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RESEARCH ARTICLE

The Specific Role of Relationship Life Events in the Onset of Depression during Pregnancy and the Postpartum

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Abstract

Background

The precipitating role of life events in the onset of depression is well-established. The present study sought to examine whether life events hypothesised to be personally salient would be more strongly associated with depression than other life events. In a sample of women making the first transition to parenthood, we hypothesised that negative events related to the partner relationship would be particularly salient and thus more strongly predictive of depression than other events.

Methods

A community-based sample of 316 first-time mothers stratified by psychosocial risk completed interviews at 32 weeks gestation and 29 weeks postpartum to assess dated occurrence of life events and depression onsets from conception to 29 weeks postpartum. Complete data was available from 273 (86.4%). Cox proportional hazards regression was used to examine risk for onset of depression in the 6 months following a relationship event versus other events, after accounting for past history of depression and other potential confounders.

Results

52 women (19.0%) experienced an onset of depression between conception and 6 months postpartum. Both relationship events (Hazard Ratio = 2.1, $p = .001$) and other life events (Hazard Ratio = 1.3, $p = .020$) were associated with increased risk for depression onset; however, relationship events showed a significantly greater risk for depression than did other life events ($p = .044$).

Conclusions

The results are consistent with the hypothesis that personally salient events are more predictive of depression onset than other events. Further, they indicate the clinical significance of events related to the partner relationship during pregnancy and the postpartum.

Introduction

Depression is one of the most prevalent and debilitating forms of psychopathology [1]. Impairment associated with the disorder can be severe, and the impact on the sufferer is amplified by the fact that depression is often recurrent, with each episode increasing the risk for experiencing a subsequent episode [2]. Epidemiological studies have consistently demonstrated a higher prevalence of depression in women, of one to three times higher than that of men [3].

Depression during pregnancy and the early postnatal period has been linked to a range of adverse outcomes for the developing foetus and child. Depression during pregnancy has been linked to risky health behaviours, such as substance use, tobacco and alcohol use and poor self-care [4] and health complications, such as preeclampsia [5,6] all of which can lead to negative obstetric and infant outcomes. Indeed, depression during pregnancy has been associated with poor obstetric outcomes, including low birth weight and preterm delivery [7–9]. Antenatal depression is also the strongest predictor of postnatal depression [10], which is associated with disturbances in mother-infant interactions [11, 12], later parenting disturbances [13] and problems in child cognitive, behavioural and social development [14–17].). Thus, the identification of mothers at risk for developing depression is essential to reduce the detrimental consequences of depression for the developing child. The frequent contact that pregnant women and new mothers have with health professionals provides an opportunity for identification and intervention. However, in order to identify those at risk, an understanding of the contributory factors to depression during this period is required.

The precipitating role of stress, particularly stressful life events, in the onset of major depression has received considerable research attention. Studies spanning several decades using both community and clinical populations have demonstrated that individuals with an onset of depression are significantly more likely to have experienced a recent stressful life event [18–21], with an average of 80% of episodes in women preceded by stressful life events [20]. The relationship between life events and depression has also been confirmed for depression during pregnancy (see review by Lancaster and colleagues [22]) and the postpartum (see review by Robertson and colleagues [10]). Indeed, birth itself may often act as a stressful life event [23].

An important yet largely neglected question regarding the nature of the life stress-depression relationship is whether certain types of life events are more depressogenic than others. A small body of research has examined this question, indicating that loss events [24–26] or events within the interpersonal domain [27, 28] are more strongly predictive of depression than other events. However, an important facet of life events that, despite possessing high face validity, has received little research attention is whether events that are meaningful or personally salient are more depressogenic. This premise was central to the development of the Life Events and Difficulties Schedule [29], a semi-structured interview in which an assessment of the ‘contextual threat’ of life events is conducted to produce a rating of their severity. This involves estimating the long-term implications of an event or difficulty for important plans, concerns, and purposes of the respondent. Brown and colleagues [30] tested this hypothesis using a prospective design by interviewing women to identify important areas of life commitment, for example, family or work. They found that the occurrence of a life event in the following year which ‘matched’ an area of commitment was almost three times more likely to result in onset of depression than a ‘non-matching’ event.

The method of matching events to commitments would suggest that these events were more personally salient to the women, and their findings support the hypothesis that salient events would be a stronger predictor of depression than other, less salient, events. However, another approach may be to examine a sample of individuals who are all making a similar life transition, allowing hypotheses to be made about events that may be particularly salient for that

group. There has been some work with adolescents taking this approach, predicting that interpersonal events, more specifically, the end of romantic relationships, would be particularly depressogenic [31]. Consistent with predictions, the authors found romantic relationship loss events to predict depression onset in adolescents after controlling for other life events, although this was only found for first onsets and not recurrences of depression.

The transition to parenthood is a period of significant change and disruption and thus represents an emotionally turbulent time for women [32]. Mothers may need to draw on all available coping resources, especially from within the partner relationship, and thus a stable and supportive relationship would seem to be particularly important. Events which threaten or disable the partner relationship, therefore, may be particularly distressing. There is substantial evidence for an association between the quality of the partner relationship and depression in perinatal women (see reviews [10, 22]). One recent cross-sectional study examined the prediction from a number of different life events to depressive symptoms in pregnancy in a sample of 693 women. They found severe marital conflict to be one of six significant predictors of depression symptoms in their multivariate model [33].

However, most previous studies have used self-report measures of psychosocial stressors and depression symptoms in cross-section. These have several limitations including that they do not provide evidence on whether or not the stressors have preceded the depressive symptoms, nor whether the reports of the stressors reflect depressive biases. Most of the studies with perinatal samples assess depression symptoms rather than diagnosis. For example, in a recent meta-analysis of 120 studies that examined associations between modifiable partner factors (e.g. conflict and partner support) and perinatal depression, only 13 assessed depression diagnosis [34]. This is the first study to make use of an established life event methodology in which the timing of events, and of onsets and offsets of depressive episodes, are assessed by interview to examine specific psychosocial risk for depression in the transition to parenthood. Using a sample of first time mothers who were interviewed during pregnancy and at 29 weeks postpartum, we tested the hypothesis that events related to the partner relationship would show a significantly higher risk for depression onset than other types of life events.

Method

Ethics statement

Ethical approval for the study was granted by the Cheshire North and West Research Ethics Committee on the 27th June 2006. The letter confirming ethical agreement for the study (reference number 05/Q1506/107) stated, 'On behalf of the Committee, I am pleased to confirm a favourable ethical approval for the above research on the basis described in the application form, protocol and supporting document as revised.' Participants gave written informed consent.

Sample

Participants were recruited into the Wirral Child Health and Development Study, a prospective epidemiological longitudinal study starting in pregnancy. The study used a two stage stratified design in which a consecutive general population sample of 1,233 first time pregnant women was recruited (the 'extensive' sample) and used to generate a subsample ('intensive' sample), for detailed study, of 316 women, stratified by psychosocial risk (see previous publication for sampling details [35]). The stratification variable, psychological abuse in current or recent relationship [36] was chosen for its known association with several risk factors for early child development. Data presented here were gathered from this intensive sample when the mothers were 32 weeks pregnant and at 29 weeks postpartum.

Socioeconomic conditions on the Wirral range between the deprived inner city and affluent suburbs, but with very low numbers from ethnic minorities. Mean age at recruitment was 27.9 years (s.d. 6.2, range 18–51) and 41.8% of the extensive sample were in the most deprived quintile of UK neighbourhoods using the English Index of Multiple Deprivation (IMD [37]).

There were 316 mothers recruited to the intensive sample at 32 weeks pregnancy, and 303 completed both the depression and life event interviews at 32 weeks, with 273 (86.4%) also completing the depression and life event interviews at 29 weeks postpartum. There were no significant differences in deprivation, risk status or maternal age between the 27 mothers who did not complete the postpartum assessment and the 273 who completed both, and they were not significantly more likely to have had a depression onset in pregnancy. The subsample of 273 was similar to the wider extensive sample in age (mean age = 27.8 years, s.d. 6.2, range 18–51) and deprivation status (37.7% in the most deprived quintile). Of the 273, 207 (75.8%) women were either married or cohabiting with a partner, 32 (11.7%) had a partner who lived elsewhere and 34 were single (12.5%). The majority, 95.2% ($N = 260$) of women described their ethnicity as White British. Within the stratified intensive sub-sample, 51% were drawn from the women with high psychosocial risk and 49% from those with low psychosocial risk.

Procedure

Mothers completed an initial brief interview at a routine antenatal visit to the hospital responsible for their antenatal care at approximately 20 weeks gestation, which included completion of the demographic questionnaire and partner psychological abuse measure. At 32 weeks gestation mothers completed a more in depth interview to gather information on mental health, recent life events, personality functioning and relationship functioning. Depression diagnosis, assessed using the Schedule for Affective Disorders and Schizophrenia-Lifetime (SADS [38]), and life events, gathered using the Life History Calendar (LHC [38]), were used for this study. The interviews were either conducted as home visits or in the study base, by graduate Research Assistants. A similar interview was then performed at 29 weeks postpartum. The mothers were recompensed for their time with high street shopping vouchers.

Measures

Demographic Questionnaire. A questionnaire was developed to collect demographic information, including marital and employment status, age and qualifications on leaving school, ethnic origin, and socio-economic status. Age and socio-economic status were included as covariates for this study. Socio-economic status was derived from post code data using the English Index of Multiple Deprivation (IMD) [37]. IMD scores were converted to quintile categories, with 1 being the 'most deprived' category, and 5 the 'least deprived'.

Psychological Abuse Questionnaire [36]. Partner psychological abuse was assessed at 20 weeks of pregnancy using a 20-item questionnaire covering humiliating, demeaning or threatening utterances in the partner relationship over the previous year. The scale is the total from 20 no/yes (coded as 0 = absent, 1 = present) items. Participants first rated these items about their own behaviour toward their partner and then about their partner's behaviour toward them. High and low psychological abuse strata were defined using the highest of the partner to participant and participant to partner scores for each family. A variable indicating whether the mother was high or low psychosocial risk allocation to the intensive sample was included as a covariate.

The Life History Calendar [39]. The LHC is a calendar-based structured interview designed to facilitate retrospective recall of life trajectories and events. The LHC comprises of a large grid, with columns representing the years and months, and rows referring to different

activities (e.g. residence, employment, life events) and is marked with key events in a respondent's life, such as birthdays. The interview begins by gathering information on life trajectories (e.g. where the respondent has lived) and then moves on to enquiring about specific events. This facilitates the recall of specific events by allowing the respondent to contextualise events by connecting them to other key life activities and events. The timing and duration of trajectories and events is then recorded on the calendar [39]. Reliability studies have reported agreement with previously collected concurrent data of 90% for a period of three years [39], 81% over a period of five years [40], and 69% to 79% across different trajectory categories over a period of one year [41].

In the present investigation, a list of 30 stressful life events was used with the LHC interview with a further open question allowing respondents to report any other stressful events they had experienced. Events reported under this category were included in analysis if they were present on other widely used life event checklists [42–44]. The two relationship events used were: end of romantic relationship and serious arguments with partner.

Schedule for Affective Disorders and Schizophrenia—Lifetime Version (SADS-L) [38]. The SADS-L is a structured diagnostic interview. The version used in this study employs an investigator-based approach and provides DSM-IV [45] based diagnoses [46]. Diagnoses of generalized anxiety disorder, panic disorder and major depression were assessed, and the major depression data used for this study. For the 32 week gestation interview, the SADS-L interview was adapted to allow ratings of DSM major depressive episodes in four lifetime periods: from one month before conception up to the 32 week interview; three years prior to conception to one month before conception; from age 16 to three years prior to conception; and before age 16. At the 29 weeks postpartum interview, ratings were made from the previous interview in pregnancy up to the current interview date. Life events and depression onsets were ascertained in the same interview and in many instances were rated by the same interviewer. However, all interviewers and coders were blind to the hypothesis that relationship life events would be more depressogenic than other types of life events.

In order to provide a constant assessment time period, depression onsets from conception to 6 months postpartum were examined for this study. A past history of depression variable was created by combining the three periods prior to pregnancy, with a positive rating in any of the periods used to indicate a past history of depression. The SADS-L was administered by trained graduate Research Assistants', 28 audio-recordings were independently rated for reliability purposes, and weighted kappa was acceptable, ranging from .83–1.00.

Statistical Analysis

To generate descriptive data regarding the life events preceding depression onset, the number and type of life events experienced in the 6 months prior to onset (based on [29, 47]) were examined for women who experienced an episode of depression ('cases'). To allow comparison to the women without a depression episode ('controls'), a random 6 month period was selected between conception and 6 months postpartum, and the number and type of life events occurring in this period was examined. The characteristics of cases and controls were estimated in SPSS.

The more formal analysis used a proportional hazards survival model [48] with time varying covariates to examine how the instantaneous risk of depression onset was associated with the relationship and non-relationship life-events in the preceding 6 months and other time fixed confounding maternal characteristics and a sample stratification factor to account for the sample design. Within the Cox proportional hazards model, the occurrence of a life event raises the risk for depression above baseline for a continuous block of 6 months after which the risk

returns to baseline (unless another event occurs within that 6 months, in which case the risk is doubled for the overlapping period), therefore in the model the woman is continuously exposed to the life event for that 6 month period. Wald tests were used to assess the significance of the log hazard ratios and to test whether there was a significant difference between the risk presented by relationship events and non-relationship events. In order to display the effects graphically four conditions were illustrated, one with no life events, and three with successive 6 months periods of risk associated with life events. These models, and associated plots, were undertaken in Stata 11 [49].

Results

Of the 273, 52 (19.0%) women experienced an onset of depression from conception to 6 months postpartum, 9 of these experienced more than one depression episode, but only first onsets were considered for this analysis. Demographic characteristics for the total sample and according to depression status are presented in [Table 1](#). There were no significant differences between cases and controls on age at consent ($t(273) = .66, p = .512$) or deprivation $\chi^2(4) = 1.41, p = .842$). There was a significant association between past history of depression and having a depression onset in the period ($\chi^2(2) = 7.49, p = .006$) with 61.2% of cases having previously experienced an episode of depression, compared to 39.7% of controls. Further, there was a significant association between sample stratification and depression onset ($\chi^2(2) = 4.68, p = .021$), 67.3% of cases were high risk allocation, compared to 50.7% of controls.

The descriptive statistics for both types of life event for cases and controls separately and the total sample are described in [Table 2](#). In the total sample, the majority of women (56.7%) experienced at least one non-relationship life event in the 6 month period, with a range of 0 to 6 events reported, and a higher proportion of cases (69.2%) experiencing at least one event than controls (53.2%). In the total sample, 12.1% of women experienced at least one relationship event, with a range of 0 to 3 events reported, and more cases (26.9%) experiencing at least one relationship event than controls (8.6%). The relationship events total score was very skewed (only 2.6% endorsed more than one event) so a binary variable was used to examine the significance of the association with depression onset. A significant association was found ($\chi^2(2) = 13.30, p = .001$) with an odds ratio of 3.92. Mann-Whitney U tests were used to examine whether there was a significant difference between cases and controls in the number of non-relationship events. Cases experienced significantly more non-relationship life events ($U(N = 273) = 6,941.50, Z = 2.67, p = .008$) than controls in the 6 month period.

[Table 3](#) presents the estimated hazard ratios (instantaneous relative risks) derived from the Cox proportional hazards survival model. A history of depression prior to pregnancy substantially and significantly raised the risk of depression, but neighbourhood deprivation, maternal age and the sample stratification factor were all non-significant. Having accounted for the effects of these potential confounders, the counts over the previous six months of non-relationship and relationship events were both estimated as having large and significant effects on the risk of depression. The Wald test of the equality for the effects of the two kinds of events indicated that relationship events were significantly more depressogenic than non-relationship events ($\chi^2(1) = 4.05, p = 0.044$). This difference is illustrated in [Fig 1](#). This shows from the time of conception, the proportion of women remaining non-depressed for four groups generated from the proportional hazards survival model: (1) women unexposed to either kind of event, (2) women continuously exposed to one non-relationship event in the preceding 6 months, (3) as (2) but for one relationship event, and (4) women continuously exposed to one of each kind of events. The graph suggests that by delivery, the proportion having experienced an onset of depression is 9% for those exposed to a non-relationship event, but 15% for those exposed to a

Table 1. Descriptive statistics for key demographic variables and past depression history for the total sample and cases and non-cases separately.

	Cases		Controls		Total	
	Mean	SD	Mean	SD	Mean	SD
Age	27.27	7.10	27.83	5.94	27.72	6.17
	N	%	N	%	N	%
Past depression						
Yes	30	61.2	85	39.7	114	43.7
No	22	38.8	136	60.3	159	56.3
Risk status^a						
High	35	67.3	112	50.7	147	53.8
Low	17	32.7	109	49.3	126	46.2
Deprivation^b						
1	18	34.6	85	38.5	103	37.7
2	10	19.2	44	19.9	54	19.8
3	14	26.9	58	26.2	73	26.7
4	5	9.6	12	5.4	18	6.6
5	5	9.6	22	10	25	9.2

^aHigh or low risk allocation to the sample

^b = Indices of multiple deprivation quintiles, 1 = most deprived

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relationship event. By 6 months postpartum, 16% of those exposed to a non-relationship event have experienced an onset of depression, whereas 30% of those exposed to a relationship event have experienced an onset.

Discussion

This study examined whether events hypothesised to be particularly salient for women making the first transition to parenthood would be a stronger predictor of depression onset than other types of life events. The results supported this hypothesis; whilst both types of events were significantly associated with risk for depression onset, the occurrence of a relationship event in the preceding six months was a significantly stronger predictor of depression onset than was the occurrence of other types of life events. This was found after accounting for past history of depression and other potential confounding variables.

Table 2. Descriptive statistics for relationship and non-relationship life events, for the total sample and for cases and non-cases separately.

	Cases		Controls		Total	
	Mean	SD	Mean	SD	Mean	SD
Relationship events	0.31	0.58	0.11	0.39	0.15	0.44
Other events	1.58	1.65	0.93	11.2	1.05	1.26
	N	% ^a	N	% ^a	N	% ^a
Relationship events	14	26.9	19	8.6	33	12.1
Other events	36	69.2	117	53.2	153	56.7

^aPercentage of women experiencing the event at least once in the 6 month period

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Table 3. The estimated hazard ratios (instantaneous relative risks) for relationship and non-relationship life events and the potential confounders.

	Hazard Ratio	P>z	95% CI
Non-relationship event	1.26	.020	1.04–1.53
Relationship event	2.13	.001	1.37–3.31
Past depression	2.24	.006	1.26–3.99
Deprivation	1.14	.225	0.92–1.42
Maternal age	.99	.643	0.94–1.04
Risk stratum	1.28	.613	0.49–3.31

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Both the non-relationship and relationship life events were associated with risk for depression, consistent with the substantial literature documenting an association between life events and depression onset [18–21]. However, the current study was novel in demonstrating that events related to the partner relationship were associated with a significantly higher risk for depression onset than were other types of events in a sample of perinatal women. The stronger prediction from relationship events is consistent with the literature which has documented associations between relationship quality and depression in perinatal women (see reviews [10,22]).

The findings are consistent with the hypothesis that life events which are more personally salient will be stronger predictors of depression onset than other life events. Previously, Brown and colleagues [30] demonstrated that the occurrence of life events which ‘matched’ an important area of life commitment were three times more likely to precipitate onset of depression than were other events. Further, Monroe and colleagues [31] hypothesised that the end of romantic relationship would be particularly meaningful for adolescents, and found that after controlling for 14 other life events from the Schedule of Recent Experiences [50] and daily hassles assessed using the Unpleasant Events Schedule [51], end of romantic relationship significantly increased the prediction of first but not recurrent onsets of depression. In the present study it was theorised that for women making the first transition to parenthood, events within the partner relationship would be particularly salient. The first transition to parenthood is a period of great emotional upheaval and so support and stability in the partner relationship would seem to be of particular importance. The results were consistent with the prediction that events which threaten the stability or mark the end of the partner relationship would be particularly depressogenic for a group of perinatal women.

The clinical significance of relationship events in perinatal depression is consistent with the theoretical underpinnings of Interpersonal Psychotherapy (IPT) for depression. IPT is rooted in attachment and interpersonal theory and is based on the premise that experiencing social disruptions increases the risk for depression [52]. A number of randomized controlled trials have provided support for the use of IPT for perinatal depression [53–55].

A strength of the current study lies in its prospective design, with two interview periods used to recall life events over a maximum period of 8 months, using the LHC, which has been demonstrated to yield reliable recall for longer periods of time [39]. This afforded some prospective prediction, with life events collected at the initial interview being used to predict onsets of depression reported at the second interview. The use of the LHC interview, which dates the occurrence of life events, also allowed that only events which were reported to have occurred before depression onset were considered for analysis. Failure to confirm this has been a problem in life event research using checklists [3] particularly in research with perinatal samples [56]. Ensuring that events precede depression onset is critical for examining the aetiological importance of relationship events, as evidence suggests that depression generates the occurrence of such events [57].

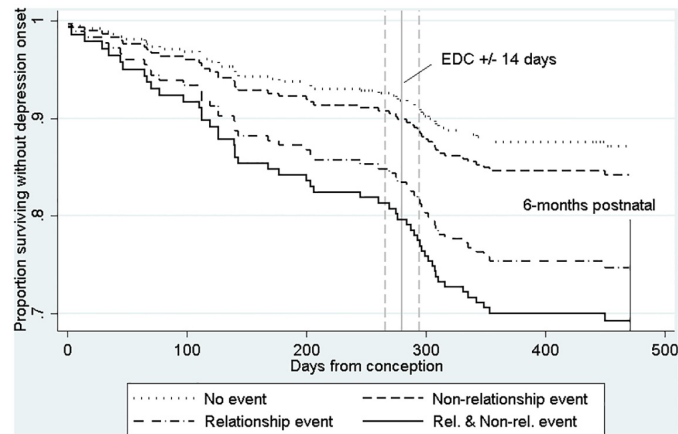


Fig 1. Survival plot showing model estimates for women with four different fixed life event exposures: 1) no events, 2) non-relationship event, 3) relationship event, 4) both relationship and non-relationship event.

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The reported findings should also be viewed in light of several limitations. Firstly, the life event and depression interviews were conducted in the same interview session, which allows the possibility of reporting bias leading to inflated associations. However, as this should affect relationship and non-relationship events equally it would not explain the different associations found for the two types of events. Secondly, as both the life event and depression interviews were administered by the same interviewer, and typically the same interviewer would administer the pregnancy and postpartum interview for each mother, the depression ratings were not always made independently of life event information, although it was ensured that all interviewers and raters were blind to the study hypothesis. Thirdly, a measure of relationship abuse administered during pregnancy was used to stratify the sample; this benefited the analysis by ensuring a sufficient number of relationship events within the time period examined. However, it does limit the generalisability of the findings, as 51% of the intensive sample were high on psychosocial risk. Stratification status was controlled for in all analyses but replication of the findings in a true general population sample would be desirable, particularly given the low representation of ethnic minority groups in the current sample. Finally, the study design does not allow us to evaluate the contribution of relationship life events in relation to other likely causes, notable genetic factors, nor does it account for heterogeneity within depressive disorders.

The identification of a specific type of life event as particularly depressogenic for women making the first transition to parenthood has important implications for practice. Pregnant and postpartum women have increased contact with health professionals, but may be less likely to seek help for depression due to the social stigma attached to depression experienced during this period [58]. The identification of specific risk factors for depression onset facilitates identification of those at risk and allows early intervention. Recent international clinical guidelines outlined the importance of conducting psychosocial risk assessments with perinatal women [59] and the current findings further underscore the importance of screening for the occurrence relationship events. As previously noted, the findings may also lend support for the use of IPT in the treatment of perinatal depression.

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Author Contributions

Conceived and designed the experiments: JH HS AP. Performed the experiments: NW. Analyzed the data: AP NW. Wrote the paper: NW AP HS JH.

References

1. Gotlib I, Hammen C. Introduction. In Gotlib I and Hammen C, editors, *Handbook of depression*. New York: Guilford Press; 2002. p.1–20.
2. American Psychiatric Association *Diagnostic and statistical manual of mental disorders*. 4th Ed. Text Revision. Washington, DC: American Psychiatric Association; 2000.
3. Kessler RC. The effects of stressful life events on depression. *Annu Rev Psychol*. 1997; 48: 191–214. PMID: [9046559](#)
4. Zuckerman B, Amaro H, Bauchner H, Cabral H. Depressive symptoms during pregnancy: Relationship to poor health behaviors. *Am J Obstet Gynecol*. 1989; 160: 1107–1111. PMID: [2729387](#)
5. Kurki T, Hiilesmaa V, Raitasalo R, Mattila H, Ylikorkla O. Depression and anxiety in early pregnancy and risk for preeclampsia. *Obstet Gynecol*. 2000; 95: 487–490. PMID: [10725477](#)
6. Schmeelk KH, Granger DA, Susman EJ, Chrousos GP. Maternal depression and risk of postpartum complications: Role of prenatal corticotropin-releasing hormone and interleukin-1 receptor antagonist. *Behav Med*. 1999; 25, 88–93. PMID: [10401538](#)
7. Grote NK, Bridge JA, Gavin AR, Melville JL, Iyengar S, Katon WJ. A meta-analysis of depression during pregnancy and the risk of preterm birth, low birth weight and intrauterine growth restriction. *Arch Gen Psychiat*. 2010; 67: 1012–1024. doi: [10.1001/archgenpsychiatry.2010.111](#) PMID: [20921117](#)
8. Hedegaard M, Henriksen TB, Sabroe S, & Secher NJ. Psychological distress in pregnancy and preterm delivery. *BMJ*. 1993; 307: 234–239. PMID: [8369684](#)
9. Hoffman S, Hatch MC. Depressive symptomatology during pregnancy: evidence for an association with decreased fetal growth in pregnancies of lower social class women. *Health Psychol*. 2000; 19: 535–543. PMID: [11129356](#)
10. Robertson E, Grace S, Wallington T, Stewart DE. Antenatal risk factors for postpartum depression: a synthesis of recent literature. *Gen Hosp Psychiat*. 2004; 26: 289–295.
11. Cohn JF, Matias R, Tronick EZ, Connell D, Lyons-Ruth K. Face-to-face interactions of depressed mothers and their infants. In Tronick EZ, Fields T, editors, *Maternal depression and infant disturbance*. San Francisco: Jossey-Bass; 1986. p. 31–45.
12. Field T, Sandberg D, Garcia R, Vega-Lahr N, Goldstein S, Guy L. Pregnancy problems, postpartum depression and early mother-infant interactions. *Dev Psychol*. 1985; 21: 1152–1156.
13. Lovejoy C, Graczyk PA, O'Hare E, Neuman G. Maternal depression and parenting behavior: A meta-analytic review. *Clin Psychol Rev*. 2000; 20: 561–592. PMID: [10860167](#)
14. Connell AM, Goodman SH. The association between psychopathology in fathers versus mothers and children's internalizing and externalizing behavior problems. *Psychol Bull*. 2002; 128: 746–773. PMID: [12206193](#)
15. Grace SL, Evindar A, Stewart DE. The effect of postpartum depression on child cognitive development and behavior: a review and critical analysis of the literature. *Arch Women Ment Health*. 2003; 6: 263–74.
16. Hammen C, Burge D, Burney E, Adrian C. Longitudinal study of diagnoses in children of women with unipolar and bipolar affective disorder. *Arch Gen Psychiat*. 1990; 47: 1112–1117. PMID: [2244796](#)
17. Luoma I., Tamminen T., Kaukonen P., Laippala P., Puura K., Salmelin R. et al. Longitudinal study of maternal depressive symptoms and child well-being. *J Am Acad Child Adolesc Psychiat*. 2001; 40: 1367–1374.
18. Brown GW, Harris TO. *Life Events and Illness*. New York: Guilford; 1989.
19. Hammen C. Stress and depression. *Annu Rev Clin Psychol*. 2005; 1: 293–319. PMID: [17716090](#)
20. Mazure CM. Life stressors as risk factors in depression. *Clin Psychol-Sci Pr*. 1998; 5: 291–313.

21. Paykel ES. Life events and affective disorders. *Acta Psychiatr Scand*. 2003; 108: 61–66.
22. Lancaster C, Gold K, Flynn H, Marcus S, Davis M. Risk factors for depressive symptoms during pregnancy: A systematic review. *Am J Obstet Gynecol*. 2010; 202: 5–14. doi: [10.1016/j.ajog.2009.09.007](https://doi.org/10.1016/j.ajog.2009.09.007) PMID: [20096252](https://pubmed.ncbi.nlm.nih.gov/20096252/)
23. Murray D, Cox JL, Chapman G, Jones P. Childbirth: Life event or start of a long-term difficulty? Further data from the Stoke-on-Trent controlled study of postnatal depression. *Brit J Psychiat*. 1995; 166: 595–600. PMID: [7620743](https://pubmed.ncbi.nlm.nih.gov/7620743/)
24. Brown GW, Harris TO, Hepworth C. Loss, humiliation and entrapment among women developing depression: a patient and non-patient comparison. *Psychol Med*. 1995; 25: 7–21. PMID: [7792364](https://pubmed.ncbi.nlm.nih.gov/7792364/)
25. Finlay-Jones R, Brown GW. Types of stressful life events and the onset of anxiety and depressive disorders. *Psychol Med*; 1981; 11: 803–815. PMID: [7323236](https://pubmed.ncbi.nlm.nih.gov/7323236/)
26. Kendler KS, Hettema JM, Butera F, Gardner CO, Prescott CA. Life event dimensions of loss, humiliation, entrapment, and danger in the prediction of onsets of major depression and generalized anxiety. *Arch Gen Psychiat*. 2003; 60: 789–96. PMID: [12912762](https://pubmed.ncbi.nlm.nih.gov/12912762/)
27. Kendler KS, Thornton LM, Prescott CA. Gender differences in the rates of exposure to stressful life events and sensitivity to their depressogenic effects. *Am J Psychiat*. 2001; 158: 587–93. PMID: [11282693](https://pubmed.ncbi.nlm.nih.gov/11282693/)
28. Zimmermann-Tansella C, Donini S, Lattanzi M, Siciliani O, Turrina C, Wilkinson G. Life events, social problems and physical health status as predictors of emotional distress in men and women in a community setting. *Psychol Med*. 1991; 21: 505–513 PMID: [1876655](https://pubmed.ncbi.nlm.nih.gov/1876655/)
29. Brown GW, Harris T. *Social origins of depression: A study of psychiatric disorder in women*. London: Tavistock; 1978.
30. Brown GW, Bifulco A, Harris TO. Life events, vulnerability and onset of depression: some refinements. *Brit J of Psychiat*. 1987; 150: 30–42.
31. Monroe SM, Rohde P, Seeley JR, Lewinsohn P. Life events and depression in adolescence: Relationship loss as a prospective risk factor for first onset of major depressive disorder. *J Abnorm Psychol*. 1999; 108: 606–614. PMID: [10609425](https://pubmed.ncbi.nlm.nih.gov/10609425/)
32. Gotlib IH, Whiffen VE, Wallace P, Mount JH. A prospective investigation of postpartum depression: Factors involved in onset and recovery. *J Abnorm Psychol*. 1991; 100: 122–132. PMID: [2040762](https://pubmed.ncbi.nlm.nih.gov/2040762/)
33. Dayan J., Creveuil C., Dreyfus M., Herlicoviez M., Baleytel J-M, & O'Keane V. Developmental model of depression applied to prenatal depression: Role of present and past life events, past emotional disorders and pregnancy stress. *PLoS One*. 2010; 5(9): e12942. doi: [10.1371/journal.pone.0012942](https://doi.org/10.1371/journal.pone.0012942) PMID: [20877652](https://pubmed.ncbi.nlm.nih.gov/20877652/)
34. Pilkington PD, Milne LC, Cairns KE, Lewis J, Whelan TA. Modifiable partner factors associated with perinatal depression and anxiety: A systematic review and meta-analysis. *J Affect Disord*. 2015; 178: 165–180. doi: [10.1016/j.jad.2015.02.023](https://doi.org/10.1016/j.jad.2015.02.023) PMID: [25837550](https://pubmed.ncbi.nlm.nih.gov/25837550/)
35. Sharp H, Pickles A, Meaney M, Marshall K, Tibu F, Hill J. Frequency of infant stroking reported by mothers moderates the effect of prenatal depression on infant behavioural and physiological outcomes. *PLoS ONE*. 2012; 7: e45446. doi: [10.1371/journal.pone.0045446](https://doi.org/10.1371/journal.pone.0045446) PMID: [23091594](https://pubmed.ncbi.nlm.nih.gov/23091594/)
36. Moffitt TE, Caspi A, Margolin G, Krueger RF, Magdol L, Silva PA et al. Do partners agree about abuse in their relationship? A psychometric evaluation of interpartner agreement. *Psychol Assessment*. 1997; 9: 47–56.
37. Noble M, Wright G, Dibben C, Smith G, McLennan D, Anttila C. *The English Indices of Deprivation 2004 (revised)*. Report to the Office of the Deputy Prime Minister. London: Neighbourhood Renewal Unit; 2004.
38. Spitzer R, Endicott J. *The schedule for affective disorders and schizophrenia, lifetime version*, 3rd ed. New York: New York State Psychiatric Institute; 1977.
39. Caspi A, Moffitt TE, Thornton A, Freedman D, Amell JW, Harrington HL et al. The life history calendar: A research and clinical assessment method for collecting retrospective event-history data. *Int J Method Psych*. 1996; 6: 101–114.
40. Freedman D, Thornton A, Camburn D, Alwin D, Young-DeMarco L. The life history calendar: A technique for collecting retrospective data. In Clogg CC, editor, (Ed.) *Sociological Methodology* Vol.18. Washington, D.C.: American Sociological Association; 1988. p.37–68.
41. Belli RF, Shay WL, Stafford FP. Event history calendars and question list surveys: A direct comparison of interviewing methods. *Public Opin Quart*. 2001; 65: 45–74.
42. Brugha T, Bebbington P, Tennant C, Hurry J. The List of Threatening Experiences: a subset of 12 life event categories with considerable long-term contextual threat. *Psychol Med*. 1985; 15: 189–194. PMID: [3991833](https://pubmed.ncbi.nlm.nih.gov/3991833/)

43. Caspi A, Karen S, Moffitt T, Taylor A, Craig IW, Harrington H, McClay L et al. Influence of life stress on depression: moderation by a polymorphism in the 5-HTT gene. *Science*. 2003; 301(5631): 386–389. PMID: [12869766](#)
44. Kendler KS, Karowski LM, Prescott CA. Causal relationship between stressful life events and the onset of major depression. *Am J Psychiat*. 1999; 156: 837–841. PMID: [10360120](#)
45. American Psychiatric Association. *Diagnostic and statistical manual of mental disorders*. 4th Ed. Washington, DC: American Psychiatric Association; 1994.
46. Harrington R, Hill J, Rutter M, John K, Fudge H, Zoccolillo M, Weissman M. The assessment of lifetime psychopathology: a comparison of two interviewing styles. *Psychol Med*. 1988; 18: 487–93. PMID: [3399595](#)
47. Surtees PG, Rennie D. Adversity and the onset of psychiatric disorder in women. *Soc Psychiat*. 1983; 18: 37–44.
48. Cox DR. Regression models and life tables. *J Roy Stat Soc B*. 1972; 34: 187–220.
49. StataCorp. *Stata Statistical Software: Release 12*. College Station, TX: StataCorp LP; 2011.
50. Holmes TH, Rahe RH. The Social Readjustment Rating Scale. *J Psychosom Res*. 1967; 11:213–8. PMID: [6059863](#)
51. Lewisohn PM, Mermelstein RM, Alexander C, MacPhillamy DJ. The Unpleasant Events Schedule: a scale for measurement of aversive events. *J C Psychol*. 1985; 41: 483–498.
52. Stuart S, O'Hara MW. Interpersonal psychotherapy for postpartum depression: a treatment program. *J Psychother Pract Res*. 1995; 4(1): 18–29. PMID: [22700210](#)
53. O'Hara MW, Stuart S, Gorman LL, Wenzel A. Efficacy of interpersonal psychotherapy for postpartum depression. *Arch Gen Psychiat*. 2000; 57(11): 1039–1045. PMID: [11074869](#)
54. Spinelli MG, Endicott J. Controlled clinical trial of interpersonal psychotherapy versus parenting education program for depressed pregnant women. *Am J Psychiat*. 2003; 160(3): 555–562. PMID: [12611838](#)
55. Spinelli MG, Endicott J, Leon AC, Goetz RR, Kalish RB, Brustman LE et al. A controlled clinical treatment trial of interpersonal psychotherapy for depressed pregnant women at 3 New York City sites. *J Clin Psychiat*. 2013; 74(4): 393–399.
56. Swendsen JD, Mazure JD. Life stress as a risk factor for postpartum depression: Current research and methodological issues. *Clin Psychol-Sci Pr*. 2000; 7: 17–31.
57. Liu RT, Alloy LB. Stress generation in depression: A systematic review of the empirical literature and recommendations for future study. *Clin Psychol Rev*. 2010; 30: 582–93. doi: [10.1016/j.cpr.2010.04.010](#) PMID: [20478648](#)
58. Beck CT. Predictors of postpartum depression: An update. *Nurs Res*. 2001; 50: 275–285. PMID: [11570712](#)
59. Austin MP, Marcé Society Position Statement Advisory Committee. Marcé International Society position statement on psychosocial assessment and depression screening in perinatal women. *Best Pract Res Cl Ob*. 2014; 28(1): 197–187.