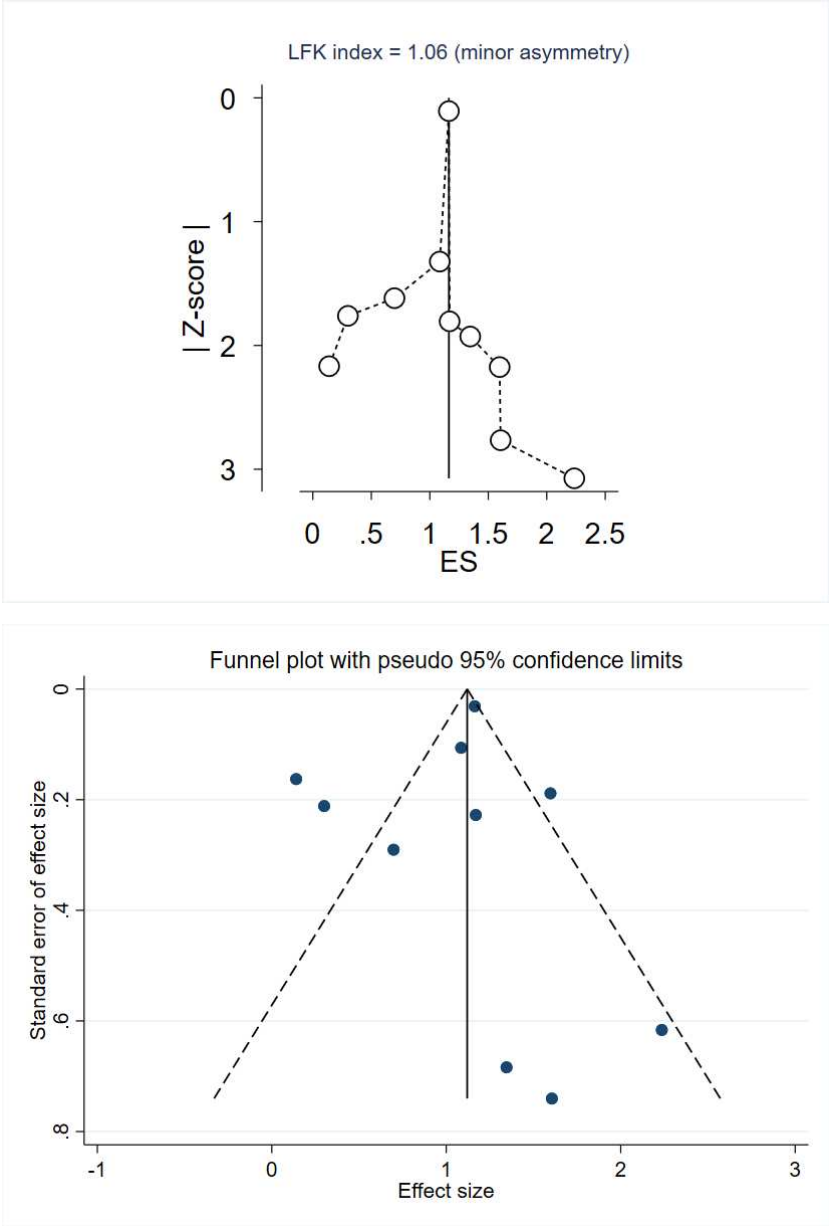
Supplementary Fig. 20: Forest plot for RDS



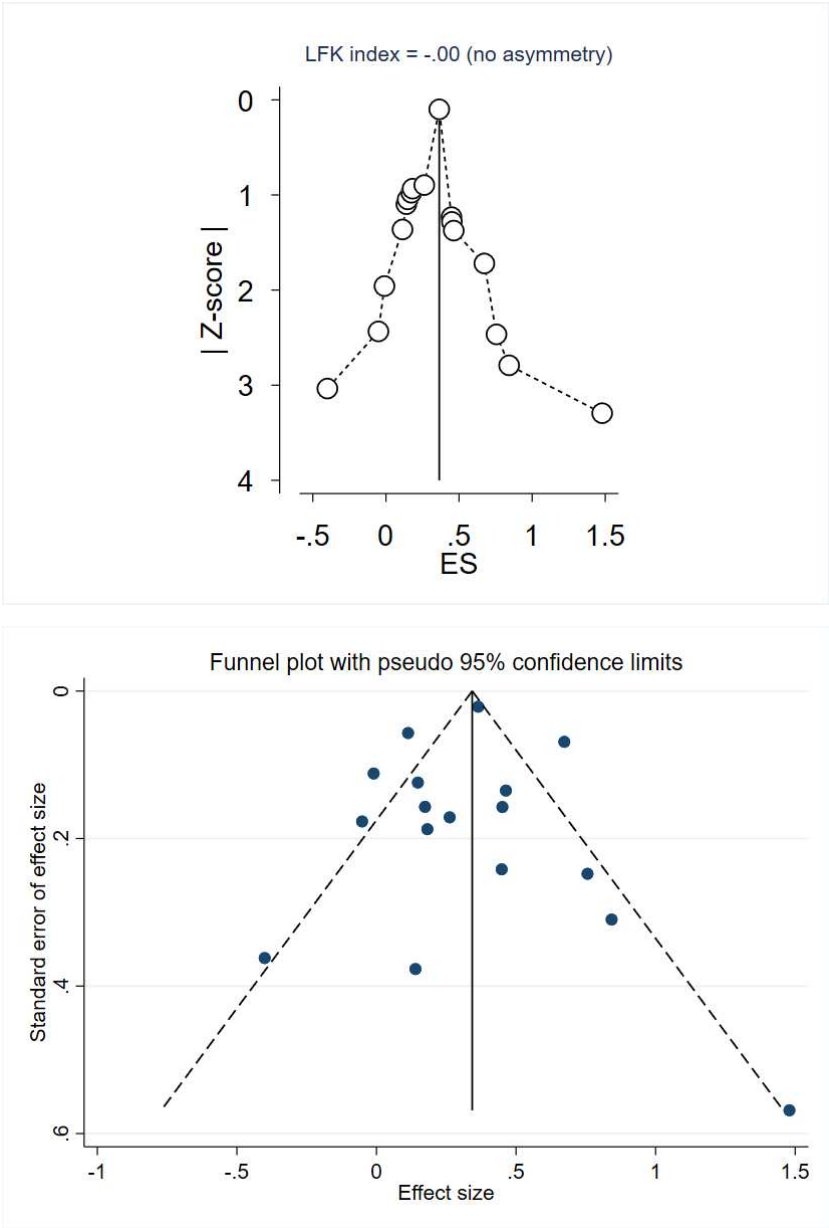
Supplementary Fig. 21: Doi and funnel plots for publication bias assessment for jaundice



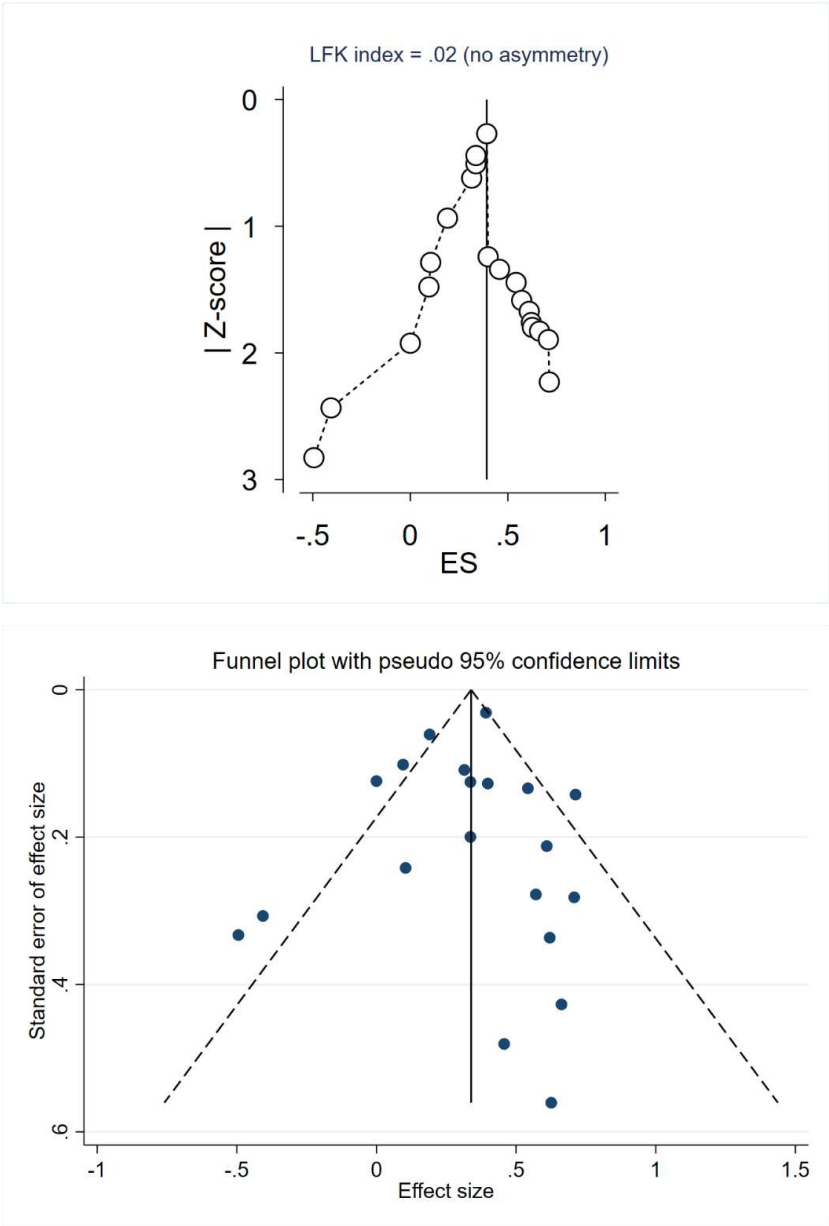
Supplementary Fig. 22: Doi and funnel plots for publication bias assessment for neonatal hypoglycemia



Supplementary Fig. 23: Doi and funnel plots for publication bias assessment for NICU admission



Supplementary Fig. 24: Doi and funnel plots for publication bias assessment for preterm birth



Supplementary Fig. 25: Doi and funnel plots for publication bias assessment for RDS

