

# **Regular Article**

# Preconception maternal posttraumatic stress and child negative affectivity: Prospectively evaluating the intergenerational impact of trauma

Danielle A. Swales<sup>1</sup>, Elysia Poggi Davis<sup>1,2</sup>, Nicole E. Mahrer<sup>3</sup>, Christine M. Guardino<sup>4</sup>, Madeleine U. Shalowitz<sup>5</sup>, Sharon L. Ramey<sup>6</sup> and Chris Dunkel Schetter<sup>7</sup>

<sup>1</sup>Department of Psychology, University of Denver, Denver, CO, USA, <sup>2</sup>Department of Psychiatry and Human Behavior, University of California, Irvine, CA, USA, <sup>3</sup>Department of Psychology, University of La Verne, La Verne, CA, USA, <sup>4</sup>Department of Psychology, Dickinson College, Carlisle, PA, USA, <sup>5</sup>Department of Pediatrics, NorthShore University HealthSystem Research Institute, Evanston, IL, USA, <sup>6</sup>Department of Psychology, Virginia Tech, Blacksburg, VA, USA and <sup>7</sup>Department of Psychology, University of California, Los Angeles, CA, USA

### **Abstract**

The developmental origins of psychopathology begin before birth and perhaps even prior to conception. Understanding the intergenerational transmission of psychopathological risk is critical to identify sensitive windows for prevention and early intervention. Prior research demonstrates that maternal trauma history, typically assessed retrospectively, has adverse consequences for child socioemotional development. However, very few prospective studies of preconception trauma exist, and the role of preconception symptoms of posttraumatic stress disorder (PTSD) remains unknown. The current study prospectively evaluates whether maternal preconception PTSD symptoms predict early child-hood negative affectivity, a key dimension of temperament and predictor of later psychopathology. One hundred and eighteen women were recruited following a birth and prior to conception of the study child and were followed until the study child was 3–5 years old. Higher maternal PTSD symptoms prior to conception predicted greater child negative affectivity, adjusting for concurrent maternal depressive symptoms and sociodemographic covariates. In exploratory analyses, we found that neither maternal prenatal nor postpartum depressive symptoms or perceived stress mediated this association. These findings add to a limited prospective literature, highlighting the importance of assessing the mental health of women prior to conception and providing interventions that can disrupt the intergenerational sequelae of trauma

Keywords: intergenerational transmission; negative affectivity; preconception; PTSD; trauma

(Received 4 March 2021; revised 8 December 2021; accepted 8 December 2021; First Published online 25 January 2022)

Maternal mental health history is strongly implicated in the developmental trajectory of child emotional development and psychopathology risk (Goodman & Gotlib, 1999; Bijl et al., 2002; Swales et al., 2020). This intergenerational link between maternal and child psychological wellbeing is especially evident among mothers who have experienced trauma and subsequent symptoms of post-traumatic stress (Lambert et al., 2014). Almost 7 out of every 10 adult women will be exposed to a traumatic event in their lifetime (Resnick et al., 1993) and approximately 4%–6% will go on to meet criteria for posttraumatic stress disorder (PTSD) (National Comorbidity Survey, 2017). Notably, PTSD rates are even higher among women living in poverty (Seng et al., 2009). Maternal trauma and subsequent PTSD symptoms have been shown to have intergenerational consequences, including deleterious effects on child socioemotional development and mental health outcomes,

 $\textbf{Corresponding author:} \ Danielle \ A. \ Swales, \ email: \\ \underline{danielle.swales@du.edu}$ 

Cite this article: Swales, D. A., et al. (2023). Preconception maternal posttraumatic stress and child negative affectivity: Prospectively evaluating the intergenerational impact of trauma. Development and Psychopathology 35: 619–629, https://doi.org/10.1017/S0954579421001760

such as elevated risk for anxiety, depression, and behavior problems (Lambert et al., 2014).

Consistent with a Developmental Origins of Health and Disease (DOHaD) framework, maternal perinatal PTSD symptoms are associated with their children's vulnerability to psychopathology via biological mechanisms of fetal programming as well as postnatal environmental cues (e.g., parenting behaviors, quality of parent-child attachments) (Doyle & Cicchetti, 2018). However, a growing literature suggests that ontogenetic pathways of risk and resilience for the development of psychopathology do not begin at conception (Keenan et al., 2018). Rather maternal mental health prior to conception, including PTSD symptoms, may also relate to developmental trajectories of mental health in the next generation via alterations in prenatal stress physiology, epigenetic pathways, and other peripartum mechanisms (Bale, 2014; Bowers & Yehuda, 2016; Buss et al., 2017; Keenan et al., 2018; Scorza et al., 2019; Swales et al., 2018). The current study expands the DOHaD framework to include preconception influences, utilizing a prospective, longitudinal design to examine how maternal PTSD symptoms prior to conception may relate to outcomes in the next generation, such as negative affectivity, which contribute to

© The Author(s), 2022. Published by Cambridge University Press. This is an Open Access article, distributed under the terms of the Creative Commons Attribution licence (http://creativecommons.org/licenses/by/4.0/), which permits unrestricted re-use, distribution and reproduction, provided the original article is properly cited.



subsequent vulnerability to later psychopathology (e.g., Clark et al., 1994; Derryberry & Rothbart, 1997; Lonigan et al., 2004; Muris & Ollendick, 2005; Nigg, 2006).

# Intergenerational consequences of preconception stress

Experimental research with animals provides compelling evidence that maternal stress exposure prior to conception exerts lasting intergenerational consequences (see Klengel et al., 2016 for review). Maternal stress exposure (e.g., chronic and unpredictable stress, overcrowding, temperature) that is limited to the preconception period adversely impacts rat offspring brain development including anatomical changes in the medial prefrontal cortex (Bock et al., 2016; Harker et al., 2015; Huang et al., 2010), altered stress physiology (Zaidan et al., 2013), and increased anxious and anhedonic behavior (Bock et al., 2016; Li et al., 2010; Zaidan et al., 2013). These experimental studies provide important evidence that severe stress exposure limited to the preconception period has intergenerational consequences.

In humans, evidence for the intergenerational transmission of trauma has been supported by epidemiological research as well as studies of collective trauma (e.g., parental exposure to genocide prior to conception) (Class et al., 2014, 2015; Flory et al., 2011; Gangi et al., 2009; Keenan et al., 2018; Perroud et al., 2014; Power et al., 2007; Solomon et al., 1988; Yehuda et al., 2001, 2008). Additionally, maternal retrospective report of traumatic and stressful life experiences, such as history of childhood maltreatment (e.g., Adverse Childhood Events Scale, Felitti et al., 1998; and Childhood Trauma Questionnaires, Bernstein et al., 1994), show that maternal trauma can have intergenerational effects. Children of women with trauma histories, assessed by maternal recall of trauma in her own childhood, are more likely to experience dysregulated temperament in infancy, including heightened negative affectivity (Bosquet Enlow et al., 2017; Lang et al., 2010) and poor emotional and behavioral outcomes in childhood (Bosquet et al., 2018; Khan & Renk, 2019; Plant et al., 2017, 2018). These studies provide epidemiological and retrospective evidence that maternal history of traumatic and potentially traumatic experiences prior to conception predicts poor emotional outcomes for the next generation. However, additional prospective research is needed to address limitations inherent to such study designs. First, retrospective recall of adverse experiences is only moderately associated with prospectively collected measures (see Baldwin et al., 2019 for meta-analysis and systematic review). Second, retrospective recall of PTSD symptoms is even more susceptible to recall biases than recall of past trauma exposure (Baldwin et al., 2019; Hardt & Rutter, 2004; Moffitt et al., 2010; Newbury et al., 2018; Reuben et al., 2016).

# Prospective assessment of preconception trauma

Given the inherent limitations of retrospective studies, there has been increasing recognition of the need for *prospective*, longitudinal studies of preconception stress (Bowers & Yehuda, 2016; Keenan et al., 2018). However, a dearth of prospective research has examined women's mental health prior to pregnancy and child outcomes due to the logistical challenges of recruiting women before conception and following them through their pregnancies. A few published prospective studies show that elevated preconception stress and exposure to stressful life experiences relate to adverse birth outcomes (i.e., shorter gestation and lower birthweight) (Harville et al., 2010; Mahrer et al., 2020) and poor sleep quality in infancy (Baird et al., 2009). A large prospective study of

women in Australia (n = 756) reported that maternal preconception symptoms of depression and anxiety predicted heightened emotional reactivity in infancy (Spry et al., 2020). Another study conducted by Hipwell et al. (2019) prospectively assessed maternal childhood trauma exposure prior to conception via standardized interviews and items from the Parent-Child Conflict Tactics Scale and found that infants of mothers exposed to childhood emotional abuse were more likely to demonstrate low emotional reactivity to the still face procedure, whereas infants of mothers exposed to childhood emotional neglect demonstrated heightened emotional reactivity, even after controlling for postpartum depressive symptoms. Collectively, these few, prospective studies are important as they reduce the influence of recall bias and affirm the need to expand research on preconception maternal trauma and mental health and child emotion regulation outcomes.

A key gap in this literature is that additional studies are needed to prospectively assess whether preconception symptoms of *post-traumatic stress* predict outcomes for the next generation. It is important to distinguish maternal *exposure* to trauma or a potentially traumatic event from posttraumatic stress *symptoms* because not all women who are exposed to a potentially traumatic event will develop subsequent PTSD symptoms. Rather mediating pathways of risk and resilience have been shown to amplify or disrupt the emergence of subsequent psychopathology following trauma exposure (McLaughlin & Lambert, 2017). Identifying whether preconception maternal PTSD symptoms relate to child outcomes is therefore important next step in exploring whether maternal mental health prior to conception confers vulnerability for subsequent psychopathology in the next generation.

Additional work is also needed to explore potential pathways that may underly this hypothesized link between preconception PTSD symptoms and child psychopathological risk. Perinatal mental health, especially maternal depression, is a plausible candidate pathway because prior work suggests that offspring may be more vulnerable to the biopsychosocial sequelae of maternal trauma during the fetal period of development (Bouvette-Turcot et al., 2020; Bowers & Yehuda, 2016; Buss et al., 2017; Davis & Narayan, 2020; Keenan et al., 2018; Swales et al., 2018; Yehuda & Meaney, 2018). Past studies demonstrate that retrospective recall of trauma history is associated with elevated internalizing symptoms (e.g., depression, anxiety, and PTSD) during pregnancy (Alvarez-Segura et al., 2014; Atzl et al., 2019; River et al., 2019), and prenatal maternal mental health predicts heightened negative emotionality in offspring during childhood (Blair et al., 2011; Davis et al., 2007; Erickson et al., 2017; Glynn et al., 2018; Werner et al., 2007). Prenatal depression is a potential pathway of particular interest because PTSD and depression diagnoses are highly comorbid (O'Donnell et al., 2004) and depressive symptoms are commonly screened for in clinical settings and thus offer implications for clinical practice. The current study provides a unique opportunity to investigate these exploratory pathways.

# The current study

The current study prospectively evaluates the association between maternal preconception symptoms of posttraumatic stress and child negative affectivity. Mothers were from a socioeconomically, racially, and ethnically diverse larger cohort of women who were studied following a birth. This subset was then followed when mothers became pregnant again with the study child and until the study child was 3–5 years of age. *Preconception* is defined in the current study as prior to conception of the study child (which

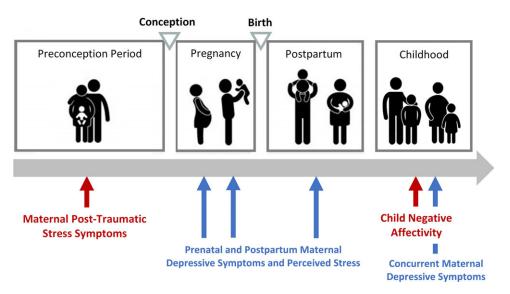


Figure 1. Overview of study measures from preconception to early childhood.

for all participants in the study sample is also an interconception period between consecutive births). As the primary aim of the current analyses, we evaluated whether preconception maternal PTSD symptoms predict child negative affectivity after accounting for sociodemographic covariates and concurrent maternal depressive symptoms (an indicator of concurrent maternal psychopathology and reporter bias). In exploratory post hoc analyses, we tested maternal prenatal and postpartum symptoms of depression as potential mediators in the relation between preconception symptoms of PTSD and child negative affectivity. Prenatal and postpartum levels of perceived stress were additionally considered as mediators in secondary, exploratory analyses.

# Method

### **Participants**

Participants were 118 mother-child dyads. Women were recruited from three study sites in the Community Child Health Network (CCHN) (i.e., North Carolina, Washington, DC, and Lake County, IL). CCHN is a multisite research network designed to investigate disparities in maternal and child mental health (see Dunkel Schetter et al., 2013; O'Campo et al., 2016; and Ramey et al., 2015 for overview of CCHN recruitment procedures, study design, data collection methods, and cohort demographics). Two hundred and forty-five women from the three selected CCHN study sites who became pregnant again between 2009 and 2013 and were assessed prospectively from preconception (i.e., between births and prior to conception of the study child) through their pregnancy and into the postpartum period. One hundred and twenty-seven women then went on to participate in a follow-up assessment when the study child was 3–5 years of age.

Additional inclusion criteria for the current analyses were mothers who provided: (a) ratings of preconception PTSD or completed at least one prenatal study visit (i.e., provided at least one rating of mood or stress during pregnancy), and (b) ratings of child negative affectivity in early childhood. One hundred and nineteen women met these criteria. One dyad was excluded from analyses due to preterm birth less than 34 weeks' gestation. See Figure 1 for overview of study design.

The 118 women in the current study analyses were more likely to be older in age (t = 4.1, p < .001), have a longer interpregnancy interval (t = 2.6, p = .002), have a lower adjusted household income (t = -2.0, p = .049), be Latina/Hispanic ( $X^2 = 8.6$ , p = .003), and be born outside of the United States ( $X^2 = 8.6$ , p = .003), in comparison to women who became pregnant during the eligibility window but were not included in the present analyses (either because they did not participate in the follow-up assessment in early childhood and/or were missing preconception or prenatal study measures). Sample attrition was not associated with maternal education (t = -1.1, p = .275), concurrent depressive symptoms (t = 0.7, t = .484), study site (t = 0.8), t = .655), cohabitation status (t = 0.4), or PTSD symptoms (t = 0.0), t = 0.326).

Mothers included in the current study sample were an average of 33 years of age at the time of child assessment (SD=5.5 years) and 39% identified as Latina/Hispanic, 32% non-Latina/Hispanic Black, and 29% non-Latina/Hispanic White (see Supplemental Table S1 for descriptive information on study measures stratified by maternal race/ethnicity). Additionally, 40% of women were born outside of the United States (46 out of the 47 of whom also identified as Latina or Hispanic). Further, 38% lived below the federal poverty line (FPL), and 21% lived near poverty at 100%–200% FPL. Children were on average 3.8 years of age at the time of assessment (SD=0.4 years) and 58% were female (see Table 1). All study procedures were approved by the Institutional Review Board for the protection of human subjects at the relevant institutions, and each mother provided written and informed consent for herself and her child.

# **Procedures**

All self-report data were collected through semistructured interviews by trained interviewers in participants' homes (see Dunkel Schetter et al., 2013). Maternal ratings of posttraumatic stress symptoms and demographic data were collected before women became pregnant with the study child. For present purposes, the preconception assessment occurred an average of 52.9 weeks prior to conception of the study child, although time from assessment to conception varied (SD = 47.3, Range: 2.0-175.0 weeks). The beginning of the pregnancy was determined using the first day of the last

Table 1. Sample characteristics

Sample characteristics (n = 118)	
Study site (N; %)	
Chicago, Illinois	74; 62.7
Rural eastern North Carolina	17; 14.4
District of Columbia	27; 22.8
Time from preconception assessment to conception (in weeks)	52.9 ± 47.3; 2-17
Maternal characteristics	
Maternal age (in years; M ± SD; Range)	33.4 ± 5.5; 24.1-44.2
Cohabitating with partner (N; %)	77; 65.3
Race/Ethnicity (N; %)	
Latina/Hispanic	46; 39.3
Non-Latina/Hispanic Black	38; 32.0
Non-Latina/Hispanic White	34; 28.7
Nativity status (N; % born outside of the U.S.)	47; 39.8
Federal poverty line (FPL) (N; %)	
Poverty (<100% FPL)	45; 38.1
Near poverty (100%–200% FPL)	25; 21.1
Above poverty (>200% FPL)	48; 40.7
Per capita household income adjusted for cost of living ( <i>M</i> ± <i>SD</i> ; <i>Range</i> )	13,757.0 ± 7,232.4; 0-89,222.0
Years of education (M ± SD; Range)	12.6 ± 2.9; 5-21
Parity (N; %)	
1	62; 52.5
2	38; 32.2
3	17; 14.4
Child characteristics	
Child age at assessment (in years; M ± SD; Range)	3.8 ± .4; 3.4–5.5
Biological sex at birth (N; % female)	68; 57.7
Gestational age at birth (in weeks) ( <i>M</i> ± <i>SD</i> ; <i>Range</i> )	38.8 ± 2.1; 32.6-42

menstrual period and/or ultrasound examinations, obtained via medical record abstraction. Standardized measures of maternal depression symptoms and perceived stress were administered twice during pregnancy in second and third trimesters, on average at 20.2 weeks gestation (SD=5.0) and 32.7 weeks gestation (SD=3.8). Maternal depression symptoms were assessed once during the postpartum period on average at 12.5 weeks after birth (SD=4.7). Additional perinatal and birth outcomes data were extracted from neonatal records. Mother-child dyads participated in an early childhood study visit when the child was 3–5 years of age, at which time child's negative affectivity was assessed. Early childhood visits were conducted in participants' homes, in the mother's preferred language (English or Spanish).

### Measures

### Maternal measures

Sociodemographic and pregnancy factors. Sociodemographic characteristics, including maternal age, years of education, per capita household income adjusted for cost of living, cohabitation status

(i.e., living with partner), parity, foreign birth (i.e., whether mother was born in a country outside of the United States), and race/ethnicity, were assessed during interviews. An outlier for adjusted household income (i.e., >8 SD above the mean) was replaced with a value 3 SD above the mean, retaining its rank as the highest value. Maternal education and per capita household income adjusted for cost of living were standardized and averaged together to create a composite of socioeconomic status (SES; Cohen et al., 2006). Maternal early life adversity was assessed by the 10-item Risky Families Questionnaire (Taylor et al., 2004), a measure of family stressors during the mother's childhood (e.g., childhood abuse, parental warmth). Mothers completed the measure during the early childhood visit, rating each item on a 5-point Likert scale, with a range of 1 (not at all) to 5 (very often). Final sum scores could range from 10 to 50, with higher scores indicating greater early life adversity. Gestational age at birth was determined from medical record abstraction.

Maternal posttraumatic stress symptoms. Maternal symptoms of posttraumatic stress were assessed prior to conception using the PTSD Checklist - Civilian Version (PCL-C; Blanchard et al., 1996). Participants provided ratings of how often they have been bothered by various symptoms of PTSD in the past month, on a 5-point Likert scale with a range of 1 (not at all) to 5 (extremely). Final sum scores could range from 17 to 85, with higher scores indicating more PTSD symptoms (see Table 2). A cut-off score of 30 was used to indicate clinically elevated symptoms and a probable diagnosis of PTSD. This threshold and scoring approach were selected because prior research demonstrates that utilizing a cut-off score of 30 maximizes the sensitivity and specificity of accurately detecting a PTSD diagnosis amongst civilian women (McDonald & Calhoun, 2010; Walker et al., 2002). The PCL-C is a reliable and well-validated measure of PTSD symptoms (Ruggiero et al., 2003), and has been previously utilized in studies of women across pregnancy and the postpartum period (e.g., Huth-Bocks et al., 2013; Thomas, Carter, et al., 2021; Thomas, Cleveland, et al., 2021). Within the current study sample, the PCL-C demonstrated excellent internal consistency ( $\alpha = .92$ ) and 18 participants (21%) reported scores above the clinical threshold. Notably, the rate of probable PTSD diagnosis was higher in this sample than in the general population (Breslau, 2009), likely reflecting the converging risk factors for trauma and PTSD symptoms experienced by many of the women in the study (e.g., poverty, low education, acculturative stress) (Breslau et al., 1995; Pole et al., 2008; Trickey et al., 2012). Indeed, women who experienced elevated symptoms of PTSD in the study sample were more likely to be born outside of the United States (t = -2.62, p = .012), to have completed fewer years of education (r = -.367, p < .001), and to have a lower per capita income adjusted for cost of living (r = -.31, p = .004). PCL-C scores were not associated with race/ethnicity or cohabitation status.

Maternal depressive symptoms. Maternal depressive symptoms were assessed using the 9-item short form of the Center for Epidemiological Studies Depression Inventory (CES-D-9; Santor & Coyne, 1997). The CES-D is a reliable and widely used self-report measure of depressive symptoms. Participants rated the frequency at which each depressive symptom occurred during the past week on a four-point Likert scale, ranging from 0 (rarely or none of the time [less than 1 day]) to 3 (most of the time [5–7 days]). Final sum scores could range from 0 to 27 (see Table 2). The CES-D scores utilized in the current analyses were assessed twice during pregnancy, once during the postpartum period,

**Table 2.** Descriptive statistics for maternal and child measures across all timepoints

Maternal and child measures	M ± SD; Range; Skewness		
Maternal preconception PTSD symptoms (PCL-C)			
Preconception PCL-C	25.1 ± 10.2; 17–63; 1.98		
Maternal depressive symptoms (CESD-D)			
Prenatal CES-D (20 weeks' GA)	8.9 ± 3.5; 2–20; 1.28		
Prenatal CES-D (33 weeks' GA)	10.0 ± 3.4; 4–18; 1.36		
Postpartum CES-D (13 weeks postpartum)	9.4 ± 3.4; 4-21; 1.61		
Concurrent CES-D (at the time of child assessment)	5.0 ± 4.7; 0–24; 1.56		
Perceived stress symptoms (PSS)			
Prenatal PSS (20 weeks' GA)	13.8 ± 6.2; 0-29; 0.00		
Prenatal PSS (33 weeks' GA)	13.9 ± 6.0; 1–33; 0.77		
Postpartum PSS (13 weeks postpartum)	13.0 ± 6.2; 3–35; 0.86		
Children's Behavior Questionnaire (CBQ)			
Negative affectivity	4.2 ± 0.8; 2.2-6.5; 0.21		

and once at the time of child assessment. CES-D scores across both prenatal timepoints were averaged to create a composite score of prenatal depressive symptoms. The CES-D-9 demonstrated fair to good internal consistency across study time points ( $\alpha$ s ranging from .76 to .85). The CES-D-9 has also been shown in prior work to be a well-validated measure of depression in adults (Corcoran & Fisher, 1987; Santor & Coyne, 1997), including pregnant and postpartum women (Mosack & Shore, 2006).

Maternal perceived stress. Maternal perceived stress symptoms were evaluated using the 10-item version of Cohen's Perceived Stress Scale (PSS; Cohen et al., 1983). The PSS is a self-report measure of generalized or nonspecific stress, which evaluates participants' feelings about how they were able to handle day-to-day problems and hassles, how often they felt nervous and stressed, and how often they felt things were going well during the past week. Responses were made on a 5-point Likert scale, ranging from 0 (never) to 4 (almost always), with a final sum score which could range from 0 to 40 and higher scores indicating greater impairment (see Table 2). The PSS scores utilized in the current analyses were assessed twice during pregnancy, once during the postpartum period, and once at the time of child assessment. PSS scores across both prenatal timepoints were averaged to create a composite score of prenatal levels of perceived stress. The PSS demonstrated good internal consistency across study time points (αs ranging from .84 to .87). The PSS has been widely used to evaluate perceived stress symptoms in women during prenatal and postpartum periods (Nast et al., 2013).

### Child negative affectivity

Child negative affectivity was assessed using the very short form of the Children's Behavior Questionnaire (CBQ-VSF), a parent report measure of child temperament (Putnam & Rothbart, 2006; Rothbart et al., 2001). The very short form of the CBQ is a 36-item questionnaire, yielding three factor scores, of which Negative Affectivity was used in the current analyses. The 12-item

Negative Affectivity scale reflects mood instability and tendency to experience dysregulated negative emotions. When completing the scale, mothers were asked the degree to which each statement (e.g., "gets quite frustrated when prevented from doing something s/he wants to do") described their child's behavior over the past six months on a 7-point Likert scale, ranging from 1 (extremely untrue of your child) to 7 (extremely true of your child) (see Table 2). An average rating was calculated for the negative affectivity scale. The CBQ-VSF has been shown to demonstrate adequate internal consistency and substantial interrater reliability (Putnam & Rothbart, 2006; Rothbart et al., 2001), and has been widely used in studies of child temperament and negative affectivity. Further, the CBQ-VSF takes advantage of mother's ability to observe her child across a wide range of contexts and relies on maternal observation of concrete behaviors rather than abstract judgments of temperament to reduce the potential influence of maternal bias in reporting. Within the study sample, the CBQ-VSF demonstrated good internal consistency on the Negative Affectivity subscale ( $\alpha = .84$ ).

# Statistical analysis

# Identification of covariates

Spearman's rank correlations (utilized to address skewness in tested covariates), t-tests, and ANOVAs were used to identify maternal (SES composite score, age, cohabitation status, race/ethnicity, foreign-born, concurrent depressive symptoms, early life adversity, parity) and child factors (gestational age at birth, sex, age at early childhood assessment) that might influence child negative affectivity. Study site and time from preconception assessment to conception also were considered as potential covariates. Variables associated with child negative affectivity at p < .10 level of significance were included as a covariate in all subsequent regression analyses. Only maternal age and the SES composite score met criterion for inclusion as a covariate (see Table 3).

### Bivariate correlations

Preliminary analyses were conducted using bivariate Pearson's correlation coefficients, assessing the relations between preconception PTSD symptoms, prenatal and postpartum depressive symptoms and perceived stress, and child negative affectivity. Pearson's correlations were utilized because tested variables were normally distributed (i.e., Skewness > -2 and < 2; see Table 2; George & Mallery, 2010). Spearman's rank correlations are also available in Supplement Table S2.

# Data imputation

To address data missingness, missing preconception PTSD data was imputed. Eighty-seven women provided PTSD symptom ratings prior to conception (as well as ratings of child negative affectivity) and 31 participants completed a prenatal mood or stress measure and completed the child negative affectivity measure but were missing PTSD ratings prior to conception (26% rate of missingness). Multiple imputations of missing data were generated using MPlus (Asparouhov & Muthén, 2010; Muthén & Muthén, 1998–2017), generating 20 imputed data sets. When running subsequent analyses, parameter estimates were computed and averaged over the 20 sets. Sensitivity tests were performed to assess the influence of imputation on study findings by repeating all regression and mediational models analyses including only participants with complete data.

**Table 3.** Associations between participant characteristics and child negative affectivity

Participant characteristics	Test statistic	<i>p</i> -value
Maternal socioeconomic status composite	−.17 <sup>a</sup> ^	.06
Maternal age	21 <sup>b</sup> *	.02
Cohabitation status	1.73 <sup>b</sup>	.32
Maternal race/ethnicity	1.46 <sup>c</sup>	.24
Foreign-born	3.10 <sup>b</sup>	.71
Maternal early life adversity	.001	.99
Parity	.46 <sup>c</sup>	.63
Time from preconception assessment to conception	16ª	.11
Gestational age at birth	.11ª	.25
Child age	.14ª	.17
Child sex	.01 <sup>b</sup>	.71
Concurrent maternal depressive symptoms	.14ª	.13
Study site	.60°	.55

Note. Reported tests statistics are aSpearman's rho,  $^{\rm b}t$ , or  $^{\rm c}F$ . Maternal childhood adversity was assessed via the Risk Families Questionnaire. Concurrent maternal depressive symptoms were assessed via the Center for Epidemiological Studies Depression Inventory (CES-D).  $^{\rm c}p < .1$ .  $^{\rm c}p < .0$ 5.  $^{\rm c}p < .0$ 1.

### Testing of study aims

Preconception maternal PTSD symptoms and child negative affectivity

Linear regression was run in Mplus to test the primary hypothesis that preconception maternal symptoms of posttraumatic stress predict child negative affectivity after inclusion of identified covariates. Concurrent maternal depressive symptoms were then added as a covariate to evaluate whether any effect of preconception maternal PTSD symptoms on child negative affectivity remains after accounting for maternal mental health at the time of child assessment.

Assessment of prenatal and postpartum depressive symptoms and perceived stress as mediating pathways

Finally, mediational pathways were added in exploratory analyses to examine whether prenatal and/or postpartum maternal depression symptoms may mediate the relation between maternal preconception PTSD symptoms and child negative affectivity. Prenatal and postpartum levels of perceived stress were also tested as mediators in secondary, exploratory analyses.

### **Results**

# **Bivariate correlations**

Preliminary bivariate correlations between maternal preconception PTSD, maternal prenatal and postpartum depressive symptoms, and child negative affectivity are presented in Table 4. Maternal preconception PTSD symptoms were positively associated with child negative affectivity on the CBQ-VSF (r=.26, p=.014). PTSD symptoms also were associated with higher prenatal and postpartum depressive symptoms (r=.46, p<.001; and r=.26, p=.020, respectively) and prenatal and postpartum perceived stress (r=.59, p<.001; and r=.48, p<.001, respectively). Prenatal and postpartum depressive symptoms in turn were

**Table 4.** Pearson's correlations between preconception PTSD symptoms, prenatal and postpartum depressive symptoms and perceived stress, and child negative affectivity

	1	2	3	4	5	6
1. Preconception PTSD	-	.46***	.26*	.59***	.48***	.26*
2. Prenatal CES-D	-	-	.40***	.55***	.31**	.20*
3. Postpartum CES-D	-	-	-	.39***	.65***	.20*
4. Prenatal PSS	-	-	-	-	.44***	.12
5. Postpartum PSS	-	-	-	-	-	.21*
6. CBQ negative affectivity	-	-	-	-	-	-

Note. \*p < .05. \*\*p < .01. \*\*\*p < .001.

**Table 5.** Regression models of preconception maternal PTSD symptoms and child negative affectivity

Model Predictors	β	SE	р			
Model 1: Regression model with identified covariates						
Preconception PTSD	.201	.087	.022*			
SES	083	.095	.383			
Maternal age	157	.096	.100			
Model 2: Regression model with identified covariates and concurrent maternal depressive symptoms						
Preconception PTSD	.187	.094	.045*			
SES	081	.095	.395			
Maternal age	153	.097	.113			
Concurrent maternal depressive symptoms	.050	.087	.566			

Note. Missing preconception PTSD data was imputed (n = 118). SES = socioeconomic status (composite of maternal education and per capita household income adjusted for cost of living). \*p < .05.

associated with higher child negative affectivity scores (r = .20, p = .043; and r = .20, p = .048, respectively). Postpartum but not prenatal levels of perceived stress positively correlated with elevated child negative affectivity (r = .21, p = .037; and r = .12, p = .210, respectively).

# Testing of study aims

Preconception maternal PTSD symptoms and child negative affectivity

Regression analyses testing the primary hypothesis revealed that greater maternal symptoms of PTSD measured prior to conception predicted heightened child negative affectivity in early childhood ( $\beta$  = 0.20, SE = 0.09, p = .022) even after inclusion of sociodemographic covariates ( $R^2$  = .093; see Table 5, Model 1). Further, this association remained after covarying concurrent symptoms of maternal depression ( $\beta$  = 0.19, SE = 0.10, p = .045;  $R^2$  = .096;  $R^2$  change = .003 see Table 5, Model 2). Sensitivity analyses were performed including only participants with complete data, and as shown in Supplemental Table S3, the effect size and significance of this finding remained.

Assessment of prenatal and postpartum depressive symptoms and perceived stress as mediating pathways

Exploratory, post hoc analyses evaluated maternal prenatal and postpartum depressive symptoms as possible mediational pathways. This mediational model is summarized in Supplementary

Figure S1. The total effect of the model was significant ( $\beta = 0.21$ , SE = 0.09, p = .022), however prenatal and postpartum depressive symptoms were not significant mediators (total indirect effect:  $\beta = 0.04$ , SE = 0.04, p = .289; specific indirect effect of prenatal depressive symptoms:  $\beta = 0.02$ , SE = 0.03, p = .511; specific indirect effect of postpartum depressive symptoms:  $\beta = 0.02$ , SE = 0.03, p = .506). Next, prenatal and postpartum levels of perceived stress were considered as mediational pathways. This mediational model is summarized in Supplementary Figure S2. The total effect of the model was significant ( $\beta = 0.20$ , SE = 0.09, p = .022) although neither prenatal nor postpartum levels of perceived mediated the relation between preconception PTSD symptoms and child negative affectivity (total indirect effect:  $\beta = 0.04$ , SE = 0.06, p = .448; specific indirect effect of prenatal levels of perceived stress:  $\beta = -0.02$ , SE = 0.04, p = .653; specific indirect effect of postpartum levels of perceived stress:  $\beta = 0.06$ , SE = 0.05, p = .250).

### **Discussion**

The current study advances our understanding of the intergenerational impact of PTSD by prospectively evaluating maternal posttraumatic stress symptoms prior to conception and negative affectivity in the next generation. Using this prospective, longitudinal approach within a socioeconomically diverse sample of women and children, we found that more severe maternal symptoms of PTSD prior to conception predicted higher negative affectivity in their children during early childhood. Notably, the relation between maternal preconception PTSD symptoms and child negative affectivity persisted when covarying SES and concurrent maternal depressive symptoms, suggesting that this association is likely not fully accounted for by postnatal shared environmental circumstances or current maternal mental health. Further, maternal early life adversity was not associated with child negative affectivity and thus did not meet covariate criteria, suggesting that maternal mental health symptoms prior to conception may be a more robust predictor of child development than history of adverse experiences. Collectively, these results build upon our understanding of the enduring effects of PTSD on development in the next generation, suggesting that elevated PTSD symptoms prior to conception are an indicator of risk for heightened negative affectivity in the offspring. Findings also underscore the crucial importance of considering the mental health of women prior to pregnancy in the intergenerational sequelae of trauma.

The current study directly builds upon the retrospective and epidemiological literature, linking maternal recall of preconception traumatic events to poor emotional outcomes in children (Bosquet Enlow et al., 2018; Briggs et al., 2014; McDonnel & Valentino, 2016) and adds to a very small number of prospective studies exploring the impact of preconception stress on emotional outcomes for the next generation (Hipwell et al., 2019; Spry et al., 2020). Current findings extend the work of Hipwell et al. (2019), who reported that in a prospective cohort, children of women exposed to traumatic events in their own childhoods were more likely to exhibit emotion dysregulation in infancy. With our direct evaluation of PTSD symptoms (rather than exposure to traumatic events), we provide novel evidence of an intergenerational association between posttraumatic stress symptoms and child negative affectivity outcomes. This finding provides evidence that PTSD symptoms are likely part of a pathway by which maternal trauma history may have a dysregulating impact on child negative affectivity; and that PTSD symptoms measured prior to conception may be a useful indicator of intergenerational risk.

Potential mediating pathways underlying the intergenerational impact of trauma

Preconception PTSD symptoms were positively associated with symptoms of depression during pregnancy. These findings align with retrospective studies linking maternal recall of trauma history to her mental health during pregnancy (Alvarez-Segura et al., 2014; Atzl et al., 2019; River et al., 2019). Further, prenatal maternal depressive symptoms were correlated with child negative affectivity, consistent with a vast fetal programming literature prospectively documenting the influence of prenatal depression on the processes of fetal development and subsequent child socioemotional outcomes (Blair et al., 2011; Davis et al., 2007; Erickson et al., 2017; Glynn et al., 2018; Letourneau et al., 2019; Werner et al., 2007). We did not find evidence that maternal depressive symptoms or perceived stress during pregnancy or the postpartum period mediated the relation between preconception PTSD and child emotional outcomes, nor did the direct effect of prenatal depressive symptoms on child negative affectivity reach statistical significance in this mediational model. It is plausible that the current study was underpowered to detect the role of perinatal depression symptoms or perceived stress in mediating this relation. Given the relatively small sample size, mediational analyses should be interpreted with caution. It is also possible that other candidate pathways may be involved in this intergenerational association. For example, other domains of maternal mental health may be more directly implicated in the intergenerational impact of PTSD, including persistent perinatal PTSD symptoms. Although depression and PTSD are often comorbid (Shalev et al., 1998), the unique features of PTSD symptomatology such as heightened vigilance may have a more potent impact on fetal development. Multiple perinatal biological mechanisms, including stress physiology (e.g., perinatal maternal and placental HPA axis activity), epigenetic mechanisms (beginning even prior to conception), immune functioning, and gut microbiota, have also been implicated in models of intergenerational stress (Bouvette-Turcot et al., 2020; Bowers & Yehuda, 2016; Buss et al., 2017; Davis & Narayan, 2020; Keenan et al., 2018; Scorza et al., 2019; Swales et al., 2018; Yehuda & Meaney, 2018). Additionally, maternal parenting (such as maternal warmth and sensitivity) and the security of attachment relationships have been frequently implicated in the intergenerational sequelae PTSD, as trauma history has been shown to be a risk factor for negative parent-child interactions (Cohen et al., 2008; Savage et al., 2019). It is important that future studies continue to explore possible mediating pathways and broaden the scope of candidate mechanisms considered.

# Strengths, limitations, and future directions

The current study has several key strengths. Due to the challenges of recruiting and assessing women prior to pregnancy, few prospective studies of preconception mental health exist. Further, the multifaceted diversity of the study sample is a notable strength. The women in the study sample came from racial, ethnic, geographical, and socioeconomic backgrounds which are underrepresented in maternal and child health research (e.g., over a third of women identified as Latina or Hispanic and approximately a third of women identified as African American or Black) (Conradt et al., 2020). Another key advantage of the current study is the use of multiple assessment timepoints. Although PTSD symptoms were only measured once in the preconception period, maternal depressive symptoms and levels of perceived stress were evaluated twice during pregnancy, once postpartum, and once during the time of

childhood assessment. This approach allowed for continued evaluation of maternal mental health across the peripartum period and into early childhood.

Despite these strengths, there also are also several limitations and key identified areas for future research. First, the sample includes only mothers who had at least one prior pregnancy at the time of recruitment. There was also a high rate of attrition amongst eligible participants (48%), reflecting the usual challenges of recruiting participants prior to conception and retaining them until 3-5 years postpartum. Participants included in the current analyses were more likely to be older, Latina, and born outside of the United States, and had a longer interpregnancy interval than those who did not participate in the present study, and notably, PTSD symptoms did not differ between the two groups. These demographic predictors of attrition should be considered in interpreting study findings. Additionally, maternal PTSD symptoms were only assessed at one time point and not during pregnancy or at any postnatal time points. Future studies should evaluate maternal PTSD symptoms over time. We also evaluated PTSD symptoms dimensionally within an at-risk community sample, with 21% of participants surpassing the clinical threshold for likely PTSD diagnosis. Future work would also benefit from exploring these associations within samples of participants recruited from clinical settings and including full diagnostic assessment of PTSD symptoms. Another limitation is that child negative affectivity was assessed via maternal report of child behavior. The likelihood of bias is reduced by the CBQ measure design which asks about behavior in concrete situations. Further, we covaried maternal depressive symptoms at the time of reporting to partially account for the influence of concurrent mental health on reporting biases; however, future studies should also seek to obtain ratings from other caregivers and through direct observations of child temperament. Finally, future studies should consider additional processes of risk and resilience that may impact the intergenerational transmission of maternal stress, such as cultural factors (considering the role of enculturation, acculturation, acculturative stress and discrimination, and community and identity based protective factors) as well as benevolent childhood experiences and other maternal characteristics which may buffer against the impact of preconception PTSD symptoms.

# Conclusion

The current findings have important implications for understanding the intergenerational impact of PTSD and stress. This study supports the continued expansion of the DOHaD model to consider how experiences prior to pregnancy may have lasting intergenerational consequences. Specifically, these findings carry important clinical significance, as heightened negative affectivity in early childhood is a key predictor of later psychopathology (Nigg, 2006), suggesting that elevated preconception PTSD symptoms may indicate risk for later mental health problems in the child. Findings underscore the critical importance of early identification of trauma and PTSD symptoms, and subsequent implementation of effective therapeutic interventions. Such screenings and therapeutic supports should be targeted not only during pregnancy and for new mothers, but also before women enter motherhood. This is important as targeted preconception interventions may confer a two-generational benefit, supporting the mental health of trauma survivors as well as disrupting intergenerational risk pathways to foster positive emotional development in children.

**Supplementary material.** To view supplementary material for this article, please visit https://doi.org/10.1017/S0954579421001760

Acknowledgments. None.

Funding statement. This research was funded by the Eunice Kennedy Shriver National Institute of Child Health and Human Development (R01HD072021; Dunkel Schetter, PI) and earlier work by the Child Community Health Network (CCHN), supported through cooperative agreements with the Eunice Kennedy Shriver National Institute of Child Health and Human Development (NICHD; U HD44207, U HD44219, U HD44226, U HD44245, U HD44253, U HD54791, U HD54019, U HD44226-05S1, U HD44245-06S1, R03 HD59584) and the National Institute for Nursing Research (U NR008929). Author N.E.M. received support from the National Institute of Mental Health (T32 MH015750).

Conflicts of interest. None.

### References

Alvarez-Segura, M., Garcia-Esteve, L., Torres, A., Plaza, A., Imaz, M. L., Hermida-Barros, L., ... Burtchen, N. (2014). Are women with a history of abuse more vulnerable to perinatal depressive symptoms? A systematic review. *Archives of Women's Mental Health*, 17(5), 343–357. https://doi.org/10.1007/s00737-014-0440-9

Asparouhov, T., & Muthén, B. (2010). Multiple imputation with Mplus. *MPlus Web Notes*. https://www.statmodel.com/download/Imputations7.pdf Atzl, V. M., Narayan, A. J., Rivera, L. M., & Lieberman, A. F. (2019). Adverse childhood experiences and prenatal mental health: Type of ACEs and age of maltreatment onset. *Journal of Family Psychology*, 33(3), 304–314. https://doi.org/10.1080/10926771.2018.1524806

Baird, J., Hill, C. M., Kendrick, T., Inskip, H. M., & SWS Study Group. (2009). Infant sleep disturbance is associated with preconceptional psychological distress: Findings from the Southampton Women's Survey. *Sleep*, 32(4), 566–568. https://doi.org/10.5665/sleep/32.4.566

Baldwin, J.R., Reuben, A., Newbury, J.B., & Danese, A. (2019). Agreement between prospective and retrospective measures of childhood maltreatment: A systematic review and meta-analysis. *JAMA Psychiatry*, 76, 584–593. https://doi.org/10.1001/jamapsychiatry.2019.0097

Bale, T. L. (2014). Lifetime stress experience: Transgenerational epigenetics and germ cell programming. *Dialogues in Clinical Neuroscience*, 16(3), 297–305. https://doi.org/10.31887/dcns.2014.16.3/tbale

Bernstein, D. P., Fink, L., Handelsman, L., Foote, J., Lovejoy, M., Wenzel, K., ... Ruggiero, J. (1994). Initial reliability and validity of a new retrospective measure of child abuse and neglect. *The American Journal of Psychiatry*, 151(8), 1132–1136. https://doi.org/10.1176/ajp.151.8.1132

Bierer, L. M., Bader, H. N., Daskalakis, N. P., Lehrner, A. L., Makotkine, I., Seckl, J. R., & Yehuda, R. (2014). Elevation of 11β-hydroxysteroid dehydrogenase type 2 activity in Holocaust survivor offspring: Evidence for an intergenerational effect of maternal trauma exposure. *Psychoneuroendocrinology*, 48, 1–10. https://doi.org/10.1016/j.psyneuen.2014.06.001

Bijl, R. V., Cuijpers, P., & Smit, F. (2002). Psychiatric disorders in adult children of parents with a history of psychopathology. *Social Psychiatry and Psychiatric Epidemiology*, 37(1), 7–12. https://doi.org/10.1007/s127-002-8208-8

Blair, M. M., Glynn, L. M., Sandman, C. A., & Davis, E. P. (2011). Prenatal maternal anxiety and early childhood temperament. *Stress*, *14*(6), 644–651. https://doi.org/10.3109/10253890.2011.594121

Blanchard, E. B., Jones-Alexander, J., Buckley, T. C., & Forneris, C. A. (1996). Psychometric properties of the PTSD Checklist (PCL). *Behaviour Research and Therapy*, 34(8), 669–673. https://doi.org/10.1016/0005-7967(96)00033-2

Bock, J., Poeschel, J., Schindler, J., Börner, F., Shachar-Dadon, A., Ferdman, N., . . . Poeggel, G. (2016). Transgenerational sex-specific impact of preconception stress on the development of dendritic spines and dendritic length in the medial prefrontal cortex. *Brain Structure and Function*, 221(2), 855–863. https://doi.org/10.1007/s00429-014-0940-4

Bosquet Enlow, M., Devick, K. L., Brunst, K. J., Lipton, L. R., Coull, B. A., & Wright, R. J. (2017). Maternal lifetime trauma exposure, prenatal cortisol,

- and infant negative affectivity. *Infancy*, 22(4), 492–513. https://doi.org/10. 1111/infa.12176
- Bosquet Enlow, M., Englund, M. M., & Egeland, B. (2018). Maternal childhood maltreatment history and child mental health: Mechanisms in intergenerational effects. *Journal of Clinical Child & Adolescent Psychology*, 47(sup1), S47–S62. https://doi.org/10.1080/15374416. 2016.1144189
- Bouvette-Turcot, A. A., Fleming, A. S., Unternaehrer, E., Gonzalez, A., Atkinson, L., Gaudreau, H., ... Meaney, M. J. (2020). Maternal symptoms of depression and sensitivity mediate the relation between maternal history of early adversity and her child temperament: The inheritance of circumstance. *Development and Psychopathology*, 32(2), 605–613. https://doi.org/10.1017/s0954579419000488
- Bowers, M. E., & Yehuda, R. (2016). Intergenerational transmission of stress in humans. *Neuropsychopharmacology*, 41(1), 232–244. https://doi.org/10. 1038/npp.2015.247
- Breslau, N. (2009). The epidemiology of trauma, PTSD, and other posttrauma disorders. *Trauma*, *Violence*, & *Abuse*, *10*(3), 198−210. https://doi.org/10. 1177/1524838009334448
- Breslau, N., Davis, G. C., & Andreski, P. (1995). Risk factors for PTSD-related traumatic events: A prospective analysis. *The American Journal of Psychiatry*, 152(4), 529–535. https://doi.org/10.1176/ajp.152.4.529
- Briggs, R. D., Silver, E. J., Krug, L. M., Mason, Z. S., Schrag, R. D., Chinitz, S., & Racine, A. D. (2014). Healthy steps as a moderator: The impact of maternal trauma on child social-emotional development. *Clinical Practice in Pediatric Psychology*, 2(2), 166–175. https://doi.org/10.1037/cpp0000060
- Buss, C., Entringer, S., Moog, N. K., Toepfer, P., Fair, D. A., Simhan, H. N., ... Wadhwa, P. D. (2017). Intergenerational transmission of maternal childhood maltreatment exposure: Implications for fetal brain development. *Journal of the American Academy of Child & Adolescent Psychiatry*, 56(5), 373–382. https://doi.org/10.1016/j.jaac.2017.03.001
- Clark, L. A., Watson, D., & Mineka, S. (1994). Temperament, personality, and the mood and anxiety disorders. *Journal of Abnormal Psychology*, 103(1), 103–116. https://doi.org/10.1037/0021-843x.103.1.103
- Class, Q. A., Abel, K. M., Khashan, A. S., Rickert, M. E., Dalman, C., Larsson, H., ... D'Onofrio, B. M. (2014). Offspring psychopathology following preconception, prenatal and postnatal maternal bereavement stress. *Psychological Medicine*, 44(1), 71–84. https://doi.org/10.1017/s0033291713000780
- Class, Q. A., Mortensen, P. B., Henriksen, T. B., Dalman, C., D'Onofrio, B. M., & Khashan, A. S. (2015). Preconception maternal bereavement and infant and childhood mortality: A Danish population-based study. *Psychosomatic Medicine*, 77(8), 863–869. https://doi.org/10.1097/psy.00000000000000229
- Cohen, L. R., Hien, D. A., & Batchelder, S. (2008). The impact of cumulative maternal trauma and diagnosis on parenting behavior. *Child Maltreatment*, 13(1), 27–38. https://doi.org/10.1177/1077559507310045
- Cohen, S., Doyle, W. J., & Baum, A. (2006). Socioeconomic status is associated with stress hormones. *Psychosomatic Medicine*, 68(3), 414–420. https://doi.org/10.1097/01.psy.0000221236.37158.b9
- Cohen, S., Kamarck, T., & Mermelstein, R. (1983). A global measure of perceived stress. *Journal of Health and Social Behavior*, 24(4), 385–396. https://doi.org/10.2307/2136404
- Conradt, E., Carter, S. E., & Crowell, S. E. (2020). Biological embedding of chronic stress across two generations within marginalized communities. *Child Development Perspectives*, 14(4), 208–214. https://doi.org/10.1111/ cdep.12382
- Corcoran, K., & Fisher, J. (1987). Measures for clinical practice: A source book. New York: Free Press.
- Davis, E. P., Glynn, L. M., Schetter, C. D., Hobel, C., Chicz-Demet, A., & Sandman, C. A. (2007). Prenatal exposure to maternal depression and cortisol influences infant temperament. *Journal of the American Academy of Child & Adolescent Psychiatry*, 46(6), 737–746. https://doi.org/10.1097/chi.0b013e318047b775
- Davis, E. P., & Narayan, A. J. (2020). Pregnancy as a period of risk, adaptation, and resilience for mothers and infants. *Development and*

- Psychopathology, 32(5), 1625–1639. https://doi.org/10.1017/s09545794200 01121
- **Derryberry, D., & Rothbart, M. K.** (1997). Reactive and effortful processes in the organization of temperament. *Development and Psychopathology*, 9(4), 633–652. https://doi.org/10.1017/s0954579497001375
- Doyle, C., & Cicchetti, D. (2018). Future directions in prenatal stress research: Challenges and opportunities related to advancing our understanding of prenatal developmental origins of risk for psychopathology. *Development and Psychopathology*, 30(3), 721–724. https://doi.org/10.1017/s095457941800069x
- Dunkel Schetter, C., Schafer, P., Lanzi, R. G., Clark-Kauffman, E., Raju, T. N., Hillemeier, M. M., & Community Child Health Network. (2013). Shedding light on the mechanisms underlying health disparities through community participatory methods: The stress pathway. Perspectives on Psychological Science, 8(6), 613–633. https://doi.org/10.1177/17456916135 06016
- Erickson, N. L., Gartstein, M. A., & Dotson, J. A. W. (2017). Review of prenatal maternal mental health and the development of infant temperament. *Journal of Obstetric, Gynecologic & Neonatal Nursing*, 46(4), 588–600. https://doi.org/10.1016/j.jogn.2017.03.008
- Felitti, V. J., Anda, R. F., Nordenberg, D., Williamson, D. F., Spitz, A. M., Edwards, V., . . . . Marks, J. S. (1998). Household dysfunction to many of the leading causes of death in adults the Adverse Childhood Experiences (ACE) Study. *American Journal of Preventive Medicine*, 14(4), 245–258. https://doi.org/10.1016/s0749-3797(98)00017-8
- Flory, J. D., Bierer, L. M., & Yehuda, R. (2011). Maternal exposure to the holocaust and health complaints in offspring. *Disease Markers*, 30(2–3), 133–139. https://doi.org/10.1155/2011/250470
- Gangi, S., Talamo, A., & Ferracuti, S. (2009). The long-term effects of extreme war-related trauma on the second generation of Holocaust survivors. Violence and Victims, 24, 687–701. https://doi.org/10.1891/0886-6708.24.5.687
- George, D., & Mallery, P. (2010). SPSS for Windows step by step. A simple study guide and reference, 17.0 update (10a ed.). Pearson Education, Inc.
- Glynn, L. M., Howland, M. A., Sandman, C. A., Davis, E. P., Phelan, M., Baram, T. Z., & Stern, H. S. (2018). Prenatal maternal mood patterns predict child temperament and adolescent mental health. *Journal of Affective Disorders*, 228, 83–90. https://doi.org/10.1016/j.jad.2017.11.065
- Goodman, S. H., & Gotlib, I. H. (1999). Risk for psychopathology in the children of depressed mothers: A developmental model for understanding mechanisms of transmission. *Psychological Review*, *106*(3), 458–490. https://doi.org/10.1037/0033-295x.106.3.458
- Hardt, J., & Rutter, M. (2004). Validity of adult retrospective reports of adverse childhood experiences: Review of the evidence. *Journal of Child Psychology and Psychiatry*, 45(2), 260–273. https://doi.org/10.1111/j.1469-7610.2004.00218.x
- Harker, A., Raza, S., Williamson, K., Kolb, B., & Gibb, R. (2015). Preconception paternal stress in rats alters dendritic morphology and connectivity in the brain of developing male and female offspring. *Neuroscience*, 303, 200–210. https://doi.org/10.1016/j.neuroscience.2015.06.058
- Harville, E. W., Boynton-Jarrett, R., Power, C., & Hyppönen, E. (2010). Childhood hardship, maternal smoking, and birth outcomes: A prospective cohort study. *Archives of Pediatrics & Adolescent Medicine*, 164(6), 533–539. https://doi.org/10.1001/archpediatrics.2010.61
- Hipwell, A. E., Tung, I., Northrup, J., & Keenan, K. (2019). Transgenerational associations between maternal childhood stress exposure and profiles of infant emotional reactivity. *Development and Psychopathology*, 31(3), 887–898. https://doi.org/10.1017/s0954579419000324
- Huang, Y., Shi, X., Xu, H., Yang, H., Chen, T., Chen, S., & Chen, X. (2010). Chronic unpredictable stress before pregnancy reduce the expression of brain-derived neurotrophic factor and *N*-methyl-D-aspartate receptor in hippocampus of offspring rats associated with impairment of memory. *Neurochemical Research*, 35(7), 1038–1049. https://doi.org/10.1007/s11064-010-0152-0
- Huth-Bocks, A. C., Krause, K., Ahlfs-Dunn, S., Gallagher, E., & Scott, S. (2013). Relational trauma and posttraumatic stress symptoms among

pregnant women. Psychodynamic Psychiatry, 41(2), 277–301. https://doi.org/10.1521/pdps.2013.41.2.277

- Keenan, K., Hipwell, A. E., Class, Q. A., & Mbayiwa, K. (2018). Extending the developmental origins of disease model: Impact of preconception stress exposure on offspring neurodevelopment. *Developmental Psychobiology*, 60(7), 753–764. https://doi.org/10.1002/dev.21773
- Khan, M., & Renk, K. (2019). Mothers' adverse childhood experiences, depressive symptoms, parenting, and attachment as predictors of young children's problems. *Journal of Child Custody*, 16(3), 268–290. https://doi.org/10.1080/15379418.2019.1575318
- Klengel, T., Dias, B. G., & Ressler, K. J. (2016). Models of intergenerational and transgenerational transmission of risk for psychopathology in mice. *Neuropsychopharmacology*, 41(1), 219–231. https://doi.org/10.1038/npp. 2015.249
- Lambert, J. E., Holzer, J., & Hasbun, A. (2014). Association between parents' PTSD severity and children's psychological distress: A meta-analysis. *Journal of Traumatic Stress*, 27(1), 9–17. https://doi.org/10.1002/jts.21891
- Lang, A. J., Gartstein, M. A., Rodgers, C. S., & Lebeck, M. M. (2010). The impact of maternal childhood abuse on parenting and infant temperament. *Journal of Child and Adolescent Psychiatric Nursing*, 23(2), 100–110. https://doi.org/10.1111/j.1744-6171.2010.0022
- Letourneau, N., Dewey, D., Kaplan, B. J., Ntanda, H., Novick, J., Thomas, J. C., ... APrON Study Team. (2019). Intergenerational transmission of adverse childhood experiences via maternal depression and anxiety and moderation by child sex. *Journal of Developmental Origins of Health and Disease*, 10(1), 88–99. https://doi.org/10.1017/s2040174418000648
- Li, H., Zhang, L., Fang, Z., Lin, L., Wu, C., & Huang, Q. (2010). Behavioral and neurobiological studies on the male progeny of maternal rats exposed to chronic unpredictable stress before pregnancy. *Neuroscience Letters*, 469(2), 278–282. https://doi.org/10.1016/j.neulet.2009.12.017
- Lonigan, C. J., Vasey, M. W., Phillips, B. M., & Hazen, R. A. (2004). Temperament, anxiety, and the processing of threat-relevant stimuli. *Journal of Clinical Child and Adolescent Psychology*, 33(1), 8–20. https://doi.org/10.1207/s15374424jccp3301\_2
- Mahrer, N. E., Guardino, C. M., Hobel, C., & Dunkel Schetter, C. (2020). Maternal stress before conception is associated with shorter gestation. *Annals of Behavioral Medicine*, 55(3), 242–252. https://doi.org/10.1093/abm/kaaa047
- McDonald, S. D., & Calhoun, P. S. (2010). The diagnostic accuracy of the PTSD checklist: A critical review. *Clinical Psychology Review*, 30(8), 976–987. https://doi.org/10.1016/j.cpr.2010.06.012
- McDonnell, C. G., & Valentino, K. (2016). Intergenerational effects of child-hood trauma: Evaluating pathways among maternal ACEs, perinatal depressive symptoms, and infant outcomes. *Child Maltreatment*, 21(4), 317–326. https://doi.org/10.1177/1077559516659556
- McLaughlin, K. A., & Lambert, H. K. (2017). Child trauma exposure and psychopathology: Mechanisms of risk and resilience. *Current Opinion in Psychology*, 14, 29–34. https://doi.org/10.1016/j.copsyc.2016.10.004
- Moffitt, T. E., Caspi, A., Taylor, A., Kokaua, J., Milne, B. J., Polanczyk, G., & Poulton, R. (2010). How common are common mental disorders? Evidence that lifetime prevalence rates are doubled by prospective versus retrospective ascertainment. *Psychological Medicine*, 40(6), 899–909. https://doi.org/10.1017/s0033291709991036
- Mosack, V., & Shore, E. R. (2006). Screening for depression among pregnant and postpartum women. *Journal of Community Health Nursing*, 23(1), 37–47. https://doi.org/10.1207/s15327655jchn2301\_4
- Muris, P., & Ollendick, T. H. (2005). The role of temperament in the etiology of child psychopathology. *Clinical Child and Family Psychology Review*, 8(4), 271–289. https://doi.org/10.1007/s10567-005-8809-y
- Muthén, L. K., & Muthén, B. O. (1998–2017). Mplus User's Guide. Eighth Edition. Muthén & Muthén.
- Narayan, A., Lieberman, A. F., Masten, A. S. (2021). Intergenerational transmission and prevention of adverse childhood experiences (ACEs). *Clinical Psychology Review*, 101997. https://doi.org/10.1016/j.cpr.2021.101997
- Nast, I., Bolten, M., Meinlschmidt, G., & Hellhammer, D. H. (2013). How to measure prenatal stress? A systematic review of psychometric instruments to assess psychosocial stress during pregnancy. *Paediatric and Perinatal Epidemiology*, 27(4), 313–322. https://doi.org/10.1111/ppe.12051

- National Comorbidity Survey (NCS). (2017). Data Table 2: 12-month prevalence DSM-IV/WMH-CIDI disorders by sex and cohort. https://www.hcp.med.harvard.edu/ncs/index.php
- Newbury, J. B., Arseneault, L., Moffitt, T. E., Caspi, A., Danese, A., Baldwin, J. R., & Fisher, H. L. (2018). Measuring childhood maltreatment to predict early-adult psychopathology: Comparison of prospective informant-reports and retrospective self-reports. *Journal of Psychiatric Research*, *96*, 57–64. https://doi.org/10.1016/j.jpsychires.2017.09.020
- Nigg, J. T. (2006). Temperament and developmental psychopathology. *Journal of Child Psychology and Psychiatry*, 47(3-4), 395–422. https://doi.org/10.1111/j.1469-7610.2006.01612.x
- O'Campo, P., Schetter, C. D., Guardino, C. M., Vance, M. R., Hobel, C. J., Ramey, S. L., . . . Community Child Health Network. (2016). Explaining racial and ethnic inequalities in postpartum allostatic load: Results from a multisite study of low to middle income woment. SSM-Population Health, 2, 850–858. https://doi.org/10.1016/j.ssmph.2016.10.014
- O'Donnell, M. L., Creamer, M., & Pattison, P. (2004). Posttraumatic stress disorder and depression following trauma: Understanding comorbidity. *American Journal of Psychiatry*, 161(8), 1390–1396. https://doi.org/10.1176/appi.ajp.161.8.1390
- Perroud, N., Rutembesa, E., Paoloni-Giacobino, A., Mutabaruka, J., Mutesa, L., Stenz, L., . . . Karege, F. (2014). The Tutsi genocide and transgenerational transmission of maternal stress: Epigenetics and biology of the HPA axis. *The World Journal of Biological Psychiatry*, 15(4), 334–345. https://doi.org/10.3109/15622975.2013.866693
- Plant, D. T., Jones, F. W., Pariante, C. M., & Pawlby, S. (2017). Association between maternal childhood trauma and offspring childhood psychopathology: Mediation analysis from the ALSPAC cohort. *The British Journal of Psychiatry*, 211(3), 144–150. https://doi.org/10.1192/bjp.bp.117.198721
- Plant, D. T., Pawlby, S., Pariante, C. M., & Jones, F. W. (2018). When one childhood meets another-maternal childhood trauma and offspring child psychopathology: A systematic review. Clinical Child Psychology and Psychiatry, 23(3), 483–500. https://doi.org/10.1177/1359104517742186
- Pole, N., Gone, J. P., & Kulkarni, M. (2008). Posttraumatic stress disorder among ethnoracial minorities in the United States. Clinical Psychology: Science and Practice, 15(1), 35–61. https://doi.org/10.1111/j.1468-2850. 2008.00109.x
- Power, C., Atherton, K., Strachan, D. P., Shepherd, P., Fuller, E., Davis, A., ... Macfarlane, G. J. (2007). Life-course influences on health in British adults: Effects of socio-economic position in childhood and adulthood. *International Journal of Epidemiology*, 36(3), 532–539. https://doi.org/10.1093/jie/dyl310
- Putnam, S. P., & Rothbart, M. K. (2006). Development of short and very short forms of the Children's Behavior Questionnaire. *Journal of Personality Assessment*, 87(1), 102–112. https://doi.org/10.1207/s15327752jpa8701\_09
- Ramey, S. L., Schafer, P., DeClerque, J. L., Lanzi, R. G., Hobel, C., Shalowitz, M., ... Raju, T. N. (2015). The preconception stress and resiliency pathways model: A multi-level framework on maternal, paternal, and child health disparities derived by community-based participatory research. *Maternal and Child Health Journal*, 19(4), 707–719. https://doi.org/10.1007/s10995-014-1581-1
- Resnick, H. S., Kilpatrick, D. G., Dansky, B. S., Saunders, B. E., & Best, C. L. (1993). Prevalence of civilian trauma and posttraumatic stress disorder in a representative national sample of women. *Journal of Consulting and Clinical Psychology*, 61(6), 984–991. https://doi.org/10.1037/0022-006x.61. 6.984
- Reuben, A., Moffitt, T. E., Caspi, A., Belsky, D. W., Harrington, H., Schroeder, F., . . . Danese, A. (2016). Lest we forget: Comparing retrospective and prospective assessments of adverse childhood experiences in the prediction of adult health. *Journal of Child Psychology and Psychiatry*, *57*(10), 1103–1112. https://doi.org/10.1111/jcpp.12621
- River, L. M., Narayan, A. J., Atzl, V. M., Rivera, L. M., & Lieberman, A. F. (2019). Past made present: The legacy of childhood maltreatment for romantic relationship quality and psychopathology during pregnancy. *Psychology of Violence*, 10(3), 324–333. https://doi.org/10.1037/vio0000273
- Rothbart, M. K., Ahadi, S. A., Hershey, K. L., & Fisher, P. (2001). Investigations of temperament at three to seven years: The Children's

- Behavior Questionnaire. *Child Development*, 72(5), 1394–1408. https://doi.org/10.1111/1467-8624.00355
- Ruggiero, K. J., Del Ben, K., Scotti, J. R., & Rabalais, A. E. (2003). Psychometric properties of the PTSD Checklist—Civilian version. *Journal of Traumatic Stress*, 16(5), 495–502. https://doi.org/10.1023/a:10257 14729117
- Santor, D. A., & Coyne, J. C. (1997). Shortening the CES-D to improve its ability to detect cases of depression. *Psychological Assessment*, 9(3), 233–243. https://doi.org/10.1037/1040-3590.9.3.233
- Savage, L. É., Tarabulsy, G. M., Pearson, J., Collin-Vézina, D., & Gagné, L. M. (2019). Maternal history of childhood maltreatment and later parenting behavior: A meta-analysis. *Development and Psychopathology*, 31(1), 9–21. https://doi.org/10.1017/s0954579418001542
- Scorza, P., Duarte, C. S., Hipwell, A. E., Posner, J., Ortin, A., Canino, G., ... Program Collaborators for Environmental influences on Child Health Outcomes. (2019). Research review: Intergenerational transmission of disadvantage: Epigenetics and parents' childhoods as the first exposure. *Journal of Child Psychology and Psychiatry*, 60(2), 119–132. https://doi. org/10.1111/jcpp.12877
- Seng, J. S., Low, L. M. K., Sperlich, M., Ronis, D. L., & Liberzon, I. (2009). Prevalence, trauma history, and risk for posttraumatic stress disorder among nulliparous women in maternity care. *Obstetrics and Gynecology*, 114(4), 839–847. https://doi.org/10.1097/aog.0b013e3181b8f8a2
- Shalev, A. Y., Freedman, S., Peri, T., Brandes, D., Sahar, T., Orr, S. P., & Pitman, R. K. (1998). Prospective study of posttraumatic stress disorder and depression following trauma. *American Journal of Psychiatry*, 155(5), 630–637. https://doi.org/10.1176/ajp.155.5.630
- Solomon, Z., Kotler, M., & Mikulincer, M. (1988). Combat-related posttraumatic stress disorder among second-generation Holocaust survivors: Preliminary findings. *The American Journal of Psychiatry*, 145(7), 865–868. https://doi.org/10.1176/ajp.145.7.865
- Spry, E., Moreno-Betancur, M., Becker, D., Romaniuk, H., Carlin, J. B., Molyneaux, E., ... Macdonald, J. A. (2020). Maternal mental health and infant emotional reactivity: A 20-year two-cohort study of preconception and perinatal exposures. *Psychological Medicine*, 50(5), 827–837. https://doi.org/10.1017/s0033291719000709
- Swales, D. A., Snyder, H. R., Hankin, B. L., Sandman, C. A., Glynn, L. M., & Davis, E. P. (2020). Maternal depressive symptoms predict general liability in child psychopathology. *Journal of Clinical Child & Adolescent Psychology*, 1–12. https://doi.org/10.1080/15374416.2020.1723598
- Swales, D. A., Stout-Oswald, S. A., Glynn, L. M., Sandman, C., Wing, D. A., & Davis, E. P. (2018). Exposure to traumatic events in childhood predicts

- cortisol production among high risk pregnant women. *Biological Psychology*, 139, 186–192. https://doi.org/10.1016/j.biopsycho.2018.10.006
- Taylor, S. E., Lerner, J. S., Sage, R. M., Lehman, B. J., & Seeman, T. E. (2004). Early environment, emotions, responses to stress, and health. *Journal of Personality*, 72(6), 1365–1394. https://doi.org/10.1111/j.1467-6494.2004. 00300.x
- Thomas, J. L., Cleveland, S., Pietrzak, R. H., Dunkel Schetter, C., & Sumner, J. A. (2021). Elucidating posttraumatic stress symptom dimensions and health correlates among postpartum women. *Journal of Affective Disorders*, 294, 314–321. https://doi.org/10.1016/j.jad.2021.07.025
- Thomas, J.L., Carter, S.E., Dunkel Schetter, C., & Sumner, J.A. (2021).
  Racial and ethnic disparities in posttraumatic psychopathology among post-partum women. *Journal of Psychiatric Research*, 137, 36–40. https://doi.org/10.1016/j.jpsychires.2021.02.030
- Trickey, D., Siddaway, A. P., Meiser-Stedman, R., Serpell, L., & Field, A. P. (2012). A meta-analysis of risk factors for post-traumatic stress disorder in children and adolescents. *Clinical Psychology Review*, 32(2), 122–138. https://doi.org/10.1016/j.cpr.2011.12.001
- Walker, E. A., Newman, E., Dobie, D. J., Ciechanowski, P., & Katon, W. (2002). Validation of the PTSD checklist in an HMO sample of women. *General Hospital Psychiatry*, 24(6), 375–380. https://doi.org/10.1016/s0163-8343(02)00203-7
- Werner, E. A., Myers, M. M., Fifer, W. P., Cheng, B., Fang, Y., Allen, R., & Monk, C. (2007). Prenatal predictors of infant temperament. *Developmental Psychobiology: The Journal of the International Society for Developmental Psychobiology*, 49(5), 474–484. https://doi.org/10.1002/dev.20232
- Yehuda, R., Bell, A., Bierer, L. M., & Schmeidler, J. (2008). Maternal, not paternal, PTSD is related to increased risk for PTSD in offspring of Holocaust survivors. *Journal of Psychiatric Research*, 42(13), 1104–1111. https://doi.org/10.1016/j.jpsychires.2008.01.002
- **Yehuda, R., Halligan, S. L., & Grossman, R.** (2001). Childhood trauma and risk for PTSD: Relationship to intergenerational effects of trauma, parental PTSD, and cortisol excretion. *Development and Psychopathology*, *13*(3), 733–753. https://doi.org/10.1017/s0954579401003170
- Yehuda, R., & Meaney, M. J. (2018). Relevance of psychological symptoms in pregnancy to intergenerational effects of preconception trauma. *Biological Psychiatry*, 83(2), 94–96. https://doi.org/10.1016/j.biopsych.2017.10.027
- Zaidan, H., Leshem, M., & Gaisler-Salomon, I. (2013). Prereproductive stress to female rats alters corticotropin releasing factor type 1 expression in ova and behavior and brain corticotropin releasing factor type 1 expression in offspring. Biological Psychiatry, 74(9), 680–687. https://doi.org/10.1016/j.biopsych.2013.04.014