



# Impact of Twin Gestation and Fetal Sex on Maternal Risk of Diabetes During and After Pregnancy

Diabetes Care 2016;39:e110–e111 | DOI: 10.2337/dc16-0825

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Pancreatic  $\beta$ -cell dysfunction is the central pathophysiological defect underlying both gestational diabetes mellitus (GDM) and the subsequent postpartum progression to type 2 diabetes mellitus (T2DM) (1). It has recently emerged that, in singleton pregnancies, carrying a boy is associated with poorer maternal  $\beta$ -cell function and hence an increased risk of GDM (1,2). Moreover, women who develop GDM while carrying a girl have a higher risk of postpartum progression to T2DM than those who develop GDM with a boy (3,4), possibly reflecting comparatively poorer  $\beta$ -cell function in the former (as they developed GDM in the absence of the adverse impact of the male fetus). It thus emerges that the sex of the fetus is associated with maternal risk of diabetes during and after a singleton pregnancy. In this context, we sought to evaluate the impact of twin gestation and the sex of both fetuses on maternal risk of diabetes during and after pregnancy.

Using population-based administrative databases, we identified all women in Ontario, Canada, with a live-birth first pregnancy between April 2000 and March 2012. There were 775,707 women with singleton pregnancies and 13,521 women

with twins (31.7% female/female, 36.0% female/male, and 32.3% male/male). The crude rate of GDM per 100 pregnancies was 5.63 in twin gestation and 3.79 with singletons. After adjustment for age, income, and region of residence, the incidence of GDM was higher in twin versus singleton pregnancies (adjusted odds ratio [OR] 1.30 [95% CI 1.21–1.40],  $P < 0.001$ ). In twin gestation, the crude rate of GDM per 100 pregnancies was 5.56 if both fetuses were female, 6.08 if one was male and one was female, and 5.20 if both were male. Upon adjustment for covariates, however, neither male/male (adjusted OR 0.92 [95% CI 0.76–1.11]) nor male/female (adjusted OR 1.02 [95% CI 0.86–1.22]) carried greater risk of GDM than female/female. Among women who developed GDM ( $n = 30,123$ ) followed over median 6 years after delivery, the incidence of postpartum progression to diabetes was 3.86 per 100 patient-years in those with a singleton pregnancy and 2.96 per 100 patient-years in those who had twins (Fig. 1A). After adjustment for age, income, and region of residence, the risk of progression to diabetes was lower in women who had GDM with twins than in those who had GDM with singletons (adjusted hazard ratio 0.76 [95% CI

0.65–0.90],  $P = 0.001$ ). However, among women who developed GDM with twins, the risk of subsequently progressing to diabetes did not differ between those with male/male, male/female, and female/female twins (Fig. 1B).

In summary, this population-based study shows that twin gestation carries an increased risk of GDM but that affected women have a lower risk of postpartum progression to T2DM than women who develop GDM with a singleton pregnancy. The sex of the twins does not appear to affect maternal risk of diabetes either during or after pregnancy. Instead, other factors (such as antepartum insulin resistance [5]) potentially may be driving the higher risk of GDM in twin pregnancy that is coupled with a lesser risk of subsequent postpartum diabetes. Overall, these data suggest that the impact of the twin gestation itself on maternal glucose metabolism supersedes that of fetal sex.

**Funding.** R.R. is supported by a Heart and Stroke Foundation of Ontario Mid-Career Investigator Award, and his research program is supported by an Ontario Ministry of Research and Innovation Early Researcher Award. B.R.S. is supported by a Canadian Institutes of Health Research New Investigator Award. The Institute for Clinical Evaluative Sciences (ICES) is a nonprofit research

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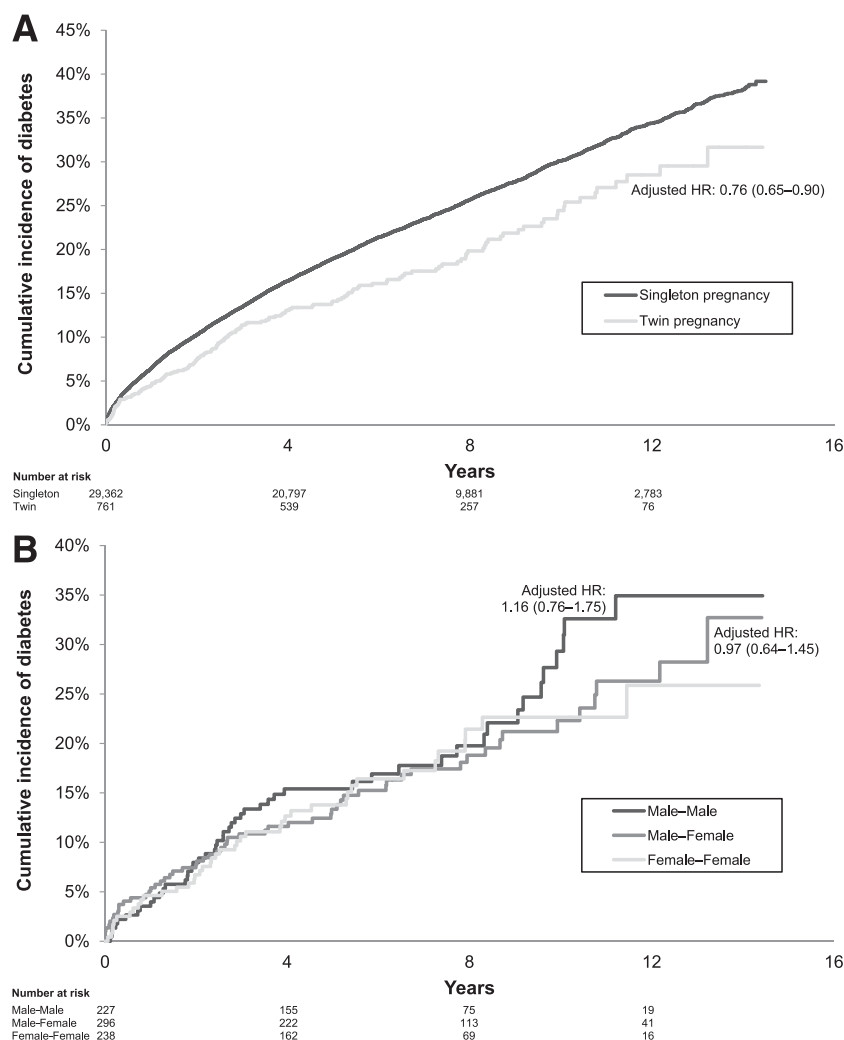
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Received 15 April 2016 and accepted 27 April 2016.

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**Figure 1—A:** Cumulative incidence of diabetes in the years after delivery in women who had GDM with a twin pregnancy vs. women who had GDM in a singleton pregnancy (reference group). **B:** Cumulative incidence of diabetes in the years after delivery in women who had GDM with a twin pregnancy, stratified according to the sex of the twins: male/male, male/female, and female/female (reference group). HR, hazard ratio.

instituted funded by the Ontario Ministry of Health and Long-Term Care (MOHLTC). The study relied on data and information compiled and provided by the Canadian Institute for Health Information (CIHI). The opinions, results, and conclusions reported in this study are those of the authors and are independent from the funding sources. No endorsement by ICES, MOHLTC, or CIHI is intended or should be inferred.

**Duality of Interest.** No potential conflicts of interest relevant to this article were reported.

**Author Contributions.** R.R. conceived of the hypothesis and wrote the manuscript. B.R.S. performed the statistical analyses. Both authors designed the analysis plan, interpreted the data, and critically revised the manuscript for important intellectual content. Both authors approved the final manuscript. B.R.S. is the guarantor of this work and, as such, had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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