Diagnostic thresholds for pregnancy hyperglycemia, maternal weight status and the risk of childhood obesity in a diverse Northern California cohort using health care delivery system data

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Abstract

Objective
To estimate the risk of childhood obesity associated with the various criteria proposed for diagnosis of gestational diabetes (GDM), and the joint effects with maternal BMI.

Methods
Cohort study of 46,396 women delivering at the Kaiser Permanente Northern California health care delivery system in 1995–2004 and their offspring, followed through 5–7 years of age. Pregnancy hyperglycemia was categorized according to the screening and oral glucose tolerance test values proposed for the diagnosis of GDM by the International Association of the Diabetes and Pregnancy Study Group (IADPSG), Carpenter Coustan (CC), and the National Diabetes Data Group (NDDG). Childhood obesity was defined by the International Obesity Task Force’s age and sex-specific BMI cut-offs. Poisson regression models estimated the risks of childhood obesity associated with each category of pregnancy glycaemia compared to normal screening, and the joint effects of maternal BMI category and GDM by the CC and the IADPSG criteria.

Results
Compared with normal screening, increased risks of childhood obesity were observed for abnormal screening [RR (95% CI): 1.30 (1.22, 1.38)], 1+ abnormal values by the IADPSG or CC [1.47 (1.36, 1.59) and 1.48 (1.37, 1.59), respectively], and 2+ values by CC or NDDG [1.52 (1.39, 1.67) and 1.60 (1.43, 1.78), respectively]. Compared to obese women without GDM, obese women with GDM defined by the CC criteria had significantly increased risk of childhood obesity [1.20 (1.07, 1.34)], which was also observed for GDM by the IADSPG.
[1.18 (1.07, 1.30)], though GDM did not significantly increase the risk of childhood obesity among normal weight or overweight women.

Conclusions
The risk of childhood obesity starts to increase at levels of pregnancy glycemia below those used to diagnose GDM and the effect of GDM on childhood obesity risk appears more pronounced in women with obesity. Interventions to reduce obesity and pregnancy hyperglycemia are warranted.

Introduction
In utero exposure to maternal hyperglycemia increases the risk of childhood obesity [1]. Gestational diabetes (GDM), or diabetes first recognized during pregnancy, affects up to 9.2% of pregnancies in the U.S. [2]. Screening for GDM is nearly universal in most clinical settings since its treatments reduces the incidence of several perinatal complications [3, 4]. However, there is a lack of consensus regarding which of several proposed criteria for the identification GDM should be used in clinical practice and an on-going clinical debate on whether reducing maternal overweight and obesity or treating pregnancy hyperglycemia is the more salient public health strategy for preventing childhood obesity. In regard to the criteria for GDM diagnosis, the American Diabetes Association [5] currently recommends either the two-step approach (i.e., screening followed by a diagnostic test) with the Carpenter and Coustan (CC) criteria [6] or the one-step approach with the International Association of the Diabetes and Pregnancy Study Groups (IADPSG) criteria [7], which is recommended by the World Health Organization [8] and uses lower glycemic thresholds than the CC criteria. The American College of Obstetricians and Gynecologists [9] currently supports the two-step approach with either the CC criteria or the less frequently used National Diabetes Data Group (NDDG) criteria [10], which uses the highest glycemic thresholds of all. Although the associations of the CC and NDDG criteria with the risk of childhood obesity have been evaluated [11, 12], no studies have evaluated the risk of childhood obesity across all of the criteria currently used for the diagnosis of GDM. In addition, although maternal overweight and obesity is a well-documented risk factor for both GDM and childhood obesity [13], the degree to which the effect of maternal overweight and obesity on childhood obesity risk is compounded by the presence of pregnancy hyperglycemia remains largely unknown.

This study sought to fill these gaps in the literature by estimating the associations of the various pregnancy glucose criteria and thresholds recommended for the diagnosis of GDM with childhood obesity at 5 to 7 years of age, both overall and across racial-ethnic groups. The study additionally sought to investigate the joint effects of maternal BMI category and pregnancy hyperglycemia on subsequent childhood obesity risk.

Materials and methods
This cohort study took place at Kaiser Permanente Northern California (KPNC), a large integrated health care delivery system with an expansive electronic health record (EHR) system, from 1995 and 2011. KPNC’s membership includes approximately 30% of the geographic area served and is representative of the surrounding population in regards to sociodemographic characteristics, except at the lower extremes of income and education [14].
The results of all blood glucose testing were obtained from the KPNC Gestational Diabetes and Pregnancy Glucose Tolerance Registry [15]; all plasma glucose measurements in this setting are performed using the hexokinase method at the KPNC regional laboratory. Women with recognized pre-gestational diabetes were identified in the KPNC Diabetes Registry [16] and excluded.

In this setting, the two-step approach is used to identify GDM: at 24–28 weeks gestation, women are screened with a 50-g, 1-hour glucose challenge test (> 95% of pregnancies are screened) and those with glucose ≥ 140 mg/dl (7.8 mmol/l) go on to a diagnostic 100-g, 3-hour oral glucose tolerance test (OGTT). During the study period (i.e., 1995–2004), women with ≥ 2 glucose values on the OGTT meeting or exceeding thresholds proposed by the National Diabetes Data Group [10] [NDDG, fasting: 105 mg/dl (5.8 mmol/l), 1-hour: 190 mg/dl (10.5 mmol/l), 2-hour: 165 mg/dl (9.1 mmol/l), and 3-hour: 145 mg/dl (8.0 mmol/l)] were diagnosed with GDM and received treatment. The lower thresholds proposed by Carpenter and Coustan [6] [CC, fasting: 95 mg/dl (5.3 mmol/l), 1-hour: 180 mg/dl (10.1 mmol/l), 2-hour: 155 mg/dl (8.7 mmol/l), and 3-hour: 140 mg/dl (7.8 mmol/l)] were implemented in this setting in 2007, thus women in the current study who met the CC criteria but not the NDDG criteria were not treated for hyperglycemia [17, 18].

The IADPSG [7] criteria were proposed in 2010 and utilize the one-step approach: a single 75-g, 2-hour OGTT with GDM diagnosed if any glucose value meets or exceeds the IADPSG thresholds [fasting: 92 mg/dl (5.1 mmol/l), 1-hour: 180 mg/dl (10.1 mmol/l), and 2-hour: 153 mg/dl (8.5 mmol/l)]. Although the one step procedure is not used in this setting, we assessed the IADPSG thresholds, which are based on a 75-g glucose load, and applied them to the 100-g OGTT to provide conservative estimates of the IADPSG with childhood obesity.

To examine associations with the diagnostic criteria for GDM, women were categorized into the following non-mutually exclusive categories: 1) abnormal screening; 2) abnormal screening and one or more 100 g, 3-hour OGTT values meeting the IADPSG thresholds for fasting, 1-hour or 2-hours; 3) abnormal screening and one or more 100 g, OGTT value meeting the CC thresholds for fasting, 1-hour, 2-hours or 3-hours; 4) abnormal screening and two or more 100 g, OGTT values meeting the CC thresholds for fasting, 1-hour, 2-hours or 3-hours; and 5) abnormal screening and two or more 100 g, OGTT values meeting the NDDG criteria for fasting, 1-hour, 2-hours or 3-hours. These categories, based on the various diagnostic criteria for GDM, were each examined in separate models and women with normal screening values (glucose < 140 mg/dl) served as the reference.

To examine pregnancy hyperglycemia categorized by the glucose thresholds for the fasting, 1-hour and 2-hour time points of the 100-g, 3-h OGTT, women were divided into non-mutually exclusive groups for time point specific analyses (i.e., separate analyses conducted for fasting, 1-hour and 2-hour): 1) abnormal screening; 2) abnormal screening and glucose meeting or exceeding the IADPSG threshold for that time point; 3) abnormal screening and glucose meeting or exceeding the CC threshold for that time point; and 4) abnormal screening and glucose meeting or exceeding the NDDG threshold for that time point. Women with normal screening (< 140 mg/dl) also served as the reference group.

The weights and heights of the children at 5 to 7 years of age were obtained from the EHR system; for children who had multiple measurements during this time frame, the measurements closest to 6 years of age were selected. BMI was calculated as weight (kilograms) divided by the height (meters) squared and the extended International Obesity Task Force’s (IOTF) BMI cut-offs [19] used to identify childhood obesity. Briefly, these age and sex specific cut-offs are based upon predicted adult BMI cut-offs for weight status [19, 20] As such, childhood obesity was defined by the age and sex specific cut-offs for a projected BMI ≥ 30 kg/m² at 18 years of age. Sensitivity analyses utilized the Centers for Disease Control and Prevention’s (CDC)
age and sex specific growth standards [21] to identify childhood obesity (i.e., ≥ 95th percentile).

Maternal age at delivery, parity, height, early pregnancy weight, and gestational age at the pregnancy weight measurement, plus child’s sex and age at the weight and height measurement, were obtained from the EHR. The early pregnancy weight measurement occurred, on average, at 16.9 weeks gestation (SD 1.3) and maternal BMI was classified <18.5 kg/m² (underweight), 18.5–24.9 kg/m² (normal weight), 25.0–29.9 kg/m² (overweight) or ≥30 kg/m² (obese). Data on self-reported race-ethnicity and educational attainment were obtained via linkage with the state of California birth certificate (99% successful linkage [22]).

We identified 48,998 pregnant women, 18–45 years of age, who were screened for GDM and delivered a singleton infant in January 1995-December 2004, and whose child was a KPNC member at 5–7 years of age (i.e., through 2011). If a woman had more than one pregnancy in this period, the first was selected. There were 2,602 women excluded for missing data on early pregnancy weight. The final analytic cohort consisted of 46,396 mother-child pairs.

This study was approved by the Kaiser Foundation Research Institute and the state of California institutional review boards. The human subjects committee of the Kaiser Foundation Research Institute waived the requirement for individual informed consent. All analyses conducted in SAS.

**Statistical analyses**

Poisson regression models with robust standard errors [23] were used to estimate the risks of childhood obesity associated with the categories of pregnancy glycemia. Two approaches were used to categorize and examine pregnancy glycemia. In the first, pregnancy glycemia was categorized into non-mutually exclusive groups based on the diagnostic criteria for GDM: abnormal screening, abnormal screening and 1+ abnormal OGTT values by the IADPSG thresholds (i.e., fasting, 1-hour or 2-hour values only), abnormal screening and 1+ abnormal OGTT value by the CC thresholds (i.e., fasting, 1-hour, 2-hour, or 3-hour values), abnormal screening and 2+ abnormal values by the CC criteria (i.e., fasting, 1-hour, 2-hour, or 3-hour values), and abnormal screening and 2+ abnormal values by the NDDG criteria (i.e., fasting, 1-hour, 2-hour, or 3-hour values). Separate models were constructed to compare each diagnostic criteria category to women with normal screening values.

For the second approach, non-mutually exclusive glucose threshold categories were determined by the time point specific thresholds of the IADPSG, CC and NDDG criteria. Time point specific analyses were then conducted: separate models were run for each glucose threshold category for the fasting, 1-hour and 2-hour time points of the OGTT, and women with normal screening values served as the reference in all models.

Interaction terms for pregnancy glucose, by both classification schemes, with maternal BMI and race-ethnicity were also examined.

The only Table 1 variable to alter risk ratio estimates by 10% or more was maternal BMI; maternal race-ethnicity and age were selected as additional adjustment variables. Multivariable models included maternal pregnancy glycemia, BMI (i.e., <18.5 kg/m², 18.5–24.9 kg/m², 25.0–29.9 kg/m² and ≥30.0 kg/m²), race-ethnicity (i.e., White, Hispanic, Asian, African American, and Other) and age (continuous).

The joint impact of maternal BMI and GDM was also examined, with GDM defined in two ways: (a) meeting the CC criteria for GDM (i.e., 2+ abnormal values by the CC), or (b) 1+ abnormal values by the IADPSG thresholds for fasting, 1-hour, or 2-hours or meeting the full CC criteria for GDM. Women with a BMI < 18.5 kg/m² (underweight) were excluded from these analyses due to the small number of women with GDM (n = 24). A Poisson
regression model was then constructed with the following categories: pregnancy BMI 18.5–24.9 kg/m$^2$ (normal weight) and no GDM (reference); pregnancy BMI 18.5–24.9 kg/m$^2$ and GDM; pregnancy BMI 25.0–29.9 kg/m$^2$ (overweight) and no GDM; pregnancy BMI 25.0–29.9 kg/m$^2$ and GDM; pregnancy BMI $\geq$30.0 kg/m$^2$ (obese) and no GDM; and pregnancy BMI $\geq$30.0 kg/m$^2$ and GDM. To arrive at estimates of the effects of GDM within the overweight and obese strata, the reference group was set to women with overweight and no GDM, and obesity and no GDM, in subsequent models.

Results

Over half of the women had early pregnancy BMI $\geq$25.0 kg/m$^2$ (Table 1). The children’s weight and height measurements at 5 to 7 years of age occurred, on average, at 6.3 years of age (SD = 0.7); 4,900 children (10.6%) were classified as obese according to the International Obesity Task Force’s cut-offs and 7,360 (15.9%) were obese according to the CDC growth standards.

In the unadjusted models evaluating pregnancy glycemia categorized into non-mutually exclusive groups based on the various *diagnostic criteria* for GDM, the risk of childhood obesity by the International Obesity Task Force’s cut-offs was increased for all categories of

| Table 1. Characteristics of the 46,396 women delivering at Kaiser Permanente Northern California in 1995–2004. |
|---------------------------------------------------------|----------------|----------------|
| N = 46,396 | N = 43,476 | N = 2,920 |
| Maternal age at delivery, years |
| 18–24 | 10,381 (22.4) | 10,122 (23.3) | 259 (8.9) |
| 25–29 | 15,161 (32.7) | 14,378 (33.1) | 783 (26.8) |
| 30–34 | 15,336 (33.1) | 14,116 (32.5) | 1,220 (41.8) |
| 35–45 | 5,581 (11.9) | 4,860 (11.2) | 658 (22.5) |
| Maternal race-ethnicity |
| White | 17,254 (37.2) | 16,486 (37.9) | 768 (26.3) |
| Hispanic | 12,740 (27.5) | 11,910 (27.4) | 830 (28.4) |
| Asian | 10,823 (23.3) | 9,751 (22.4) | 1,072 (36.7) |
| African American | 4,051 (8.7) | 3,883 (8.9) | 168 (5.8) |
| Other | 1,528 (3.3) | 1,446 (3.3) | 82 (2.8) |
| Maternal education* |
| Some High School or High School graduate | 15,955 (35.4) | 14,990 (35.5) | 965 (33.8) |
| Some college | 13,424 (29.8) | 12,570 (29.8) | 854 (29.9) |
| College graduate or beyond | 15,719 (34.9) | 14,685 (34.8) | 1,034 (36.2) |
| Parity* |
| 0 | 27,654 (59.6) | 26,061 (60.0) | 1,593 (54.6) |
| 1 | 11,592 (25.0) | 10,848 (25.0) | 744 (25.5) |
| 2+ | 7,134 (15.4) | 6,551 (15.1) | 583 (20.0) |
| Maternal BMI Category, kg/m$^2$ |
| <18.5 | 853 (1.8) | 829 (1.9) | 24 (0.8) |
| 18.5–24.9 | 21,273 (45.9) | 20,371 (46.9) | 902 (30.9) |
| 25.0–29.9 | 14,211 (30.6) | 13,293 (30.6) | 918 (31.4) |
| $\geq$30.0 | 10,059 (21.7) | 8,983 (20.7) | 1,076 (36.9) |

* There were n = 1,298 women missing data on education and n = 16 missing data on parity

GDM: gestational diabetes, defined by either the Carpenter and Coustan or National Diabetes Data Group criteria, BMI: body mass index

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pregnancy hyperglycemia (Table 2). As compared to normal screening, the risk of childhood obesity was statistically significantly elevated among women with abnormal screening [RR = 1.30 (95% CI 1.22, 1.38)]; abnormal screening plus one or more abnormal OGTT values by the IADPSG thresholds [RR = 1.47 (95% CI 1.36, 1.59)]; abnormal screening plus one or more abnormal OGTT values by the CC thresholds [RR = 1.47 (95% CI 1.37, 1.59)]; abnormal screening plus two or more abnormal OGTT values by the CC criteria [RR = 1.52 (95% CI 1.39, 1.67)]; and abnormal screening plus two or more abnormal OGTT values by the NDDG criteria [RR = 1.60 (95% CI 1.43, 1.78); Table 2]. Following adjustment for maternal BMI, race-ethnicity, and age, all risk estimates were attenuated but remained statistically significant (Table 2). Risk estimates were attenuated but remained statistically significant with childhood obesity defined according to the CDC growth standards (S1 Table).

Table 3 shows the risk estimates for pregnancy hyperglycemia categorized into non-mutually exclusive groups based on the glucose thresholds for the fasting, 1-hour and 2-hour time points of the 100-g OGTT. In the unadjusted models, the risk of childhood obesity by the International Obesity Task Force’s cut-offs was significantly increased for women with abnormal screening as compared to normal screening, as well as for all higher degrees of pregnancy hyperglycemia at each time point. For the fasting, 1-hour and 2-hour time points, women with abnormal screening who additionally met any of the time point specific glucose thresholds had significantly increased risks for childhood obesity (RR ranges: 1.86–2.68 for fasting, 1.57–1.61 for 1-hour, and 1.42–1.47 for 2-hour).

Following adjustment for maternal BMI, race-ethnicity, and age, the risk estimates for the glucose threshold categories were all attenuated but remained statistically significant (Table 3). The adjusted risk estimates for the association of fasting glycemia with childhood obesity were 1.27 (95% CI 1.15, 1.40) for those meeting or exceeding the IADPSG threshold (i.e., fasting glucose ≥92 mg/dl); 1.32 (95% CI 1.18, 1.48) for those meeting or exceeding the CC threshold (i.e., fasting glucose ≥95 mg/dl); and 1.59 (95% CI 1.35, 1.86) for those meeting or exceeding the NDDG threshold (i.e., fasting glucose ≥105 mg/dl) compared to normal screening. For 1-hour glycemia, the adjusted risk of childhood obesity remained significantly elevated for those who met the CC/IADPSG threshold for 1-hour glucose [i.e., 1-hour glucose ≥180 mg/dl, RR = 1.23 (95% CI 1.13, 1.35)] or those who met or exceeded the NDDG threshold [i.e.,

| Diagnostic Criteria for GDM | N women | n cases of childhood obesity | Unadjusted | Adjusted |
|----------------------------|---------|-----------------------------|------------|----------|
| Normal screening           | 38,184  | 3,830                       | Reference  | Reference |
| Abnormal screening         | 8,212   | 1,070                       | 1.30 (1.22, 1.38) | 1.13 (1.06, 1.20) |
| Abnormal screening and 1+ abnormal OGTT values by IADPSG | 4,431 | 654 | 1.47 (1.36, 1.59) | 1.18 (1.09, 1.27) |
| Abnormal screening and 1+ abnormal OGTT values by CC | 4,392 | 650 | 1.48 (1.37, 1.59) | 1.19 (1.10, 1.29) |
| Abnormal screening and 2+ abnormal OGTT values by CC | 2,731 | 417 | 1.52 (1.39, 1.67) | 1.20 (1.09, 1.32) |
| Abnormal screening and 2+ abnormal OGTT values by NDDG | 1,825 | 292 | 1.60 (1.43, 1.78) | 1.25 (1.12, 1.39) |

Adjusted for maternal age, race-ethnicity, and BMI category
OGTT: 100g, 3-hr oral glucose tolerance test, IADPSG: International Association of Diabetes in Pregnancy Study Groups, CC: Carpenter and Coustan, NDDG: National Diabetes Data Group
Note that the diagnostic criteria categories are not mutually exclusive, RR estimates obtained from separate models

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For 2-hour glycemia, the adjusted risk of childhood obesity remained significantly elevated among women who met the IADPSG threshold (i.e., 2-hour glucose ≥153 mg/dl; RR = 1.17 (95% CI 1.07, 1.27); the CC threshold [i.e., 2-hour glucose ≥155 mg/dl: RR = 1.20 (95% CI 1.09, 1.31)] or who met or exceeded the NDDG threshold [i.e., 2-hour glucose ≥165 mg/dl: RR = 1.18 (95% CI 1.06, 1.30)]. With childhood obesity defined according to the CDC growth standards, risk estimates were attenuated but remained statistically significant (S2 Table).

The risk of childhood obesity by the International Obesity Task Force’s cut-offs was also increased among women with overweight or obesity, and among African American and Hispanic women. In a crude model with just maternal BMI, the risk of childhood obesity was RR = 2.36 (95% CI 2.19, 2.54) for maternal BMI 25.0–29.9 kg/m² and RR = 4.44 (95% CI 4.14, 4.76) for maternal BMI ≥ 30 kg/m² compared to BMI 18.5–24.9 kg/m²; for maternal BMI < 18.5 kg/m², the risk was RR = 0.43 (95% CI 0.27, 0.68). In a crude model with just maternal race-ethnicity, the risk was RR = 1.92 (95% CI 1.76, 2.11) for African American women, RR = 2.08 (95% CI 1.95, 2.22) for Hispanic women, and RR = 0.99 (95% CI 0.91, 1.08) for Asian women compared to White women. Interaction terms for pregnancy glucose with maternal BMI and race-ethnicity were added to the models presented in Tables 2 and 3. The interaction term for abnormal screening and maternal BMI attained statistical significance (P

### Table 3. Risk Ratio estimates and 95% confidence intervals for the associations of the fasting, 1-hour and 2-hour Glucose Threshold Categories with childhood obesity at 5–7 years of age among 46,396 women delivering at Kaiser Permanente Northern California in 1995–2004.

| Glucose Threshold Categories | N women | n cases of childhood obesity | Unadjusted | Adjusted† |
|-----------------------------|---------|-----------------------------|------------|-----------|
| **Fasting Glucose Thresholds** |         |                             |            |           |
| Normal screening             | 38,184  | 3,830                       | Reference  | Reference |
| Abnormal screening           | 8,212   | 1,070                       | 1.30 (1.22, 1.38) | 1.13 (1.06, 1.20) |
| Abnormal screening and fasting glucose ≥92 mg/dl† | 1,751 | 334 | 1.86 (1.69, 2.06) | 1.27 (1.15, 1.40) |
| Abnormal screening and fasting glucose ≥95 mg/dl‡ | 1,277 | 259 | 2.02 (1.81, 2.26) | 1.32 (1.18, 1.48) |
| Abnormal screening and fasting glucose ≥105 mg/dl§ | 439 | 118 | 2.68 (2.29, 3.14) | 1.59 (1.35, 1.86) |
| **1-hour Glucose Thresholds** |         |                             |            |           |
| Normal screening             | 38,184  | 3,830                       | Reference  | Reference |
| Abnormal screening           | 8,212   | 1,070                       | 1.30 (1.22, 1.38) | 1.13 (1.06, 1.20) |
| Abnormal screening, 1-hour glucose ≥180 mg/dl¶ | 3,044 | 478 | 1.57 (1.43, 1.71) | 1.23 (1.13, 1.35) |
| Abnormal screening, 1-hour glucose ≥190 mg/dl§ | 2,183 | 353 | 1.61 (1.46, 1.78) | 1.24 (1.12, 1.37) |
| **2-hour Glucose Thresholds** |         |                             |            |           |
| Normal screening             | 38,184  | 3,830                       | Reference  | Reference |
| Abnormal screening           | 8,212   | 1,070                       | 1.30 (1.22, 1.39) | 1.13 (1.06, 1.20) |
| Abnormal screening, 2-hour glucose ≥153 mg/dl† | 3,290 | 478 | 1.42 (1.30, 1.55) | 1.17 (1.07, 1.27) |
| Abnormal screening, 2-hour glucose ≥155 mg/dl‡ | 3,134 | 462 | 1.47 (1.34, 1.61) | 1.20 (1.09, 1.31) |
| Abnormal screening, 2-hour glucose ≥165 mg/dl§ | 2,239 | 325 | 1.45 (1.30, 1.61) | 1.18 (1.06, 1.30) |

† Adjusted for maternal age, race-ethnicity, and BMI category
‡ Meeting the Carpenter and Coustan threshold
§ Meeting National Diabetes Data Group threshold
¶ Meeting the International Association of Diabetes in Pregnancy Study Groups/Carpenter and Coustan thresholds, which are identical for the 1-hour time point

IADPSG: International Association of Diabetes in Pregnancy Study Groups, CC: Carpenter and Coustan, NDDG: National Diabetes Data Group

Note that the glucose threshold categories are not mutually exclusive, RR estimates obtained from separate models

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< .10 for BMI categorical and \( P < .01 \) for BMI continuous). In analyses stratified by maternal BMI (underweight and normal weight versus overweight and obese, presented in S3 and S4 Tables), similar findings were observed, though several adjusted risks estimates did not attain statistical significance among the underweight and normal weight women. The interaction between abnormal screening and race-ethnicity also attained statistical significance (\( P < .05 \)).

In analyses stratified by maternal race-ethnicity, similar findings were observed for each racial-ethnic group (S5, S6, S7 and S8 Tables).

Fig 1 displays the risk estimates for the joint effects of maternal BMI category and GDM status defined by the CC criteria (i.e. \( 2+ \) CC abnormal values, Panel A) or expanded to include the IADPSG criteria (i.e. \( 1+ \) IADPSG abnormal value by the fasting, 1-hour, or 2-hour time points or \( 2+ \) CC abnormal values, Panel B). When GDM was defined by the CC criteria alone (n = 2,896 with GDM by the CC criteria alone, Panel A), as compared to women with normal weight and without GDM, those classified as overweight without GDM had an increased risk of childhood obesity \( [RR = 2.27 \ (95\% CI 2.10, 2.45)] \) and the risk increased further for those classified as overweight with GDM \( [RR = 2.62 \ (95\% CI 2.19, 3.12)] \), as obese without GDM \( [RR = 4.17 \ (95\% CI 3.86, 4.49)] \) and as obese with GDM \( [RR = 4.98 \ (95\% CI 4.41, 5.62); \text{Fig 1} \) . The risk of childhood obesity in women with normal weight or overweight and GDM did not differ from that observed among women in these same weight categories without GDM \( [RR = 1.15 \ (95\% CI 0.88, 1.52) \text{ for normal weight, Fig 1, and } RR = 1.15 \ (95\% CI 0.97, 1.37) \text{ for overweight, from a separate model}] \). However, as compared to women with obesity who did not have GDM, those with obesity and GDM had a significantly increased risk of childhood obesity \( [RR = 1.20 \ (95\% CI 1.07, 1.34) \text{ from a separate model}] \).

When the definition of GDM was expanded to include \( 1+ \) abnormal value by the IADPSG thresholds for the fasting, 1-hour or 2-hour time points (n = 4,414 with GDM by the CC criteria or meeting \( 1+ \) IADPSG thresholds, Panel B, Fig 1), similar risk estimates were obtained for each maternal BMI and GDM status subgroup, and the impact of GDM on childhood obesity risk within strata of maternal weight status was unaltered. The risk of childhood obesity in women with normal or overweight did not differ from that observed among women in the same weight category without GDM \( [RR = 1.12 \ (95\% CI 0.97, 1.31) \text{ for normal weight, Fig 1, and } RR = 1.15 \ (95\% CI 0.99, 1.32) \text{ for overweight, from a separate model}] \), though women with obesity and GDM had significantly increased risk of childhood obesity as compared to women with obesity alone \( [RR = 1.18 \ (95\% CI 1.07, 1.30) \text{ for obese, from a separate model}] \).

**Discussion**

The results of this study suggest that the risk of childhood obesity starts to increase with pregnancy hyperglycemia detected by an abnormal screening value alone. The risk among women with a single abnormal value by the IADPSG thresholds was nearly the same as that observed among women with a single abnormal value by the CC thresholds and similar to risk observed among women with two or more abnormal values by the CC criteria, though only the latter are typically diagnosed and treated for GDM. The risk of childhood obesity was further increased among women who met the full NDDG criteria, the only group to receive treatment for GDM in this setting during the study period. These findings suggest that in a clinical setting, an abnormal screening test and/or a single abnormal value by either the IADPSG or the CC thresholds could be used to identify infants at increased risk of childhood obesity. However, it should be noted that the prevalence of GDM would increase by at least 60.6% (22) if a single abnormal value by either the CC or the IADPSG thresholds were used to diagnose GDM. Therefore, the benefits of treating women meeting these lower thresholds must be considered alongside the increased burden on the health care system and cost of treatment.
Following adjustment for maternal age, race-ethnicity and BMI, the risk estimates for childhood obesity were attenuated but remained statistically significant, suggesting that, etiologically, pregnancy hyperglycemia increases the risk of childhood obesity independently of
recognized maternal risk factors. Importantly, similar associations between the pregnancy glucose thresholds and criteria and risk of childhood obesity were observed across all racial-ethnic groups. The risk of childhood obesity increased with increasing maternal BMI and the presence of GDM (whether defined by the CC or IADPSG criteria) further increased the risk among women with obesity. Although there were small numbers of women with GDM among the normal or overweight women, the risk of childhood obesity in those with normal weight or overweight was not significantly increased by the presence of GDM. Taken together, the results of this study suggest that maternal BMI may be the more salient, modifiable risk factor for childhood obesity than GDM.

Consistent with the results of the current study, a follow-up study to the Hyperglycemia and Adverse Pregnancy Outcome (HAPO) study [from which the IADPSG thresholds originated [24]], conducted at 10 of the 15 original HAPO field centers recently reported that the frequency of childhood obesity at 10 to 14 years of age progressively increased across mothers without GDM, those with IADPSG-defined GDM (i.e., meeting the IADPSG but not the CC), and those with CC-defined GDM (i.e., 2 to 3 glucose values at or above the CC thresholds applied to the 75-g OGTT), and that the trend attained statistical significance [25]. As in the current study, GDM defined by the IADPSG criteria was statistically significantly associated with childhood obesity following adjustment for pregnancy BMI [25]. A follow-up study conducted at a single HAPO field center, the Belfast Centre [26], found that fasting pregnancy hyperglycemia defined by the IADPSG threshold was associated with increased risk of childhood obesity in a sample of 1,320 children, 5 to 7 years of age, but the association was no longer significant following adjustment for pregnancy BMI [26]. A follow-up study of 970 ethnic Chinese women conducted at the Hong Kong HAPO field center reported that GDM defined by the IADPSG criteria was significantly associated with offspring overweight or obesity at 7 years of age and that the results remained significant following adjustment for prepregnancy BMI [27]. Yet a large population-based cohort study in Tianjin, China of 27,155 mother-child pairs found that GDM defined by the IADPSG criteria was significantly associated with a higher mean BMI for age $Z$-score through 6 years of age; the difference remained statistically significant at 5 years of age but disappeared for 4 and 6 years of age with adjustment for prepregnancy BMI [28]. These inconsistent findings may be explained by a number of factors, including sample size, population differences (e.g., genetic susceptibility) and/or the use of different methods to assess pregnancy hyperglycemia, alternately, they could reflect true variation of effects over the course of childhood.

Although treatment for GDM bestows benefits in terms of neonatal outcomes (e.g., infant birth weight, macrosomia, and neonatal fat mass [3, 4]), the evidence pertaining to childhood obesity is less clear. The Eunice Kennedy Shriver National Institute of Child Health and Human Development Maternal-Fetal Medicine Units (MFMU) Network randomized clinical trial assessing the effects of treatment on perinatal adverse outcomes among women with mild GDM (i.e., fasting glucose $<95$ mg/dl and two of three timed measurements exceeding the CC thresholds) conducted a follow-study and found no difference in obesity at 5 to 10 years of age by treatment status [29]. In the current study, those who met the NDDG criteria for GDM and received treatment similarly did not show a reduced risk of childhood obesity. However, in a large observational study, Hillier et al. [11] found that women with GDM by the CC criteria who were not treated had an increased odds of childhood obesity at 5 to 7 years of age, but those with GDM by the NDDG criteria who received treatment were not at statistically significant increased risk [i.e., OR = 1.38 (95% CI 0.84, 2.27)]. Additional studies are needed to assess whether treatment of GDM may reduce the risk of childhood obesity.

The current study was conducted in a large, diverse cohort of pregnant women representing the complete range pregnancy glycemia, a clear strength. A limitation of the current study is
the lack of data on gestational weight gain, which is associated with both GDM [30] and child-
hood obesity [31] thus may confound the association of interest. We also assessed the IADPSG
thresholds with a two-step procedure and a 100-g OGTT instead of a 75-g OGTT, as intended
[7], thereby providing conservative estimates of the IADPSG glucose thresholds’ associations
with childhood obesity due to the use of a higher glucose load and the fact that some of the
women with normal screening (our reference group) may have had fasting, 1-hour or 2-hour
glucose values meeting the IADPSG thresholds but could not be captured since they received
only the screening test.

The results of this study suggest that the risk of childhood obesity is present at pregnancy
glycemia levels below those recommended for the diagnosis of GDM and it increases with
increasing level of pregnancy glycemia. The association between pregnancy hyperglycemia
and childhood obesity is in large part explained by maternal overweight and obesity status.
Examination of the joint impact of maternal BMI and GDM revealed that GDM may only
compound the risk of childhood obesity in women with obesity. In terms of the upstream pre-
vention of childhood obesity, interventions that identify women at risk of GDM early in preg-
nancy and aim to reduce pregnancy hyperglycemia as well as interventions that aim to at
reduce obesity among reproductive aged women are warranted.

Supporting information

S1 Table. Risk ratio estimates and 95% Confidence Intervals for associations of the Diag-
nostic Criteria for gestational diabetes with childhood obesity at 5–7 years of age, identi-
fied by the Centers for Disease Control and Prevention’s growth standards, among 46,396
women delivering at Kaiser Permanente Northern California in 1995–2004. Adjusted for
maternal age, race-ethnicity, and BMI category. OGTT: 100g, 3-hr oral glucose tolerance test,
IADPSG: International Association of Diabetes in Pregnancy Study Groups, CC: Carpenter
and Coustan, NDDG: National Diabetes Data Group. Note that the diagnostic criteria catego-
ries are not mutually exclusive, RR estimates obtained from separate models.

S2 Table. Risk ratio estimates and 95% Confidence Intervals for the associations of the
Fasting, 1-hour and 2-hour Glucose Threshold Categories with childhood obesity at 5–7
years of age, identified by the Centers for Disease Control and Prevention’s growth stan-
dards, among 46,396 women delivering at Kaiser Permanente Northern California in
1995–2004. Adjusted for maternal age, race-ethnicity, and BMI category. Meeting the Inter-
national Association of Diabetes in Pregnancy Study Groups threshold. Meeting the Carpen-
ter and Coustan threshold. Meeting National Diabetes Data Group threshold Meeting the
International Association of Diabetes in Pregnancy Study Groups/Carpenter and Coustan
thresholds, which are identical for the 1-hour time point. IADPSG: International Association
of Diabetes in Pregnancy Study Groups, CC: Carpenter and Coustan, NDDG: National Diabe-
tes Data Group. Note that the glucose threshold categories are not mutually exclusive, RR esti-
mates obtained from separate models.

S3 Table. Risk ratio estimates and 95% Confidence Intervals for the associations of the
GDM Diagnostic Criteria and Glucose Threshold Categories with childhood obesity at 5–7
years of age, identified by International Obesity Task Force’s cut-offs, among underweight
and normal weight women (n = 22,126), Kaiser Permanente Northern California, 1995–
2011. Multivariable models include the respective pregnancy glycemia variable, maternal age
and BMI category (<18.5 kg/m² and 18.5–24.9 kg/m²). Meeting the International
Association of Diabetes in Pregnancy Study Groups threshold. § Meeting the Carpenter and Coustan threshold. ¶ Meeting the International Association of Diabetes in Pregnancy Study Groups/Carpenter and Coustan thresholds, which are identical for the 1-hour time point. OGTT: 100g, 3-hr oral glucose tolerance test, IADPSG: International Association of Diabetes in Pregnancy Study Groups, CC: Carpenter and Coustan, NDDG: National Diabetes Data Group, CC: Carpenter and Coustan, NDDG: National Diabetes Data Group, BMI: body mass index Note that glucose categories are not mutually exclusive, RR estimates obtained from separate models.

(DOCX)

S4 Table. Risk ratio estimates and 95% Confidence Intervals for the associations of the GDM Diagnostic Criteria and Glucose Threshold Categories with childhood obesity at 5–7 years of age, identified by International Obesity Task Force’s cut-offs, among women with overweight or obesity (n = 24,270), Kaiser Permanente Northern California, 1995–2011. Multivariable models include the respective pregnancy glycemia variable, maternal age and BMI category (25.0–29.9 kg/m² and ≥30.0 kg/m²) † Meeting the International Association of Diabetes in Pregnancy Study Groups threshold. ² Meeting the Carpenter and Coustan threshold. ³ Meeting National Diabetes Data Group threshold. ¶ Meeting the International Association of Diabetes in Pregnancy Study Groups/Carpenter and Coustan thresholds, which are identical for the 1-hour time point. OGTT: 100g, 3-hr oral glucose tolerance test, IADPSG: International Association of Diabetes in Pregnancy Study Groups, CC: Carpenter and Coustan, NDDG: National Diabetes Data Group, CC: Carpenter and Coustan, NDDG: National Diabetes Data Group, BMI: body mass index. Note that glucose categories are not mutually exclusive, RR estimates obtained from separate models.

(DOCX)

S5 Table. Risk ratio estimates and 95% Confidence Intervals for the associations of the GDM Diagnostic Criteria and Glucose Threshold Categories with childhood obesity at 5–7 years of age, identified by International Obesity Task Force’s cut-offs, among White women (n = 17,254), Kaiser Permanente Northern California, 1995–2011. Multivariable models include the respective pregnancy glycemia variable, maternal age and BMI category (<18.5 kg/m², 18.5–24.9 kg/m², 25.0–29.9 kg/m², and ≥30.0 kg/m²). † Meeting the International Association of Diabetes in Pregnancy Study Groups threshold ² Meeting the Carpenter and Coustan threshold. ³ Meeting National Diabetes Data Group threshold. ¶ Meeting the International Association of Diabetes in Pregnancy Study Groups/Carpenter and Coustan thresholds, which are identical for the 1-hour time point. OGTT: 100g, 3-hr oral glucose tolerance test, IADPSG: International Association of Diabetes in Pregnancy Study Groups, CC: Carpenter and Coustan, NDDG: National Diabetes Data Group, CC: Carpenter and Coustan, NDDG: National Diabetes Data Group, BMI: body mass index. Note that glucose categories are not mutually exclusive, RR estimates obtained from separate models.

(DOCX)

S6 Table. Risk ratio estimates and 95% Confidence Intervals for the associations of the GDM Diagnostic Criteria and Glucose Threshold Categories with childhood obesity at 5–7 years of age, identified by International Obesity Task Force’s cut-offs, among Hispanic women (n = 12,740), Kaiser Permanente Northern California, 1995–2011. Multivariable models include the respective pregnancy glycemia variable, maternal age and BMI category (<24.9 kg/m², 25.0–29.9 kg/m², and ≥30.0 kg/m²). † Meeting the International Association of Diabetes in Pregnancy Study Groups threshold ² Meeting the Carpenter and Coustan threshold. ³ Meeting National Diabetes Data Group threshold. ¶ Meeting the International
Association of Diabetes in Pregnancy Study Groups/Carpenter and Coustan thresholds, which are identical for the 1-hour time point. OGTT: 100g, 3-hr oral glucose tolerance test, IADPSG: International Association of Diabetes in Pregnancy Study Groups, CC: Carpenter and Coustan, NDDG: National Diabetes Data Group, CC: Carpenter and Coustan, NDDG: National Diabetes Data Group, BMI: body mass index. Note that glucose categories are not mutually exclusive, RR estimates obtained from separate models.

S7 Table. Risk ratio estimates and 95% Confidence Intervals for the associations of the GDM Diagnostic Criteria and Glucose Threshold Categories with childhood obesity at 5–7 years of age, identified by International Obesity Task Force’s cut-offs, among Asian women (n = 10,823), Kaiser Permanente Northern California, 1995–2011. Multivariable models include the respective pregnancy glycemia variable, maternal age and BMI category (<18.5 kg/m², 18.5–24.9 kg/m², 25.0–29.9 kg/m², and ≥30.0 kg/m²). † Meeting the International Association of Diabetes in Pregnancy Study Groups threshold. § Meeting the Carpenter and Coustan threshold.

S8 Table. Risk ratio estimates and 95% Confidence Intervals for the associations of the GDM Diagnostic Criteria and Glucose Threshold Categories with childhood obesity at 5–7 years of age, identified by International Obesity Task Force’s cut-offs, among African American women (n = 4,051), Kaiser Permanente Northern California, 1995–2011. Multivariable models include the respective pregnancy glycemia variable, maternal age and BMI category (<18.5 kg/m², 18.5–24.9 kg/m², 25.0–29.9 kg/m², and ≥30.0 kg/m²). † Meeting the International Association of Diabetes in Pregnancy Study Groups threshold. § Meeting the Carpenter and Coustan threshold.

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