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## Differential Nongenetic Impact of Birth Weight Versus Third-Trimester Growth Velocity on Glucose Metabolism and Magnetic Resonance Imaging Abdominal Obesity in Young Healthy Twins

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**Context:** Low birth weight is associated with type 2 diabetes, which to some extent may be mediated via abdominal adiposity and insulin resistance. Fetal growth velocity is high during the third trimester, constituting a potential critical window for organ programming. Intra-pair differences among monozygotic twins are instrumental in determining nongenetic associations between early environment and adult metabolic phenotype.

**Objective:** Our objective was to investigate the relationship between size at birth and third-trimester growth velocity on adult body composition and glucose metabolism using intra-pair differences in young healthy twins.

**Methods:** Fifty-eight healthy twins (42 monozygotic/16 dizygotic) aged 18–24 yr participated. Insulin sensitivity was assessed using hyperinsulinemic-euglycemic clamps. Whole-body fat was assessed by dual-energy x-ray absorptiometry scan, whereas abdominal visceral and sc fat (L1–L4) were assessed by magnetic resonance imaging. Third-trimester growth velocity was determined by repeated ultrasound examinations.

**Results:** Size at birth was nongenetically inversely associated with adult visceral and sc fat accumulation but unrelated to adult insulin action. In contrast, fetal growth velocity during third trimester was not associated with adult visceral or sc fat accumulation. Interestingly, third-trimester growth was associated with insulin action in a paradoxical inverse manner.

**Conclusions:** Abdominal adiposity including accumulation of both sc and visceral fat may constitute primary nongenetic factors associated with low birth weight and reduced fetal growth before the third trimester. Reduced fetal growth during vs. before the third trimester may define distinct adult trajectories of metabolic and anthropometric characteristics influencing risk of developing type 2 diabetes. (*J Clin Endocrinol Metab* 96: 2835–2843, 2011)

There is a well-established association between low birth weight (LBW) and risk of developing insulin resistance and type 2 diabetes (T2D) (1) as well as cardiovascular disease (2, 3). LBW represents the composite end-

point of growth velocities and trajectories during gestation including first, second, and third trimesters. Fetal growth velocity (FGV) is at its maximum in the third trimester, constituting a potentially important critical win-

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Abbreviations: BMI, Body mass index; DXA, dual-energy x-ray absorptiometry; FGV, fetal growth velocity; FPIR, first-phase insulin response; IVGTT, iv glucose tolerance test; LBW, low birth weight; MR, magnetic resonance; MRI, MR imaging; SDS, sd score; T2D, type 2 diabetes.

dow for organ programming. Indeed, third-trimester exposure to famine during the Dutch hunger winter of 1944–1945 was linked to decreased glucose tolerance in adult life (4).

There is compelling evidence for detrimental effects of abdominal obesity on all components in the metabolic syndrome (5, 6). LBW was associated with central obesity in some (7) but not all (8) studies. The inconsistency in the literature may arise from many different factors including age-dependent impact of birth weight on adult body composition, genetic admixture, and the nature and timing of fetal insults as well as the use of indirect measures of abdominal obesity in most studies.

Visceral and sc fat has not previously been assessed directly using the multi-slice magnetic resonance (MR) scanning technology in human study cohorts addressing the developmental origin of obesity and T2D. Furthermore, only few studies in the area included detailed ultrasound measurements of growth velocities throughout third trimester (9).

At least two of the currently more than 40 identified T2D gene variants are associated with reduced birth weight, providing proof of principle for a genetic link between LBW and T2D (10–12). Nevertheless, the vast majority of T2D genes are not associated with weight at birth, and studies of genetically identically monozygotic twins have revealed a significant nongenetic impact of size at birth on T2D and associated metabolic changes in adult life (13, 14). Indeed, intra-pair correlations among genetically identical twins has been documented to represent a very sensitive and powerful tool to detect nongenetic associations between markers of the fetal environment including birth weight on one side and metabolic and anthropometric outcome variables relevant to T2D pathophysiology on the other.

We aimed to investigate the nongenetic relationship between size at birth and third-trimester FGV, respectively, and state of the art measures of adult body composition and glucose metabolism in a unique cohort of young adult twins with extensive ultrasound determinations of FGV during the third trimester.

## Subjects and Methods

### Subjects

Participants were offspring of a cohort of women with twin pregnancies and admitted to Herlev Hospital, Denmark, in the period 1983–1996 ( $n = 457$ ). Gestational age was determined by ultrasound at wk 18 of gestation. Additional ultrasound examinations were performed with 2-wk intervals from wk 28 of gestation until delivery. Fetal weight was calculated from standard biometric ultrasound examinations (15). Coefficients of variation of ultrasound determinations have in singleton preg-

nancies been reported to be in the range of 6.5% and the difference between estimated fetal weight and actual birth weight to be  $-2.2 \pm 8.7\%$  (16). Weight estimates were transformed to SD scores (SDS) using an age- and sex-specific reference of singleton fetal growth (16). FGV in the third trimester was determined by linear regression using a minimum of three examinations (range, 3–8) and was expressed as  $\Delta$ SDS per 28 d. Birth weight SDS was calculated using a large Swedish singleton growth reference (17).

A random sample of 58 young healthy twins (30 male and 28 female) participated, of which a subgroup of eight participants did not participate in the clamp protocol. None received medication known to interfere with glucose homeostasis. Zygosity was determined by comparison of 10 microsatellite DNA polymorphisms showing very high heterozygosity in the population (18). The protocol was approved by the regional ethics committee, and procedures were performed according to the principles of the Helsinki Declaration. After thorough written and oral explanation of the study, all participants gave their written consent.

### Study design

#### ***Hyperinsulinemic-euglycemic clamp combined with stable isotope infusion, iv glucose tolerance test (IVGTT), and indirect calorimetry***

The study procedure was performed as previously described (19). Female participants who did not use oral contraceptives were examined on d 2–5 of their menstrual period. Participants were instructed to consume a diet rich in carbohydrate for 2 d and to abstain from strenuous exercise activity for 24 h before the examination. After an overnight fast, the participants underwent standard blood testing, anthropometric measurements, blood pressure, and a dual-energy x-ray absorptiometry (DXA) scan. At 0730 h, a polyethylene catheter was placed in each antecubital vein for blood sampling and test infusions. One hand was kept in a heated plexiglas box to ensure arterialization of the venous blood. Immediately after taking the background samples, a primed constant infusion of [6,6- $^2$ H5]glucose (priming bolus of 20  $\mu$ mol/kg; continuous infusion rate of 0.220  $\mu$ mol/min  $\cdot$  kg) was started (time 0 min) and maintained for 150 min to determine glucose kinetics in the basal state. Blood samples for measuring plasma glucose enrichments were drawn at baseline (time 0 min) and in the basal steady-state period (time 120–150 min) when the tracer equilibria of [ $^2$ H5]glucose was expected. Isotopes were purchased from Cambridge Isotope Laboratories (Andover, MA). A 30-min IVGTT (time 150–180 min) was performed for the assessment of  $\beta$ -cell function. A glucose bolus of 0.3 g/kg was infused over 1 min, and blood samples for glucose, insulin, and C-peptide were collected at 0, 2, 4, 6, 8, 10, 15, 20, and 30 min. After the IVGTT, a primed-continuous insulin infusion was initiated and fixed at 80 mU/m $^2$   $\cdot$  min through the 180-min clamp (time 180–360 min). The insulin-stimulated steady-state period was defined as the last 30 min of the insulin clamp period. A variable infusion of unlabeled glucose (180 g/liter) was used to maintain euglycemia at 5 mM. Plasma glucose concentration was monitored every 5 min during clamp using a OneTouch (LifeScan, Milpitas, CA) blood glucose meter. Oxygen consumption ( $V_{O_2}$ ) and carbon dioxide production ( $V_{CO_2}$ ) were measured during steady-state using indirect calorimetry with a flow-through canopy gas analyzer system (Deltatrac; Datex, Helsinki, Finland) as previously described (20).

### Biochemical and tracer analyses

Blood samples were centrifuged immediately at 4°C, and plasma samples were stored at –80°C. Plasma insulin and C-peptide concentrations were determined by AutoDELPHIA time-resolved fluoroimmunoassay (PerkinElmer Wallac Oy, Turku, Finland). Stable isotope enrichments were measured as previously described (21).

### Magnetic resonance

T1 weighted MR images were acquired with a 3-T Philips Achieva whole-body scanner. For each patient, between 15 and 26 slices were used to cover the abdominal region bounded by L1 and L4 (pixel spacing  $0.8984 \times 0.8984 \text{ mm}^2$ , slice thickness 7 mm, gap 1 mm). The images were automatically processed to quantify the abdominal adiposity using image analysis techniques (22, 23). The applied method distinguished between superficial sc, deep sc, and visceral fat, reporting the quantities as percentages of total abdominal volume.

### Calculations

#### IVGTT

The area under the curve was calculated using a trapezoidal method for insulin during the first-phase insulin response (FPIR), 0–10 min of the IVGTT.

#### Hyperinsulinemic-euglycemic clamp

For stable isotope tracer calculations, the total rate of glucose appearance was calculated as  $R_{a(\text{endogenous})} = R_d = F_{\text{total}}/E_{\text{glucose}}$ , where  $R_a$  and  $R_d$  are the respective rates of appearance and disappearance (micromoles per kilogram fat-free mass per minute), and  $F_{\text{total}}$  is the total infusion rate of glucose tracer (micromoles per kilogram fat-free mass per minute).  $E_{\text{glucose}}$  is the enrichment of glucose in plasma expressed as tracer-to-tracee ratio. The  $R_a$  of glucose is a measure of endogenous glucose production and represents hepatic glucose production in the basal state (24). To measure insulin action, M-value was calculated as the mean glucose infusion rate during the insulin-stimulated steady-state period (milligrams per kilogram fat-free mass per minute). During the predefined clamp steady-state periods, the coefficients of variation of plasma insulin and glucose levels were 0.11 and 0.06, respectively. Basal and insulin-stimulated glucose and lipid oxidation rates were calculated according to the methods of Frayn (25). The  $\beta$ -cell function was assessed by calculating the disposition index ( $D_i$ ) expressing the inverse hyperbolic relationship between insulin secretion and insulin action and calculated as  $\text{FPIR} \times \text{M-value}$ .

### Statistics

The comparison of males and females (see Tables 2 and 3) were performed using mixed ANOVA (PROC MIXED procedure; SAS Institute, Cary, NC). monozygotic twins share their entire genome, whereas dizygotic twins on average share half of their segregating genes. Therefore, observations in neither monozygotic nor dizygotic twin pairs can be regarded as independent observations. Accordingly, we adjusted for the intra-twin pair relationship in the analyses by including a random-effect term for twin pair membership and fixed-effect terms for zygosity.

Percent-wise impact of birth weight, birth weight SDS and third-trimester FGV (per 1 SD) on adult measures of metabolism

and body composition were studied in a similar PROC MIXED procedure adjusted for gender. The analyses on MR-derived measures of abdominal obesity were adjusted for current BMI (see Table 4).

To quantify the nongenetic impact of birth weight, birth weight SDS, and third-trimester FGV on adult body composition and glucose homeostasis, multivariate linear regression analyses were made using intra-pair differences (twin A – twin B) exclusively in the 42 monozygotic twins. This approach adjusts for common environmental factors (such as the maternal, placental, and common postnatal environmental effects) and most importantly genetic effects. Consequently, any observed association using this approach is of environmental origin. The multivariate linear regression analyses were made with intra-pair differences of adult anthropometry and body composition as response variable and intra-pair differences of birth weight, birth weight SDS, and third-trimester FGV as explanatory variable. All analyses were adjusted for sex.

The designation of a member in a twin pair is arbitrary; *i.e.*, there is no consistency in which of the twins in a pair is assigned A and which is assigned B. The correlation coefficient may differ dependent on the assignment of the twins, so intra-pair differences were calculated using  $2n$ , as previously recommended (26). All analyses were carried out in SAS (version 9.1; SAS Institute);  $P < 0.05$  was considered significant.

## Results

### Clinical characteristics

A total of 42 monozygotic (21 pairs; 26 males and 16 females) and 16 dizygotic twins (eight pairs; four males and 12 females) were included. Subject characteristics stratified by gender and zygosity is provided in Table 1. The median gestational age was 263 d (range, 208–279 d).

### Gender differences

Females had a higher total fat percentage and a 2-fold increase of the deep and superficial sc fat contents compared with males (Table 2). However, females and males had similar volume percentage of visceral fat (Table 2). Females had lower fasting plasma glucose but a higher level of free fatty acids compared with males (Table 3). During the hyperinsulinemic-euglycemic clamps, females had a higher rate of glucose appearance from the liver and a higher glucose oxidation rate compared with males (Table 3).

### Impact of birth weight, birth weight SDS, and FGV on body composition (Table 4)

The quantitative impact of 1 SD increase in birth weight SDS on visceral, deep sc, and superficial sc adipose tissue was –12.5, –32.3, and –8.1%, respectively, when accounting for current BMI. In comparison, birth weight *per se* was inversely associated with visceral [–14.7% (95% confidence interval = –26.3 to –1.4)] but not deep or superficial sc adipose tissue. Third-trimester FGV was not associated with visceral or sc adipose tissue accumulation.

In supplementary analysis, BMI was replaced as explanatory variable by body fat percentage, whole-body fat

**TABLE 1.** Subject characteristics

	Males		Females	
	MZ	DZ	MZ	DZ
Pre- and perinatal growth and condition				
n	26	4	16	12
Gestational age (d)	257 (253–267)	268 (266–270)	265 (263–266)	248 (237–259)
Fetus				
FGV	−0.88 (−1.56 to −0.55)	−0.76 (−1.01 to −0.42)	−0.35 (−0.62–0.12)	−0.46 (−1.31–0.41)
Birth				
Birth weight (g)	2450 ± 442	3389 ± 331	2502 ± 317	2185 ± 390
Birth weight Z-score	−1.30 ± 1.24	0.24 ± 0.88	−1.65 ± 1.01	−0.97 ± 1.24
Birth length (cm)	48.0 (46.5–48.5)	51.0 (49.0–52.5)	47.8 (45.5–49.0)	46.0 (44.0–47.5)
Birth length Z-score	−0.50 ± 0.99	−0.50 ± 1.11	−0.61 ± 1.35	−0.35 ± 1.08
Ponderal index	22.8 ± 2.1	26.0 ± 2.2	23.5 ± 2.2	22.8 ± 2.1
Head circumference (cm)	32.8 ± 1.9	34.4 ± 0.95	32.9 ± 1.3	32.0 ± 1.3
APGAR 1	9 (8–10)	9 (7–10)	10 (10–10)	10 (9–10)
APGAR 5	10 (10–10)	10 (10–10)	10 (10–10)	10 (10–10)
Adulthood				
Age (yr)	21.7 ± 2.1	18.9 ± 0.2	21.6 ± 1.4	23.2 ± 1.7
Height (m)	1.79 ± 0.05	1.84 ± 0.03	1.66 ± 0.05	1.69 ± 0.03
Weight (kg)	71.5 ± 8.2	75.4 ± 2.8	54.9 ± 9.2	58.4 ± 8.6
BMI (kg/m <sup>2</sup> )	22.4 ± 2.3	22.4 ± 1.3	20.0 ± 2.8	20.6 ± 3.0
Waist circumference (cm)	79.1 ± 5.8	80.3 ± 2.1	68.4 ± 6.4	69.2 ± 7.9
Hip circumference (cm)	94.0 ± 6.1	96.5 ± 1.7	91.7 ± 6.8	95.4 ± 7.1
Waist/hip ratio	0.84 ± 0.05	0.83 ± 0.02	0.75 ± 0.04	0.72 ± 0.04
Total cholesterol	3.8 ± 0.6	3.2 ± 0.5	3.8 ± 0.6	4.3 ± 0.72
HDL	1.11 ± 0.25	1.32 ± 0.23	1.72 ± 0.42	1.58 ± 0.26
LDL	2.26 ± 0.57	1.48 ± 0.62	1.71 ± 0.38	2.28 ± 0.68
Triglycerides	0.88 (0.65–1.06)	0.71 (0.59–0.95)	0.67 (0.51–0.94)	0.82 (0.71–1.24)
Systolic blood pressure	123 ± 9	135 ± 5	118 ± 9	111 ± 7
Diastolic blood pressure	63 ± 8	73 ± 7	69 ± 6	64 ± 7

Data are provided as mean ± sd or when not normally distributed as median (interquartile range). DZ, Dizygotic; HDL, high-density lipoprotein; LDL, low-density lipoprotein; MZ, monozygotic.

content (kilograms), or lean body mass (kilograms) (data not shown). However, substitution of BMI with DXA-derived measures of body composition made only very subtle changes to the results presented.

### Impact of birth weight, birth weight SDS, and FGV on insulin action and insulin secretion

Birth weight and birth weight SDS were not related to insulin action or insulin secretion. In contrast, FGV was inversely associated with M-value but not related to other measures of glucose metabolism (Table 4).

### Nongenetic impact of birth weight, birth weight SDS, and FGV on body composition

Intra-pair differences in birth weight, birth weight SDS, and FGV were correlated with intra-pair differences in adult body composition and glucose metabolism (Table 5). The quantitative nongenetic impact of 1 sd increase in birth weight within a twin pair was 2.0 cm in adult height and 1370 g in adult lean body mass (Table 5). Similar nongenetic relationships were seen for 1 sd increase in intra-twin pair birth weight SDS; however, birth weight SDS was additionally associated with decreased visceral, deep sc, and superficial sc fat accumulation of 1.9, 0.5, and 0.9 volume percent per 1 sd (Table 5). Third-trimester FGV was

associated with adult height but not with measures of abdominal obesity.

## Discussion

This study identifies novel and apparently independent differential contributions from third-trimester FGV and birth weight SDS, respectively, on abdominal obesity and insulin action. Birth weight SDS was inversely associated with both sc and visceral obesity in a nongenetic manner but not related to insulin action and insulin secretion. In contrast, third-trimester FGV was inversely associated with insulin action but not significantly with visceral or sc fat accumulation.

Previous studies have reported somewhat conflicting positive, inverse, or no associations between birth weight and different measures of adult obesity. Some of the diversity in the literature may arise from age-dependent associations (8), ethnic differences, and the nature and timing of growth restriction (27) as well as methodological differences related to the definition of obesity. Epidemiological studies have reported a positive association be-

**TABLE 2.** Adult anthropometrics determined by DXA and MR scan

	Males	Females
DXA		
Total fat (%)	14.2 (11.5–16.6)	26.2 (21.7–28.8) <sup>c</sup>
Total fat (kg)	10.71 (8.38–13.58)	14.95 (11.38–17.24) <sup>b</sup>
Total lean body mass (kg)	62.1 (57.7–65.4)	41.9 (37.7–45.0) <sup>c</sup>
Trunk fat (kg)	3.94 (3.30–5.31)	5.21 (3.53–6.60)
Trunk fat (%)	11.5 (9.75–15.2)	19.4 (15.5–24.1) <sup>c</sup>
Leg fat (kg)	2.06 (1.51–2.56)	2.93 (2.64–3.80) <sup>c</sup>
Leg fat (%)	16.2 (12.7–18.4)	32.0 (27.7–35.2) <sup>c</sup>
Trunk to leg ratio (kg)	2.02 (0.74–2.20)	1.39 (1.25–1.81) <sup>b</sup>
Trunk to leg ratio (%)	0.76 (0.68–0.86)	0.60 (0.53–0.69) <sup>b</sup>
Trunk to total fat ratio (kg)	0.38 (0.36–0.40)	0.34 (0.31–0.39) <sup>a</sup>
Leg to total fat ratio (kg)	0.38 (0.36–0.41)	0.47 (0.42–0.49) <sup>c</sup>
MR		
Superficial sc fat (vol%)	8.48 (7.06–10.79)	15.5 (12.0–20.1) <sup>c</sup>
Deep sc fat (vol%)	1.08 (0.70–2.10)	2.86 (1.77–3.43) <sup>b</sup>
Visceral fat (vol%)	5.62 (4.43–6.75)	4.95 (3.56–6.48)
Total sc (vol%)	10.3 (8.5–12.1)	18.7 (14.0–23.1) <sup>c</sup>
Abdominal fat (vol%)	15.9 (13.2–19.5)	22.6 (17.6–28.8) <sup>b</sup>

Data from the MR scan are provided in volume percent (vol%) (95% confidence interval).

<sup>a</sup>  $P < 0.05$ .

<sup>b</sup>  $P < 0.01$ .

<sup>c</sup>  $P < 0.001$ .

tween birth weight and BMI in young (28), middle-aged (29), and elderly individuals (30). However, a smaller study, using DXA scans, reported that the positive association of birth weight with adult BMI was explained by its association with lean body mass (31). A study in 32 elderly men from England reported higher fat mass and higher trunk-to-limb fat ratio after adjusting for total fat mass in men born with a low birth weight (32). Similarly, we have previously observed an elevated trunk-to-leg fat ratio among young healthy singletons born with low birth weight (7). The present study used state-of-art multi-slice MR imaging (MRI) and found birth weight SDS to be associated with both visceral and sc fat deposition but not with whole-body fat percentage, BMI, or waist circumference.

Visceral fat accumulation is considered to be detrimental for insulin action (33), which potentially can be explained by a greater release of free fatty acids from the visceral than from the sc adipose tissue (34). Others have suggested that the quantitatively higher abundance of sc fat may represent the most critical adipose tissue compart-

**TABLE 3.** Hyperinsulinemic-euglycemic clamp

	Males	Females
Plasma glucose (mmol/liter)		
Basal	5.28 ± 0.21	4.82 ± 0.36 <sup>c</sup>
Serum insulin (pmol/liter)		
Basal	34.9 ± 12.6	33.2 ± 13.7
Free fatty acids (μmol/liter)		
Basal	269 (217–383)	420 (351–511) <sup>a</sup>
Insulin stimulated	6.0 (4.0–10.0)	9.0 (6.5–12.0)
Glucose oxidation (mg/min · kg FFM)		
Insulin stimulated	4.45 ± 0.56	5.21 ± 0.73 <sup>b</sup>
Fat oxidation (mg/min · kg FFM)		
Insulin stimulated	0.10 ± 0.22	−0.02 ± 0.24
Non-Ox GM (mg/min · kg FFM)	7.8 (6.9–9.6)	8.3 (7.2–9.3)
$R_a$ glucose (mg/min · kg FFM)	2.52 ± 0.54	3.23 ± 0.85 <sup>b</sup>
M-value (mg/min · kg FFM)	12.2 (11.5–13.8)	13.0 (11.8–15.0)
$D_i$ peripheral (10 <sup>−3</sup> pmol/liter · min × mg/min · kg FFM)	19.5 (13.6–32.2)	24.4 (19.1–30.7)

Data are shown as mean ± SD or median (interquartile range) when data not normally distributed. FFM, Fat-free mass;  $R_a$ , rate of appearance. Non-Ox, Non-oxidative glucose metabolism; GM, glucose metabolism;  $D_i$ , disposition index.

<sup>a</sup>  $P < 0.05$ .

<sup>b</sup>  $P < 0.01$ .

<sup>c</sup>  $P < 0.001$ .

ment adversely influencing insulin action (35, 36). A recent study in obese Pima Indians assessed abdominal obesity by MRI and found both ip and deep sc fat to predict peripheral and hepatic insulin action (37). Likewise, both visceral and sc adipose tissue were associated with blood pressure, diabetes, and the metabolic syndrome in the Framingham Heart Study (38). Finally, a prospective study among Japanese Americans found intraabdominal fat to predict T2D independent of BMI and sc fat area (39). The significant inverse intra-twin pair associations between birth weight SDS on one side, and both visceral as well as abdominal sc fat as determined using MRI on the other, demonstrates that an adverse intrauterine environment predisposes to abdominal obesity by influencing both visceral and sc abdominal fat accumulation with no impact on total-body fat content in young adult life. Importantly, the significant associations observed among the genetically identical monozygotic twins provide strong evidence in favor of a non-genetic component contributing to the association between low birth weight and abdominal fat accumulation.

In contrast to birth weight SDS scores, third-trimester FGV was not associated with visceral or sc fat volume

**TABLE 4.** Percent-wise impact of size and birth and FGV per 1 SD on measures of metabolism and body composition

	Birth weight	Birth weight SDS	FGV
MR			
Superficial sc fat (vol%) <sup>a</sup>	−2.9 (−8.8–3.3)	−8.1 (−12.7 to −3.3) <sup>c</sup>	−3.6 (−8.9–2.0)
Deep sc fat (vol%) <sup>a</sup>	−7.52 (−25.3–14.4)	−32.3 (−43.2 to −21.7) <sup>d</sup>	7.0 (−14.8–34.4)
Visceral fat (vol%) <sup>a</sup>	−14.7 (−26.3 to −1.4) <sup>b</sup>	−12.5 (−23.4 to −0.0) <sup>b</sup>	3.3 (−10.6–19.4)
Total sc (vol%) <sup>a</sup>	−4.2 (−10.5–2.6)	−10.3 (−15.3 to −5.00) <sup>d</sup>	−3.2 (−9.5–3.4)
Abdominal fat (vol%) <sup>a</sup>	−7.3 (−13.8 to −0.2) <sup>b</sup>	−12.0 (−17.1 to −6.5) <sup>d</sup>	−2.5 (−9.3–4.8)
DXA			
Total fat (%)	−3.76 (−8.59–1.33)	0.07 (−4.55–4.90)	−1.69 (−6.39–3.24)
Total fat (kg)	−3.08 (−10.31–4.69)	2.77 (−4.06–10.1)	−1.17 (−7.79–5.93)
Total lean body mass (kg)	1.66 (−0.47–3.91)	1.82 (−0.10–3.78)	−0.98 (−2.68–0.76)
Trunk to leg ratio (kg)	−1.47 (−6.79–4.15)	−0.56 (−5.44–4.56)	3.65 (−1.55–9.13)
Trunk to total fat ratio (kg)	−1.20 (−4.11–1.79)	−0.00 (−2.67–2.73)	1.68 (−1.15–4.59)
Hyperinsulinemic-euglycemic clamp			
Fasting plasma glucose (mmol/liter)	0.54 (1.21–1.02)	−0.37 (1.97–1.25)	0.65 (−1.05–2.37)
Fasting serum insulin (pmol/liter)	7.38 (−3.80–19.9)	4.17 (−5.97–15.4)	1.15 (−8.97–12.4)
Free fatty acids (μ mol/liter)			
Fasting	1.71 (−9.35–14.1)	6.29 (−5.08–19.0)	11.54 (−1.84–26.7)
Insulin stimulated	2.23 (−19.6–18.9)	−8.36 (−23.9–10.4)	6.90 (−11.6–29.2)
Glucose oxidation (mg/min · kg FFM)			
Insulin stimulated	0.25 (−3.77–4.45)	−1.79 (−5.52–2.08)	−3.30 (−7.51–1.10)
Fat oxidation (mg/min · kg FFM)			
Insulin stimulated	0.34 (−32.8–49.9)	−4.86 (35.4–40.1)	4.79 (−26.8–50.0)
Non-Ox GM (mg/min · kg FFM)	2.38 (−5.18–10.54)	−4.30 (−10.1–1.88)	−6.66 (−13.2–0.46)
FPIR	−0.84 (−16.1; 24.3)	6.48 (−8.76; 24.3)	7.97 (−7.43; 25.9)
R <sub>a</sub> glucose (mg/min · kg FFM)	0.02 (−6.71–7.14)	−3.59 (−9.48–2.68)	−3.84 (−9.69–2.39)
M-value (mg/min · kg FFM)	1.92 (−3.85–8.03)	−3.24 (−8.28–2.09)	−5.41 (−9.99 to −0.66) <sup>b</sup>
D <sub>i</sub> peripheral (10 <sup>−3</sup> pmol/liter · min × mg/min · kg FFM)	0.41 (−13.3–16.3)	3.24 (−9.79–18.2)	−0.24 (−13.0–14.4)

Mean change (95% confidence interval) in variables per 1 SD change in birth weight (473 g), birth weight z-score (1.23), or FGV (1.14 Δ z-score). All data were adjusted for sex. FFM, Fat-free mass; R<sub>a</sub>, rate of appearance; vol%, volume percent.

<sup>a</sup> Data were additionally adjusted for current BMI.

<sup>b</sup>  $P < 0.05$ .

<sup>c</sup>  $P < 0.01$ .

<sup>d</sup>  $P < 0.001$ .

percent, indicating that central fat accumulation represents a consequence of fetal programming before the third trimester. In accordance, a study in 805 human embryos and fetuses suggested the critical window for adipose tissue development to be the period from the 14th to 23rd week of gestation (40). Additionally, maternal undernutrition during the first and second but not third trimester was related to offspring obesity at age 19 in the Dutch Hunger Winter Study (27). We have previously investigated the impact of FGV *vs.* birth weight SDS on body composition assessed by DXA in a cohort of singletons recruited from the same Danish pregnancy clinic (9). In our previous study, birth weight SDS among singletons was inversely associated with DXA fat mass as well as trunk fat percentage, the latter finding being consistent with the current data. In further support of the current twin data, we did not observe any significant association between FGV and body composition among singletons (9). Taken together, the current and previous data support a key role of the intrauterine environment during the first or second trimester in the origin of abdominal obesity. The

minor difference between this and our previous study with regard to the associations between birth weight and total fat mass may be explained by the use of intra-pair differences among twins *vs.* singletons as well as the use of DXA *vs.* MRI in the two studies to assess regional and total fat mass. In terms of methodologies, it may be emphasized that intra-pair correlations among in particular monozygotic twins as well as the use of MRI to assess regional fat mass may represent more distinct, sensitive, and powerful assessments of the association between fetal growth and abdominal obesity as compared with the use of singletons and DXA scans. Finally, comparison between twins and singletons may be complicated by the different intrauterine challenges and shorter gestation or the higher risk of insulin resistance and T2D in twins *per se* (41).

We observed no associations between birth weight SDS and insulin action in this cohort of young healthy twins. In contrast, FGV during the third trimester was associated with insulin action in a somewhat paradoxical inverse manner with a high third-trimester FGV being associated with reduced insulin action. In our previous study of

**TABLE 5.** Intra-twin pair regression analyses of size at birth and FGV on measures of metabolism and body composition in 42 monozygotic twins

	$\Delta$ Birth weight	$\Delta$ Birth weight SDS	$\Delta$ FGV
Weight (kg)	2.06 (0.47–3.65) <sup>a</sup>	1.60 (–0.05–3.24)	0.66 (–1.22–2.55)
Height (cm)	2.0 (1.3–2.6) <sup>c</sup>	1.91 (1.22–2.60) <sup>c</sup>	1.17 (0.21–2.14) <sup>a</sup>
BMI (kg/m <sup>2</sup> )	0.24 (–0.29–0.78)	0.08 (–0.46–0.62)	–0.06 (–0.066–0.54)
Waist (cm)	1.73 (0.35–3.11) <sup>a</sup>	1.13 (–0.31–2.57)	0.03 (–1.65–1.70)
Hip (cm)	1.46 (–0.13–3.04)	1.18 (–0.43–2.79)	–0.31 (–2.15–1.53)
Waist/hip ratio	0.01 (–0.00–0.01)	0.00 (–0.01–0.01)	0.00 (–0.01–0.01)
Total fat (g)	104 (–628–836)	274 (–452–999)	103 (–706–911)
Total fat (%)	–0.57 (–1.34–0.21)	0.02 (–0.77–0.82)	–0.09 (–0.94–0.76)
Lean body mass (g)	1370 (624–2117) <sup>c</sup>	923 (117–1730) <sup>a</sup>	653 (–222–1527)
Trunk to leg ratio	0.00 (–0.08–0.08)	0.02 (–0.06–0.10)	0.04 (–0.06–0.13)
Trunk to total fat ratio	–0.00 (–0.01–0.01)	0.00 (–0.00–0.01)	0.00 (–0.01–0.01)
Superficial sc fat (vol%)	0.03 (–0.71–0.76)	–0.88 (–1.56 to –0.20) <sup>a</sup>	–0.44 (–1.24–0.35)
Deep sc fat (vol%)	–0.39 (–0.65 to –0.13) <sup>b</sup>	–0.46 (–0.70 to –0.21) <sup>c</sup>	–0.10 (–0.41–0.21)
Visceral fat (vol%)	–1.17 (–2.44–0.10)	–1.89 (–3.06 to –0.71) <sup>b</sup>	–0.65 (–2.08–0.78)
Total sc (vol%)	–0.36 (–1.25–0.53)	–1.34 (–2.13 to –0.55) <sup>c</sup>	–0.54 (–1.52–0.43)
Abdominal fat (vol%)	–1.53 (–3.39–0.33)	–3.22 (–4.84–1.60) <sup>c</sup>	–1.19 (–3.27–0.88)
FPIR (pmol/liter · min)	1.5 (–326–329)	50 (–137–237)	1.5 (–200–203)
R <sub>a</sub> glucose (mg/min · kg FFM)	–0.12 (–0.29–0.04)	–0.06 (–0.23–0.11)	–0.12 (–0.29–0.05)
M-value (mg/min · kg FFM)	–0.06 (–1.07–0.94)	–0.28 (–0.96–0.10)	–0.67 (–1.36–0.01)
D <sub>i</sub> peripheral (10 <sup>–3</sup> pmol/liter · min × mg/min · kg FFM)	33 (–1835–1900)	188 (–1660–2036)	–1104 (–3080–873)

Intra-twin pair change (95% confidence interval) per 1 sd change in intra-twin pair birth weight (406 g), birth weight SDS (1.24), and FGV (1.33). Analyses were adjusted for sex. FFM, Fat-free mass; R<sub>a</sub>, rate of appearance; vol%, volume percent.

<sup>a</sup>  $P < 0.05$ .

<sup>b</sup>  $P < 0.01$ .

<sup>c</sup>  $P < 0.001$ .

young healthy singletons using different methodologies, we found no association of FGV on insulin action (9). The inverse relationship between FGV and insulin action is in accordance with our previous report of an age-dependent nongenetic influence of birth weight on *in vivo* clamp insulin action with a negative association among young twins and a positive association among elderly twins (42). The notion of a time- or age-dependent effect of birth weight on glucose tolerance and insulin action is supported by studies of offspring of protein-malnourished rats (43, 44). At 3 months of age, the young offspring of protein-restricted mothers had an improved glucose tolerance; however, at 15 month of age, they were less glucose tolerant compared with controls (44).

Following the line of considering distinct trajectories of metabolic and anthropometric alterations associated with impaired fetal growth, we speculate that abdominal obesity proceeds, and may subsequently contribute to, the development of overt insulin resistance.

Caution is warranted in extrapolating whole-body insulin action determinations to represent accurate measurements of insulin action in the primary target tissues of insulin including muscle, liver, and fat. Thus, data are accumulating that insulin action may differ significantly within each of these tissues with hepatic insulin action being distinct from peripheral insulin action (19, 45, 46)

as well as adipose tissue insulin action being distinct from muscle insulin action as recently documented in bed-rest experiments from our group (47). Indeed, we have previously shown that young LBW men exhibit significant defects of muscle insulin action despite normal whole-body insulin action (48, 49). Although the notion of organ-specific trajectories of insulin action is in line with the idea of impaired muscle insulin action preceding overt whole-body insulin resistance among young LBW men, more knowledge is required to dissect the distinct effects of third-trimester growth velocity, as opposed to growth before or after the third trimester, on both whole-body as well as organ-specific insulin action. All together, we are facing a hitherto unrecognized complexity of the associations as well as underlying mechanisms linking the intrauterine environment to the development of overt whole-body and organ-specific insulin resistance, which needs to be established to understand, and subsequently to prevent, the development of T2D in subjects experiencing impaired fetal growth.

The twin approach is unique in regard to determination of nongenetic contributions to phenotypic traits. We found solid evidence for a nongenetic inverse association between birth weight SDS and visceral and sc adipose tissue. Family studies have suggested that 42–56% of the variability in visceral and sc fat may be due to genetic

causes (50, 51). In addition, recent genome-wide association studies have identified common variants in the *FTO* gene to be associated with obesity and provided further evidence for a genetic influence on common obesity (52). Altogether, central obesity appears to be determined by both genetic and nongenetic factors.

In conclusion, abdominal adiposity including both sc and visceral fat may constitute primary nongenetic defects associated with reduced fetal growth before the third trimester, preceding overt defects of insulin secretion and action. Reduced FGV during *vs.* before the third trimester may define distinct adult trajectories of metabolic and anthropometric characteristics influencing risk of developing T2D.

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## References

- Hales CN, Barker DJ, Clark PM, Cox LJ, Fall C, Osmond C, Winter PD 1991 Fetal and infant growth and impaired glucose tolerance at age 64. *BMJ* 303:1019–1022
- Barker DJ, Bull AR, Osmond C, Simmonds SJ 1990 Fetal and placental size and risk of hypertension in adult life. *BMJ* 301:259–262
- Barker DJ, Osmond C, Winter PD, Margetts B, Simmonds SJ 1989 Weight in infancy and death from ischaemic heart disease. *Lancet* 334:577–580
- Ravelli AC, van der Meulen JH, Michels RP, Osmond C, Barker DJ, Hales CN, Bleker OP 1998 Glucose tolerance in adults after prenatal exposure to famine. *Lancet* 351:173–177
- Ford ES, Williamson DF, Liu S 1997 Weight change and diabetes incidence: findings from a national cohort of US adults. *Am J Epidemiol* 146:214–222
- Hubert HB, Feinleib M, McNamara PM, Castelli WP 1983 Obesity as an independent risk factor for cardiovascular disease: a 26-year follow-up of participants in the Framingham Heart Study. *Circulation* 67:968–977
- Rasmussen EL, Malis C, Jensen CB, Jensen JE, Storgaard H, Poulsen P, Pilgaard K, Schou JH, Madsbad S, Astrup A, Vaag A 2005 Altered fat tissue distribution in young adult men who had low birth weight. *Diabetes Care* 28:151–153
- Monrad RN, Grunnet LG, Rasmussen EL, Malis C, Vaag A, Poulsen P 2009 Age-dependent nongenetic influences of birth weight and adult body fat on insulin sensitivity in twins. *J Clin Endocrinol Metab* 94:2394–2399
- Vielwerth SE, Jensen RB, Larsen T, Holst KK, Mølgaard C, Greisen G, Vaag A 2008 The effect of birthweight upon insulin resistance and associated cardiovascular risk factors in adolescence is not explained by fetal growth velocity in the third trimester as measured by repeated ultrasound fetometry. *Diabetologia* 51:1483–1492
- Andersson EA, Pilgaard K, Pisinger C, Harder MN, Grarup N, Faerch K, Poulsen P, Witte DR, Jørgensen T, Vaag A, Hansen T, Pedersen O 2010 Type 2 diabetes risk alleles near *ADCY5*, *CDKAL1* and *HHEX-IDE* are associated with reduced birthweight. *Diabetologia* 53:1908–1916
- Freathy RM, Bennett AJ, Ring SM, Shields B, Groves CJ, Timpson NJ, Weedon MN, Zeggini E, Lindgren CM, Lango H, Perry JR, Pouta A, Ruukonen A, Hyppönen E, Power C, Elliott P, Strachan DP, Jarvelin MR, Smith GD, McCarthy MI, Frayling TM, Hattersley AT 2009 Type 2 diabetes risk alleles are associated with reduced size at birth. *Diabetes* 58:1428–1433
- Pulizzi N, Lyssenko V, Jonsson A, Osmond C, Laakso M, Kajantie E, Barker DJ, Groop LC, Eriksson JG 2009 Interaction between prenatal growth and high-risk genotypes in the development of type 2 diabetes. *Diabetologia* 52:825–829
- Bo S, Cavallo-Perin P, Scaglione L, Ciccone G, Pagano G 2000 Low birthweight and metabolic abnormalities in twins with increased susceptibility to type 2 diabetes mellitus. *Diabet Med* 17:365–370
- Poulsen P, Vaag AA, Kyvik KO, Møller Jensen D, Beck-Nielsen H 1997 Low birth weight is associated with NIDDM in discordant monozygotic and dizygotic twin pairs. *Diabetologia* 40:439–446
- Warsof SL, Gohari P, Berkowitz RL, Hobbins JC 1977 The estimation of fetal weight by computer-assisted analysis. *Am J Obstet Gynecol* 128:881–892
- Larsen T, Petersen S, Greisen G, Larsen JF 1990 Normal fetal growth evaluated by longitudinal ultrasound examinations. *Early Hum Dev* 24:37–45
- Niklasson A, Ericson A, Fryer JG, Karlberg J, Lawrence C, Karlberg P 1991 An update of the Swedish reference standards for weight, length and head circumference at birth for given gestational age (1977–1981). *Acta Paediatr Scand* 80:756–762
- Eiberg H, Hansen L, Kjer B, Hansen T, Pedersen O, Bille M, Rosenberg T, Tranebjaerg L 2006 Autosomal dominant optic atrophy associated with hearing impairment and impaired glucose regulation caused by a missense mutation in the *WFS1* gene. *J Med Genet* 43:435–440
- Alibegovic AC, Højbjerg L, Sonne MP, van Hall G, Stallknecht B, Dela F, Vaag A 2009 Impact of 9 days of bed rest on hepatic and peripheral insulin action, insulin secretion, and whole-body lipolysis in healthy young male offspring of patients with type 2 diabetes. *Diabetes* 58:2749–2756
- Ferrannini E 1988 The theoretical bases of indirect calorimetry: a review. *Metabolism* 37:287–301
- van HG, Sacchetti M, Radegran G, Saltin B 2002 Human skeletal muscle fatty acid and glycerol metabolism during rest, exercise and recovery. *J Physiol* 543(Pt 3):1047–1058
- Nielsen TL, Hagen C, Wraae K, Bathum L, Larsen R, Brixen K, Andersen M 2010 The impact of the CAG repeat polymorphism of the androgen receptor gene on muscle and adipose tissues in 20–29-year-old Danish men: Odense Androgen Study. *Eur J Endocrinol* 162:795–804
- Jørgensen P, Larsen R, Wraae K 2009 Unsupervised assessment of subcutaneous and visceral fat by MRI. In: Salberg AB, Hardeberg J, Jenssen R, eds. *Image analysis*. Vol. 5575. Berlin and Heidelberg: Springer; 179–188
- Wolfe RR, Chinkes D 2005 *Isotope tracers in metabolic research: principles and practice of kinetic analysis*. Hoboken, NJ: Wiley and Sons
- Frayn KN 1983 Calculation of substrate oxidation rates in vivo from gaseous exchange. *J Appl Physiol* 55:628–634
- Bring J, Wernroth L 1999 Inefficient analysis of twin data: is there an association between diabetes and birth weight? *Diabetologia* 42: 898–899
- Ravelli GP, Stein ZA, Susser MW 1976 Obesity in young men after famine exposure in utero and early infancy. *N Engl J Med* 295:349–353
- Rasmussen F, Johansson M 1998 The relation of weight, length and

- ponderal index at birth to body mass index and overweight among 18-year-old males in Sweden. *Eur J Epidemiol* 14:373–380
29. **Pilgaard K** 2010 Impact of size at birth and prematurity on adult anthropometry in 4744 middle-aged Danes: the Inter99 study. *J Dev Origins Health Dis* 1:319–328
  30. **Eriksson J, Forsén T, Osmond C, Barker D** 2003 Obesity from cradle to grave. *Int J Obes Relat Metab Disord* 27:722–727
  31. **Yliharsila H, Kajantie E, Osmond C, Forsén T, Barker DJ, Eriksson JG** 2007 Birth size, adult body composition and muscle strength in later life. *Int J Obes (Lond)* 31:1392–1399
  32. **Kensara OA, Wootton SA, Phillips DI, Patel M, Jackson AA, Elia M** 2005 Fetal programming of body composition: relation between birth weight and body composition measured with dual-energy x-ray absorptiometry and anthropometric methods in older Englishmen. *Am J Clin Nutr* 82:980–987
  33. **Kissebah AH** 1996 Intra-abdominal fat: is it a major factor in developing diabetes and coronary artery disease? *Diabetes Res Clin Pract* 30(Suppl):25–30
  34. **Richelsen B, Pedersen SB, Møller-Pedersen T, Bak JF** 1991 Regional differences in triglyceride breakdown in human adipose tissue: effects of catecholamines, insulin, and prostaglandin E<sub>2</sub>. *Metabolism* 40:990–996
  35. **Abate N, Garg A, Peshock RM, Stray-Gundersen J, Grundy SM** 1995 Relationships of generalized and regional adiposity to insulin sensitivity in men. *J Clin Invest* 96:88–98
  36. **Goodpaster BH, Thaete FL, Simoneau JA, Kelley DE** 1997 Subcutaneous abdominal fat and thigh muscle composition predict insulin sensitivity independently of visceral fat. *Diabetes* 46:1579–1585
  37. **Koska J, Stefan N, Votruba SB, Smith SR, Krakoff J, Bunt JC** 2008 Distribution of subcutaneous fat predicts insulin action in obesity in sex-specific manner. *Obesity (Silver Spring)* 16:2003–2009
  38. **Fox CS, Massaro JM, Hoffmann U, Pou KM, Maurovich-Horvat P, Liu CY, Vasan RS, Murabito JM, Meigs JB, Cupples LA, D'Agostino RB Sr, O'Donnell CJ** 2007 Abdominal visceral and subcutaneous adipose tissue compartments: association with metabolic risk factors in the Framingham Heart Study. *Circulation* 116:39–48
  39. **Boyko EJ, Fujimoto WY, Leonetti DL, Newell-Morris L** 2000 Visceral adiposity and risk of type 2 diabetes: a prospective study among Japanese Americans. *Diabetes Care* 23:465–471
  40. **Poissonnet CM, Burdi AR, Bookstein FL** 1983 Growth and development of human adipose tissue during early gestation. *Early Hum Dev* 8:1–11
  41. **Poulsen P, Grunnet LG, Pilgaard K, Storgaard H, Alibegovic A, Sonne MP, Carstensen B, Beck-Nielsen H, Vaag A** 2009 Increased risk of type 2 diabetes in elderly twins. *Diabetes* 58:1350–1355
  42. **Poulsen, Levin, Beck N, Vaag A** 2002 Age-dependent impact of zygosity and birth weight on insulin secretion and insulin action in twins. *Diabetologia* 45:1649–1657
  43. **Ozanne SE, Wang CL, Coleman N, Smith GD** 1996 Altered muscle insulin sensitivity in the male offspring of protein-malnourished rats. *Am J Physiol* 271(6 Pt 1):E1128–E1134
  44. **Hales CN, Desai M, Ozanne SE, Crowther NJ** 1996 Fishing in the stream of diabetes: from measuring insulin to the control of fetal organogenesis. *Biochem Soc Trans* 24:341–350
  45. **Brons C, Vaag A** 2009 Skeletal muscle lipotoxicity in insulin resistance and type 2 diabetes. *J Physiol* 587(Pt 16):3977–3978
  46. **Faerch K, Brons C, Alibegovic AC, Vaag A** 2010 The disposition index: adjustment for peripheral vs. hepatic insulin sensitivity? *J Physiol* 588(Pt 5):759–764
  47. **Højbjerg L, Sonne MP, Alibegovic AC, Dela F, Vaag A, Meldgaard JB, Christensen KB, Stallknecht B** 2010 Impact of physical inactivity on subcutaneous adipose tissue metabolism in healthy young male offspring of patients with type 2 diabetes. *Diabetes* 59:2790–2798
  48. **Hermann TS, Rask-Madsen C, Ihlemann N, Domínguez H, Jensen CB, Storgaard H, Vaag AA, Kober L, Torp-Pedersen C** 2003 Normal insulin-stimulated endothelial function and impaired insulin-stimulated muscle glucose uptake in young adults with low birth weight. *J Clin Endocrinol Metab* 88:1252–1257
  49. **Sonne MP, Højbjerg L, Alibegovic AC, Vaag A, Stallknecht B, Dela F** 2010 Diminished insulin-mediated forearm blood flow and muscle glucose uptake in young men with low birth weight. *J Vasc Res* 47:139–147
  50. **Hong Y, Rice T, Gagnon J, Després JP, Nadeau A, Pérusse L, Bouchard C, Leon AS, Skinner JS, Wilmore JH, Rao DC** 1998 Familial clustering of insulin and abdominal visceral fat: the HERITAGE Family Study. *J Clin Endocrinol Metab* 83:4239–4245
  51. **Pérusse L, Després JP, Lemieux S, Rice T, Rao DC, Bouchard C** 1996 Familial aggregation of abdominal visceral fat level: results from the Quebec family study. *Metabolism* 45:378–382
  52. **Dina C, Meyre D, Gallina S, Durand E, Körner A, Jacobson P, Carlsson LM, Kiess W, Vatin V, Lecoer C, Delplanque J, Vaillant E, Pattou F, Ruiz J, Weill J, Levy-Marchal C, Horber F, Potoczna N, Hercberg S, Le Stunff C, Bougnères P, Kovacs P, Marre M, Balkau B, Cauchi S, Chèvre JC, Froguel P** 2007 Variation in FTO contributes to childhood obesity and severe adult obesity. *Nat Genet* 39:724–726