BMJ Open

Psychosocial deprivation in women with gestational diabetes mellitus is associated with poor fetomaternal prognoses: an observational study

Journal:	BMJ Open
Manuscript ID:	bmjopen-2014-007120
Article Type:	Research
Date Submitted by the Author:	05-Nov-2014
Complete List of Authors:	Cosson, Emmanuel; AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Endocrinology-Diabetology-Nutrition; Sorbonne Paris Cité, UMR U1153 Inserm / U1125 Inra / Cnam / Université Paris 13, Bihan, Hélène; AP-HP, Avicenne Hospital, Paris 13 University, Sorbonne Paris Cité, Diabetology, Metabolic Diseases; Sorbonne Paris Cité, UMR U1153 Inserm / U1125 Inra / Cnam / Université Paris 13, Reach, GTrard; APHP, Endocrinology Vittaz, Laurence; Ballanger Hospital, Carbillon, Lionel; Assistance Publique Hopitaux de Paris, Paris 13 University, Department of Obstetrics and Gynaecology Valensi, Paul; AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Endocrinology-Diabetology-Nutrition
Primary Subject Heading :	Diabetes and endocrinology
Secondary Subject Heading:	Global health, Obstetrics and gynaecology, Occupational and environmental medicine
Keywords:	Diabetes in pregnancy < DIABETES & ENDOCRINOLOGY, OBSTETRICS, SOCIAL MEDICINE

SCHOLARONE[™] Manuscripts

BMJ Open

Psychosocial deprivation in women with gestational diabetes mellitus is associated with poor fetomaternal prognoses: an observational study

E Cosson *professor*¹², H Bihan *investigator*²³, G Reach *professor*³, L Vittaz *investigator*⁴, L Carbillon *professor*⁵, P Valensi *professor*¹

¹ AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Endocrinology-Diabetology-Nutrition, CRNH-IdF, CINFO, 93143 Bondy, France; ² Sorbonne Paris Cité, UMR U1153 Inserm / U1125 Inra / Cnam / Université Paris 13, 93000 Bobigny, France; ³ AP-HP, Avicenne Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Diabetology, Metabolic Diseases, CRNH-IdF, 93000 Bobigny, France; ⁴ Ballanger Hospital, Department of Endocrinology-Diabetology, 93600 Aulnay-Sous-Bois, France; ⁵ AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Obstetrics and Gynecology, 93143 Bondy, France

Corresponding author:

Emmanuel Cosson

Department of Endocrinology-Diabetology-Nutrition

Avenue du 14 juillet, Hôpital Jean Verdier, 93143 Bondy Cedex. FRANCE

Tel: +33 1 48-02-65-80, Fax: +33 1 48-02-65-79

Sorbonne Paris Cité, UMR U1153 Inserm / U1125 Inra / Cnam / Université Paris 13, 93000 Bobigny, France

e-mail: emmanuel.cosson@jvr.aphp.fr

professor

- Hélène Bihan

² Sorbonne Paris Cité, UMR U1153 Inserm / U1125 Inra / Cnam / Université Paris 13, 93000
Bobigny, France;

³AP-HP, Avicenne Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Diabetology, Metabolic Diseases, CRNH-IdF, 93000 Bobigny, France

helene.bihan@avc.aphp.fr

investigator

- Gérard Reach

³AP-HP, Avicenne Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Diabetology, Metabolic Diseases, CRNH-IdF, 93000 Bobigny, France

gerard.reach @avc.aphp.fr

professor

- Laurence Vittaz

⁴Ballanger Hospital, Department of Endocrinology-Diabetology, 93600 Aulnay-Sous-Bois, France;

laurence.vittaz@ch-aulnay.fr

investigator

- Lionel Carbillon

⁵ AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Obstetrics and Gynecology, 93143 Bondy, France

lionel.carbillon@jvr.aphp.fr

professor

- Paul Valensi

BMJ Open

¹ AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Endocrinology-Diabetology-Nutrition, CRNH-IdF, CINFO, 93143 Bondy, France

paul.valensi@jvr.aphp.fr

professor

rds, main u Word counts: abstract 250 words, main text 2800 words

3 tables and 1 figure

1 supplemental file

50 references

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

What is already known on this topic

Psychosocial deprivation is associated with more gestational diabetes mellitus.

What this study adds

Among women with gestational diabetes mellitus, psychosocial deprivation is associated with large for gestational age infants, independently of obesity, gestational weight gain and other confounders.

BMJ Open

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

Abstract

Objective To evaluate the prognoses associated with psychosocial deprivation in women with gestational diabetes mellitus (GDM).

Design Observational study considering the 1498 multiethnic women with GDM who gave birth between January 2009 and February 2012.

Setting Four largest maternity units in the northeastern suburban area of Paris.

Participants The 994 women who completed the Evaluation of Precarity and Inequalities in Health Examination Centers [EPICES] questionnaire.

Main outcome measure Main complications of GDM (large infant for gestational age (LGA), shoulder dystocia, cesarean section, preeclampsia).

deprivation (EPICES score \geq 30.17) affected 577 women (56%) and was **Results** Psychosod with overweight/obesity, parity, and non-European origin and negatively positively associat associated with far y history of diabetes, fruit and vegetable consumption, and working status. The psychosociall eprived women were diagnosed with GDM earlier, received insulin treatment during p gnancy more often and were more likely to have LGA infants (15.1 vs. 10.6%, odds ratio [95% confidence interval 1.02-2.2], p<0.05) and shoulder dystocia (3.1 vs. 1.2%, OR 2.7[0.9] .2], p<0.05). In addition to psychosocial deprivation, LGA was associated besity, history of GDM, ethnicity, excessive gestational weight gain and with greater parity ultivariate analysis using these covariates revealed that EPICES score was insulin therapy. A independently asso ated with LGA infants (per 10 units, OR 1.12[1.03-1.20], p<0.01).

Conclusions In our area, psychosocial deprivation is common in women with GDM and is associated with earlier GDM diagnoses and greater insulin treatment, an increased likelihood of

luat
es n
ona
200
est 1
99
on (
ieas
, ce
cial
ted
mil
ly d
oreg
1.5
7-7.
y, ol
mu
ocia
ur a
rlie

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

shoulder dystocia and, independently of obesity, gestational weight gain and other confounders with LGA infants.

Strengths and limitations:

- It is known that psychosocial deprivation is associated with more gestational diabetes and we report for the first time that, among women with gestational diabetes mellitus, psychosocial deprivation is associated with a poor prognosis.
- Our large, multicenter and diverse cohort and the adjustments for the relevant confounding factors, such as body mass index and gestational weight gain, ensure the robustness of our findings.
- We defined psychosocial vulnerability using the Evaluation of Precarity and Inequalities in Health Examination Centers [EPICES] questionnaire, which has been validated during pregnancy or not. This tool evaluates at an individual level several domains, including material goods, money, friendship and family networks, healthcare and leisure.
- However, the EPICES questionnaire was retrospectively fulfilled (6 to 24 months after pregnancy).

Key words: gestational diabetes mellitus, psychosocial deprivation, prognoses, large infant for gestational age

Introduction

Socioeconomic status reflects access to resources to prosper, and psychosocial deprivation is associated, across countries and over time,¹² with higher mortality and morbidity, including type 2 diabetes.³ The main drivers in more incident type 2 diabetes appear to be higher body mass index (BMI) and impaired health behaviors.⁴ The American Diabetes Association recommends the inclusion of assessments of patients' psychological and social situations as an ongoing part of the medical management of diabetes.⁵ Indeed, psychosocial deprivation in patients with diabetes has been reported to be associated with increased obesity,⁶ worse glycemic control,⁷ poorer adherence,^{8 9} more diabetic complications,^{6 7 10 11} and perhaps greater mortality.¹²⁻¹⁴ During pregnancy, psychosocial deprivation is also associated with poor outcomes that include increased rates of maternal¹⁵ and neonatal^{15 16} hospitalization, stillbirth,¹⁷ postnatal death,¹⁸ preterm delivery^{17 19} and changes in fetal growth.¹⁷⁻²⁰

Gestational diabetes mellitus (GDM) is defined as any degree of glucose intolerance with onset or first recognition during pregnancy and is now very common, with a prevalence ranging from 9.3 in Israel to 25.5% in California, US.²¹ Although GDM is also more frequent in cases of psychosocial deprivation,^{18 19 22 23} its prognosis in case of poor psychological and social conditions is currently unknown. We hypothesized that psychosocial deprivation might be associated with poor prognoses in women with GDM when confounding factors, such as obesity,¹⁹ gestational weight gain (GWG)¹⁶ and smoking habits,²⁰ are considered.

The four largest maternity units in the Northeastern suburban area of Paris, France participated in the IMPACT initiative, which aimed to improve postpartum screening for dysglycemia after GDM.^{24 25} The women who attended these maternity units responded to the Evaluation of Precarity and Inequalities in Health Examination Centers [EPICES] questionnaire,

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

a questionnaire which evaluates psychosocial deprivation.^{67 19 24 26} Therefore, for the first time, we had the opportunity to investigate the fetomaternal prognoses of these women with GDM according to their individual psychosocial statuses.

Methods

Patients

This study is a secondary analysis of the IMPACT study.^{24 25} Briefly, the IMPACT initiative began in March 2011 and was a mobilization campaign for women with GDM and their community caregivers that sought to increase postpartum screening for dysglycemia. We aimed to evaluate the effect of this initiative by comparing the postpartum screening rates between the women who delivered before (between January 2009 and December 2010) and after this initiative (between April 2011 to February 2012). We systematically included women who were at least 18 years of age, free of known pregestational diabetes, had GDM and were followed during pregnancy in one of the four largest maternity units of the Seine-Saint Denis area of France during these periods of time. GDM was detected by oral glucose tolerance test and was defined by fasting blood glucose values ≥ 5.3 mmol/l and/or a 2-hour blood glucose value ≥ 7.8 mmol/l between January 2009 and December 2010,^{24 25} and thereafter according to the International Association of Diabetes and Pregnancy Study Groups criteria, adopted in France in 2010.²⁷ In the primary analyses, we included the women who could be contacted by telephone and provided self-reports that indicated whether they had undergone postpartum screening tests during the six months following their deliveries.^{24,25} For the current analysis, we included all of the women who delivered between January 2009 and February 2012 and who retrospectively

BMJ Open

completed the [EPICES] questionnaire by phone regardless of their report concerning the postpartum screening.

Data collection and assessment of outcomes

One single investigator extracted the following data from hospital records: age at conception, origin/ethnicity, family history of diabetes, history of previous GDM, gestational age at the time of GDM diagnosis (three classes: <24 weeks of gestation, between 24 and 28 weeks of gestation and ≥ 28 weeks of gestation), insulin treatment during pregnancy, and GWG. Excessive GWG was defined according to the recommendations of the Institute of Medicine; *i.e.*, $GWG \ge 16$ kg in women with pregravid BMIs $< 25 \text{ kg/m}^2, \ge 11.5 \text{ kg}$ in overweight women (BMIs between 25 and 29.9 kg/m²) and \geq 9 kg in obese women (BMI \geq 30 kg/m²). We also collected information about events during pregnancy, including offspring birth weight in comparison to the standard French population (large for gestational age (LGA) was defined by a birth weight exceeding the 90th percentile)²⁸, preeclampsia (blood pressure > 140/90 mmHg on two recordings 4 h apart and proteinuria of at least 300 mg/24 h or 3+ or higher upon dipstick testing of a random urine sample), shoulder dystocia (defined as the use of obstetrical maneuvers *i.e.*, McRoberts episiotomy after delivery of the fetal head, suprapubic pressure, posterior arm rotation to an oblique angle, rotation of the infant by 180 degrees, delivery of the posterior arm) and caesarian section.

The investigator conducted semi-structured interviews by phone between January and November 2011 for the women who delivered before the IMPACT campaign (maximum delay of time since delivery 24 months) and at least six months after delivery for the women who delivered after the IMPACT initiative. The investigator requested information about the subjects' current weights, heights, waist circumferences, professional statuses, smoking statuses, number

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

of children, antihypertensive and lipid-lowering treatments, family histories of diabetes and daily consumptions of fruits and vegetables. All these data were therefore declarative. Waist circumference was deducted from current waist size of trousers (waist circumference <80 cm: 6-14 (UK) or 34-42 (France), 80-88 cm: 16-20 (UK) or 44-48 (France); >88 cm: 22 (UK) or 48 (France) or more).

The investigator also conducted interviews to assess psychosocial deprivation using the EPICES score, which is a French deprivation score that is calculated based on responses to 11 questions that consider both socioeconomic conditions and family environment (supplementary material).^{7 26 29} It evaluates at an individual level several domains, including material goods, money, friendship and family networks, healthcare and leisure. As previously reported, the EPICES score is a continuous variable, and increasing quintiles are associated with increased risks for poor health conditions such as obesity, diabetes in women, higher rates of smoking, poorer access to dental and gynecological care, and poorer perceived health statuses.^{24 26} However, psychosocial deprivation can be defined by a score ≥ 30.17 ,²⁹ which was the threshold used here.

Statistical analyses

Sample size calculations were based on the main criterion of the IMPACT study, *i.e.* a postpartum screening test performed six months following delivery.^{24 25} Results reported in this manuscript were prespecified, exploratory endpoints. Continuous variables are expressed as means \pm SD, and normality was assessed with Kolmogorov-Smirnov tests. There were no missing data concerning psychosocial deprivation and main outcomes. Comparisons of two independent groups were performed using the Student t-test if the variable was normally

distributed; otherwise, the Wilcoxon Mann-Whitney test was used. The significance of differences in proportions (*i.e.*, qualitative variables) were tested with the x^2 test, and the odds ratios (ORs) and 95% confidence intervals (CIs) were calculated in cases of statistical significance (p<0.05). We defined EPICES score tertiles in our cohort: first tertile: EPICES score <23.71 (mean 11.7±6.2; n=296); second tertile: score between 23.71 and 51.5 (mean 35.0±8.5; n=355); and third tertile: score \geq 51.5 (mean 69.9±12.6; n=343). The factors associated with having a LGA infant were assessed with a univariate logistic regression method. For multivariate analyses, we included all factors that were associated with LGA infants with p values \leq 0.05 in the univariate analyses. SAS Statistics version (9.2) (Cary, USA) was used to conduct all statistical analyses.

Results

Characteristics of the women

A total of 1498 women gave birth following GDM between January 2009 and February 2012 in our maternity units. Of these women, 994 retrospectively responded by phone to the EPICES questionnaire. Table 1 illustrates the characteristics of these women. The characteristics of the 994 women who responded to the EPICES questionnaire were similar to those of the 504 who did not respond with the exception of greater daily consumptions of fruits and vegetable (66.1 vs. 59.0%, respectively, p<0.01) and a trend toward being older (33.3 ± 5.2 vs. 32.7 ± 5.5 years, respectively, p=0.06). The EPICES questionnaire could not be completed by the women who could not be reached by phone and those with French language proficiencies that were insufficient for answering the questions.

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

Psychosocial deprivation affected 577 women (56%) and was positively associated with parity, overweight and obesity, greater waist circumference, and non-European origin. Psychosocial deprivation was negatively associated with daily fruit and vegetable consumption, reduced family history of diabetes and working status (Table 1).

Pregnancy outcomes

Table 2 shows that the psychosocially deprived women were more likely to have been diagnosed with GDM prior to 24 weeks of gestation and more likely to have been treated with insulin during pregnancy than the non-psychosocially deprived women. The psychosocially deprived women were more likely to have LGA infants and infants with shoulder dystocia, but no differences in cesarean section or preeclampsia were found. Figure 1 (panels A to D) shows that the prevalences of insulin treatment during pregnancy (Figure 1A), LGA infants (Fig 1C) and shoulder dystocia (Figure 1D) increased with increasing EPICES score tertile.

Table 3 shows that, in addition to psychosocial deprivation (OR 1.5 [95%CI 1.02-2.22]), LGA was associated with higher parity, greater BMI and obesity of the mother (OR 2.1 [1.4–3.1]), increased incidence of GDM history (OR 2.0 [1.4–3.1]), ethnicity/origin, greater EPICES score (per 10 units: OR 1.50 [1.22-2.22]), greater GWG and excessive GWG during pregnancy (OR 2.8 [1.9-4.1]) and insulin treatment during pregnancy (OR 1.6 [1.1–2.4]). A multivariate analysis that considered parity, obesity, personal history of GDM, ethnicity, EPICES score, excessive GWG and insulin therapy during pregnancy revealed that the EPICES score remained independently associated with LGA infants (Table 3). In a model that was identical to the aforementioned model with the exception that psychosocial deprivation (*i.e.*, EPICES score

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

 \geq 30.17) was used in the place of the EPICES score, a trend toward an association between psychosocial deprivation and LGA infants remained (OR 1.53 [0.98-2.39], p=0.06).

Discussion

In this study, psychosocial deprivation in women with GDM was associated with earlier GDM diagnoses and more extensive insulin treatment. Moreover, we show for the first time that, independent of confounding factors, psychosocial deprivation was associated with increases in adverse outcomes, particularly LGA infants. We report that psychosocial deprivation (*i.e.*, an EPICES score above 30.17) affected 56% of the women with GDM in our study; another study reported a prevalence of 48% (11/23 women with GDM) in another area of France using the same definition of deprivation.¹⁹ This high prevalence is due to the prevalence of precarity³⁰ and multiethnicity³¹ in the Northeastern suburban area of Paris and to the roles played by these conditions in the rate of GDM. Indeed, the prevalence of GDM has been reported to be 1.7- to 2.9-fold higher among patients with high EPICES scores,¹⁹ low educational statuses^{22 23} or low family incomes^{18 23} compared to their counterparts without these conditions. Notably, 23% of pregnant women in France have been reported to have high EPICES scores regardless of GDM status,¹⁹ and 17.5% have been coded as psychosocially deprived by social workers¹⁵ in two other areas in France. Together, our results advocate for screening for deprivation among pregnant women with GDM.

As previously reported for women with and without GDM,^{19 32} we found that psychosocially deprived women with GDM were more likely to be obese. These women were also more likely to be unemployed and less likely to be daily consumers of fruits or vegetables; the latter

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

association is likely due to the cost of these foods. An association between socioeconomic status and healthy eating status, including fruits and vegetable consumption, has previously been reported.^{33 34} We also observed a link between ethnicity/origin and deprivation; similar links have previously been described as complex relationships.³⁵⁻³⁸ The women with and without psychosocial deprivation reported similar prevalences of prepregnancy antihypertensive and lipid-lowering treatments although metabolic disorders are often associated with elevated EPICES scores^{39 40} and stress.⁴¹ The lack of association observed in our study might be specific to women of reproductive age or might be attributable to reduced numbers of medical visits prior to pregnancy due to precarity.¹⁶ The latter supposition would also result in undiagnosed metabolic syndrome prior to pregnancy, which would be in accordance with the greater prevalence of GDM diagnoses prior to 24 weeks of gestation among psychosocially deprived women. These findings suggest the possibility that these women might actually have had undiagnosed pregravid type 2 diabetes. Indeed, precarity is a risk factor for undiagnosed type 2 diabetes even in women of reproductive age.⁴² However, we do not have access to the results of postpartum glycemic assessments that would be needed to confirm this hypothesis.

We also studied the association between psychosocial deprivation and adverse pregnancy outcomes in women with GDM for the first time. Compared to those without precarity, the women with precarity were more likely to have LGA infants and infants with shoulder dystocia. The association between EPICES scores and LGA infants was independent of obesity, which suggests that this relationship was only partially driven by the increased prevalence of overweight/obesity among the deprived women.^{43 44} GWG and the prevalence of excessive GWG, which are other confounding factors regarding LGA infants,^{43 44} were comparable between the women with and without precarity. We have recently shown that, compared to

Page 15 of 37

BMJ Open

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

women from Sub-Saharan Africa, European women experience more GDM-related events.³¹ Furthermore, racial/ethnic differences in the clinical outcomes of GDM, including macrosomia, are commonly reported (for review).⁴⁵ Here, the association of LGA with precarity remained significant after adjusting for origin/ethnicity while we did not find any association between precarity and offspring with birth weights greater than 4000 g or 4250 g. The association between psychosocial deprivation and shoulder dystocia, which was not adjusted for confounding factors because the rate of dystocia events was low, was most likely driven by the prevalence of LGA infants. In a population-based study, the risk for shoulder dystocia significantly increased with BMI category in an unadjusted analysis, but this significance disappeared after adjusting for GDM.³⁵ As previously reported for pregnant women regardless of GDM status,¹⁵ we found that the women in our cohort with GDM underwent cesarean section at similar rates regardless of the presence of psychosocial deprivation. The vulnerable women were diagnosed with GDM earlier, which suggests that unknown pregravid dysglycemia might partially explain the increased rate of LGA infants.⁴⁶ In a recent German study, the groups that were found to at high risk for GDM were women of low socio-economic status, migrants and obese women. An elevated risk of fetal malformations was found among the women who had been diagnosed with GDM, which suggests that many of these women might have had high glucose levels by the first trimester.⁴⁷

The present study has limits and strengths. Our large, multicenter and diverse cohort and the adjustments for the relevant confounding factors ensure the robustness of our findings. However, we did not have access to data about glycemic control, diet, physical activity or the numbers of visits during pregnancy. Thus, the adverse outcomes observed for the women with precarity might have been due to these factors based on the following arguments: *(i)* poor glycemic control

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

has been reported in vulnerable patients with diabetes⁷ and was likely present in our population with GDM and psychosocial insecurity because insulin treatment was more often necessary during GDM among this population; *(ii)* fruit and vegetable consumption was lower among the vulnerable women following pregnancy, which might be indicative of poorer nutritional habits during pregnancy;^{33 34 48} *(iii)* exercise during late pregnancy has been reported to vary with the education level of the mother;^{33 34 49} and *(vi)* access to health care might depend on socioeconomic status,¹⁹ but it is unlikely that access to health care influenced our results because health care is free of charge within the French healthcare system. We used the EPICES score, which is an individual index that has been validated during pregnancy¹⁹ and appears to be more strongly linked to the risk of adverse materno-fetal outcomes than neighborhood-level socioeconomic status.¹⁷ However, the EPICES questionnaire was retrospectively fulfilled (6 to 24 months after pregnancy).

Conclusions

To conclude, our results from a large multiethnic multicenter European cohort from an area in which precarity is common demonstrate that psychosocial deprivation affected more than half of the women with GDM. Psychosocial deprivation was associated with higher BMIs and earlier GDM diagnoses among the vulnerable women, which suggests that GDM likely corresponded to unknown type 2 diabetes mellitus in these women and that prenatal diagnosis of type 2 diabetes should be reinforced in them, with weight control intervention and adherence to healthy lifestyle before pregnancy.⁵⁰ The vulnerable women were also more likely to be treated with insulin, but they gained as much weight during pregnancy as did the non-vulnerable women. Independent of the gestational age at GDM diagnosis, insulin use, overweight/obesity, GWG and other confounders, these women were also more likely to have LGA infants. This finding suggests that

BMJ Open

the routine screening of women with GDM for psychosocial vulnerability may be an important tool for improving the prognoses of these women and their children. For example, specific follow-up and psychosocial support might be beneficial in these women.

Contributors: EC researched data, directed research and wrote the manuscript; HB researched data and wrote the manuscript; GR directed research and reviewed/edited the manuscript; LV researched data and contributed to discussion; LC researched data and reviewed/edited the manuscript; PV directed research and reviewed/edited the manuscript. All authors contributed to the interpretation of the results and the revision of the manuscript for intellectual content and approved the final version of the manuscript. Delphine Dubois, Umanis, Paris, is the guarantor of this work and, as such, had fully access to all the data in the study and takes the responsibility for the integrity of the data and the accuracy of the data analyses.

Funding: This research was supported by a grant from Novo Nordisk® France. Novo Nordisk was not involved in study design, analyses and interpretation of results and article writing. It provided money to pay the investigator, to perform statistical analyses and for English language editing service.

Competing interest: All authors have completed the ICMJE uniform disclosure form at www.icmje.org/coi_disclosure.pdf . Dr. COSSON reports grants from Novo Nordisk France, during the conduct of the study; personal fees from Novo Nordisk France, outside the submitted work. Dr. Bihan reports personal fees from Novonordisk, during the conduct of the study. Dr Reach reports no conflict of interest in the case of this procedure. Dr. Carbillon reports personal

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

fees from Novonordisk during the conduct of the study. Dr. VALENSI reports grants from Novo Nordisk France during the conduct of the study; personal fees from Novo Nordisk France, outside the submitted work.

Ethical approval: The study protocol was approved by the National Ethics Committee

(CCTIRS: *Comité Consultatif sur le Traitement de l'Information en Matière de Recherche*; advisory committee on research information processing).

Transparency declaration: The lead author, EC, affirms that the manuscript is an honest, accurate, and transparent account of the study being reported; that no important aspects of the study have been omitted; and that any discrepancies from the study as planned (and, if relevant, registered) have been explained.

Clinical trial registration: observational study.

Data sharing: No additional data available.

Acknowledgments: The authors acknowledge the statistical assistance of Bénédicte Borsik, Delphine Dubois and Anne Ourliac (Umanis, Paris, France – funding source: Novo Nordisk®). The authors also thank Dr Faranaz Faghfouri (AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Obstetrics and Gynecology, Bondy, France) who was the study investigator (funding source: Novo Nordisk®); and Drs H Dauphin (Ballanger Hospital, Department of Obstetrics and Gynecology, Aulnay-Sous-Bois, France); Chafika Khiter (De La Fontaine Hospital, Department of Diabetology, Saint Denis, France), Dominique Leboeuf (Seine-Saint-Denis Private Hospital, Department of Obstetrics and Gynecology, Le Blanc Mesnil, France) and Astrid Lepagnol (De La Fontaine Hospital, Department of Obstetrics and Gynecology, Saint Denis, France) who recruited the women. This program was sponsored by the Société Francophone de Diabétologie, the réseau pour la prise en charge et la prévention de

BMJ Open

l'obésité en pédiatrie 93-Seine-Saint-Denis (REPOP 93), L'Assurance Maladie Seine-Saint-Denis, l'Ordre National Des Pharmaciens and Université Paris 13.

 e-Saint-Denis (k.

 cs Pharmaciens and Unive

EXCLUSIVE LICENCE

BM Group

BMJ Licence to BMJ Publishing Group Limited ("BMJ Group") for Publication To be agreed to by the corresponding author or guarantor on behalf of all authors ("Corresponding Author"). All authors collectively are referred to as the "Contributors"

In consideration of the BMJ Group ("the Publishers") considering to publish the article contained within the original manuscript which includes without limitation any diagrams, photographs, other illustrative material, video, film or any other material howsoever submitted by the Contributor(s) at any time and related to the Contribution ("the Contribution") in the BMJ ("the Journal"), certain rights are required to be granted by each different category of author(s), which are as follows:

- For employees of the UK Crown acting in the course of their employment- a non exclusive Licence, as set out below. All provisions of this document apply. The non exclusivity relates to the original submitted manuscript video, films, images, photographs, diagrams and/or illustrative material only).
- For employees of the US Federal Government employees acting in the course of their employment, no copyright exists and the Contribution is in the public domain so no licence is required to be granted. The Author Warranties below apply (excluding 1.iii).
- For all other authors, an exclusive Licence, as set out below. All provisions of this document apply.

NB where a Contribution is a multi authored work, each author's element of the Contribution will be dealt with in accordance with 1, 2 or 3 above, as applicable.

The Licence

The Licence granted in accordance with 1 or 3 above is:

A worldwide, licence, to the Publishers and its licensees in perpetuity (subject to the Reversion of Rights set out below), in all forms, formats and media (whether known now or created in the future), to i) publish, reproduce, distribute, display and store the Contribution, ii) translate the Contribution into other languages, create adaptations, reprints, include within collections and create summaries, extracts and/or, abstracts of the Contribution and convert or allow conversion into any format including without limitation audio, iii) create any other derivative work(s) based in whole or part on the Contribution, v) to exploit all subsidiary rights that currently exist or as may exist in the future in the Contribution, v) the inclusion of electronic links from the Contribution to third party material where-ever it may be located; and, vi) licence any third party to do any or all of the above.

If you and/or any co-author's employer owns the copyright to your contribution you must obtain in writing the relevant employers' consent to grant the licence and agree to all obligations herein. The author(s) hereby agree that in the event that the BMJ Group sell, the whole or part of its journal business to any third party, the benefit and the burden of the Licence contained herein shall be assigned to that third party.

Additional Rights and Obligations

The author(s) (and their employers as applicable), hereby authorise the Publishers to take such steps as they consider necessary at their own expense in the copyright owners name and on their behalf, if they believe that a third party is infringing or is likely to infringe copyright or the rights granted to the Publishers herein in the Contribution without further recourse to the copyright owner(s).

For Original Research articles and Open Access Funded Articles (as both defined below), the Publishers expressly agree to place the published Contribution for display on PubMed Central (including its international mirror sites) promptly without charge to the authors or their employers (provided Pubmed Central does not charge the Publishers), which will include any Publishers' supplied amendments or retractions.

BMJ Author Licence March 2013 doc

"Original Research" means an article reporting a research study, with a research structured abstract and normally appearing in the Research section of the BMJ. "Open Access Funded Articles" means articles funded in whole or part by a research grant from a government and/or charitable organisation(s) that requires open access deposit in PubMed Central. Such articles are identified by the reference to a Creative Commons licence.

The author(s) acknowledge and accept that BMJ Group may make additional changes to the contribution as considered necessary in accordance with standard editorial processes whether before or after publication. The Corresponding Author will usually see proofs for their Contribution s and every effort will be made to consult with the Corresponding Author if substantial alterations are made. The BMJ Group may also retract or publish a correction or other notice when it considers this appropriate for legal or editorial reasons and this shall be at its absolute discretion which shall be exercised reasonably.

Reversion of Rights

If the Contribution is not published in either the print or electronic versions of the Journal or any other Publisher(s) products, within 12 months of final acceptance by the BMJ Group, (or as otherwise agreed), any Licence granted herein shall automatically terminate and all rights shall revert to the copyright owner. The Publishers may keep a copy of the Contribution as a record (including via any contractor).

Rights Granted to Owners of the Contribution

Ownership of copyright remains with the author(s) or their employers. All rights not expressly granted are, subject to the Licence terms, reserved by the Publishers. In return for the grant of the Licence herein, the copyright owner(s) shall have the following rights for <u>non-Commercial Use</u> (unless otherwise stated) of the Contribution:

1.The right to reproduce a reasonable number (no more than 100) print copies of the final Contribution, by copying or downloading from the BMJ Group website, for personal use and to send copies to colleagues in print or electronic form provided no fee is charged and this is not done on a systematic basis (which includes via mass e-mailings).

2. The right to include the Contribution in a compilation for classroom use (course packs) to be distributed free of charge (other than for direct photocopying cost) to students at the Contributor(s)'s institution or to be stored in digital format in data rooms for access by students as part of their course work and for in house training programmes of the Contributor(s)'s employer or at seminars or conferences subject to a limit of 100 copies per conference or seminar.

3. For all articles (excluding articles commissioned by the Publishers), the right to post a version of the final published version of the Contribution, or any abstract of the final published Contribution on the Contributor(s)'s own and/or his/her institution's website after the Publisher's publication.

4. For all Publisher commissioned articles, the right to post a version of the final published version of the Contribution, or any abstract of the final published Contribution on the Contributor(s)'s own and/or his/her institution's website 12 months after publication.

5. The following statement must accompany the articles posted on the Contributor(s)'s and/or his/her institution's website:

"This article has been published in the BMJ [insert full citation reference] and can also be viewed on the journal's website at <u>www.bmj.com</u>"

6. In addition, for Original Research articles and Open Access Funded Articles copyright owners (and the Publishers) may and may allow third parties to use the Contribution in accordance one of the following Creative Commons licences depending on the source of the research funding as per below:

a) where the Original Research article and/or Open Access Funded Articles is not funded by the Wellcome Trust or UK Research Council, the articles may be re-used under the terms of the Creative Commons Attribution-Non Commercial 3.0 Unported (CC BY-NC 3.0) see:

http://creativecommons.org/licenses/by-nc/3.0/

BMJ Author Licence March 2013 doc

R

and

http://creativecommons.org/licenses/by-nc/3.0/legalcode

or any updated versions as determined by the Publisher from time to time.

or

b) where the Original Research article and/or Open Access Funded Articles is funded by the Wellcome Trust or UK Research Council, the Contribution may be re-used under the terms of the Creative Commons Attribution 3.0 Unported Licence (CC BY 3.0) see:

http://creativecommons.org/licenses/by/3.0/

and

http://creativecommons.org/licenses/by/3.0/legalcode

or any updated versions as determined by the Publisher from time to time

subject to ensuring the Publishers and the Journal are referenced (including a full citation) as set out above; all third party rights within all images, diagrams, photograph, other illustrative material or films, not owned by the authors or BMJ Group are cleared independently and appropriately; and all the Publishers trademarks are removed from any derivative works and ensuring any translations, for which a prior translation agreement with BMJ Group has not been established, must prominently display the statement: "This is an unofficial translation of an article that appeared in a BMJ Group publication. BMJ Group has not endorsed this translation".

7. The right to publish with the necessary acknowledgement of the Publishers and the Journal, all or part of the material from the published Contribution in a book, essay, position paper, or other non peer reviewed publication authored or edited by the Contributor(s)'s (which may be a Commercial Use). This does not apply to multiple Contributions in the same journal, for which permission from the Publishers must be sought.

8. The right to use selected figures and tables and (of which the author or his employer owns or has licensed) and selected text (up to 300 words) from the Contribution for incorporation within another work published in print or digital format by a third party, so long as full credit is given to the Publishers and use of the parts of the Contribution is non Commercial Use.

9. Subject to it not being contrary to English law to do so (such as for example where the UK has trading or other bans with the country of the Corresponding Authors origin or certain groups of people within), the right to receive a royalty for up to 5 years from publication of 10% of any net receipts less sales commission on single orders in excess of £2000 received by the Publisher for any single Contribution reprint or translation sales to a single third party, subject however to any fee being determined (if charged) at the absolute discretion of the Publishers as may be altered from time to time. If the Publishers receive such an order for reprint sales of the Contribution to find out to whom payment should be made. Corresponding Authors have the responsibility to ensure that all authors have agreed what should be done with any such royalty payment and to keep the Publisher updated with current contact details.

For permission to use materials that are beyond permitted here, visit http://www.bmj.com/aboutbmj/resources-readers/permissions

"Commercial Use" includes:

- copying or downloading of documents, or linking to such postings, for further redistribution, sale or licensing, for a fee;
- copying, downloading or posting by a site or service that incorporates advertising with such content;
- the inclusion or incorporation of document content in other works or services (other than for legally permitted quotations with an appropriate citation) that is then available for sale or licensing, for a fee.
- use of documents or document content (other than for legally permitted quotations with appropriate citations) by organisations for any promotional or advertising

BMJ Author Licence March 2013 doc

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

2

9

10

11

12

13

14

15

16

17

18 19

20

21

22

23

24

25

26

27

28

29

30

31

32

33

34

35

36 37

38

39

40

41

42

43

44

45

46

47

48

49

50

51 52 53

purposes whether direct or indirect, whether for a fee or otherwise. Distribution by or on behalf of pharmaceutical organisations is considered in all cases as Commercial Use:

use for the purposes of monetary reward by means of sale, resale, license, loan, hire transfer or other form of commercial exploitation.

Author warranties

1. The author(s) warrant that: i) they are the sole author(s) of the Contribution which is an original work; ii) the whole or a substantial part of the Contribution has not previously been published; iii) they or their employers are the copyright owners of the Contribution; iv) to the best of their knowledge that the Contribution does not contain anything which is libellous, illegal or infringes any third party's copyright or other rights; v) that they have obtained all necessary written consents for any patient information which is supplied with the Contribution and vi) that they have declared or will accurately declare all competing interests to the Publisher.

Anti Bribery

As a service provider to the BMJ Group, you agree that you shall: (a)comply with all applicable laws, statutes, regulations and codes relating to anti-bribery and anti-corruption including but not limited to the Bribery Act 2010 (Relevant Requirements); b) not engage in any activity, practice or conduct which would constitute an offence under sections 1, 2 or 6 of the Bribery Act 2010 (as amended) if such activity, practice or conduct had been carried out in the UK; (c) comply with any Publisher Ethics and Anti-bribery Policy supplied to you from time to time including as contained as follows (Relevant Policies):

http://group.bmj.com/group/about/corporate/Anti-Bribery%20and%20Corruption%20Policy%20-August%202012.pdf;

(d) promptly report to the Chief Executive Officer or Chairman of the Publisher any request or demand for any undue financial or other advantage of any kind received by you in connection with the performance of this Agreement; Breach of this Clause shall be deemed a material breach of this Agreement.

Law and Jurisdiction

To the fullest extent permitted by law, this Agreement will be governed by the laws of England and shall be governed and construed in accordance with the laws of England whose courts shall have exclusive jurisdiction, unless as at the date of formation of this Agreement either i) an English judgement could not be enforced in the Corresponding Author's stated country location; or ii) it would take six months or more for the BMJ Group to enforce an English judgement in the Corresponding Author's stated country location, then it is hereby agreed that this Agreement shall be governed by the laws of the Corresponding Author's stated country (or state if applicable) and their courts shall have jurisdiction. Notwithstanding any of the above, this clause is governed by the laws of England.

The following statement must be included in your manuscript, together with the relevant tick box line below:

"I [insert full name] The Corresponding Author of this article contained within the original manuscript which includes any diagrams & photographs within and any related or stand alone film submitted (the Contribution") has the right to grant on behalf

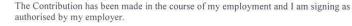
of all authors and does grant on behalf of all authors, a licence to the BMJ Publishing Group Ltd and its licencees, to permit this Contribution (if accepted) to be published in the BMJ and any other BMJ Group products and to exploit all subsidiary rights, as set out in our licence set out at: http://www.bmj.com/about-bmj/resources-authors/forms-policies-and-checklists/copyright-openaccess-and-permission-reuse.'

IF YOU ARE A NATIONAL INSTITUTE OF HEALTH ("NIH") EMPLOYEE, CONTRACTOR OR TRAINEE the following cover sheet will be accepted by the BMJ Group and NIH and incorporated into the above Licence.

Please tick one or more boxes as appropriate:

I am the sole author of the Contribution. ×

I am one author signing on behalf of all co-owners of the Contribution.



BMJ Author Licence March 2013 doc

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

I am a US Federal Government employee acting in the course of my employment.
 I am not a US Federal Government employee, but some or all of my co-authors are.
 I am an employee of the UK Crown* acting in the course of my employment
 I am a US Federal Government employee acting in the course of my employment.
 I am not a US Federal Government employee, but some or all of my co-authors are.
 I am not a US Federal Government employee, but some or all of my co-authors are.
 I am not a US Federal Government employee, but some or all of my co-authors are.
 I am not an employee of the UK Crown acting in the course of my employment but some/all of my co-authors are.*

*Such authors should consult guidance and if necessary return any completed form.

HOPITAL CAN VERDIER UE DU 11 JUIL ET. VS143 BONDY CEDEX SRUUE CU DY P VALENSI SRUUE CU DY P VALENSI ieur des Unwersites - Praticien Hospitalier AVENUEDO ENDOCRINOL OGIE-DI Professeur des Universités - Praticien Hospitalier

BMJ Open

References

- 1. Clarke R, Emberson J, Fletcher A, Breeze E, Marmot M, Shipley MJ. Life expectancy in relation to cardiovascular risk factors: 38 year follow-up of 19,000 men in the Whitehall study. *BMJ* 2009;339:b3513.
- 2. Chandola T, Ferrie J, Sacker A, Marmot M. Social inequalities in self reported health in early old age: follow-up of prospective cohort study. *BMJ* 2007;334(7601):990.
- Kivimaki M, Virtanen M, Kawachi I, Nyberg ST, Alfredsson L, Batty GD, et al. Long working hours, socioeconomic status, and the risk of incident type 2 diabetes: a metaanalysis of published and unpublished data from 222 120 individuals. *Lancet Diabetes Endocrinol* 2014.
- 4. Stringhini S, Tabak AG, Akbaraly TN, Sabia S, Shipley MJ, Marmot MG, et al. Contribution of modifiable risk factors to social inequalities in type 2 diabetes: prospective Whitehall II cohort study. *BMJ* 2012;345:e5452.
- 5. American Diabetes Association. Standards of medical care in diabetes--2014. *Diabetes Care* 2014;37 Suppl 1:S14-80.
- 6. Bihan H, Ramentol M, Fysekidis M, Auclair C, Gerbaud L, Desbiez F, et al. Screening for deprivation using the EPICES score: a tool for detecting patients at high risk of diabetic complications and poor quality of life. *Diabetes Metab* 2012;38(1):82-5.
- 7. Bihan H, Laurent S, Sass C, Nguyen G, Huot C, Moulin JJ, et al. Association among individual deprivation, glycemic control, and diabetes complications: the EPICES score. *Diabetes Care* 2005;28(11):2680-5.
- 8. Brown AF, Ettner SL, Piette J, Weinberger M, Gregg E, Shapiro MF, et al. Socioeconomic position and health among persons with diabetes mellitus: a conceptual framework and review of the literature. *Epidemiologic reviews* 2004;26:63-77.
- 9. Wamala S, Merlo J, Bostrom G, Hogstedt C, Agren G. Socioeconomic disadvantage and primary non-adherence with medication in Sweden. *Int J Qual Health Care* 2007;19(3):134-40.
- Chaturvedi N, Jarrett J, Shipley MJ, Fuller JH. Socioeconomic gradient in morbidity and mortality in people with diabetes: cohort study findings from the Whitehall Study and the WHO Multinational Study of Vascular Disease in Diabetes. *Bmj* 1998;316(7125):100-5.
- 11. Nicolucci A, Carinci F, Ciampi A. Stratifying patients at risk of diabetic complications: an integrated look at clinical, socioeconomic, and care-related factors. SID-AMD Italian Study Group for the Implementation of the St. Vincent Declaration. *Diabetes Care* 1998;21(9):1439-44.
- 12. Saydah SH, Imperatore G, Beckles GL. Socioeconomic status and mortality: contribution of health care access and psychological distress among U.S. adults with diagnosed diabetes. *Diabetes Care* 2013;36(1):49-55.
- 13. Booth GL, Bishara P, Lipscombe LL, Shah BR, Feig DS, Bhattacharyya O, et al. Universal drug coverage and socioeconomic disparities in major diabetes outcomes. *Diabetes Care* 2012;35(11):2257-64.
- 14. Gnavi R, Petrelli A, Demaria M, Spadea T, Carta Q, Costa G. Mortality and educational level among diabetic and non-diabetic population in the Turin Longitudinal Study: a 9-year follow-up. *International journal of epidemiology* 2004;33(4):864-71.

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

15. Gayral-Taminh M, Daubisse-Marliac L, Baron M, Maurel G, Reme JM, Grandjean H. [Social and demographic characteristics and perinatal risks for highly deprived mothers]. *J Gynecol Obstet Biol Reprod (Paris)* 2005;34(1 Pt 1):23-32.

- 16. Lejeune VN, Chaplet VM, Carbonne B, Jannet DJ, Milliez JM. Precarity and pregnancy in Paris. *Eur J Obstet Gynecol Reprod Biol* 1999;83(1):27-30.
- 17. Luo ZC, Wilkins R, Kramer MS. Effect of neighbourhood income and maternal education on birth outcomes: a population-based study. *CMAJ* 2006;174(10):1415-20.
- Joseph KS, Liston RM, Dodds L, Dahlgren L, Allen AC. Socioeconomic status and perinatal outcomes in a setting with universal access to essential health care services. *CMAJ* 2007;177(6):583-90.
- 19. Convers M, Langeron A, Sass C, Moulin JJ, Augier A, Varlet MN, et al. [Is the socioeconomic deprivation EPICES score useful in obstetrics?]. *Gynecol Obstet Fertil* 2012;40(4):208-12.
- 20. Brooke OG, Anderson HR, Bland JM, Peacock JL, Stewart CM. Effects on birth weight of smoking, alcohol, caffeine, socioeconomic factors, and psychosocial stress. *BMJ* 1989;298(6676):795-801.
- 21. Sacks DA, Hadden DR, Maresh M, Deerochanawong C, Dyer AR, Metzger BE, et al. Frequency of gestational diabetes mellitus at collaborating centers based on IADPSG consensus panel-recommended criteria: the Hyperglycemia and Adverse Pregnancy Outcome (HAPO) Study. *Diabetes Care* 2012;35(3):526-8.
- 22. Bo S, Menato G, Bardelli C, Lezo A, Signorile A, Repetti E, et al. Low socioeconomic status as a risk factor for gestational diabetes. *Diabetes Metab* 2002;28(2):139-40.
- 23. Hunsberger M, Rosenberg KD, Donatelle RJ. Racial/ethnic disparities in gestational diabetes mellitus: findings from a population-based survey. *Womens Health Issues* 2010;20(5):323-8.
- 24. Bihan H, Cosson E, Khiter C, Vittaz L, Faghfouri F, Leboeuf D, et al. Factors associated with screening for glucose abnormalities after gestational diabetes mellitus: Baseline cohort of the interventional IMPACT study. *Diabetes Metab* 2014;40:151-7.
- 25. Cosson E, Bihan H, Vittaz L, Khiter C, Carbillon L, Faghfouri F, et al. Improving postpartum glucose screening following gestational diabetes mellitus: the IMPACT multicenter initiative. A cohort study. *Diabet Med* 2014:in press.
- 26. Sass C, Gueguen R, Moulin JJ, Abric L, Dauphinot V, Dupre C, et al. [Comparison of the individual deprivation index of the French Health Examination Centres and the administrative definition of deprivation]. *Sante publique (Vandoeuvre-les-Nancy, France)* 2006;18(4):513-22.
- 27. Gestational diabetes. Summary of expert consensus. Diabetes Metab 2010;36(6 Pt 2):695-9.
- 28. Leroy B, Lefort F. [The weight and size of newborn infants at birth]. *Rev Fr Gynecol Obstet* 1971;66(6):391-6.
- 29. Sass C, Belin S, Chatain C, Moulin JJ, Debout M, Duband S. [Social vulnerability is more frequent in victims of interpersonal violence: value of the EPICES score]. *Presse Med* 2009;38(6):881-92.
- 30. Fiche 6.1 Pauvreté, précarité, Observatoire Régional de la Santé 2011.
- 31. Cosson E, Cussac-Pillegand C, Benbara A, Pharisien I, Jaber Y, Banu I, et al. The diagnostic and prognostic performance of a selective screening strategy for gestational diabetes mellitus according to ethnicity in Europe. *J Clin Endocrinol Metab* 2014;99:996-1005.

BMJ Open

		27
32. Pa	k JH, Lee BE, Park HS, Ha EH, Lee SW, Kim YJ. Association between pre-pre	
	body mass index and socioeconomic status and impact on pregnancy outcomes <i>J Obstet Gynaecol Res</i> 2011;37(2):138-45.	in Korea.
33. Bi	aveman PA, Cubbin C, Egerter S, Williams DR, Pamuk E. Socioeconomic dispation health in the United States: what the patterns tell us. <i>American journal of public</i> 2010;100 Suppl 1:S186-96.	
34. B	policy. Annu Rev Public Health 2012;33:7-40.	ss, and
35. O	esen P, Rasmussen S, Kesmodel U. Effect of prepregnancy maternal overweigh obesity on pregnancy outcome. <i>Obstet Gynecol</i> 2011;118(2 Pt 1):305-12.	t and
36. N	zroo JY. The structuring of ethnic inequalities in health: economic position, raci discrimination, and racism. <i>American journal of public health</i> 2003;93(2):277-	
37. Ta	naka M, Jaamaa G, Kaiser M, Hills E, Soim A, Zhu M, et al. Racial disparity in hypertensive disorders of pregnancy in New York State: a 10-year longitudinal population-based study. <i>American journal of public health</i> 2007;97(1):163-70.	
38. Sı	ith GD. Learning to live with complexity: ethnicity, socioeconomic position, an Britain and the United States. <i>American journal of public health</i> 2000;90(11):1	
39. Ja	fiol C, Thomas F, Bean K, Jego B, Danchin N. Impact of socioeconomic status diabetes and cardiovascular risk factors: results of a large French survey. <i>Diabe</i> 2013;39(1):56-62.	
40. La	Rosa E, Le Clesiau H, Valensi P. Metabolic syndrome and psychosocial deprivation collected from a Paris suburb. <i>Diabetes Metab</i> 2008;34(2):155-61.	ation. Data
41. R	smond R. Role of stress in the pathogenesis of the metabolic syndrome. <i>Psychoneuroendocrinology</i> 2005;30(1):1-10.	
42. Pa	Ilweber B, Valensi P, Lindstrom J, Lalic NM, Greaves CJ, McKee M, et al. A E evidence-based guideline for the prevention of type 2 diabetes. <i>Horm Metab Re</i> Suppl 1:S3-36.	
43. B	ack MH, Sacks DA, Xiang AH, Lawrence JM. The relative contribution of preproverweight and obesity, gestational weight gain, and IADPSG-defined gestation diabetes mellitus to fetal overgrowth. <i>Diabetes Care</i> 2013;36(1):56-62.	
44. Bo	wers K, Laughon SK, Kiely M, Brite J, Chen Z, Zhang C. Gestational diabetes, pregnancy obesity and pregnancy weight gain in relation to excess fetal growth variations by race/ethnicity. <i>Diabetologia</i> 2013;56(6):1263-71.	-
45. G	Iden SH, Brown A, Cauley JA, Chin MH, Gary-Webb TL, Kim C, et al. Health in endocrine disorders: biological, clinical, and nonclinical factorsan Endocrin scientific statement. <i>J Clin Endocrinol Metab</i> 2012;97(9):E1579-639.	-
46. Bo	ulot P, Chabbert-Buffet N, d'Ercole C, Floriot M, Fontaine P, Fournier A, et al. multicentric survey of outcome of pregnancy in women with pregestational dia <i>Diabetes Care</i> 2003;26(11):2990-3.	
47. So	nneider S, Hoeft B, Freerksen N, Fischer B, Roehrig S, Yamamoto S, et al. Neor complications and risk factors among women with gestational diabetes mellitus <i>Obstet Gynecol Scand</i> 2011;90(3):231-7.	
48. Bi	an H, Castetbon K, Mejean C, Peneau S, Pelabon L, Jellouli F, et al. Sociodem factors and attitudes toward food affordability and health are associated with fr vegetable consumption in a low-income French population. <i>J Nutr</i> 2010;140(4)	uit and

- 49. Foxcroft KF, Rowlands IJ, Byrne NM, McIntyre HD, Callaway LK. Exercise in obese pregnant women: the role of social factors, lifestyle and pregnancy symptoms. *BMC Pregnancy Childbirth* 2011;11:4.
- 50. Zhang C, Tobias DK, Chavarro JE, Bao W, Wang D, Ley SH, et al. Adherence to healthy lifestyle and risk of gestational diabetes mellitus: prospective cohort study. BMJ 2014;349:g5450. to peet eview only

BMJ Open

Table1: Characteristics of the total cohort of women by psychosocial status

	Total	No psychosocial	Psychosocial	OR [95% CI]	р
		deprivation	deprivation	•[/•/·•-]	r
	n = 994	n = 417	n = 577		
EPICES score, unit	40.1 ± 25.5	15.6 ± 8.2	57.7 ± 18.1		< 0.001
Age, years	33.3 ± 5.2	33.5 ± 5.0	33.2 ± 5.4		NS
Parity, n	2.4 ± 1.3	2.3 ± 1.2	2.6 ± 1.3		< 0.001
Nulliparity (%)	266 (26.8)	123 (29.6)	143 (24.8)		0.093
Body mass index, kg/m ²	27.8 ± 5.4	27.2 ± 5.3	28.2 ± 5.4		< 0.001
Weight status					< 0.01
Normal weight (%)	307 (31.7)	153 (37.4)	154 (27.5)	REF	
Overweight (%)	374 (38.6)	150 (36.7)	224 (40.1)	1.5 [1.1-2.0]	< 0.05
Obesity (%)	287 (29.6)	106 (25.9)	181 (32.4)	1.7 [1.2-2.4]	< 0.005
Waist circumference					<0.01
<80 cm (%)	505 (51.8)	240 (58.3)	265 (47.2)	REF	
80-88 cm (%)	414 (42.5)	154 (37.4)	260 (46.3)	1.5 [1.2 – 2.0]	< 0.01
>88 cm (%)	55 (5.6)	18 (4.4)	37 (6.6)	1.8 [1.03 – 3.36]	< 0.05
Family history of diabetes	545 (55.3)	247 (59.8)	298 (52.0)	0.7 [0.6 – 0.9]	< 0.05
(%)					
Non daily fruits and	336 (33.9)	108 (25.9)	228 (39.7)	1.9 [1.4 – 2.5]	< 0.001
vegetable consumption (%)					
Anti-hypertensive	62 (6.3)	20 (4.8)	42 (7.3)		NS
treatment (%)					
Lipid lowering treatment	8 (0.8)	2 (0.5)	6 (1.1)		NS
(%)					
Smoking (%)	76 (7.7)	36 (8.7)	40 (6.9)		NS
History of GDM (%)	184 (20.6)	71 (18.9)	113 (21.8)		NS
Ethnicity / origin					< 0.001
Europe (%)	229 (23.7)	140 (34.2)	89 (16.0)	REF	
Antilla (%)	19 (2.0)	8 (2.0)	11 (2.0)		NS
North Africa (%)	382 (39.5)	183 (44.7)	199 (35.7)	1.7 [1.2 – 2.4]	< 0.01
Sub Saharan Africa (%)	145 (15.0)	22 (5.4)	122 (22.1)	8.8 [5.2 – 14.9]	< 0.001
Middle East (%)	25 (2.6)	8 (2.0)	17 (3.1)	3.3 [1.4 – 8.1]	< 0.01
India Pakistan (%)	74 (7.7)	26 (6.4)	48 (8.6)	2.9 [1.7 – 5.0]	< 0.001
Asia (%)	92 (9.5)	22 (5.4)	70 (12.6)	5.0 [2.9 -8.7]	< 0.001
Working status (%)	376 (38.1)	212 (53.4)	154 (26.9)	0.3 [0.2 – 0.4]	< 0.001

The data are expressed as n (%) or as the means \pm the SDs.

GDM: gestational diabetes mellitus; EPICES: Evaluation of Precarity and Inequalities in Health

Examination Centers; OR: odds ratio; REF: reference; 95% CI: 95% confidence interval.

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright.

1
2
3
4
5
6
/ 0
0 Q
10
11
12
13
14 15
15 16
17
18
19
20
21
$\begin{array}{c}2\\3\\4\\5\\6\\7\\8\\9\\10\\11\\23\\14\\15\\16\\17\\8\\9\\021\\223\\24\\25\\27\\28\\9\\30\\132\\33\\4\\56\\37\\38\end{array}$
24
25
26
27
28
29 30
31
32
33
34
30 36
37
38
39
40
41
42 43
44
45
46
47
48 49
49 50
50 51
52
53
54
55 56
50 57
57 58
59
60

60

Table 2: Events during pregnancy by psychosocial stat	us
---	----

	Total	No psychosocial deprivation	Psychosocial deprivation	OR [95% CI]	р
	n = 994	n = 417	n = 577		
GDM diagnosis					0.024
<24 weeks gestation (%)	122 (15.1)	41 (12.1)	81 (17.3)	REF	
24-28 weeks gestation (%)	350 (43.3)	141 (41.5)	209 (44.7)	0.8 [0.5 – 1.2]	NS
\geq 28 weeks gestation (%)	336 (41.6)	158 (46.5)	178 (38.0)	0.6 [0.4 -0.9]	0.011
Insulin therapy during	260 (29.4)	80 (21.8)	180 (34.8)	1.9 [1.4 – 2.6]	< 0.001
pregnancy (%)					
GWG, kg	9.9 ± 6.1	9.9 ± 5.7	9.9 ± 6.4		NS
Excessive GWG (%)	265 (27.4)	109 (26.6)	156 (27.9)		NS
Birth weight, kg	3.4 ± 0.6	3.4 ± 0.5	3.4 ± 0.5		NS
Large for gestational age	131 (13.2)	44 (10.6)	87 (15.1)	1.5 [1.02 – 2.2]	0.037
infants (%)					
Birth weight \geq 4000 g (%)	107 (11.7)	39 (10.1)	68 (12.9)		NS
Birth weight \geq 4250 g (%)	42 (4.6)	17 (4.4)	25 (4.7)		NS
Shoulder dystocia (%)	23 (2.3)	5 (1.2)	18 (3.1)	2.7 [0.97 – 7.2]	0.047
Cesarean section (%)	256 (25.8)	104 (24.9)	152 (26.3)		NS
Preeclampsia (%)	18 (1.8)	11 (2.6)	7 (1.2)	0.5 [0.2 – 1.2]	0.096

The data are expressed as n (%) or as the means \pm the SDs.

GDM: gestational diabetes mellitus; GWG: gestational weight gain; OR: odds ratio; REF: reference; 95%

CI: 95% confidence interval.

BMJ Open

Table 3: Factors associated with lar	rge-for-gestational-age infants
--------------------------------------	---------------------------------

	No LGA infant	LGA infant	Univariate analysis	Multivariate analysis	
	n=863	n=131	p	OR [95% CI]*	p *
Age, years	33.3 ± 5.2	33.5 ± 5.2	NS		-
Parity, n	2.4 ± 1.3	2.7 ± 1.2	< 0.01	1.10 [0.93-1.31]	NS
Body mass index, kg/m ²	27.5 ± 5.4	29.8 ± 5.0	< 0.001		-
Obesity (%)	231 (27.4)	56 (44.4)	< 0.001	1.53 [0.998-2.45]	0.06
Family history of diabetes (%)	470 (55.0)	75 (57.3)	NS		-
Non daily fruits and	284 (33.0)	52 (39.7)	NS		-
vegetable consumption (%)					
Smoking (%)	66 (7.7)	10 (7.6)	NS		-
History of GDM (%)	143 (18.7)	41 (31.8)	< 0.001	1.73 [1.09-2.75]	< 0.05
Ethnicity / origin			< 0.05		
Europe (%)	207 (24.8)	22 (16.8)		REF	
Antilla (%)	17 (2.0)	2 (1.0)		0.90 [0.18-4.38]	NS
North Africa (%)	314 (37.6)	68 (51.9)		1.63 [0.93-2.87]	0.09
Sub Saharan Africa (%)	122 (14.6)	23 (17.6)		1.11 [0.54-2.32]	NS
Middle East (%)	24 (2.9)	1 (0.8)		0.32 [0.04-2.55]	NS
India Pakistan (%)	66 (7.9)	8 (6.1)		1.02 [0.40-2.59]	NS
Asia (%)	85 (10.2)	7 (5.3)	•	0.59 [0.22-1.61]	NS
Working (%)	499 (39.0)	41 (31.3)	0.09		-
EPICES score, unit	39.1 ± 25.4	46.5 ± 25.3	0.002	1.12 [1.03-1.20]**	< 0.01
Psychosocial deprivation (%)	490 (56.8)	87 (66.4)	0.037		-
GDM diagnosis			NS		-
<24 gestational weeks (%)	101 (14.9)	21 (16.4)			-
24-28 gestational weeks (%)	290 (42.6)	60 (46.9)	C		-
>28 gestational weeks (%)	289 (42.5)	47 (36.7)			-
GWG, kg	9.7 ± 6.1	10.9 ± 5.8	< 0.05		-
Excessive GWG (%)	205 (24.3)	60 (47.6)	< 0.001	2.34 [1.54-3.55]	< 0.0001
Insulin therapy during pregnancy (%)	210 (27.8)	50 (38.8)	<0.05	1.32 [0.86-2.04]	NS

The data are expressed as n (%) or as the means \pm the SDs

EPICES: Evaluation of Precarity and Inequalities in Health Examination Centers; GDM: gestational diabetes mellitus; GWG: gestational weight gain; LGA: large for gestational age; OR: odds ratio; REF: reference; 95% CI: 95% confidence interval.

*Multivariate analysis considering parity, obesity, personal history of GDM, ethnic origin, EPICES score, excessive GWG during pregnancy and insulin therapy during pregnancy; **per 10 units.

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

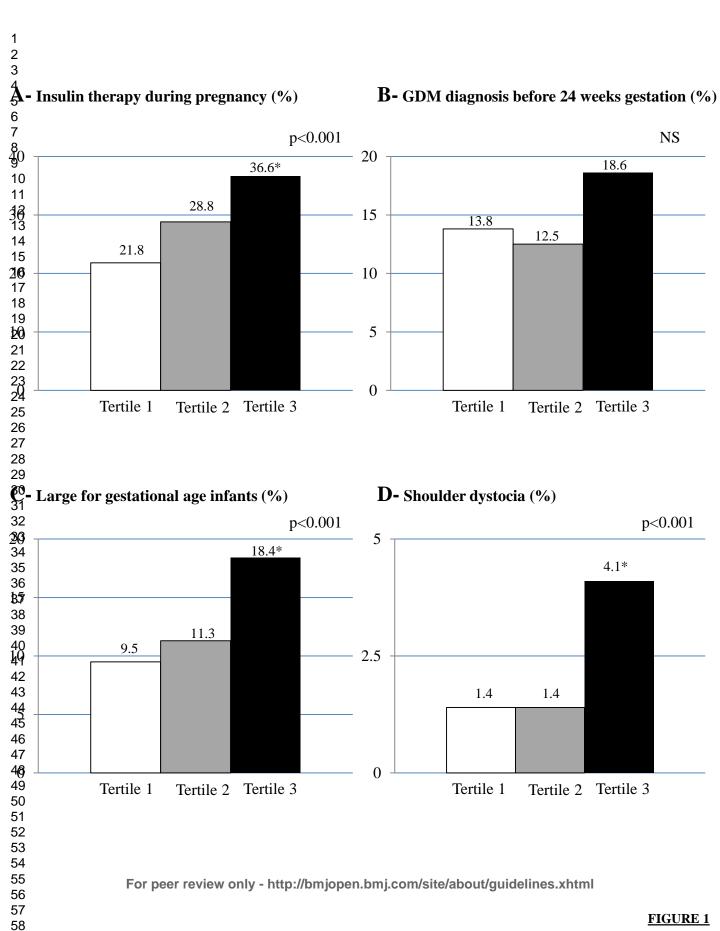
Figure 1: Prevalence of events according to Evaluation of Precarity and Inequalities in Health Examination Centers (EPICES) score tertiles

* p<0.05 versus the first tertile.

GDM: gestational diabetes mellitus; tertile 1: EPICES score <23.71 (mean 11.7±6.2); tertile 2:

EPICES score between 23.71 and 51.5 (mean 35.0 ± 8.5) and tertile 3: EPICES score ≥ 51.5 (mean

69.9±12.6).



Appendix 1 Self-monitoring questionnaire to evaluate deprivation status.

Questions	Coefficient	
1. Do you sometimes meet a social worker (welfare worker, educator)?	10.06	
2. Do you have a complementary health insurance* (mutual insurance)?	-11.83	
3. Do you live as a couple?	-8.28	
4. Are you a homeowner or will you be one in the near future?	-8.28	
5. Are there periods in the month when you have real financial difficulties to face your	14.80	
needs (food, rent, electricity)?		
6. Have you done any sport activities in the last 12 months?	-6.51	
7. Have you gone to any shows (cinema, theatre) over the last 12 months?		
8. Have you gone on holiday over the last 12 months?	-7.10	
9. Have you seen any family member over the last six months (other than your parents or	-9.47	
children)?		
10. If you have difficulties (financial, family or health), is there anyone around you who	-9.47	
could take you in for a few days?		
11. If you have difficulties (financial, family or health), is there anyone around you who		
could help you financially (material aid such as money lending)?		
Intercept	75.14	

EPICES: Evaluation de la Précarité et des Inégalités de santé dans les Centres d'Examen de Santé. Score calculation: each question coefficient is added to intercept whenever the answer is "yes". A score equal to zero corresponds to non-deprivation, a score equal to 100 corresponds to maximum deprivation.

Questions were translated from French to English.

* In France, about 95% of the population is under the general French social security scheme. It gives right to the basic health insurance coverage that reimburses only part of medical expenses. The remainder of the medical cost not reimbursed by the French social security scheme remind on charge of people. Subscription to a complementary private insurance permits to cover partly or completely the percentage of medical costs not paid by the general social security scheme.

STROBE

Psychosocial deprivation in women with gestational diabetes mellitus is associated with poor fetomaternal prognoses: an observational study

	Item No	Recommendation			
Title and abstract					
Page 1		(a) Indicate the study's design with a commonly used term in the title or the abstract			
Page 5	1	(b) Provide in the abstract an informative and balanced summary of what was done and what was found			
Introduction					
Background/rationale Page 7	2	Explain the scientific background and rationale for the investigation being reported			
Objectives Page 7-8	3	State specific objectives, including any prespecified hypotheses			
Methods					
Study design Page 8	4	Present key elements of study design early in the paper			
Setting Page 8-10	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection			
Participants Page 8	6	(a) Cohort study?Give the eligibility criteria, and the sources and methods of selection of participants. Describe methods of follow-upCase-control study?Give the eligibility criteria, and the sources and methods of case ascertainment and control selection. Give the rationale for the choice of cases and controlsCross sectional study?Give the eligibility criteria, and the sources and methods of selection of participants			
		(b) Cohort study?For matched studies, give matching criteria and number of exposed and unexposedCase- control study?For matched studies, give matching criteria and the number of controls per case			
Variables Page 8-10	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable			
Data sources/ measurement Page 8-10	8*	For each variable of interest, give sources of data and details of methods of assessment (measurement). Describe comparability of assessment methods if ther is more than one group			
Bias Page 10-11	9	Describe any efforts to address potential sources of bias			
Study size Page 11	10	Explain how the study size was arrived at			
Quantitative variables Page 11	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why			

	Item No	Recommendation
		(a) Describe all statistical methods, including those used to control for confounding
		(b) Describe any methods used to examine subgroups and interactions
		(c) Explain how missing data were addressed
Statistical methods Page 10-11	12	(d) Cohort study?If applicable, explain how loss to follow-up was addressedCase-control study?If applicable, explain how matching of cases and controls was addressedCross sectional study?If applicable, describe analytical methods taking account of sampling strategy
		(e) Describe any sensitivity analyses
Results		
Participants Page 11	13*	(a) Report numbers of individuals at each stage of study?eg numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study, completing follow-up, and analysed
		(b) Give reasons for non-participation at each stage
		(c) Consider use of a flow diagram
		(a)Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders
Descriptive data Page 11 and Table 1	14*	(b) Indicate number of participants with missing data for each variable of interest
		(c) Cohort study?Summarise follow-up time (eg average and total amount)
		Cohort study?Report numbers of outcome events or summary measures over time
Outcome data Page 11 and Table 2	15*	Case-control study?Report numbers in each exposure category, or summary measures of exposure
		Cross sectional study?Report numbers of outcome events or summary measures
Main results Page 11 and Table 1-	16	(a) Report the numbers of individuals at each stage of the study?eg numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study, completing follow-up, and analysed
3 and figure 1		(b) Give reasons for non-participation at each stage
		(c) Consider use of a flow diagram
Other analyses Non applicable	17	Report other analyses done?eg analyses of subgroups and interactions, and sensitivity analyses
Discussion		
Key results Page 13-15	18	Summarise key results with reference to study objectives
Limitations Page 15	19	Discuss limitations of the study, taking into account sources of potential bias or imprecision. Discuss both direction and magnitude of any potential bias
Interpretation	20	Give a cautious overall interpretation of results

	Item No	Recommendation
Page 13-16		considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence
Generalisability Page 15	21	Discuss the generalisability (external validity) of the study results
Other information		
Funding Page 17	22	Give the source of funding and the role of the funders for the present study and, if applicable, for the original study on which the present article is based

*Give information separately for cases and controls in case-control studies and, if applicable, for exposed and unexposed groups in cohort and cross sectional studies.

Т

BMJ Open

Psychosocial deprivation in women with gestational diabetes mellitus is associated with poor fetomaternal prognoses: an observational study

Journal:	BMJ Open
Manuscript ID:	bmjopen-2014-007120.R1
Article Type:	Research
Date Submitted by the Author:	18-Jan-2015
Complete List of Authors:	Cosson, Emmanuel; AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Endocrinology-Diabetology-Nutrition; Sorbonne Paris Cité, UMR U1153 Inserm / U1125 Inra / Cnam / Université Paris 13, Bihan, Hélène; AP-HP, Avicenne Hospital, Paris 13 University, Sorbonne Paris Cité, Diabetology, Metabolic Diseases; Sorbonne Paris Cité, UMR U1153 Inserm / U1125 Inra / Cnam / Université Paris 13, Reach, GTrard; APHP, Endocrinology Vittaz, Laurence; Ballanger Hospital, Carbillon, Lionel; Assistance Publique Hopitaux de Paris, Paris 13 University, Department of Obstetrics and Gynaecology Valensi, Paul; AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Endocrinology-Diabetology-Nutrition
Primary Subject Heading :	Diabetes and endocrinology
Secondary Subject Heading:	Global health, Obstetrics and gynaecology, Occupational and environmental medicine
Keywords:	Diabetes in pregnancy < DIABETES & ENDOCRINOLOGY, OBSTETRICS, SOCIAL MEDICINE

SCHOLARONE[™] Manuscripts

BMJ Open

Psychosocial deprivation in women with gestational diabetes mellitus is associated with poor fetomaternal prognoses: an observational study

E Cosson *professor*¹², H Bihan *investigator*²³, G Reach *professor*³, L Vittaz *investigator*⁴, L Carbillon *professor*⁵, P Valensi *professor*¹

¹ AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Endocrinology-Diabetology-Nutrition, CRNH-IdF, CINFO, 93143 Bondy, France; ² Sorbonne Paris Cité, UMR U1153 Inserm / U1125 Inra / Cnam / Université Paris 13, 93000 Bobigny, France; ³ AP-HP, Avicenne Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Diabetology, Metabolic Diseases, CRNH-IdF, 93000 Bobigny, France; ⁴ Ballanger Hospital, Department of Endocrinology-Diabetology, 93600 Aulnay-Sous-Bois, France; ⁵ AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Obstetrics and Gynecology, 93143 Bondy, France

Corresponding author:

Emmanuel Cosson

Department of Endocrinology-Diabetology-Nutrition

Avenue du 14 juillet, Hôpital Jean Verdier, 93143 Bondy Cedex. FRANCE

Tel: +33 1 48-02-65-80, Fax: +33 1 48-02-65-79

Sorbonne Paris Cité, UMR U1153 Inserm / U1125 Inra / Cnam / Université Paris 13, 93000 Bobigny, France

e-mail: emmanuel.cosson@jvr.aphp.fr

professor

- Hélène Bihan

² Sorbonne Paris Cité, UMR U1153 Inserm / U1125 Inra / Cnam / Université Paris 13, 93000 Bobigny, France;

³AP-HP, Avicenne Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Diabetology, Metabolic Diseases, CRNH-IdF, 93000 Bobigny, France

helene.bihan@avc.aphp.fr

investigator

- Gérard Reach

³AP-HP, Avicenne Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Diabetology, Metabolic Diseases, CRNH-IdF, 93000 Bobigny, France

gerard.reach @avc.aphp.fr

professor

- Laurence Vittaz

⁴Ballanger Hospital, Department of Endocrinology-Diabetology, 93600 Aulnay-Sous-Bois, France;

laurence.vittaz@ch-aulnay.fr

investigator

- Lionel Carbillon

⁵ AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Obstetrics and Gynecology, 93143 Bondy, France

lionel.carbillon@jvr.aphp.fr

professor

- Paul Valensi

BMJ Open

¹ AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Endocrinology-Diabetology-Nutrition, CRNH-IdF, CINFO, 93143 Bondy, France

paul.valensi@jvr.aphp.fr

professor

ds, main tex. Word counts: abstract 250 words, main text 2800 words

3 tables and 1 figure

1 supplemental file

50 references

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

What is already known on this topic

Psychosocial deprivation is associated with more gestational diabetes mellitus.

What this study adds

Among women with gestational diabetes mellitus, psychosocial deprivation is associated with large for gestational age infants, independently of obesity, gestational weight gain and other confounders.

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

Abstract

Objective To evaluate the prognoses associated with psychosocial deprivation in women with gestational diabetes mellitus (GDM).

Design Observational study considering the 1498 multiethnic women with GDM who gave birth between January 2009 and February 2012.

Setting Four largest maternity units in the northeastern suburban area of Paris.

Participants The 994 women who completed the Evaluation of Precarity and Inequalities in Health Examination Centers [EPICES] questionnaire.

Main outcome measure Main complications of GDM (large infant for gestational age (LGA), shoulder dystocia, cesarean section, preeclampsia).

Results Psychosocial deprivation (EPICES score \geq 30.17) affected 577 women (56%) and was positively associated with overweight/obesity, parity, and non-European origin and negatively associated with family history of diabetes, fruit and vegetable consumption, and working status. The psychosocially deprived women were diagnosed with GDM earlier, received insulin treatment during pregnancy more often and were more likely to have LGA infants (15.1 vs. 10.6%, odds ratio 1.5[95% confidence interval 1.02-2.2], p<0.05) and shoulder dystocia (3.1 vs. 1.2%, OR 2.7[0.97-7.2], p<0.05). In addition to psychosocial deprivation, LGA was associated with greater parity, obesity, history of GDM, ethnicity, excessive gestational weight gain and insulin therapy. A multivariate analysis using these covariates revealed that EPICES score was independently associated with LGA infants (per 10 units, OR 1.12[1.03-1.20], p<0.01).

Conclusions In our area, psychosocial deprivation is common in women with GDM and is associated with earlier GDM diagnoses and greater insulin treatment, an increased likelihood of

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

shoulder dystocia and, independently of obesity, gestational weight gain and other confounders with LGA infants.

Strengths and limitations:

- It is known that psychosocial deprivation is associated with more gestational diabetes and we report for the first time that, among women with gestational diabetes mellitus, psychosocial deprivation is associated with a poor prognosis.
- Our large, multicenter and diverse cohort and the adjustments for the relevant confounding factors, such as body mass index and gestational weight gain, ensure the robustness of our findings.
- We defined psychosocial vulnerability using the Evaluation of Precarity and Inequalities in Health Examination Centers [EPICES] questionnaire, which has been validated during pregnancy or not. This tool evaluates at an individual level several domains, including material goods, money, friendship and family networks, healthcare and leisure.
- However, the EPICES questionnaire was retrospectively fulfilled (6 to 24 months after pregnancy).

Key words: gestational diabetes mellitus, psychosocial deprivation, prognoses, large infant for gestational age

BMJ Open

Introduction

Socioeconomic status reflects access to resources to prosper, and psychosocial deprivation is associated, across countries and over time,¹² with higher mortality and morbidity, including type 2 diabetes.³ The main drivers in more incident type 2 diabetes appear to be higher body mass index (BMI) and impaired health behaviors.⁴ The American Diabetes Association recommends the inclusion of assessments of patients' psychological and social situations as an ongoing part of the medical management of diabetes.⁵ Indeed, psychosocial deprivation in patients with diabetes has been reported to be associated with increased obesity,⁶ worse glycemic control,⁷ poorer adherence,^{8 9} more diabetic complications,^{6 7 10 11} and perhaps greater mortality.¹²⁻¹⁴ During pregnancy, psychosocial deprivation is also associated with poor outcomes that include increased rates of maternal¹⁵ and neonatal^{15 16} hospitalization, stillbirth,¹⁷ postnatal death,¹⁸ preterm delivery^{17 19} and small for gestational age infants.¹⁷⁻²⁰

Gestational diabetes mellitus (GDM) is defined as any degree of glucose intolerance with onset or first recognition during pregnancy and is now very common, with a prevalence ranging from 9.3 in Israel to 25.5% in California, US.²¹ Although GDM is also more frequent in cases of psychosocial deprivation,^{18 19 22 23} its prognosis in case of poor psychological and social conditions is currently unknown. We hypothesized that psychosocial deprivation might be associated with poor prognoses in women with GDM when confounding factors, such as obesity,¹⁹ gestational weight gain (GWG)¹⁶ and smoking habits,²⁰ are considered.

The four largest maternity units in the Northeastern suburban area of Paris, France participated in the IMPACT initiative, which aimed to improve postpartum screening for dysglycemia after GDM.^{24 25} During this study, the women who attended these maternity units responded to the Evaluation of Precarity and Inequalities in Health Examination Centers

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

[EPICES] questionnaire, a questionnaire which evaluates psychosocial deprivation.^{6 7 19 24 26} Therefore, for the first time, we had the opportunity to investigate the fetomaternal prognoses of these women with GDM according to their individual psychosocial statuses.

Methods

Patients

This study is a secondary analysis of the IMPACT study.^{24 25} Briefly, the IMPACT initiative began in March 2011 and was a mobilization campaign for women with GDM and their community caregivers that sought to increase postpartum screening for dysglycemia. We aimed to evaluate the effect of this initiative by comparing the postpartum screening rates between the women who delivered before (between January 2009 and December 2010) and after this initiative (between April 2011 to February 2012). We systematically included women who were at least 18 years of age, free of known pregestational diabetes, had GDM and were followed during pregnancy in one of the four largest maternity units of the Seine-Saint Denis area of France during these periods of time. GDM was detected by oral glucose tolerance test and was defined by fasting blood glucose values ≥ 5.3 mmol/l and/or a 2-hour blood glucose value ≥ 7.8 mmol/l between January 2009 and December 2010,^{24 25} and thereafter according to the International Association of Diabetes and Pregnancy Study Groups criteria, adopted in France in 2010.²⁷ GDM screening was universal in the four centers. In the primary analyses, we included the women who could be contacted by telephone and provided self-reports that indicated whether they had undergone postpartum screening tests during the six months following their deliveries.²⁴ ²⁵ For the current analysis, we included all of the women who delivered between January 2009

BMJ Open

and February 2012 and who retrospectively completed the [EPICES] questionnaire by phone regardless of their report concerning the postpartum screening.

Data collection and assessment of outcomes

One single investigator extracted the following data from hospital records: age at conception, origin/ethnicity, family history of diabetes, history of previous GDM, gestational age at the time of GDM diagnosis (three classes: <24 weeks of gestation, between 24 and 28 weeks of gestation and ≥ 28 weeks of gestation), insulin treatment during pregnancy, and GWG. Excessive GWG was defined according to the recommendations of the Institute of Medicine; *i.e.*, $GWG \ge 16$ kg in women with pregravid BMIs $< 25 \text{ kg/m}^2$, $\geq 11.5 \text{ kg}$ in overweight women (BMIs between 25) and 29.9 kg/m²) and \geq 9 kg in obese women (BMI \geq 30 kg/m²). We also collected obstetrical and neonatal outcomes, including offspring birth weight in comparison to the standard French population (large for gestational age (LGA) was defined by a birth weight exceeding the 90th percentile)²⁸, preeclampsia (blood pressure \geq 140/90 mmHg on two recordings 4 h apart and proteinuria of at least 300 mg/24 h or 3+ or higher upon dipstick testing of a random urine sample), shoulder dystocia (defined as the use of obstetrical maneuvers *i.e.*, McRoberts, episiotomy after delivery of the fetal head, suprapubic pressure, posterior arm rotation to an oblique angle, rotation of the infant by 180 degrees, delivery of the posterior arm and acute or elective cesarian section.

The investigator conducted semi-structured interviews by phone between January and November 2011 for the women who delivered before the IMPACT campaign (maximum delay of time since delivery 24 months) and at least six months after delivery for the women who delivered after the IMPACT initiative. The investigator requested information about the subjects' current weights, heights, waist circumferences, professional statuses, smoking statuses, number

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

of children, antihypertensive and lipid-lowering treatments, family histories of diabetes and daily consumptions of fruits and vegetables. All these data were therefore declarative. Waist circumference was deducted from current waist size of trousers (waist circumference <80 cm: 6-14 (UK) or 34-42 (France), 80-88 cm: 16-20 (UK) or 44-48 (France); >88 cm: 22 (UK) or 48 (France) or more).

The investigator also conducted interviews to assess psychosocial deprivation using the EPICES score, which is a French deprivation score that is calculated based on responses to 11 questions that consider both socioeconomic conditions and family environment (appendix 1).^{7 26} ²⁹ It evaluates at an individual level several domains, including material goods, money, friendship and family networks, healthcare and leisure. As previously reported, the EPICES score is a continuous variable, and increasing quintiles are associated with increased risks for poor health conditions such as obesity, diabetes in women, higher rates of smoking, poorer access to dental and gynecological care, and poorer perceived health statuses.^{24 26} However, psychosocial deprivation can be defined by a score ≥ 30.17 ,²⁹ which was the threshold used here.

Statistical analyses

Sample size calculations were based on the main criterion of the IMPACT study, *i.e.* a postpartum screening test performed six months following delivery.^{24 25} Results reported in this manuscript were prespecified, exploratory endpoints. Continuous variables are expressed as means \pm SD, and normality was assessed with Kolmogorov-Smirnov tests. There were no missing data concerning psychosocial deprivation and main outcomes. Comparisons of two independent groups were performed using the Student t-test if the variable was normally distributed; otherwise, the Wilcoxon Mann-Whitney test was used. The significance of

differences in proportions (*i.e.*, qualitative variables) were tested with the x^2 test, and the odds ratios (ORs) and 95% confidence intervals (CIs) were calculated in cases of statistical significance (p<0.05). We defined EPICES score tertiles in our cohort: first tertile: EPICES score <23.71 (mean 11.7±6.2; n=296); second tertile: score between 23.71 and 51.5 (mean 35.0±8.5; n=355); and third tertile: score \geq 51.5 (mean 69.9±12.6; n=343). The factors associated with having a LGA infant were assessed with a univariate logistic regression method. For multivariate analyses, we included all factors that were associated with LGA infants with p values ≤0.05 in the univariate analyses. SAS Statistics version (9.2) (Cary, USA) was used to conduct all statistical analyses.

Results

Characteristics of the women

A total of 1498 women gave birth following GDM between January 2009 and February 2012 in our maternity units. Of these women, 994 retrospectively responded by phone to the EPICES questionnaire. Table 1 illustrates the characteristics of these women. The characteristics of the 994 women who responded to the EPICES questionnaire were similar to those of the 504 who did not respond with the exception of greater daily consumptions of fruits and vegetable (66.1 vs. 59.0%, respectively, p<0.01) and a trend toward being older (33.3 ± 5.2 vs. 32.7 ± 5.5 years, respectively, p=0.06). The EPICES questionnaire could not be completed by the women who could not be reached by phone and those with French language proficiencies that were insufficient for answering the questions.

Psychosocial deprivation affected 577 women (56%) and was positively associated with parity, overweight and obesity, greater waist circumference, and non-European origin.

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

Psychosocial deprivation was negatively associated with daily fruit and vegetable consumption, reduced family history of diabetes and working status (Table 1).

Pregnancy outcomes

Table 2 shows that the psychosocially deprived women were more likely to have been diagnosed with GDM prior to 24 weeks of gestation and more likely to have been treated with insulin during pregnancy than the non-psychosocially deprived women. The psychosocially deprived women were more likely to have LGA infants and infants with shoulder dystocia, but no differences in cesarean section or preeclampsia were found. Figure 1 (panels A to D) shows that the prevalences of insulin treatment during pregnancy (Figure 1A), LGA infants (Fig 1C) and shoulder dystocia (Figure 1D) increased with increasing EPICES score tertile.

Table 3 shows that, in addition to psychosocial deprivation (OR 1.5 [95%CI 1.02-2.22]), LGA was associated with higher parity, greater BMI and obesity of the mother (OR 2.1 [1.4– 3.1]), increased incidence of GDM history (OR 2.0 [1.4–3.1]), ethnicity/origin, greater EPICES score (per 10 units: OR 1.50 [1.22-2.22]), greater GWG and excessive GWG (OR 2.8 [1.9-4.1]) and insulin treatment during pregnancy (OR 1.6 [1.1–2.4]). A multivariate analysis that considered parity, obesity, personal history of GDM, ethnicity, EPICES score, excessive GWG and insulin therapy during pregnancy revealed that the EPICES score remained independently associated with LGA infants (Table 3). In a model that was identical to the aforementioned model with the exception that weight and height were used in the place of the obesity, an association between psychosocial deprivation and LGA infants remained (per 10 units: OR 1.11 [1.02-1.20, p<0.05). In another model that was identical to the aforementioned model with the exception that psychosocial deprivation (*i.e.*, EPICES score \geq 30.17) was used in the place of the

BMJ Open

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

EPICES score, a trend toward an association between psychosocial deprivation and LGA infants remained (OR 1.53 [0.98-2.39], p=0.06). The prevalence of shoulder dystocia was too low to allow multivariate analyses.

Discussion

In this study, psychosocial deprivation in women with GDM was associated with earlier GDM diagnoses and more extensive insulin treatment. Moreover, we show for the first time that, independent of confounding factors, psychosocial deprivation was associated with increases in adverse outcomes, particularly LGA infants. We report that psychosocial deprivation (*i.e.*, an EPICES score above 30.17) affected 56% of the women with GDM in our study; another study reported a prevalence of 48% (11/23 women with GDM) in another area of France using the same definition of deprivation.¹⁹ This high prevalence is due to the prevalence of precarity³⁰ and multiethnicity³¹ in the Northeastern suburban area of Paris and to the roles played by these conditions in the rate of GDM. Indeed, the prevalence of GDM has been reported to be 1.7- to 2.9-fold higher among patients with high EPICES scores,¹⁹ low educational statuses^{22 23} or low family incomes^{18 23} compared to their counterparts without these conditions. Notably, 23% of pregnant women in France have been reported to have high EPICES scores regardless of GDM status,¹⁹ and 17.5% have been coded as psychosocially deprived by social workers¹⁵ in two other areas in France. Together, our results advocate for screening for deprivation among pregnant women with GDM.

As previously reported for women with and without GDM,^{19 32} we found that psychosocially deprived women with GDM were more likely to be obese. These women were also more likely to be unemployed and less likely to be daily consumers of fruits or vegetables; the latter

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

association is likely due to the cost of these foods. An association between socioeconomic status and healthy eating status, including fruits and vegetable consumption, has previously been reported.^{33 34} We also observed a link between ethnicity/origin and deprivation; similar links have previously been described as complex relationships.³⁵⁻³⁸ The women with and without psychosocial deprivation reported similar prevalences of prepregnancy antihypertensive and lipid-lowering treatments although metabolic disorders are often associated with elevated EPICES scores^{39 40} and stress.⁴¹ The lack of association observed in our study might be specific to women of reproductive age or might be attributable to reduced numbers of medical visits prior to pregnancy due to precarity.¹⁶ The latter supposition would also result in undiagnosed metabolic syndrome prior to pregnancy, which would be in accordance with the greater prevalence of GDM diagnoses prior to 24 weeks of gestation among psychosocially deprived women. These findings suggest the possibility that these women might actually have had undiagnosed pregravid type 2 diabetes. Indeed, precarity is a risk factor for undiagnosed type 2 diabetes even in women of reproductive age.⁴² However, we do not have access to the results of postpartum glycemic assessments that would be needed to confirm this hypothesis.

We also studied the association between psychosocial deprivation and adverse pregnancy outcomes in women with GDM for the first time. Compared to those without precarity, the women with precarity were more likely to have LGA infants and infants with shoulder dystocia. The association between EPICES scores and LGA infants was independent of obesity, which suggests that this relationship was only partially driven by the increased prevalence of overweight/obesity among the deprived women.^{43 44} GWG and the prevalence of excessive GWG, which are other confounding factors regarding LGA infants,^{43 44} were comparable between the women with and without precarity. We have recently shown that, compared to

Page 15 of 33

BMJ Open

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

women from Sub-Saharan Africa, European women experience more GDM-related events.³¹ Furthermore, racial/ethnic differences in the clinical outcomes of GDM, including macrosomia, are commonly reported (for review).⁴⁵ Here, the association of LGA with precarity remained significant after adjusting for origin/ethnicity while we did not find any association between precarity and offspring with birth weights greater than 4000 g or 4250 g. The association psychosocial deprivation. Our large, multicenter and diverse cohort and the adjustments for the

between psychosocial deprivation and shoulder dystocia, which was not adjusted for confounding factors because the rate of dystocia events was low, was most likely driven by the prevalence of LGA infants. In a population-based study, the risk for shoulder dystocia significantly increased with BMI category in an unadjusted analysis, but this significance disappeared after adjusting for GDM.³⁵ As previously reported for pregnant women regardless of GDM status,¹⁵ we found that the women in our cohort with GDM underwent cesarean section at similar rates regardless of the presence of psychosocial deprivation. The vulnerable women were diagnosed with GDM earlier, which suggests that unknown pregravid dysglycemia might partially explain the increased rate of LGA infants.⁴⁶ In a recent German study, the groups that were found to at high risk for GDM were women of low socio-economic status, migrants and obese women. An elevated risk of fetal malformations was found among the women who had been diagnosed with GDM, which suggests that many of these women might have had high glucose levels by the first trimester.⁴⁷ The present study has limits and strengths. The public hospital recruitment and the area we cover probably included a higher proportion of women living with vulnerable conditions, precluding a generalization of our results. On the other hand, we could only include women who could fulfill the EPICES instrument and this may have underestimated the prevalence of

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

relevant confounding factors ensure the robustness of our findings. However, we did not have access to data about glycemic control, diet, physical activity or the numbers of visits during pregnancy. Thus, the adverse outcomes observed for the women with precarity might have been due to these factors based on the following arguments: (i) poor glycemic control has been reported in vulnerable patients with diabetes⁷ and was likely present in our population with GDM and psychosocial insecurity because insulin treatment was more often necessary during GDM among this population; (ii) fruit and vegetable consumption was lower among the vulnerable women following pregnancy, which might be indicative of poorer nutritional habits during pregnancy;^{33 34 48} (iii) exercise during late pregnancy has been reported to vary with the education level of the mother;^{33 34 49} and (vi) access to health care might depend on socioeconomic status,¹⁹ but it is unlikely that access to health care influenced our results because health care is free of charge within the French healthcare system. Compliance may also differ according to psychosocial vulnerability status. Some data were self-reported, such as current weight, height, waist circumference class, professional status, smoking status, number of children, antihypertensive and lipid-lowering treatments, family history of diabetes and daily consumptions of fruits and vegetables. We used the EPICES score, which is an individual index that has been validated during pregnancy¹⁹ and appears to be more strongly linked to the risk of adverse materno-fetal outcomes than neighborhood-level socioeconomic status.¹⁷ However, the EPICES questionnaire was retrospectively fulfilled (6 to 24 months after pregnancy).

Conclusions

To conclude, our results from a large multiethnic multicenter European cohort from an area in which precarity is common demonstrate that psychosocial deprivation affected more than half of the women with GDM. Psychosocial deprivation was associated with higher BMIs and earlier

GDM diagnoses among the vulnerable women, which suggests that GDM likely corresponded to unknown type 2 diabetes mellitus in these women and that prenatal diagnosis of type 2 diabetes should be reinforced in them, with weight control intervention and adherence to healthy lifestyle before pregnancy.⁵⁰ The vulnerable women were also more likely to be treated with insulin, but they gained as much weight during pregnancy as did the non-vulnerable women. Independent of the gestational age at GDM diagnosis, insulin use, overweight/obesity, GWG and other confounders, these women were also more likely to have LGA infants. This finding suggests that the routine screening of women with GDM for psychosocial vulnerability may be an important tool for improving the prognoses of these women and their children. For example, specific follow-up and psychosocial support might be beneficial in these women.

Contributors: EC researched data, directed research and wrote the manuscript; HB researched data and wrote the manuscript; GR directed research and reviewed/edited the manuscript; LV researched data and contributed to discussion; LC researched data and reviewed/edited the manuscript; PV directed research and reviewed/edited the manuscript. All authors contributed to the interpretation of the results and the revision of the manuscript for intellectual content and approved the final version of the manuscript. Delphine Dubois, Umanis, Paris, is the guarantor of this work and, as such, had fully access to all the data in the study and takes the responsibility for the integrity of the data and the accuracy of the data analyses.

Funding: This research was supported by a grant from Novo Nordisk® France. Novo Nordisk was not involved in study design, analyses and interpretation of results and article writing. It

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

provided money to pay the investigator, to perform statistical analyses and for English language editing service.

Competing interest: All authors have completed the ICMJE uniform disclosure form at www.icmje.org/coi_disclosure.pdf . Dr. COSSON reports grants from Novo Nordisk France, during the conduct of the study; personal fees from Novo Nordisk France, outside the submitted work. Dr. Bihan reports personal fees from Novonordisk, during the conduct of the study. Dr Reach reports no conflict of interest in the case of this procedure. Dr. Carbillon reports personal fees from Novo Nordisk France, Dr. VALENSI reports grants from Novo Nordisk France, outside the submitted work is france during the conduct of the study; personal fees from Novo Nordisk France, outside the submitted work.

Ethical approval: The study protocol was approved by the National Ethics Committee (CCTIRS: *Comité Consultatif sur le Traitement de l'Information en Matière de Recherche*; advisory committee on research information processing).

Transparency declaration: The lead author, EC, affirms that the manuscript is an honest, accurate, and transparent account of the study being reported; that no important aspects of the study have been omitted; and that any discrepancies from the study as planned (and, if relevant, registered) have been explained.

Clinical trial registration: observational study.

Data sharing: No additional data available.

Acknowledgments: The authors acknowledge the statistical assistance of Bénédicte Borsik, Delphine Dubois and Anne Ourliac (Umanis, Paris, France – funding source: Novo Nordisk®). The authors also thank Dr Faranaz Faghfouri (AP-HP, Jean Verdier Hospital, Paris 13 University, Sorbonne Paris Cité, Department of Obstetrics and Gynecology, Bondy, France) who was the

study investigator (funding source: Novo Nordisk®); and Drs H Dauphin (Ballanger Hospital, Department of Obstetrics and Gynecology, Aulnay-Sous-Bois, France); Chafika Khiter (De La Fontaine Hospital, Department of Diabetology, Saint Denis, France), Dominique Leboeuf (Seine-Saint-Denis Private Hospital, Department of Obstetrics and Gynecology, Le Blanc Mesnil, France) and Astrid Lepagnol (De La Fontaine Hospital, Department of Obstetrics and Gynecology, Saint Denis, France) who recruited the women. This program was sponsored by the Société Francophone de Diabétologie, the réseau pour la prise en charge et la prévention de l'obésité en pédiatrie 93-Seine-Saint-Denis (REPOP 93), L'Assurance Maladie Seine-Saint-Denis, l'Ordre National Des Pharmaciens and Université Paris 13. BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

References

- 1. Clarke R, Emberson J, Fletcher A, Breeze E, Marmot M, Shipley MJ. Life expectancy in relation to cardiovascular risk factors: 38 year follow-up of 19,000 men in the Whitehall study. *BMJ* 2009;339:b3513.
- 2. Chandola T, Ferrie J, Sacker A, Marmot M. Social inequalities in self reported health in early old age: follow-up of prospective cohort study. *BMJ* 2007;334(7601):990.
- 3. Kivimaki M, Virtanen M, Kawachi I, Nyberg ST, Alfredsson L, Batty GD, et al. Long working hours, socioeconomic status, and the risk of incident type 2 diabetes: a metaanalysis of published and unpublished data from 222 120 individuals. *Lancet Diabetes Endocrinol* 2014.
- 4. Stringhini S, Tabak AG, Akbaraly TN, Sabia S, Shipley MJ, Marmot MG, et al. Contribution of modifiable risk factors to social inequalities in type 2 diabetes: prospective Whitehall II cohort study. *BMJ* 2012;345:e5452.
- 5. American Diabetes Association. Standards of medical care in diabetes-2014. *Diabetes Care* 2014;37 Suppl 1:S14-80.
- Bihan H, Ramentol M, Fysekidis M, Auclair C, Gerbaud L, Desbiez F, et al. Screening for deprivation using the EPICES score: a tool for detecting patients at high risk of diabetic complications and poor quality of life. *Diabetes Metab* 2012;38(1):82-5.
- 7. Bihan H, Laurent S, Sass C, Nguyen G, Huot C, Moulin JJ, et al. Association among individual deprivation, glycemic control, and diabetes complications: the EPICES score. *Diabetes Care* 2005;28(11):2680-5.
- 8. Brown AF, Ettner SL, Piette J, Weinberger M, Gregg E, Shapiro MF, et al. Socioeconomic position and health among persons with diabetes mellitus: a conceptual framework and review of the literature. *Epidemiologic reviews* 2004;26:63-77.
- 9. Wamala S, Merlo J, Bostrom G, Hogstedt C, Agren G. Socioeconomic disadvantage and primary non-adherence with medication in Sweden. *Int J Qual Health Care* 2007;19(3):134-40.
- Chaturvedi N, Jarrett J, Shipley MJ, Fuller JH. Socioeconomic gradient in morbidity and mortality in people with diabetes: cohort study findings from the Whitehall Study and the WHO Multinational Study of Vascular Disease in Diabetes. *Bmj* 1998;316(7125):100-5.
- Nicolucci A, Carinci F, Ciampi A. Stratifying patients at risk of diabetic complications: an integrated look at clinical, socioeconomic, and care-related factors. SID-AMD Italian Study Group for the Implementation of the St. Vincent Declaration. *Diabetes Care* 1998;21(9):1439-44.
- 12. Saydah SH, Imperatore G, Beckles GL. Socioeconomic status and mortality: contribution of health care access and psychological distress among U.S. adults with diagnosed diabetes. *Diabetes Care* 2013;36(1):49-55.
- 13. Booth GL, Bishara P, Lipscombe LL, Shah BR, Feig DS, Bhattacharyya O, et al. Universal drug coverage and socioeconomic disparities in major diabetes outcomes. *Diabetes Care* 2012;35(11):2257-64.

BMJ Open

14. Gnavi R, Petrelli A, Demaria M, Spadea T, Carta Q, Costa G. Mortality and educational level among diabetic and non-diabetic population in the Turin Longitudinal Study: a 9-year
follow-up. International journal of epidemiology 2004;33(4):864-71.
15. Gayral-Taminh M, Daubisse-Marliac L, Baron M, Maurel G, Reme JM, Grandjean H.
[Social and demographic characteristics and perinatal risks for highly deprived mothers]. <i>J Gynecol Obstet Biol Reprod (Paris)</i> 2005;34(1 Pt 1):23-32.
16. Lejeune VN, Chaplet VM, Carbonne B, Jannet DJ, Milliez JM. Precarity and pregnancy in Paris. <i>Eur J Obstet Gynecol Reprod Biol</i> 1999;83(1):27-30.
17. Luo ZC, Wilkins R, Kramer MS. Effect of neighbourhood income and maternal education on birth outcomes: a population-based study. <i>CMAJ</i> 2006;174(10):1415-20.
 Joseph KS, Liston RM, Dodds L, Dahlgren L, Allen AC. Socioeconomic status and perinatal outcomes in a setting with universal access to essential health care services. <i>CMAJ</i> 2007;177(6):583-90.
 Convers M, Langeron A, Sass C, Moulin JJ, Augier A, Varlet MN, et al. [Is the socioeconomic deprivation EPICES score useful in obstetrics?]. <i>Gynecol Obstet Fertil</i> 2012;40(4):208-12.
20. Brooke OG, Anderson HR, Bland JM, Peacock JL, Stewart CM. Effects on birth weight of smoking, alcohol, caffeine, socioeconomic factors, and psychosocial stress. <i>BMJ</i> 1989;298(6676):795-801.
21. Sacks DA, Hadden DR, Maresh M, Deerochanawong C, Dyer AR, Metzger BE, et al. Frequency of gestational diabetes mellitus at collaborating centers based on IADPSG consensus panel-recommended criteria: the Hyperglycemia and Adverse Pregnancy Outcome (HAPO) Study. <i>Diabetes Care</i> 2012;35(3):526-8.
22. Bo S, Menato G, Bardelli C, Lezo A, Signorile A, Repetti E, et al. Low socioeconomic status as a risk factor for gestational diabetes. <i>Diabetes Metab</i> 2002;28(2):139-40.
23. Hunsberger M, Rosenberg KD, Donatelle RJ. Racial/ethnic disparities in gestational diabetes mellitus: findings from a population-based survey. <i>Womens Health Issues</i> 2010;20(5):323-8.
24. Bihan H, Cosson E, Khiter C, Vittaz L, Faghfouri F, Leboeuf D, et al. Factors associated with screening for glucose abnormalities after gestational diabetes mellitus: Baseline cohort of the interventional IMPACT study. <i>Diabetes Metab</i> 2014;40:151-7.
25. Cosson E, Bihan H, Vittaz L, Khiter C, Carbillon L, Faghfouri F, et al. Improving postpartum glucose screening following gestational diabetes mellitus: the IMPACT multicenter initiative. A cohort study. <i>Diabet Med</i> 2014:in press.
26. Sass C, Gueguen R, Moulin JJ, Abric L, Dauphinot V, Dupre C, et al. [Comparison of the individual deprivation index of the French Health Examination Centres and the administrative definition of deprivation]. <i>Sante publique (Vandoeuvre-les-Nancy, France)</i> 2006;18(4):513-22.
 27. Gestational diabetes. Summary of expert consensus. <i>Diabetes Metab</i> 2010;36(6 Pt 2):695-9. 28. Leroy B, Lefort F. [The weight and size of newborn infants at birth]. <i>Rev Fr Gynecol Obstet</i>

- 1971;66(6):391-6.
- 29. Sass C, Belin S, Chatain C, Moulin JJ, Debout M, Duband S. [Social vulnerability is more frequent in victims of interpersonal violence: value of the EPICES score]. *Presse Med* 2009;38(6):881-92.
- 30. Fiche 6.1 Pauvreté, précarité, Observatoire Régional de la Santé 2011.

h	r
7	7

- 31. Cosson E, Cussac-Pillegand C, Benbara A, Pharisien I, Jaber Y, Banu I, et al. The diagnostic and prognostic performance of a selective screening strategy for gestational diabetes mellitus according to ethnicity in Europe. *J Clin Endocrinol Metab* 2014;99:996-1005.
- 32. Park JH, Lee BE, Park HS, Ha EH, Lee SW, Kim YJ. Association between pre-pregnancy body mass index and socioeconomic status and impact on pregnancy outcomes in Korea. *J Obstet Gynaecol Res* 2011;37(2):138-45.
- 33. Braveman PA, Cubbin C, Egerter S, Williams DR, Pamuk E. Socioeconomic disparities in health in the United States: what the patterns tell us. *American journal of public health* 2010;100 Suppl 1:S186-96.
- 34. Bleich SN, Jarlenski MP, Bell CN, LaVeist TA. Health inequalities: trends, progress, and policy. *Annu Rev Public Health* 2012;33:7-40.
- 35. Ovesen P, Rasmussen S, Kesmodel U. Effect of prepregnancy maternal overweight and obesity on pregnancy outcome. *Obstet Gynecol* 2011;118(2 Pt 1):305-12.
- 36. Nazroo JY. The structuring of ethnic inequalities in health: economic position, racial discrimination, and racism. *American journal of public health* 2003;93(2):277-84.
- 37. Tanaka M, Jaamaa G, Kaiser M, Hills E, Soim A, Zhu M, et al. Racial disparity in hypertensive disorders of pregnancy in New York State: a 10-year longitudinal population-based study. *American journal of public health* 2007;97(1):163-70.
- 38. Smith GD. Learning to live with complexity: ethnicity, socioeconomic position, and health in Britain and the United States. *American journal of public health* 2000;90(11):1694-8.
- 39. Jaffiol C, Thomas F, Bean K, Jego B, Danchin N. Impact of socioeconomic status on diabetes and cardiovascular risk factors: results of a large French survey. *Diabetes Metab* 2013;39(1):56-62.
- 40. La Rosa E, Le Clesiau H, Valensi P. Metabolic syndrome and psychosocial deprivation. Data collected from a Paris suburb. *Diabetes Metab* 2008;34(2):155-61.
- 41. Rosmond R. Role of stress in the pathogenesis of the metabolic syndrome. *Psychoneuroendocrinology* 2005;30(1):1-10.
- 42. Paulweber B, Valensi P, Lindstrom J, Lalic NM, Greaves CJ, McKee M, et al. A European evidence-based guideline for the prevention of type 2 diabetes. *Horm Metab Res* 2010;42 Suppl 1:S3-36.
- 43. Black MH, Sacks DA, Xiang AH, Lawrence JM. The relative contribution of prepregnancy overweight and obesity, gestational weight gain, and IADPSG-defined gestational diabetes mellitus to fetal overgrowth. *Diabetes Care* 2013;36(1):56-62.
- 44. Bowers K, Laughon SK, Kiely M, Brite J, Chen Z, Zhang C. Gestational diabetes, prepregnancy obesity and pregnancy weight gain in relation to excess fetal growth: variations by race/ethnicity. *Diabetologia* 2013;56(6):1263-71.
- 45. Golden SH, Brown A, Cauley JA, Chin MH, Gary-Webb TL, Kim C, et al. Health disparities in endocrine disorders: biological, clinical, and nonclinical factors--an Endocrine Society scientific statement. *J Clin Endocrinol Metab* 2012;97(9):E1579-639.
- 46. Boulot P, Chabbert-Buffet N, d'Ercole C, Floriot M, Fontaine P, Fournier A, et al. French multicentric survey of outcome of pregnancy in women with pregestational diabetes. *Diabetes Care* 2003;26(11):2990-3.
- 47. Schneider S, Hoeft B, Freerksen N, Fischer B, Roehrig S, Yamamoto S, et al. Neonatal complications and risk factors among women with gestational diabetes mellitus. *Acta Obstet Gynecol Scand* 2011;90(3):231-7.

BMJ Open

- 48. Bihan H, Castetbon K, Mejean C, Peneau S, Pelabon L, Jellouli F, et al. Sociodemographic factors and attitudes toward food affordability and health are associated with fruit and vegetable consumption in a low-income French population. *J Nutr* 2010;140(4):823-30.
- 49. Foxcroft KF, Rowlands IJ, Byrne NM, McIntyre HD, Callaway LK. Exercise in obese pregnant women: the role of social factors, lifestyle and pregnancy symptoms. *BMC Pregnancy Childbirth* 2011;11:4.
- 50. Zhang C, Tobias DK, Chavarro JE, Bao W, Wang D, Ley SH, et al. Adherence to healthy lifestyle and risk of gestational diabetes mellitus: prospective cohort study. *BMJ* 2014;349:g5450.

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

₽

Total	No psychosocial deprivation	Psychosocial deprivation	OR [95% CI]	р
n = 994	n = 417	n = 577		
0.1 ± 25.5	15.6 ± 8.2	57.7 ± 18.1		< 0.001
3.3 ± 5.2	33.5 ± 5.0	33.2 ± 5.4		NS
2.4 ± 1.3	2.3 ± 1.2	2.6 ± 1.3		< 0.001
66 (26.8)	123 (29.6)	143 (24.8)		0.093
4.3±15.1	72.2±14.5	75.7±155		< 0.001
163±6	163±6	164±7		0.073
7.8 ± 5.4	27.2 ± 5.3	28.2 ± 5.4		< 0.001
(21.7)	152 (27.4)	154 (27.5)	DEE	< 0.01
07(31.7)	153 (37.4)	154 (27.5)	REF	<0.05
74 (38.6)	150 (36.7)	224 (40.1)	1.5 [1.1-2.0]	< 0.05
87 (29.6)	106 (25.9)	181 (32.4)	1.7 [1.2-2.4]	< 0.005
(51.0)	240 (59 2)	265(472)	REF	< 0.01
05 (51.8) 14 (42.5)	240(58.3)	265 (47.2)		< 0.01
55 (5.6)	154 (37.4) 18 (4.4)	260 (46.3) 37 (6.6)	$ \begin{array}{c} 1.5 [1.2 - 2.0] \\ 1.8 [1.03 - 3.36] \end{array} $	< 0.01
45 (55.3)	247 (59.8)	298 (52.0)	0.7 [0.6 - 0.9]	< 0.05
		· · · ·		
6 (33.9)	108 (25.9)	228 (39.7)	1.9 [1.4 – 2.5]	< 0.001
62 (6.3)	20 (4.8)	42 (7.3)		NS
8 (0.8)	2 (0.5)	6 (1.1)		NS
76 (7.7)	36 (8.7)	40 (6.9)		NS
84 (20.6)	71 (18.9)	113 (21.8)		NS
				< 0.001
29 (23.7)	140 (34.2)	89 (16.0)	REF	
19 (2.0)	8 (2.0)	11 (2.0)		NS
82 (39.5)	183 (44.7)	199 (35.7)	1.7 [1.2 – 2.4]	< 0.01
45 (15.0)	22 (5.4)	122 (22.1)	8.8 [5.2 – 14.9]	< 0.001
25 (2.6)	8 (2.0)	17 (3.1)	3.3 [1.4 – 8.1]	< 0.01
74 (7.7)	26 (6.4)	48 (8.6)	2.9 [1.7 – 5.0]	< 0.001
92 (9.5)	22 (5.4)	70 (12.6)	5.0 [2.9 -8.7]	< 0.001
76 (38.1)	212 (53.4)	154 (26.9)	0.3 [0.2 – 0.4]	< 0.001

Table1: Characteristics of the

The data are expressed as n (%) or as the means \pm the SDs.

GDM: gestational diabetes mellitus; EPICES: Evaluation of Precarity and Inequalities in Health Examination Centers; OR: odds ratio; REF: reference; 95% CI: 95% confidence interval.

1

EPICES score, unit

Body mass index, kg/m²

Waist circumference

Non daily fruits and

Anti-hypertensive

treatment (%)

Smoking (%)

Family history of diabetes

vegetable consumption (%)

Lipid lowering treatment

History of GDM (%)

Working status (%)

Ethnicity / origin

Normal weight (%)

Overweight (%)

Obesity (%)

<80 cm (%)

>88 cm(%)

Europe (%)

Antilla (%)

Asia (%)

North Africa (%)

Middle East (%)

India Pakistan (%)

Sub Saharan Africa (%)

80-88 cm (%)

Age, years

Nulliparity (%)

Weight (kg)

Height (cm)

Weight status

(%)

(%)

Parity, n

	Total	No psychosocial deprivation	Psychosocial deprivation	OR [95% CI]	
	n = 994	n = 417	n = 577		
GDM diagnosis					
<24 weeks gestation (%)	122 (15.1)	41 (12.1)	81 (17.3)	REF	
24-28 weeks gestation (%)	350 (43.3)	141 (41.5)	209 (44.7)	0.8 [0.5 – 1.2]	
≥ 28 weeks gestation (%)	336 (41.6)	158 (46.5)	178 (38.0)	0.6 [0.4 -0.9]	
Insulin therapy during	260 (29.4)	80 (21.8)	180 (34.8)	1.9 [1.4 – 2.6]	<
pregnancy (%)					
GWG, kg	9.9 ± 6.1	9.9 ± 5.7	9.9 ± 6.4		
Excessive GWG (%)	265 (27.4)	109 (26.6)	156 (27.9)		
Birth weight, kg	3.4 ± 0.6	3.4 ± 0.5	3.4 ± 0.5		
Large for gestational age	131 (13.2)	44 (10.6)	87 (15.1)	1.5 [1.02 – 2.2]	
infants (%)					
Birth weight \geq 4000 g (%)	107 (11.7)	39 (10.1)	68 (12.9)		
Birth weight \geq 4250 g (%)	42 (4.6)	17 (4.4)	25 (4.7)		
Shoulder dystocia (%)	23 (2.3)	5 (1.2)	18 (3.1)	2.7 [0.97 – 7.2]	
Cesarean section (%)	256 (25.8)	104 (24.9)	152 (26.3)		
Preeclampsia (%)	18 (1.8)	11 (2.6)	7 (1.2)	0.5 [0.2 – 1.2]	

The data are expressed as n (%) or as the means \pm the SDs.

... ouds ratio; REF: re GDM: gestational diabetes mellitus; GWG: gestational weight gain; OR: odds ratio; REF: reference; 95%

CI: 95% confidence interval.

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

Table 3: Factors associated w	No LGA	LGA infant	Univariate	Multivariate an	alveie
	infant	LGA IIIiaiit	analysis		alysis
	n=863	n=131	p	OR [95% CI]*	p*
Age, years	33.3 ± 5.2	33.5 ± 5.2	ŃS		-
Parity, n	2.4 ± 1.3	2.7 ± 1.2	< 0.01	1.10 [0.93-1.31]	NS
Weight (kg)	73.1±14.8	82.1±15.3	< 0.001		
Height (cm)	163±6	167±6	< 0.001		
Body mass index, kg/m ²	27.5 ± 5.4	29.8 ± 5.0	< 0.001		-
Obesity (%)	231 (27.4)	56 (44.4)	< 0.001	1.53 [0.998-2.45]	0.06
Family history of diabetes (%)	470 (55.0)	75 (57.3)	NS	i	-
Non daily fruits and	284 (33.0)	52 (39.7)	NS		-
vegetable consumption (%)					
Smoking (%)	66 (7.7)	10 (7.6)	NS		-
History of GDM (%)	143 (18.7)	41 (31.8)	< 0.001	1.73 [1.09-2.75]	< 0.05
Ethnicity / origin		. ,	< 0.05		
Europe (%)	207 (24.8)	22 (16.8)		REF	
Antilla (%)	17 (2.0)	2 (1.0)		0.90 [0.18-4.38]	NS
North Africa (%)	314 (37.6)	68 (51.9)		1.63 [0.93-2.87]	0.09
Sub Saharan Africa (%)	122 (14.6)	23 (17.6)		1.11 [0.54-2.32]	NS
Middle East (%)	24 (2.9)	1 (0.8)		0.32 [0.04-2.55]	NS
India Pakistan (%)	66 (7.9)	8 (6.1)		1.02 [0.40-2.59]	NS
Asia (%)	85 (10.2)	7 (5.3)	٠	0.59 [0.22-1.61]	NS
Working (%)	499 (39.0)	41 (31.3)	0.09		-
EPICES score, unit	39.1 ± 25.4	46.5 ± 25.3	0.002	1.12 [1.03-1.20]**	< 0.01
Psychosocial deprivation (%)	490 (56.8)	87 (66.4)	0.037	•	-
GDM diagnosis			NS		-
<24 gestational weeks (%)	101 (14.9)	21 (16.4)			-
24-28 gestational weeks (%)	290 (42.6)	60 (46.9)			-
>28 gestational weeks (%)	289 (42.5)	47 (36.7)			-
GWG, kg	9.7 ± 6.1	10.9 ± 5.8	< 0.05		-
Excessive GWG (%)	205 (24.3)	60 (47.6)	< 0.001	2.34 [1.54-3.55]	< 0.0001
Insulin therapy during	210 (27.8)	50 (38.8)	< 0.05	1.32 [0.86-2.04]	NS

The data are expressed as n (%) or as the means \pm the SDs

EPICES: Evaluation of Precarity and Inequalities in Health Examination Centers; GDM: gestational diabetes mellitus; GWG: gestational weight gain; LGA: large for gestational age; OR: odds ratio; REF: reference; 95% CI: 95% confidence interval.

*Multivariate analysis considering parity, obesity, personal history of GDM, ethnic origin, EPICES score, excessive GWG during pregnancy and insulin therapy during pregnancy; **per 10 units.

BMJ Open

Current weights, heights, professional statuses, smoking statuses, number of children, family histories of diabetes and daily consumptions of fruits and vegetables were self-reported.

BMJ Open: first published as 10.1136/bmjopen-2014-007120 on 6 March 2015. Downloaded from http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protected by copyright

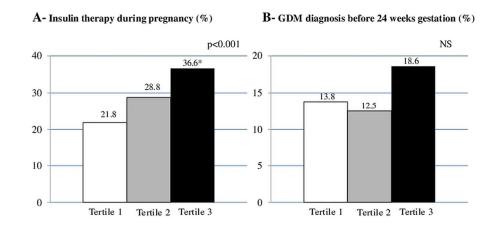
Figure 1: Prevalence of events according to Evaluation of Precarity and Inequalities in Health Examination Centers (EPICES) score tertiles

* p<0.05 versus the first tertile.

GDM: gestational diabetes mellitus; tertile 1: EPICES score <23.71 (mean 11.7±6.2); tertile 2:

EPICES score between 23.71 and 51.5 (mean 35.0 \pm 8.5) and tertile 3: EPICES score \geq 51.5 (mean

69.9±12.6).



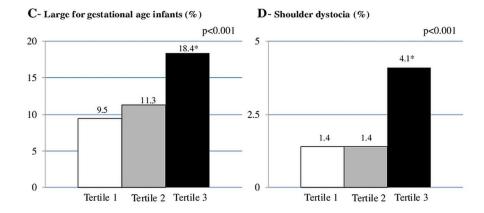


FIGURE 1

90x119mm (300 x 300 DPI)

Appendix 1 <u>Self-monitoring questionnaire to evaluate deprivation status.</u>

Questions	Coefficient
1. Do you sometimes meet a social worker (welfare worker, educator)?	10.06
2. Do you have a complementary health insurance* (mutual insurance)?	-11.83
3. Do you live as a couple?	-8.28
4. Are you a homeowner or will you be one in the near future?	-8.28
5. Are there periods in the month when you have real financial difficulties to face your	14.80
needs (food, rent, electricity)?	
6. Have you done any sport activities in the last 12 months?	-6.51
7. Have you gone to any shows (cinema, theatre) over the last 12 months?	-7.10
8. Have you gone on holiday over the last 12 months?	-7.10
9. Have you seen any family member over the last six months (other than your parents or	-9.47
children)?	
10. If you have difficulties (financial, family or health), is there anyone around you who	-9.47
could take you in for a few days?	
11. If you have difficulties (financial, family or health), is there anyone around you who	-7.10
could help you financially (material aid such as money lending)?	
Intercept	75.14

EPICES: Evaluation de la Précarité et des Inégalités de santé dans les Centres d'Examen de Santé. Score calculation: each question coefficient is added to intercept whenever the answer is "yes". A score equal to zero corresponds to non-deprivation, a score equal to 100 corresponds to maximum deprivation.

Questions were translated from French to English.

* In France, about 95% of the population is under the general French social security scheme. It gives right to the basic health insurance coverage that reimburses only part of medical expenses. The remainder of the medical cost not reimbursed by the French social security scheme remind on charge of people. Subscription to a complementary private insurance permits to cover partly or completely the percentage of medical costs not paid by the general social security scheme.

STROBE

Psychosocial deprivation in women with gestational diabetes mellitus is associated with poor fetomaternal prognoses: an observational study

	Item No	Recommendation
Title and abstract		
Page 1	1	(a) Indicate the study's design with a commonly used term in the title or the abstract
Page 5	Ţ	(b) Provide in the abstract an informative and balanced summary of what was done and what was found
Introduction		
Background/rationale Page 7	2	Explain the scientific background and rationale for the investigation being reported
Objectives Page 7-8	3	State specific objectives, including any prespecified hypotheses
Methods		
Study design Page 8	4	Present key elements of study design early in the paper
Setting Page 8-10	5	Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection
Participants Page 8	6	(a) Cohort study?Give the eligibility criteria, and the sources and methods of selection of participants. Describe methods of follow-upCase-control study?Give the eligibility criteria, and the sources and methods of case ascertainment and control selection. Give the rationale for the choice of cases and controlsCross sectional study?Give the eligibility criteria, and the sources and methods of selection of participants
		(b) Cohort study?For matched studies, give matching criteria and number of exposed and unexposedCase- control study?For matched studies, give matching criteria and the number of controls per case
Variables Page 8-10	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable
Data sources/ measurement Page 8-10	8*	For each variable of interest, give sources of data and details of methods of assessment (measurement). Describe comparability of assessment methods if there is more than one group
Bias Page 10-11	9	Describe any efforts to address potential sources of bias
Study size Page 11	10	Explain how the study size was arrived at
Quantitative variables Page 11	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen and why

₽
Ş
2
8
ĕ
÷
5
Ť
p
Ъ
blished as 10
÷
Ð
0
as `
0.1136/bmj
-
36/b
×
mq
<u>ج</u>
R
õ
Ļ
-2014-0
2
4
6
007120
4
12
õ
0
o
<
larc
5
÷
N
2
5
5 Г
U
ş
own
ownlc
ownloa
ownloade
ownloaded
ownloaded fr
ownloaded fro
<u>_</u>
ownloaded from h
ownloaded from http
ownloaded from http:/
http://
http://bm
http://bm
http://bm
http://bm
http://
http://bm
http://bmjopen.bmj.com/ on A
http://bmjopen.bmj.com/ on Apr
http://bmjopen.bmj.com/ on Apr
http://bmjopen.bmj.com/ on Apr
http://bmjopen.bmj.com/ on April 19,
http://bmjopen.bmj.com/ on April 19, 2024 by guest.
http://bmjopen.bmj.com/ on April 19,
http://bmjopen.bmj.com/ on April 19, 2024 by guest.
http://bmjopen.bmj.com/ on April 19, 2024 by guest. Pr
http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protec
http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protecte
http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protec
http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protecte
http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protecte
http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protecte
http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protecte
http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protecte
http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protecte
http://bmjopen.bmj.com/ on April 19, 2024 by guest. Protecte

	Item No	Recommendation
Statistical methods Page 10-11	12	(a) Describe all statistical methods, including those used to control for confounding
		(b) Describe any methods used to examine subgroups and interactions
		(c) Explain how missing data were addressed
		(d) Cohort study? If applicable, explain how loss to follow-up was addressed Case-control study? If applicable, explain how matching of cases and control was addressed Cross sectional study? If applicable, describe analytical methods taking account of samplin strategy
		(e) Describe any sensitivity analyses
Results		
Participants Page 11	13*	(a) Report numbers of individuals at each stage of study?eg numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study, completing follow-up, and analysed
		(b) Give reasons for non-participation at each stage
		(c) Consider use of a flow diagram
Descriptive data Page 11 and Table 1	14*	(a)Give characteristics of study participants (eg demographic, clinical, social) and information on exposures and potential confounders
		(b) Indicate number of participants with missing data for each variable of interest
		(c) Cohort study?Summarise follow-up time (eg average and total amount)
Outcome data Page 11 and Table 2	15*	Cohort study?Report numbers of outcome events or summary measures over time
		Case-control study?Report numbers in each exposure category, or summary measures of exposure
		Cross sectional study?Report numbers of outcome event or summary measures
Main results Page 11 and Table 1- 3 and figure 1	16	(a) Report the numbers of individuals at each stage o the study?eg numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study, completing follow-up, and analysed
		(b) Give reasons for non-participation at each stage
		(<i>c</i>) Consider use of a flow diagram
Other analyses Non applicable	17	Report other analyses done?eg analyses of subgroups and interactions, and sensitivity analyses
Discussion		
Key results Page 13-15	18	Summarise key results with reference to study objectives
Limitations Page 15	19	Discuss limitations of the study, taking into account sources of potential bias or imprecision. Discuss bot direction and magnitude of any potential bias
Interpretation	20	Give a cautious overall interpretation of results

	Item No	Recommendation
Page 13-16		considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence
Generalisability Page 15	21	Discuss the generalisability (external validity) of the study results
Other information		
Funding Page 17	22	Give the source of funding and the role of the funders for the present study and, if applicable, for the original study on which the present article is based

*Give information separately for cases and controls in case-control studies and, if applicable, for exposed and unexposed groups in cohort and cross sectional studies.

Т