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## Abstract

*Objective:* The study aimed to evaluate the impact of prenatal maternal stress on birth weight using a large cohort of predominantly Caucasian women living in an urban area. *Method:* Women were recruited between 2005 and 2010. Data collection took place between the 24th and the 28th week of gestation. The Measure of Psychological Stress (MSP-9), a validated tool to assess stress symptoms, was used to collect data on prenatal maternal stress (independent variable). Birth weight (dependent variable) was classified as low birth weight (LBW; <2,500 g), normal birth weight (NBW; 2,500-4,000 g), and macrosomia (>4,000 g). Adjusted odd ratios (aOR) were obtained after performing multivariate logistic regressions adjusted for potential cofounders. At the final stage, 5,721 women were included in analysis. *Results:* When compared with women experiencing low stress, participants with high-stress scores were at increased risk of delivering an infant with LBW before adjustment [OR = 2.06, 95% CI (1.04-4.09)], but, after adjustment, only a nonsignificant trend remained. However, women experiencing intermediate and high levels of stress were at increased risk of delivering a baby with macrosomia, even after adjustment [aOR = 1.23 ; 95% CI (1.02-1.49)] and [aOR = 1.76 ; 95% CI (1.11-2.77)] compared to those who scored low on the psychological stress scale. *Conclusion:* Women exposed to high self-reported psychological stress during second trimester (24<sup>th</sup> to 28<sup>th</sup> weeks) of pregnancy have a 1.7-fold increased risk for delivering a baby with macrosomia when compared to women exposed to low psychological stress.

*Keywords:* prenatal psychological stress, pregnancy, fetal macrosomia, low birth weight infant

### **Impact of Maternal Psychological Stress on Birth weight**

Birth weight is an important predictor of the newborn short and long-term health. Low birth weight (LBW) and macrosomia increase the risk of morbidity and mortality (Ay et al., 2009; Boulet, Alexander, Salihu, & Pass, 2003; Canadian Institute for Health Information, 2009). Newborns weighing under 2,500 grams, defined as low birth weight (LBW) by the World Health Organization (WHO) (United Nations Children's Fund and World Health Organization, 2004) have a higher risk, especially if they have very LBW (<1,500 g), to suffer from cerebral palsy, deafness, blindness, epilepsy, chronic lung disease, learning disabilities, and attention deficit disorders (Bergman et al., 1985; Dunn, Robertson, & Crichton, 1986; Ellenberg & Nelson, 1979; Gallo & Lennerstrand, 1991; Kraybill, Bose, & D'ercole, 1987; McCormick, Gortmaker, & Sobol, 1990). On the other hand, while there is no consensus regarding the definition of macrosomia (Boulet et al., 2003; Ye et al., 2015), newborns weighing more than 4,000 g are generally defined as having macrosomia since this threshold could indicate higher risk of adverse obstetric and neonatal outcomes (Dubois, Girard, & Tatone-Tokuda, 2007; Wang et al., 2016). Macrosomia increases the risk of shoulder dystocia (Benedetti & Gabbe, 1978), clavicular fracture, and brachial palsy (Boyd, Usher, & Mclean, 1983), prolongation of the second and third stage of labor, instrumented vaginal delivery, emergency caesarean section, Apgar score lower than four, neonatal hypoglycemia, and admission at the neonatal intensive care unit (Jolly, Sebire, Harris, Regan, & Robinson, 2003; Mohammadbeigi et al., 2013), to name only a few. Adverse outcomes related to both extremes of the birth weight spectrum thus represent an obstetrical challenge.

In Canada over the last fourteen years, the rate of LBW increased, from 5.6% in 2000 to 6.4% in 2015 (Statistics Canada, 2016). Neonates weighing >4,500 g and large for gestational age (LGA) represented 1.5% and 9.5% of all Canadian births in 2015 (Statistics Canada, 2016). Several factors are known to contribute to LBW and macrosomia. LBW is

more common in women with unfavorable socioeconomic level, chronic hypertension or nephropathy, tobacco, alcohol or drug consumption (de Bernabé et al., 2004), whereas risk factors for macrosomia include mothers with a prepregnancy body mass index (BMI) >30, parity >4, age >40, and preexisting diabetes or gestational diabetes mellitus (GDM) (Jolly et al., 2003).

In the past years, psychological stress has been proposed to play a role in birth weight, especially LBW (E. L. Bussières et al., 2015). Various hypotheses have been proposed, but research has yielded few stable findings. Some studies observed a link between maternal stress and LBW (Dolatian et al., 2016; Hashim & Moawed, 2000; Rondó et al., 2003), while others did not (Sable & Wilkinson, 2000; Wing et al., 2017). Hashim and Moawed found that an intermediate score on a life stressor scale doubled the risk of giving birth to a baby with LBW [OR=2.0, 95% CI (1.2-3.2)] in a cohort of 500 Saudi women (Hashim & Moawed, 2000). Similarly, a study conducted by Rondó et al. involving 865 Brazilian participants showed that, when compared to women experiencing low levels of stress, high-stress mothers were twice as likely to deliver a baby with LBW [OR=1.97, 95% CI 1.12-3.47] (Rondó et al., 2003). Liou et al. found that stress measured after 24 weeks of gestation could not predict the occurrence of LBW (Liou, Wang, & Cheng, 2016). Wadhwa et al. had previously found that perceived stress was not related with birth weight in a group of mostly married, employed, Caucasian women with annual family income > \$50,000 (Wadhwa, Sandman, Porto, Dunkel-Schetter, & Garite, 1993). Bussières et al. showed that the relation between maternal prenatal stress and LBW was modest but significant, and that this link was related to different and significant moderators such as type of stress and type of population (high or low risk) (E. L. Bussières et al., 2015).

In contrast, the link between prenatal maternal psychological stress and macrosomia is not well studied. The quality of life is impaired by decreased self-confidence, anxiety, stress

and depression. Jakubauskiene et al. found that lower score of quality of life (evaluated by mental and physical health scores) during the first trimester are associated with LGA newborns (Jakubauskiene et al., 2019). Tegethoff et al. observed that life stress, defined as perceptions of important difficulties in major areas such as finances, relationships, work, pregnancy, housing, and health, was associated with increased birth weight after controlling for duration of gestation. (Tegethoff, Greene, Olsen, Meyer, & Meinlschmidt, 2010a). A Canadian study founded that the relationship between social disparities and macrosomia varied by geographic region (Dubois et al., 2007).

A frequently cited mechanism to explain the relation between prenatal stress and birth weight involves the hypothalamic-pituitary-adrenal (HPA) axis (Beijers, Buitelaar, & de Weerth, 2014; Gilles et al., 2018). Stress can induce dysfunction of the HPA axis. Prenatal maternal psychological distress may alter molecular mechanisms that regulate fetal exposure to maternal cortisol (Togher, O'Keeffe, Khashan, Clarke, & Kenny, 2018). Overexposure to maternal cortisol has been associated with reduced birth weight (Field et al., 2006).

The placenta is an important mediator for intrauterine fetal development and growth (Nelissen, van Montfoort, Dumoulin, & Evers, 2011). Tegethoff *et al.* have observed that maternal life stress during the pregnancy was associated with an increased placenta weight (Tegethoff, Greene, Olsen, Meyer, & Meinlschmidt, 2010b). Some evidence suggests that maternal cortisol can stimulate the placental release of the corticotropin-releasing hormone (CRH) in fetal and maternal circulation, which is associated with decreased fetal growth and preterm delivery, the two major factors that cause LBW (Hobel & Culhane, 2003; Rakers et al., 2017). It is also known that norepinephrine can indirectly affect the fetus by reducing uteroplacental perfusion (Rakers et al., 2017). Reduced placental perfusion may lead to fetal undernutrition, reducing fetal growth (Baschat & Hecher, 2004). Finally, stress may be

linked to other behaviors that may affect birthweight, such as smoking, alcohol and drug use, poor nutrition, and loss of sleep (Gennaro & Hennessy, 2003).

Also, imprinted genes have been demonstrated to play important roles prenatally regulating fetal growth, placental development and signaling between the mother and her fetus (John, 2017). It has been hypothesized that aberrant imprinting in the placenta contributes to the co-occurrence of complications of pregnancy including LBW and maternal mood disorders (John, 2017). A recent genome-wide association study highlighted maternal genetic effects on the birth weight that are independent of fetal genetics (Beaumont et al., 2018). They have identified variants in the maternal genome that are robustly associated with newborn birth weight (Beaumont et al., 2018). This study aimed to evaluate the impact of prenatal maternal psychological stress on birthweight using a large cohort of predominantly Caucasian women living in an urban area. We hypothesized that high maternal psychological stress results in an unfavorable impact on birth weight. Many studies that have investigated the prenatal effects of stress on birth weight have been done in low socioeconomic status populations. The relevance of this study lies in the fact that it was performed from a large-scale prospective cohort of pregnant women ( $n=5,721$ ) and in a socioeconomic context that is mostly favorable.

## **Method**

### **Study Design**

This study is based on a large prospective cohort on pregnancies complications that includes 7,866 pregnant women recruited at the Centre Hospitalier Universitaire (CHU) de Quebec-Université Laval from April 2005 to March 2010 (A comprehensive Healthy Pregnancy Initiative from the Institute for Human Development, Child and Youth Health, Canadian Institutes of Health Research). Details of the original study design can be found

elsewhere (Bernard et al., 2019; Giguere et al., 2015; Theriault, Giguere, Masse, Girouard, & Forest, 2016; Woolcott et al., 2016). Pregnant women aged to 18 years or older and without chronic hepatic or renal diseases were invited to participate in the biobank. Women whose pregnancies included major fetal anomalies or that were terminated prior to 20 weeks of gestation because of miscarriage, voluntary pregnancy termination or fetal death were excluded of the present study (figure 1). Furthermore, women lost to follow-up or with missing delivery data, as well as women with a multiple pregnancy, were excluded. This left a sample of 7492 women with singleton pregnancy of more than 20 weeks. From this sample, 1771 women were excluded either because of missing data, the psychological stress questionnaire was not completed or the child was born with major congenital anomaly or was stillborn. The final sample included 5,721 women (figure 1). Participants gave written informed consent and the study was approved by the CHU de Quebec-Université Laval Ethics Review Board (initial approval date: 9 November 2004, Project 5-04-10-01 [95.05.17], SC12-01-159).

Infant birth weight was classified within three categories: LBW (<2,500 g), normal birth weight (NBW; 2,500-4,000 g), and macrosomia (>4,000 g).

### **Data Collection**

Data collection took place during the 50 g glucose challenge test performed between the 24<sup>th</sup> and the 28<sup>th</sup> week of gestation. Women were invited to complete a questionnaire that included information about their past medical and family history, life habits (tobacco, alcohol, drugs, physical activity, nutrition, etc.), sociodemographic characteristics (ethnicity, marital status, annual household income, highest level of education, employment status, etc.), anthropometric measures (age, height, weight), and perceived level of stress.

Infant anthropometric measures and obstetrical and neonatal complications were collected by research nurses from participant medical records following delivery. Medical

history, including gynecological and obstetrical history during the pregnancy, was also collected from the medical records.

### **Measure of Psychological Stress (MSP)**

The MSP-9 questionnaire was used to collect data on prenatal maternal stress (Langevin, François, Boini, & Riou, 2012). The MSP-9 is a validated questionnaire used to assess stress symptoms felt by someone in the last four to five days. Developed by Lemyre and Tessier, this unidimensional scale is divided in three distinctive domains: cognitive-affective, somatic, and behavioral (Lemyre & Tessier, 2003). The original version contains 45 items. A shorter nine-items version (MSP-9) was created in 2002 to meet research needs. This version meets the same reliability and validity criteria as the original measure and has an internal consistency of .89 (Lemyre & Tessier, 2003). A four point likert scale is used to answer each of the nine stress items: (1) “*Never*”, (2) “*Sometimes*”, (3) “*Often*”, (4) “*Very Often*”. A result from 9 to 15 was classified as low stress (perceived stress  $\leq$  the general population average), between 16 and 25 as intermediate stress (perceived stress above the general population average), and above 26 as high stress (extreme perceived stress) (Lemyre & Tessier, 2003).

### **Statistical Analyses**

Continuous variables were expressed as mean  $\pm$  one standard deviation (SD). Characteristics of mothers who gave birth to an infant with LBW or macrosomia and those who did not (i.e. NBW newborns) were compared using z-test for categorical variables and Mann-Whitney U-test for continuous variables. Odds ratios and their 95% confidence intervals were obtained after performing multinomial logistic regressions with infant birth weight (LBW, NBW, Macrosomia) as the dependent variable and the level of stress following the classification obtained from the MSP-9 questionnaire as the independent variable. As a first step, to determine which confounders will be added to the multivariate model, we

performed univariate logistic regression for each potential variable. A  $p$ -value of  $<.2$  in univariate analysis was set for entry of the variable in multivariate analysis. The following potential confounders were tested in univariate analysis: gestational age at delivery, parity (nulliparous, multiparous), maternal age, prepregnancy BMI, weight gain during pregnancy, ethnicity (2 Caucasian parents, Not 2 Caucasians parents, Unknown), level of education (University, Collegial, High school, None, Unknown), annual household income ( $>60,000$ \$, between  $40,000-59,999$ \$,  $<40,000$ \$, Unknown), marital status (Married/commun law, Others, Unknown), alcohol intake (0/week,  $\leq 1$ /week,  $>1$ /week, Unknown), drug consumption during pregnancy (No, Yes, Unknown), smoking status (Smoker, Ex-smoker, Nonsmoker, Unknown), antidepressants or anxiolytics use during pregnancy (No, Yes, Unknown), gender of the baby, hypertension (No, Preeclampsia, Gestational hypertension), and glucose disorders during pregnancy (No, GDM, IGT), past history of preterm delivery (Yes, No), past history of pregnancy termination (Yes, No) or miscarriage (Yes, No). As second step, the selected variables were added at the multivariate logistic regression. If the variable did not contribute significantly to the model, it was remove. In the final model, the odds ratios were adjusted for maternal age, gestational age at delivery, parity, prepregnancy BMI, weight gain during pregnancy, presence of hypertension during pregnancy, smoking status and gender of the baby. Results were considered significant if the  $p$  value was  $< .05$ . Data were analysed using XLSTAT 2018.6 software.

## Results

### Characteristics of the study population

Of the 5,721 pregnant women included in the final analysis, 175 babies weighed less than 2,500 grams (3.1%) and 611 weighed more than 4,000 grams (10.7%). Participant anthropometric and sociodemographic characteristics are described in Table 1. Table 2 refers to obstetrical characteristics for the three groups of mothers. Infants with LBW, when

compared to newborns with NBW, were more prone to be born from a mother who smoked, suffered from preeclampsia, delivered by caesarean section, experienced a previous preterm birth, and had a previous psychiatric diagnosis. Newborns with LBW were also more likely to be female. Newborns over 4,000 g, when compared to babies with normal birth weight, were more likely to be males, to be born from an older mother with a higher BMI, and delivered by caesarean section, but less likely to be born from nulliparous, smoking mothers who drank alcohol more than once per week, although the percentage of subjects in this latter group was low (1.1% in NBW vs 0.34% in the macrosomia subgroups).

### **Association between psychological stress and birth weight**

#### **Low birth weight**

Women who obtained an intermediate score on the MSP-9 did not present a higher risk of giving birth to an infant with LBW, both before and after adjustment for confounders [aOR =1.37, 95% CI (0.88-2.12)] (Table 3). Those who scored high were at increased risk to deliver a LBW baby before adjustment [OR = 2.06, 95% CI (1.04-4.09)], but the association did not remain significant after adjustment [aOR = 2.09, 95% CI (0.83-5.26)] (Table 3). However, it was observed that as the level of stress of the participants increased, the odds ratios, even if not statistically significant, increased.

#### **Macrosomia**

Expecting mothers who scored intermediate or high on the MSP-9 were at greater risk of giving birth to a newborn with macrosomia, both before and after corrections for confounders. After adjustment, risk almost doubled for women experiencing high levels of stress in comparison to those experiencing low levels of stress [aOR = 1.76 ; 95% CI (1.11-2.77)]. Risk for macrosomia was 23% greater in women at intermediate-stress levels [aOR = 1.23 ; 95% CI (1.02-1.49)] (Table 3).

### Discussion

Our study aimed to evaluate the impact of prenatal maternal psychological stress during the second trimester on birth weight in a population of mostly Caucasian women from relatively favorable socioeconomic contexts. Results indicate that maternal prenatal stress is not significantly related to LBW after confounding variables were accounted for, even though a two-fold increase in the rates of LBW infants was noted for women who scored high on the MSP-9 stress measure. The most intuitive explanation for the absence of a statistically significant association is the insufficient analytical power. Only 175 out of more than 5,000 women gave birth to an infant with LBW (3.1%), which is lower than what would be expected (6.4% in 2015)(Statistics Canada, 2016). Some studies did not find a statistically significant relationship between prenatal psychological stress and LBW (Wadhwa et al., 1993; Wang et al., 2016), unlike other studies (A. E. Borders, W. A. Grobman, L. B. Amsden, & J. L. Holl, 2007; Hashim & Moawed, 2000; Rondó et al., 2003; Stylianou-Riga et al., 2018). There are explanations for apparent inconsistent results relative to the relationship between maternal prenatal stress and infant birth weight. In their meta-analysis, Bussières et al. observed that overall there is a significant but low-level inverse relation between maternal prenatal stress and infant birth weight and gestational age. Also, they observed that the effect on birth weight increased if maternal prenatal stress was pregnancy-specific and if studies were conducted outside of North America and Europe (E. L. Bussières et al., 2015).

A number of studies that found an association between maternal stress and LBW were conducted in low-income populations (A. E. B. Borders, W. A. Grobman, L. B. Amsden, & J. L. Holl, 2007; Hashim & Moawed, 2000; Nkansah-Amankra, Luchok, Hussey, Watkins, & Liu, 2010), which was not the case with the current sample, where 63% of women had experienced college-level education or had a university degree, and 51% had a family income over 60,000 \$(CDN). Of note, our results were consistent with Wadhwa et al. who founded

that perceived stress was not related with birth weight in a favorable socioeconomic population (Caucasian women mostly married, employed, and with annual family income > \$50, 000) (Wadhwa et al., 1993). In the various previous studies, the classification was essentially based on birthweight of less than 2,500 g compared to those weighing more than 2,500 g, leading to a dichotomous approach (LBW vs the rest) (A. E. Borders et al., 2007; Hashim & Moawed, 2000; Nkansah-Amankra et al., 2010). This classification of birth weight into only two groups that includes macrosomic newborns, as opposed to the classification we used, could possibly influence the association of prenatal stress with the LBW.

Interestingly, our findings show that intermediate and high levels of maternal prenatal stress is significantly associated with the delivery of infants weighing more than 4,000 g. Women exposed to high prenatal stress had a 1.8-fold increased risk of giving birth to a newborn with macrosomia. To our knowledge, no previous studies conducted on prenatal stress and birth weight had distinguished between infants with normal birth weight and with macrosomia. When using birth weight as a continuous variable, only one other study reported a link between a higher level of perceived life stress and increased birth weight (Tegethoff et al., 2010a). Similar to our study, their Danish population sample was mainly of favorable SES. They suggested that the placenta is a key candidate linking maternal life stress and increased fetal growth. The placenta has two main functions; it provides the fetus with the nutrients and the oxygen it requires for tissue accretion and energy metabolism. It also produces hormones and reacts to maternal hormones and neutrophins that are known to affect fetal growth and development (Fowden, Forhead, Coan, & Burton, 2008). A neutrophin, brain-derived neurotrophic factor (BDNF), is suggested to play an important role in placental and fetal growth and development. A possible hypothesis explaining the relationship between maternal stress and birth weight is that altered intrauterine concentrations of BDNF may lead

to abnormal fetal growth. In nondiabetic macrosomic neonates, cord blood BDNF was significantly lower than in neonates with normal birth weight (Cai et al., 2017). Moreover, some studies reported that cord blood BDNF levels were lower in infants born to mothers with major depression and generalized anxiety disorder during pregnancy (Sonmez et al., 2019; Uguz et al., 2013).

Although the biological processes remain unclear, they hypothesized, based on animal studies, that perceived prenatal stress in early pregnancy could lower the concentration of the interleukin 10 (IL10). In mice, lower levels of stress were associated with lower IL10 and increased placental size by 28% in one study (Roberts, White, Wiemer, Ramsay, & Robertson, 2003). They also suggested that the pregnancy-associated plasma protein-A (PAPP), which can be activated by stress-activated signaling pathways (Resch, Oxvig, Bale, & Conover, 2006), may stimulate the insulin-like growth system, potentially leading to an increase in placental weight.

Research on growth factors linked to stress experience in mice increasingly suggests that stress may affect both placental volume and birth weight through different epigenetic and other mechanisms. Maternal genetic factors may influence fetal growth indirectly through the intra-uterine environment or directly through inheritance by the fetus (Beaumont et al., 2018). The identified loci highlighted potentially relevant maternal traits including fasting glucose, blood glucose, blood pressure, sex hormone levels and immune function (Beaumont et al., 2018). Adaptation to stress is influenced by different factors, such as ...

It is also possible that the heterogeneity of results in the literature may be linked to the diversity of methods used across studies and, in particular, to the definition and measurement of stress. In their meta-analysis, Bussières et al. revealed that studies concerned with pregnancy-related stress yield greater effect sizes than studies that used other definitions of

stress (E. L. Bussieres et al., 2015). It is therefore difficult to identify the kinds of stress that may affect neonatal outcomes such as LBW or macrosomia. The prenatal stress literature has conceptualized stress as negative life events, psychological distress, physical and psychological strain, perceived stress or daily hassles. The measure of psychological stress used in our study was not used in other studies, which could explain that we found an association between higher maternal prenatal stress and macrosomia.

Several researchers have also suggested that the timing of the stress assessment also needs to be considered. Roy-Matton et al. observed that women with pregnancy complications had significantly higher perceived stress at 10 to 20 weeks of gestation compared to 25 to 30 weeks of gestation, whereas the difference between women with and without pregnancy complications was not significant (Roy-Matton, Moutquin, Brown, Carrier, & Bell, 2011). Several carefully conducted studies have shown that stress experienced in the first or second trimesters may have more impact on infant birth weight than stress experienced later during pregnancy (Khashan et al., 2014; Lobel et al., 2008). Finally, other researchers have shown that study design also may affect results. Some studies measure stress retrospectively, or examine stress as it might have been experienced by mothers over a long period of time (e.g., one year), whereas other studies examine the prenatal stress-birth weight association prospectively. Such design-based variations may induce differences in results (E.-L. Bussieres et al., 2015; Gennaro & Hennessy, 2003; Tarabulsky et al., 2014).

### **Strengths and Limitations**

Our study has numerous strengths. First, this is a large-scale study involving 5,721 participants and which included 175 infants with LBW and 611 neonates with macrosomia. Our design permitted the categorization of birth weight in three groups for analysis, LBW, NBW and macrosomia, which compares favorably to most other studies that relied on a dichotomous approach with respect to birth weight (LBW vs infants > 2,500 g). Additionally,

stress was measured directly, i.e. women self-reported their perceived stress level rather than using a diagnosis of depression or anxiety. According to Gennaro and Hennessy (2003), asking women directly about perceived psychological stress, rather than anxiety or depression, could be a more accurate approach to identify acute stress (Gennaro & Hennessy, 2003). Finally, the prospective design of the study excluded the possibility of recall bias in the measure of maternal stress experience.

There were, however, several limits to the current study. Stress perception was measured only once during pregnancy, between the 24<sup>th</sup> and 28<sup>th</sup> week of gestation. Since it is not clear when stress impacts obstetrical outcomes, it could have been useful to obtain repeated measures of stress across trimesters to examine relative contributions to predicting birth weight. Likewise, it would have been useful to operationalize stress in different ways, also to see how different angles on prenatal stress may contribute to predicting infant birth weight.

### **Conclusion**

In conclusion, in this specific, relatively educated socioeconomically middle-class population of Caucasian women, maternal prenatal psychological stress measured between the 24<sup>th</sup> and the 28<sup>th</sup> week of gestation was associated with a higher risk of delivering a baby with macrosomia. This association requires corroboration by future research. There was also a two-fold, nonsignificant trend of delivering a newborn with LBW with increased maternal stress, after controlling for confounding variables. Considering the consequences of both macrosomia and LBW on the newborn's immediate and future health, healthcare prevention programs that help lower stress during pregnant women would probably be beneficial to improving infant birth weight.

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### Tables

Table 1

*General Characteristics of the Study Population*

Characteristics	LBW ( <i>n</i> =175)	NBW ( <i>n</i> =4935)	Macrosomia ( <i>n</i> =611)
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Age, years, <i>mean ± SD</i>	29.69 ± 4.55	29.52 ± 4.30	29.86 ± 4.02†
BMI before pregnancy, kg/m <sup>2</sup> , <i>mean ± SD</i>	23.79 ± 5.68	24.16 ± 5.14	26.36 ± 6.05†
Ethnicity, <i>n (%)</i>			
2 Caucasian parents	165 (98.21)	4570 (97.01)	560 (96.72)
Not 2 Causasian parents	3 (1.79)	141 (2.99)	19 (3.28)
Smoking status, <i>n (%)</i>			
Smoker	46 (26.29)†	616 (12.49)	43 (7.06)†
Ex-smoker	33 (18.86)	1213 (24.60)	164 (26.93)
Never smoked	96 (54.86)†	3101 (62.90)	402 (66.01)
Alcohol consumption, <i>n (%)</i>			
0	145 (91.19)	4398 (93.53)	556 (94.56)
≤ 1 per week	13 (8.18)	252 (5.36)	30 (5.10)
> 1 per week	1 (0.63)	52 (1.11)	2 (0.34)†
Drug consumption, <i>n (%)</i>			
Yes	6 (3.53)	113 (2.35)	10 (1.70)
No	164 (96.47)	4690 (97.65)	579 (98.30)
Marital status, <i>n (%)</i>			
Married/Common law	159 (91.38)	4581 (93.11)	566 (92.79)
Single	15 (8.62)	339 (6.89)	44 (7.21)
Highest education level, <i>n (%)</i>			
No diploma	12 (6.90)	187 (3.80)	22 (3.61)
High school	52 (29.89)	1154 (23.44)	135 (22.17)
Collegial	47 (27.01)	1606 (32.62)	199 (32.68)
University	63 (36.21)	1976 (40.14)	253 (41.54)
Annual family income, \$CAN, <i>n (%)</i>			
< 40 000	46 (28.57)	996 (21.75)	107 (18.90)
40 000-59 999	32 (19.88)	1021 (22.29)	119 (21.02)
≥60 000	83 (51.55)	2563 (55.96)	340 (60.07)
Previous psychiatric	23 (14.74)†	377 (7.93)	38 (6.33)

diagnosis, *n* (%)

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*Note.* LBW = low birth weight; NBW = normal birth weight; BMI = body mass index.

†*p* < .05 when compared to NBW

Table 2

*Obstetrical Characteristics of the Study Population*

Characteristics	LBW ( <i>n</i> =175)	NBW ( <i>n</i> =4935)	Macrosomia ( <i>n</i> =611)
Gestational age at delivery, <i>mean ± SD</i>	35.47±2.98†	39.43±1.23	40.10±0.92†
Baby's sex, <i>n</i> (%)			
Male	72 (41.14)†	2519 (51.08)	403 (65.96)†
Female	103 (58.86)	2414 (48.92)	208 (34.04)
Mode of delivery, <i>n</i> (%)			
Vaginal	109 (62.29)	3918 (79.39)	229 (73.49)
Caesarean section	66 (37.71) †	1017 (20.61)	162 (26.51) †
Parity, <i>n</i> (%)			
Nulliparous	96 (54.86)	2404 (48.71)	208 (34.04)†
Multiparous	79 (45.14)	2531 (51.29)	403 (65.96)
Hypertensive disorders, <i>n</i> (%)			
Preeclampsia	24 (13.71) †	70 (1.42)	11 (1.80)
Gestational hypertension	12 (6.86)	139 (2.82)	21 (3.44)
Glucose disorders, <i>n</i> (%)			

GDM	13	(7.43)	219	(4.44)	38	(6.22)
IGT	8	(4.57)	106	(2.15)	17	(2.78)
Previous preterm birth, <i>n</i> (%)	28	(16.00)†	234	(4.74)	24	(3.93)
Previous miscarriage, <i>n</i> (%)	40	(22.86)	1087	(22.03)	136	(22.26)
Previous pregnancy termination, <i>n</i> (%)	37	(21.14)	950	(19.25)	112	(18.33)

*Note.* LBW = low birth weight; NBW = normal birth weight; † =  $p < .05$  when compared to NBW.

Table 3

*Association Between Psychological Stress and Birth weight*

MSP-9 score	LBW (n=175)			Macrosomia (n=611)		
	n (%)	Non-adjusted OR (CI 95%)	Adjusted OR* (CI 95%)	n (%)	Non-adjusted OR (CI 95%)	Adjusted OR* (CI 95%)
Low	65 (37.14)	1 (Reference)	1 (Reference)	229 (37.48)	1 (Reference)	1 (Reference)
Intermediate	100 (57.17)	1.24 (0.90-1.70) $p = .185$	1.37 (0.88-2.12) $p = .116$	353 (57.77)	<b>1.24 (1.04-1.48)</b> $p = .016$	<b>1.23 (1.02-1.49)</b> $p = .027$
High	10 (5.71)	<b>2.06 (1.04-4.09)</b> $p = .038$	2.09 (0.83-5.26) $p = .116$	29 (4.74)	<b>1.70 (1.12-2.58)</b> $p = .013$	<b>1.76 (1.11-2.77)</b> $p = .016$

\*adjustments done for gestational age at delivery, parity, maternal age, pre-pregnancy body mass index, weight gain during pregnancy, , smoking status, sex of the baby, hypertensive disorders during pregnancy. Boldface = significant result. LBW = low birth weight ; MSP = Measure of Psychological Stress; OR = odds ratios.

Figure

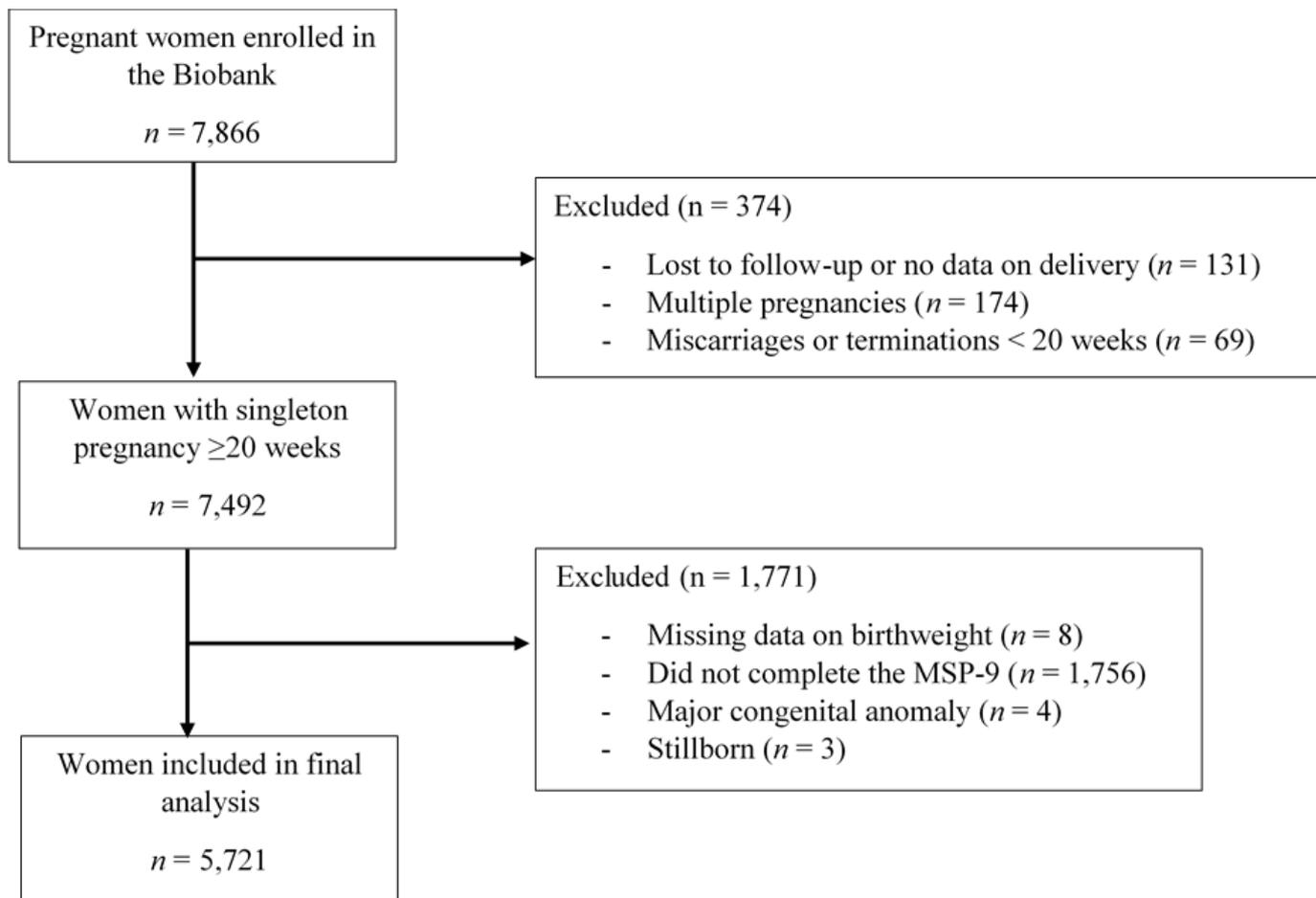


Figure 1. Flow chart of the sample selection for the study of the impact of the maternal psychological stress on birth. MSP = Measure of Psychological Stress.

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