Common mental disorders in the perinatal period in Primary Healthcarein Latin America: A Systematic review and meta-analysis Trastornos mentales comunes en el período perinatal en Atención Primaria de Salud en América Latina: una revisión sistemática y metanálisis. Transtornos mentais comuns no período perinatal na Atenção Primária à Saúde na América Latina: uma revisão sistemática e meta-análise

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Abstract

There is a statistically higher risk for developing mental comorbidities in the perinatal period. There is a scientific gap in the analysis of the occurrence of mental health problems in the perinatal period among middle and low-income countries in the context of Primary Healthcare. Thus, the objective of this study was to review the occurrence of common mental disorders in the perinatal period in Primary Healthcare in Latin America. An overall estimate of 37% (95%CI: 32%-43%) of common mental disorders was identified for the entire perinatal period. Despite the high frequencies identified for mental comorbidity symptoms, there were no studies in other Latin American countries with the exception of Brazil, preventing an analysis on the mental health of Latino women in the perinatal period.

Key Words: perinatal, primary care, mental disorders, public health

Resumen:

Existe un riesgo estadísticamente mayor de desarrollar comorbilidades mentales en el período perinatal. Existe un vacío científico en el análisis de la ocurrencia de problemas de salud mental en el período perinatal entre países de ingresos medios y bajos en el contexto de la Atención Primaria de Salud. Así, el objetivo de este estudio fue revisar la ocurrencia de trastornos mentales comunes en el período perinatal en la Atención Primaria de Salud en América Latina. Se identificó una estimación global del 37 % (IC 95 %: 32 %-43 %) de trastornos mentales comunes durante todo el período perinatal. A pesar de las altas frecuencias identificadas para los síntomas de comorbilidad mental, no hubo estudios en otros países de América Latina con excepción de Brasil, lo que impide un análisis sobre la salud mental de las mujeres latinas en el período perinatal.

Palabras clave: trastornos mentales, atención primaria perinatal, salud publica

Resumo:

Existe um risco estatisticamente maior de desenvolver comorbidades mentais no período perinatal. Existe uma lacuna científica na análise da ocorrência de problemas de saúde mental no período perinatal entre famílias de média e baixa renda países no contexto da Atenção Primária à Saúde. Assim, o objetivo deste estudo foi revisar a ocorrência de transtornos mentais comuns no período perinatal em Atenção Primária à Saúde na América Latina. Uma estimativa geral de 37% (95% CI:32%-43%) de transtornos mentais comuns foram identificados durante todo o período perinatal período. Apesar das altas frequências identificadas para sintomas de comorbidade mental, não houve estudos em outros países latino-americanos com exceção de Brasil, impossibilitando uma análise sobre a saúde mental das mulheres latinas no período perinatal.

Palavras chave: perinatal, atenção primária, transtornos mentais, saude publica

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Common mental disorders (CMDs) are defined as a suspected mental comorbidity for mood, anxiety, and somatization disorders, being characterized by non-psychotic manifestations of depressive symptoms, anxiety symptoms and somatization, with manifestations of difficulty concentrating, forgetfulness, insomnia, fatigue and irritability, in addition to non-specific somatic complaints. Although the initial symptoms are not aggressively characterized, they can generate extreme consternation and lead to dysfunction in the person's daily activities (World Health Organization, 2017).

Women at all stages of life are susceptible to manifesting CMDs. The risk for developing CMDs and/or other mental health problems is significantly increased (Austin et al., 2017) during the perinatal period, from the beginning of pregnancy to 12 months after delivery (Lara-Cinisomo et al., 2018; O'Hara & Wisner, 2014). Systematic reviews have pointed to high occurrences of gestational CMDs in the world, with a prevalence between 10-15% being estimated in high-income countries, while the occurrence in low and middle-income countries is between 10-41% (Fisher et al.,

2012; Gelaye et al., 2016).

The occurrence of CMDs in high-income countries is well established, and although there are studies with alarming data for low- and middle-income countries (Silva et al., 2020), such systematic reviews analyzed the occurrence of CMDs in different contexts from Primary Healthcare (PHC) to specialized services. This configures in considerable heterogeneity of screening and diagnostic methods and instruments used to achieve the outcome of interest (Fisher et al., 2012; Gelaye et al., 2016). Conceptual differences in relation to CMDs and the perinatal period have also been found (Silva et al., 2020).

A previous study reported 13% of postpartum depression in a global review and meta-analysis (O'Hara & Swain, 1996), but no review studies on CMDs in the perinatal period in Latin American countries were identified. The most recent systematic review on this phenomenon is from 2012 (Fisher et al., 2012), which investigated the prevalence and determinants of CMDs in women in low- and middle-income countries, but with the inclusion of only a few studies from Latin America and it did not have PHC as a scenario of interest.

Given the gap presented above, the relevance of CMDs in the perinatal period, as well as the possible negative outcomes for the woman, the child and the family, the aim of this study is to review the occurrence of CMDs in the perinatal period in PHC in Latin America.

Methods

Study design

This is a systematic review carried out in accordance with the guidelines of the Preferred Reporting Items for Systematic Reviews and Meta-analyses (PRISMA), Figure 1, for conducting systematic reviews (Galvão et al., 2015; Moher et al., 2009). The question of this review is the following: What is the occurrence of CMDs in the perinatal period in women treated in PHC in Latin America? The PICOS strategy was applied in developing the research question: Population (P): women in the perinatal period, from pregnancy to 12 months after delivery, attended in PHC; Intervention or exposure (I): occurrence of CMDs in the perinatal period; Comparison or control (C): none; Outcomes or outcome measures (O): prevalence and incidence of CMDs (depressive and anxiety symptoms, depressive symptoms and anxiety symptoms during pregnancy, postpartum and both periods); Studies (S): cross-sectional (prevalence) and cohort (prevalence and incidence). The searches were carried out in the following information resources: Cumulative Index to Nursing and Allied Health Literature (CINAHL), Embase, PubMed, PsycINFO, Latin American and

Caribbean Health Science Literature (LILACS) and Scientific Electronic Library Online (Scielo). A specific search strategy was used for each resource, as shown in Table 1. The search results were organized for screening in the EndNote software program which excluded duplicate articles, and the Rayyan software program (Ouzzani et al., 2016) was used with "blind on" reading by two researchers (BPS and TSD) and by a third (RAAZ) to break a tie. Studies which met the following inclusion criteria were occurrrence of CMDs (WHO, 2017) in the perinatal period (Lara-Cinisomo et al., 2018); (VI) conducted in PHC; and (VII) of Latin American countries. As exclusion criteria: (I) review articles; (II) articles which did not answer the guiding question; (III) articles which did not cover the selected theme; and (IV) studies developed in countries other than Latin America.

Titles and abstracts were initially read with the application of eligibility criteria in order to refine the sample with the inclusion of studies that responded to the objective of the review. Next, each selected article was read in full, subsidizing reflections on the phenomenon studied and the organization of articles for data collection. Finally, a critical analysis was carried out to identify the issues related to each article, different methodologies, samples and data collection.

The initial search was performed by two independent reviewers with "blind on" (BPS and TSD), with a standardized protocol for applying the inclusion and exclusion criteria. The final decision to include disagreement cases between independent reviewers was made by a third reviewer (RAAZ).

Data extraction

The data extracted from the selected studies in this review was done using a structured form that contains the following information: authors, date of publication, country in which the study was conducted, language, diagnosis of the patients studied, number of patients included, study design, and incidence/prevalence (Table

Quality analysis
The study quality analysis was performed according to the criteria of the Mixed Methods Appraisal Tool (MMAT), a strategy for evaluating the quality of studies when a review includes different designs. This tool was developed in 2006 and updated in 2018 by researchers at McGill University (Hong et al., 2018).

The MMAT scoring system enables reviewers and readers to view the global quality for each study assessed. The MMAT was designed to appraise the methodological quality of the studies, not the quality of their reporting (writing). This is an important feature because good research may not be well reported. Each article was independently assessed by all three reviewers for methodological quality using the MMAT to critically appraise the quality of the research studies (Table 3). The 2018 version of the MMAT enables describing and appraising the methodological quality of five types of studies: i) qualitative; ii) quantitative randomized controlled trials; iii) quantitative non-randomized; iv) quantitative descriptive; and v) mixed methods. Each type has its own set of quality criteria. The criteria are scored 'yes', 'no' or 'can't tell', followed by comments (Hong et al., 2018).

Qualitative synthesis

A qualitative synthesis was conducted with the studies that met the inclusion criteria. The synthesis included the study design, the period of the occurrence (pregnancy and/or post-partum), and the type of mental disorder screened by the original authors (common mental disorders, anxiety, and/or depression).

Statistical analysis

The number of women diagnosed with common mental disorders,

depression and/or anxiety and the total sample size were extracted from the original literature to calculate the pooled prevalence. Data were analyzed using the Stata statistical software program, and the command metaprop described by Nyaga et al. (2014). The pooled estimates of binomial data, the confidence intervals, continuity correct, and the Freeman-Tukey transformation the PROSERRO protocol of this systematic review was registered in the PROSPERO platform with record number CRD42021249557.

Results

General description

All 12 included studies were developed in PHC in Brazil. Of these, nine used a cross-sectional design (Cardillo et al., 2016; Fernandes Moll et al., 2019; Hassan et al., 2016; Shu et al., 2016; Silva & Cavalcante Neto, 2015; Silva et al., 2017; Silva et al., 2016; Silva et al., 2015; Tabb et al., 2015) and three were of the cohort type (Lima et al., 2017; Maclean et al., 2015; Paskulin et al., 2017). Sample sizes ranged from 66 (Fernandes Moll et al., 2019) to 831 participants (Shu et al., 2016). Three articles (Silva et al., 2017; Silva et al., 2016; Silva et al., 2015) were presumably developed with the same sample, with different objectives and published in different journals. The same occurred with studies conducted by Shu et al. (2016) and Tabb et al. (2015) which relied on data from a study conducted by Faisal-Cury et al. (2009) to carry out their analyzes.

Screening instruments and cut-off points

Several screening instruments were identified: the Edinburgh Postnatal Depression Scale (EPDS) was used for depression with scores ≥12 and ≥13 (Cardillo et al., 2016; Fernandes Moll et al., 2019; Lima et al., 2017); the General Health Questionnaire (GHQ-12) with a control of the control of 12) with a score ≥9 (Hassan et al., 2016); the Hospital Depression Subscale (HADS-D), part of the Hospital Anxiety and Depression Scale (HADS), with a cutoff point ≥9 (Silva et al., 2016; Silva et al., 2015); and the Primary Care Evaluation of Mental Distriction (PRIME-MD), diagnostic criteria according to the DSM-III-R for major depression (Paskulin et al., 2017).

The Hamilton Rating Scale for Depression (HAM-D) with a cutoff point ≥25 (Cardillo et al., 2016), and the Hospital Anxiety Subscale (HADS-A) with a cut-off point ≥9 (Silva et al., 2017; Silva et al., 2015) were used for anxiety screening; and the Primary Care Evaluation of Mental Disorders (PRIME-MD) with diagnostic criteria according to the DSM-III-R was used to assess generalized

anxiety (Paskulin et al., 2017).

The instruments used for CMD screening were the Self-Report Questionnaire (SRQ-20) (Maclean et al., 2015; Shu et al., 2016; Silva & Cavalcante Neto, 2015; Tabb et al., 2015) with two cut-off points of \geq 7 and \geq 8; and the General Health Questionnaire (GHQ-12) (Hassan et al., 2016), with a cut-off point ≥ 3 .

Groups (CMD, SD, SA, Pregnancy, Postpartum and both) and

study designs (cross-sectional and cohort)

Most studies were performed during pregnancy. Three screened CMDs (Shu et al., 2016; Silva & Cavalcante Neto, 2015; Tabb et al., 2015), two screened depressive symptoms (Lima et al., 2017; Silva et al., 2016) and two others depressive and anxious symptoms (Paskulin et al., 2017; Silva et al., 2015). Three studies investigated occurrences of the disorders only in the postpartum period, with screening for depressive and anxious symptoms (Cardillo et al., 2016), CMDs and depressive symptoms (Hassan et al., 2016), and only depressive symptoms (Formandes Moll et al., 2010). Only one only depressive symptoms (Fernandes Moll et al., 2019). Only one study screened CMDs during both pregnancy and postpartum (Maclean et al., 2015).

The gestational moments divided into weeks or trimesters assumed by the studies were quite diverse. Three cross-sectional studies considered screening for depressive and anxiety symptoms (Silva et al., 2015), depressive symptoms (Silva et al., 2016) and anxiety symptoms (Silva et al., 2017) in all gestational trimesters, evaluating the disorders as the participants sought care at the PHC. In a cohort study, Lima et al. (2017) defined three moments (20th, 28th and 36th weeks) for screening depressive symptoms, as well as Paskulin et al. (2017) in the 16th and 36th weeks of pregnancy. Shu et al. (2016) and Tabb et al. (2015) screened CMDs in women at 20 and 30 gestational weeks, and Silva and Cavalcante Neto (2015) in the 12th and 34th gestational weeks.

Among the studies that screened the occurrence of disorders in the postpartum period, a cross-sectional study screened depressive and anxiety symptoms among women at 0 to 4 months postpartum (Cardillo et al., 2016); another cross-sectional study screened CMDs and depressive symptoms in participants at 6 months postpartum (Hassan et al., 2016), whereas Fernandes Moll et al. (2019) identified the occurrence of depressive symptoms among participants at two weeks and 6 months after delivery. Only one cohort study screened the occurrence of CMDs during both pregnancy (between 20 and 30 weeks) and postpartum (between 6 and 18 months) (Maclean et al., 2015).

Qualitative synthesis

A qualitative synthesis was conducted with the studies that met the inclusion criteria. The synthesis included the period of the occurrence (pregnancy and/or post-partum), and the type of mental disorder (common mental disorders, anxiety, and/or depression).

The prevalence of depressive symptoms during pregnancy ranged from 21.6% (Paskulin et al., 2017) to 25.4% (Lima et al., 2017) at the 36th gestational week, as evaluated by the PRIM-MD/PHQ and EPDS, respectively. The occurrences of anxious symptoms observed in the studies were 19.8% (Paskulin et al., 2017) evaluated with the PRIM-MD/PHQ, 26.8% (Silva et al., 2015) and 28.6% (Silva et al., 2017), both with the HADS, score \geq 9. A high prevalence of CMDs was observed (61.4%) screened with the SRQ-20, cut-off point \geq 7, between the 12th and 34th gestational weeks (Silva & Cavalcante Neto, 2015).

High prevalence was also observed in a cohort study that evaluated the occurrence of depressive symptoms (EPDS≥13) at the 20th, 28th and 36th weeks of pregnancy of 27.2%, 21.7% and 25.4%, respectively, in one of the evaluation moments. The prevalence for the entire pregnancy was 7% (Lima et al. 2017)

the entire pregnancy was 7% (Lima et al., 2017). In the study by Cardillo et al. (2016), the occurrence of depressive symptoms screened with EPDS and adopting a cut-off point \geq 12 from 0 to 4 months postpartum was 20.8%. On the other hand, Hassan et al. (2016) found a prevalence of 8.3% for depressive symptoms using the GHQ-12 with a score \geq 9 at 6 months after delivery. On the other hand, these same authors found a high prevalence of CMDs (39.9%) for the same sample and period screened with the GHQ-12 with a score \geq 3.

Finally, a cohort study carried out between the 20th and 30th weeks of pregnancy and between the 6th and 18th months postpartum adopting a score ≥8 for positive CMDs found a prevalence of 36% CMDs screened only during pregnancy.

Meta-analyses (overall estimate)

The prevalence of disorders (CMDs, depressive symptoms or anxiety symptoms) in the first group of analyses for studies carried out in the gestational period was 31% (95% CI: 24%-38%); the estimate in the postpartum period was 22% (95% CI: 6%-38%) and heterogeneity of p=0.365 between groups. The overall estimate of occurrences was 28% (95%CI 22%-34%).

The overall prevalence for each of the disorders (CMDs, depressive symptoms and anxiety symptoms) in the second group of analyses was 37% (95%CI: 32%-43%) for CMDs, 21% (95%CI: 12%-31%)

for depressive symptoms and 20% for anxiety symptoms (95%CI: 16%-23%), with statically significant heterogeneity (p<0.0001). Finally, a prevalence of 37% (95%CI: 30%-43%) of CMDs during pregnancy was found in the third group of analyses (type of disorder by perinatal period), 27% (95%CI: 13%-41%) for depressive symptoms during pregnancy, and 20% (95%CI: 16%-23%) for anxiety symptoms during pregnancy. The prevalence of CMDs in the postpartum period was 40% (95%CI: 34%-46%) and 15% for depressive symptoms (95%CI: 95%-6%, 25%) depressive symptoms (95%CI: 95%:6%-25%).

Discussion

All included studies were conducted in Brazil between 2015 and 2019, applying different instruments and cut-off points. As a result of the differences in the study designs used by the authors, different

instruments were applied in different perinatal periods.

The lowest and highest prevalences for depressive symptoms were 7% and 27.2%, respectively, observed in the same study. The lowest prevalence was estimated for all pregnancy periods and the highest for the 20th gestational week. The instrument used was the EPDS with a score ≥13 (Lima et al., 2017). The lowest prevalence for anxiety disorders was 19.8% assessed with the PRIME-MD/PHQ (Paskulin et al., 2017); the highest was 28.6% (Silva et al., 2017) screened with the HADS-A≥9. The prevalence for CMDs ranged from 32% (Tabb et al., 2015) and 32.5% (Shu et al., 2016), with the use of the SRQ-20 score ≥ 8 , and the highest prevalence of 61.4% for CMDs screened with SRQ-20, cut-off point ≥7 (Silva et al., 2015). Such differences in occurrences are related to the perinatal period evaluated, the instruments used and the cut-off points used. Study designs and populations included are quite different.

In a systematic review study (Gelaye et al., 2016) performed with studies developed in low- and middle-income countries, the pooled prevalence for the gestational period was 25.3% (95%CI 21.4-29-6%), with significant heterogeneity between studies (p<0.001). The pooled prevalence of depression in a sensitivity analysis after excluding a study with a larger sample size was 25.8% (95%CI 22.8-

Gelaye et al. (2016) found estimates of 19% for the postpartum period (95%CI 15.5-23.0%) in a grouped analysis of the studies, with significant heterogeneity (p<0.001). The prevalence in the sensitivity analysis was 19.6% (95%CI 16.8-22.6%).

The overall estimate for depression in the present study was 21% (95%CI 12%-31%), with a higher pooled prevalence for the studies included in this review. Our analysis considered the occurrences of depression (pooled analysis) for the entire perinatal period.

In a systematic review with meta-analysis which summarized the evidence on the prevalence and determinants of CMDs in the perinatal period in low- and middle-income countries, Fisher et al. (2012) found a prevalence of 15.9% in the gestational period (95%CI 15.0%-16.8%). The prevalence of CMDs for the postpartum period was 19.8% (95%CI 19.2%-20.6%). The overall estimate in the present review was 37% (95%CI 32%-43%) for CMDs for the

entire perinatal period.

Through the high frequencies found for depression, anxiety and CMDs throughout the perinatal, pregnancy and postpartum period, this systematic review supports the importance of actions to detect mental disorders in the perinatal period in primary healthcare, with emphasis on the need for screening starting from the first prenatal visit. All studies were developed with the population of Brazil, which indicates the absence of investigations analyzing the population of other Latin American countries and shows a gap that should mobilize researchers, scholars and professionals to analyze the conditions and outcomes of maternal mental health in the perinatal period. From the care point of view, the results of the analysis point to the need to create and implement clinical protocols for tracking and monitoring mental health in the perinatal period, with a view to preventing maternal and child mental health problems.

The leadership and work of nurses comprise a central role in qualifying mental healthcare in the prenatal and postpartum periods. Due to the prenatal and postpartum follow-up carried out by nurses, the decrease in maternal mortality rates is central to the leadership and performance of Primary Health Care nurses in the Brazilian context. The leadership and work of nurses comprise a central role in qualifying mental healthcare in the prenatal and postpartum periods.

As a limitation of the present study, only studies available electronically in certain databases in English, Portuguese and Spanish were included. Only observational studies of the cohort and cross-sectional types were considered. These choices may have led to omitting studies which include other inclusion criteria related to the descriptors used. Furthermore, the heterogeneity of the study designs, screening instruments and scores used in each of them and the evaluation moments of the occurrences, both during pregnancy and in the postpartum period. The use of the metrapop command used for the statistical analysis does not allow differentiating prevalence from incidence. Therefore, only the prevalence measures from the cohort studies were considered.

Conclusion

High frequencies were found for depression, anxiety and CMDs throughout the perinatal period, pregnancy and postpartum, which demonstrate the need to focus attention/care on maternal mental health in prenatal care. There is a very diverse population and methodological set, even though all the studies were carried out in the same country, making it difficult to compile the results around a single estimate.

Research is needed to determine paths for professional training and the establishment of care protocols within the scope of PHC, calling for special attention from researchers and health professionals to mental health during the perinatal period.

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Table 1. Search strategies modulated according to each of the information resources, 2021.

Database	Search strategy
CINAHL	(((MH "Pregnancy") OR TI Pregnancy OR AB Pregnancy) OR ((MH "Depression, Postpartum") OR TI ("postpartum depression" OR "postnatal depression" OR ppd) OR AB ("postpartum depression" OR "postnatal depression" OR ppd)) AND ("common mental disorder" OR "common mental disorders"))
Embase	("common mental disorder"/exp OR "common mental disorder" OR "common mental disorders") AND ("postnatal depression"/exp OR "pregnancy"/exp) AND [Embase]/lim
LILACS	tw: (("transtorno mental comum" OR "transtornos mentais comuns" OR "common mental disorder" OR "common mental disorders")) AND (mh: ("Gravidez" OR "Depressão Pós-Parto"))
PubMed/MEDLINE	(("common mental disorder" OR "common mental disorders")) AND ((("Pregnancy" [Mesh] OR "Pregnancy" [tiab])) OR ("Depression, Postpartum" [Mesh] OR "Depression, Postpartum" [tiab]))
PsycINFO	"COMMON MENTAL DISORDER" OR Any Field: "COMMON MENTAL DISORDERS" AND (Any Field: PREGNANCY OR Any Field: "POSTPARTUM DEPRESSION")
Scielo	((Transtomos mentais comuns) AND (Gavidez) OR (Depressão Pós-Parto))

Platform registration number PROSPERO: CRD42021249557

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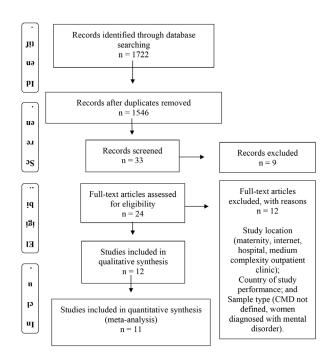


Figure 1. PRISMA flowchart of the inclusion process in the systematic review.

Table 2. Characterization of included studies according to authorship, study site, design, sample, assessed perinatal period, instrument and score tracked symptom and occurrence.

Author, Year	Country and publication language	Design	Sample	Perinatal period	Screening instrument and score	Outcome measured	Frequen cy
Cardillo et		Cross-			EPDS ≥12	Depressive	20.8%
al., (2016)	Brazil, English	sectional	72	0 to 4 months postpartum	HAM- D≥25*	Anxiety	17.3%*
Hassan et	D1	C			GHQ-12	CMDs	39.9%
Hassan et al., (2016)	Brazil, Portuguese	Cross- sectional	288	6 months postpartum	≥3 and ≥9	Depressive	8.3%
Silva et al., 2015	Brazil, Portuguese	Cross- sectional	209	Pregnancy (all gestational trimesters)	HADS ≥9	Anxiety Depressive	26.8% 14.8%
Silva et al., (2016)	Brazil, English	Cross- sectional	209	Pregnancy (all gestational trimesters)	HADS-D ≥9	Depressive	14.8%
Silva et al., (2017)	Brazil, Portuguese	Cross- sectional	209	Pregnancy (all gestational trimesters)	HADS-A ≥9	Anxiety	28.6%
Lima et al.,	Brazil,	Cohort	272	20th, 28th and 36th	EPDS ≥13	Depressive (all gestation period)	7%
(2017)	Portuguese		gestational weeks	gestational weeks		20th week 28th week 36th week	27.2% 21.7% 25.4%
MacLean et al., (2015)	Brazil, English	Cohort	270 (adolescent	Pregnancy (between 20 and 30 weeks) and postpartum (between 6 and 18 months)	SRQ-20 ≥8	CMDs	36%**

			s and adults)				
Fernandes Moll et al., (2019)	Brazil, English	Cross- sectional	66	2 weeks to 6 months postpartum	EPDS ≥12	Depressive	19.7%
Paskulin et				Pregnancy	PRIME-	Major depression	21.6%
al., (2017)	Brazil, English	Cohort	712	(16th and 36th weeks)	MD/PHQ**	General anxiety	19.8%
Shu et al., (2016)	Braziil, English	Cross- sectional	831 (adolescent s and adults)	Pregnancy (between 20 and 30 weeks)	SRQ-20 ≥8	CMDs	32.5%
Silva e Cavalcante Neto, (2015)	Braziil, Portuguese	Cross- sectional	74	Pregnancy (between 12 and 34 weeks)	SRQ-20 ≥7	CMDs	61.4%
Tabb et al., (2015)	Braziil, English	Cross- sectional	811 (adolescent s and	Pregnancy (between 20 and 30 weeks)	SRQ-20 ≥8	CMDs	32%

adults)

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Table 3. MMAT tool quality evaluation.

	Screening questions		Quantitative studies						
Authors, Year	S1. Are there clear research questions? S2. Do the collected data address the research questions?		4.1. Is the sampling strategy relevant to address the research question?	4.2. Is the sample representative of the target population?	4.3. Are the measurements appropriate?	4.4. Is the risk of nonrespons e bias low?	4.5. Is the statistical analysis appropriate answer the research question?		
Cardillo et al., (2016)	Yes	Yes	Can't tell	Yes	Yes	Yes	Yes		
Hassan et al., (2016)	Yes	Yes	Yes	Yes	Yes	Yes	Yes		
Silva et al., (2015)	Yes	Yes	Yes	Yes	Yes	Yes	Yes		
Silva et al., (2016)	Yes	Yes	Yes	Yes	Yes	Yes	Yes		
Silva et al., (2017)	Yes	Yes	Yes	Yes	Yes	Yes	Yes		
Fernandes Moll et al., (2019)	Yes	Yes	Yes	Yes	Yes	Yes	Yes		
Paskulin et al., (2017)	Yes	Yes	Yes	Yes	Yes	Yes	Yes		
Shu et al., (2016) Silva e	Yes	Yes	Yes	Yes	Yes	Yes	Yes		
Cavalcante Neto, (2015)	Yes	Yes	Can't tell	Yes	Yes	Yes	Yes		

Screening questions				Non-				
Authors, Year	S1. Are there clear research questions?	S2. Do the collected data address the research questions?	3.1. Are the participants representative of the target population?	3.2. Are measurements appropriate regarding both the outcome and intervention (or exposure)?	3.3. Are there complete outcome data?	3.4. Are the confounders accounted for in the design and analysis?	3.5. During the study period, is the intervention administered (or exposure occurred) as intended?	
Lima et al., (2017)	Yes	Yes	Yes	Yes	Yes	Yes	Yes	
MacLean et al., (2015)	Yes	Yes	Yes	Yes	Yes	Yes	Yes	
Tabb et al., (2015)	Yes	Yes	Yes	Yes	Yes	Yes	Yes	

^{*}only evaluated the anxiety symptom in adolescents with positive depressive symptoms.

^{***}diagnostic criteria according to the DSM-III-R.

Table 4. Meta-analyses of common mental disorders, depressive symptoms and anxiety symptoms in the perinatal period.

	Cub anama	Study	Estimation	959	6CI	Weight			
	Sub-groups	Study	Prevalence	Inf	Sup	%			
		MacLean et al., (2015)	0.33	0.27	0.38	8.49			
		Shu et al., (2016)	0.32	0.29	0.36	8.82			
		Silva e Cavalcante Neto, (2015)	0.59	0.48	0.70	7.28			
	Pregnancy	Tabb et al., (2015)	0.32	0.29	0.36	8.82			
		Silva et al., (2015)	0.15	0.11	0.20	8.61			
		Silva et al., (2016)	0.15	0.11	0.20	8.61			
		Silva et al., (2017)	0.27	0.21	0.33	8.42			
Period		Lima et al., (2017)	0.39	0.33	0.45	8.46			
	Subtotal (I^2	=93.15%, p<0.001)	0.31	0.24	0.38	67.50			
_		Hassan et al., (2016)	0.40	0.34	0.46	8.35			
	Post-partum	Hassan et al., (2016)	0.08	0.05	0.13	8.78			
		Fernandes Moll et al., (2019)	0.20	0.12	0.31	7.66			
		Cardillo et al., (2016)	0.21	0.13	0.32	7.71			
_	Subtotal (I^2	0.22	0.06	0.38	32.50				
	Heterogeneity between groups: p=0.365								
	Overall (I^2	=95.09%, p<0.001)	0.28	0.22	0.34	100.00			
		MacLean et al., (2015)	0.33	0.27	0.38	7.84			
		Shu et al., (2016)	0.32	0.29	0.36	8.16			
	CMD	Silva e Cavalcante Neto, (2015)	0.59	0.48	0.70	6.65			
		Tabb et al., (2015)	0.32	0.29	0.36	8.16			
		Hassan et al., (2016)	0.40	0.34	0.46	7.70			
_	Subtotal (I^2	0.37	0.32	0.43	38.52				
_		Silva et al., (2015)	0.27	0.21	0.33	7.77			
		Silva et al., (2016)	0.15	0.11	0.20	7.96			
Mental	D	Lima et al., (2017)	0.39	0.33	0.45	7.80			
Disorders	Depression	Hassan et al., (2016)	0.08	0.05	0.13	8.12			
		Fernandes Moll et al., (2019)	0.20	0.12	0.31	7.03			
		Cardillo et al., (2016)	0.21	0.13	0.32	7.08			
_	Subtotal (I^2	=94.25%, p<0.001)	0.21	0.12	0.31	45.75			
_		Silva et al., (2015)	0.15	0.11	0.20	7.96			
	Anxiety	Silva et al., (2017)	0.27	0.21	0.33	7.77			
_	Subtotal (I	^2=0%, p<0.001)	0.20	0.16	0.23	15.73			
		eity between groups: p<							
		=95.09%, p<0.001)	0.28	0.22	0.34	100.00			

		MacLean et al., (2015)	0.33	0.27	0.38	7.84
	CMD Pregnancy	Shu et al., (2016)	0.32	0.29	0.36	8.16
		Silva e Cavalcante Neto, (2015)	0.59	0.48	0.70	6.65
		Tabb et al., (2015)	0.32	0.29	0.36	8.16
	Subtotal (I^2=	86.16%, p<0.001)	0.37	0.30	0.43	30.82
	Depression	Silva et al., (2015)	0.27	0.21	0.33	7.77
	Pregnancy	Silva et al., (2016)	0.15	0.11	0.20	7.96
Period		Lima et al., (2017)	0.39	0.33	0.45	7.80
and	Subtotal (I^2	0.27	0.13	0.41	23.53	
Mental	Anxiety Pregnancy	Silva et al., (2015)	0.15	0.11	0.20	7.96
Disorders		Silva et al., (2017)	0.27	0.21	0.33	7.77
	Subtotal (I^2	0.20	0.16	0.23	15.73	
	CMD Post-partum	Hassan et al., (2016)	40.00	0.34	0.46	7.70
	Sul	40.00	0.34	0.46	7.70	
		Hassan et al., (2016)	0.08	0.05	0.13	8.12
	Depression Post- partum	Fernandes Moll et al., (2019)	0.20	0.12	0.31	7.03
		Cardillo et al., (2016)	0.21	0.13	0.32	7.08
	Subtotal (I^2	2=0%, p<0.001)	0.15	0.06	0.25	22.22
	Overall (I^2=9	0.28	0.22	0.34	100.00	