

GYNECOLOGY

Waist circumference in relation to outcomes of infertility treatment with assisted reproductive technologies



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BACKGROUND: Many studies have documented a lower likelihood of live birth with increasing body mass index among women undergoing assisted reproductive technology, but few have examined the association with waist circumference, an anthropometric measure that allows assessment of central adiposity.

OBJECTIVE: To examine the relation between baseline waist circumference and infertility treatment outcomes among women undergoing treatment with assisted reproductive technology.

MATERIALS AND METHODS: We followed up 264 women who underwent 445 assisted reproductive technology cycles for infertility treatment at the Massachusetts General Hospital between 2010 and 2017. Waist circumference was assessed at enrollment. We used cluster-weighted generalized estimating equation models to estimate the probability of live birth by tertiles of waist circumference (<77, 77–86, >86 cm), while accounting for multiple treatment cycles per woman and adjusting for age, race, smoking, infertility diagnosis, day 3 follicle-stimulating hormone, body mass index, and height.

RESULTS: Mean (standard deviation) waist circumference and body mass index were 83.6 (12.6) cm and 24.1 (4.3) kg/m², respectively. Waist

circumference and body mass index were positively correlated ($r = 0.69$, $P < .0001$). Waist circumference was inversely related to the probability of live birth after adjusting for BMI and other confounders. The multivariable adjusted probability of live birth (95% confidence interval) for women in increasing tertiles of waist circumference were 53% (42–65%), 42% (32–53%), and 38% (28–50%) (P , trend = .04). When women were classified in joint categories of body mass index and waist circumference, women with a body mass index ≥ 25 kg/m² and a waist circumference ≥ 77 cm had the lowest live birth rate (38% [27–50%]), whereas women with a body mass index between 18.5 and 25 kg/m² and a waist circumference <77 cm had the highest (54% [42–66%]). The results were similar using different waist circumference cut-off values.

CONCLUSION: Waist circumference was inversely related to the probability of live birth among women undergoing assisted reproductive technology independently of body mass index.

Key words: assisted reproductive technology, body mass index, ICSI, in vitro fertilization, intracytoplasmic sperm injection, IVF, waist circumference

Obesity has become a major public health problem as its prevalence has reached epidemic proportions. Between 2007 and 2016, the prevalence of obesity and severe obesity among women in the United States increased from 35.4% to 41.1% and from 7.3% to 9.7%, respectively.¹ Obesity has been identified as a risk factor for subfertility, especially with ovulation disorders.^{2,3} Several studies have shown the increased risk for delayed time to pregnancy among obese women.^{4–9} Obesity also predicts a lower probability of live birth among women undergoing assisted reproductive technology (ART).^{10,11} The

most updated systematic review and meta-analysis concluded that both overweight (body mass index [BMI] ≥ 25 kg/m²) and obese (BMI ≥ 30 kg/m²) women have a significantly lower live birth rate, compared with women with a normal BMI (18.5–24.9 kg/m²).¹²

A known limitation of BMI as a marker of adiposity is that, despite being highly correlated with fat mass,¹³ it is also positively related to lean mass.^{14,15} Hence, BMI can misclassify muscular individuals as obese, although misclassification of individuals with high adiposity as “normal” is a more frequent problem.¹⁶ Women presenting to fertility centers tend to have a lower BMI than women of reproductive age in the general population, and thus misclassification may occur more often in studies conducted in fertility centers than in other settings,^{1,11} although the extent of this issue is unknown. Waist circumference (WC) is an anthropometric measure that is not only highly correlated with fat mass but is also used allow for

classification of central obesity.¹⁷ In fact, WC can predict risk of multiple chronic conditions when considered alone or in combination with BMI.^{18–24} However, few studies have examined the association of WC with fertility or outcomes of infertility treatment. To address this gap, we examined the relation between WC and ART outcomes, when considered alone or in combination with BMI, in an ongoing prospective cohort study of women in couples presenting for infertility evaluation and treatment.

Material and Methods

Study population

This study consisted of a subset of women enrolled in the Environment and Reproductive Health (EARTH) Study, an ongoing prospective cohort started in 2004 to explore environmental and lifestyle determinants of fertility among couples presenting for evaluation or treatment of infertility to the Massachusetts General Hospital (MGH) Fertility Center (Boston, MA). Women

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AJOG at a Glance

Why was this study conducted?

Many studies have identified overweight and obesity as risk factors for infertility and failed infertility treatment. However, the role abdominal obesity independent of overall adiposity, as measured by body mass index (BMI), has received little attention.

Key findings

The probability of achieving a live birth as a result of assisted reproductive technology decreased with increasing waist circumference independently of BMI. Overweight women with a high waist circumference had the lowest probability of live birth, whereas overweight women with low waist circumference had probabilities of live birth similar to those of women with normal BMI.

What does this add to what is known?

Waist circumference provides information regarding a woman's chances of infertility treatment success with assisted reproductive technology above and beyond her BMI.

aged 18–45 years who were planning to use their own gametes for ART were eligible. Women remained eligible to participate if their physician later determined that using donor oocytes or embryos was clinically necessary. Among women referred to the study, approximately 60% of those approached by the research staff enrolled in the study.

Waist circumference assessments were introduced to the study in 2010. At study entry, participants underwent an anthropometric assessment by trained study staff. Waist circumference was measured using a Gullick II Plus Measuring Tape and recorded to the nearest millimeter. Participants were asked to stand, discover their abdomen, and hold their shirt or examination gown above their abdomen with their crossed arms. Study staff then placed the measuring tape at the level of the umbilicus on a plane parallel to the floor and applying pressure based on the tape's pressure indicator. All participants completed a staff-administered brief questionnaire aimed at collecting basic demographic, medical, and lifestyle information. Participants were also asked to allow investigators access to their medical records, and to complete a detailed questionnaire addressing lifestyle and medical history at home. Women were then followed up during their in vitro fertilization or intracytoplasmic sperm injection treatment cycles for clinical outcomes until a live

birth was achieved or until discontinuation of treatment.

The current analysis included 264 women with available WC measurements who underwent 445 ART cycles between 2010 and 2017. The minimum and maximum number of ART cycles contributed per woman were 1 ($n = 142$) and 5 ($n = 10$). This study was approved by the Institutional Review Boards of the MGH and the Harvard T.H. Chan School of Public Health. All participants provided written informed consent after the study procedures were explained by study staff.

Clinical management and assessment of outcomes

Trained study staff abstracted clinical information from the patients' electronic medical records. Details of patient clinical management have been described elsewhere.¹⁰ In brief, for fresh ART cycles, patients underwent 1 of 3 stimulation protocols as clinically indicated: 1) luteal-phase gonadotropin-releasing hormone (GnRH) agonist protocol; 2) follicular-phase GnRH-agonist/flare protocol; or 3) GnRH-antagonist protocol. Patients were monitored by clinical staff during gonadotropin stimulation for serum estradiol (E2), follicle size and counts, and endometrial thickness, and were administered human chorionic gonadotropin (hCG) approximately 35 hours before the scheduled oocyte retrieval to

induce oocyte maturation. Embryologists classified oocytes as germinal vesicle, metaphase I, metaphase II, or degenerated, and then determined fertilization rate as the number of oocytes with 2 pronuclei divided by the number of metaphase II oocytes at 17–20 hours after either conventional in vitro fertilization or intracytoplasmic sperm injection. For cryo-thaw or donor-egg recipient cycles, patients underwent endometrial preparation protocols as clinically indicated. Early ART end points referred to those preceding embryo transfer, including markers of ovarian responses to stimulation (peak E2 levels, endometrial thickness, MII, and total oocytes), and fertilization rate. Clinical outcomes per initiated ART cycle were assessed, including the probabilities of implantation (a serum β -hCG level >6 mIU/mL, measured approximately 17 days after oocyte retrieval), clinical pregnancy (presence of intrauterine gestational sac[s] on transvaginal ultrasonography at 6 weeks), and live birth (the birth of a neonate on or after 24 weeks' gestation). The denominator for all clinical outcomes was the total number of ART cycles started.

Statistical analysis

Participants were categorized by tertiles of WC measured in centimeters. The associations of baseline personal and reproductive characteristics with WC were evaluated using Kruskal–Wallis tests for continuous variables and χ^2 or Fisher' exact tests for categorical variables. We used cluster-weighted generalized estimating equations models²⁵ to evaluate the associations of WC with ART outcomes, while accounting for correlation among multiple treatment cycles per woman and the unbalanced design. A normal distribution and identity link were specified for continuous outcomes (peak E2 levels, endometrial thickness, and day 3 follicle-stimulating hormone [FSH] levels); a Poisson distribution and log link for count outcomes (number of total and mature oocytes retrieved); and a log-binomial distribution for proportions (fertilization and clinical outcomes). Tests for linear trend across tertiles of

TABLE 1
Demographic and baseline reproductive characteristics of study participants (N = 264 women)

Characteristics	All participants		Waist circumference, cm						
	61–158		Tertile 1 (<77.0)		Tertile 2 (77.0–86.0)		Tertile 3 (>86.0)		P value ^a
Range, cm	61–158		Tertile 1 (<77.0)		Tertile 2 (77.0–86.0)		Tertile 3 (>86.0)		P value ^a
Waist circumference, cm, mean (SD)	83.6	(12.6)	71.7	(4.0)	81.0	(3.0)	97.3	(10.6)	
Number of participants	264		85		88		91		
Personal characteristics									
Age, y, mean (SD)	34.9	(4.2)	34.6	(4.3)	34.5	(4.3)	35.4	(4.1)	.17
BMI, kg/m ² , mean (SD)	24.1	(4.3)	21.5	(3.5)	22.8	(2.4)	27.8	(3.9)	<.0001
Height, cm, mean (SD)	165.6	(6.8)	164.6	(6.5)	166.1	(7.0)	166.0	(6.7)	.32
Smoker, n (%)									
Never	201	(76)	66	(78)	67	(76)	68	(75)	.74
Past	59	(22)	19	(22)	19	(22)	21	(23)	
Current	4	(2)	0	(0)	2	(2)	2	(2)	
White, n (%)	212	(80)	64	(75)	69	(78)	79	(87)	.13
College degree or higher, n (%)	261	(99)	83	(98)	87	(99)	91	(100)	.34
Reproductive characteristics									
Initial primary infertility diagnosis, n (%)									
Male factor	71	(27)	21	(25)	28	(32)	22	(24)	.39
Female factor	71	(27)	23	(27)	18	(20)	30	(33)	
Unexplained	122	(46)	41	(48)	42	(48)	39	(43)	
Initial treatment protocol, n (%)									
Antagonist	41	(16)	11	(13)	16	(18)	14	(15)	.75
Flare	30	(11)	9	(10)	7	(8)	14	(15)	
Luteal phase agonist	180	(68)	60	(71)	61	(69)	59	(65)	
Endometrial preparation (donor/cryo cycles)	13	(5)	5	(6)	4	(5)	4	(5)	
Ever pregnant, n (%)	109	(41)	32	(38)	37	(42)	40	(44)	.59
Type of fertilization procedure, n (%) ^b									
In vitro fertilization	115	(48)	34	(45)	42	(51)	39	(48)	.80
Intracytoplasmic sperm injection	125	(52)	41	(55)	41	(49)	43	(52)	
Day 3 FSH, IU/L, mean (SD)	7.21	(2.23)	7.48	(2.48)	7.07	(2.02)	7.09	(2.19)	.36

BMI, body mass index; FSH, follicle-stimulating hormone; SD, standard deviation.

Values are mean (SD) or n (%) and are standardized to the age distribution of the study population.

^a From Kruskal–Wallis test for continuous variables and χ^2 tests for categorical variables or Fisher test when cells have <5 individuals; ^b A total of 24 women were not included because of not reaching fertilization.

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WC were conducted using the median WC in each tertile as a continuous variable in the regression models. Primary analyses were performed using measured WC (divided into tertiles) and included terms for WC-adjusted BMI (BMI residual). Confounding was evaluated based on previous knowledge and taking into consideration the

associations between WC with baseline characteristics and the probability of live birth. Multivariable adjusted models were included terms for age (continuous, years), race/ethnicity (white vs. nonwhite), smoking (ever vs. never), infertility diagnosis (male factor, female factor, unexplained), day 3 FSH levels (IU/L), BMI (continuous, kg/m²), and

height (continuous, cm). Population marginal means were used to present population averages adjusted for the covariates (at the mean level for continuous variables and at a value weighted according to their frequencies for categorical variables) in the model.²⁶ We also assessed the relation of WC and ART using the World Health Organization

TABLE 2
Association between waist circumference and intermediate assisted reproductive technology outcomes

Adjusted means (95% CI) ^a	Cycles	Terile of waist circumference, cm						
		Terile 1 (<77.0 cm)		Terile 2 (77.0–86.0 cm)		Terile 3 (>86.0 cm)		<i>P</i> trend
Endometrial thickness, mm	422	10.4	(9.6–11.1)	10.4	(9.8–11.0)	9.9	(9.3–10.5)	
E2 trigger levels, pmol/L	353	2319	(2116–2521)	2118	(1936–2301)	2011 ^c	(1838–2185)	.01
Day 3 FSH, IU/L ^b	445	7.59	(6.98–8.20)	7.23	(6.76–7.70)	7.05	(6.54–7.56)	.11
Total oocyte yield, n	350	11.8	(10.5–13.2)	10.9	(9.6–12.3)	11.8	(10.5–13.2)	.84
M2 oocyte yield, n	350	9.6	(8.5–10.9)	9.1	(8.0–10.4)	9.7	(8.7–10.8)	.78
Fertilization, rate	349	0.71	(0.64–0.76)	0.74	(0.69–0.79)	0.72	(0.67–0.77)	.81

CI, confidence interval; FSH, follicle-stimulating hormone.

^a All models were adjusted for age, race (white vs. nonwhite), smoking (ever vs. never), infertility diagnosis (male factor, female factor, unexplained), day 3 FSH, body mass index, and height; ^b Not adjusted for day 3 FSH; ^c *P* < .05 compared with terile 1.

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(WHO) suggested cut-off values for WC (80 and 88 cm)²⁷ rather than distribution-based cutoffs. Because WC and BMI tend to be highly correlated, we calculated BMI-adjusted WC values (residuals of the regression of WC on BMI) and WC-adjusted BMI values (residuals of the regression of BMI on WC) to allow for the computation of more efficient estimates in multivariable models.¹⁷ We performed sensitivity analyses to evaluate the robustness of the findings and modeling assumptions, which included modeling WC as a restricted cubic spline and adjusting models with different strategies. We tested whether the association between WC and ART outcomes differed between fresh and cryo cycles or by primary infertility diagnosis by introducing cross-product terms to the multivariable adjusted models. When there was a suggestive interaction effect (*P* < .10), stratified analysis was further conducted. All statistical analyses were performed using SAS 9.4 (SAS Institute, Cary, NC).

Results

The study population included 264 women who underwent 445 ART cycles, of which 158 (35.5%) resulted in a live birth. Mean (standard deviation) age, WC, and BMI of participants were 34.9 (4.2) years, 83.6 (12.6) cm, and 24.1 (4.3) kg/m², respectively. The majority of women were white (80%), had a college degree (99%), and had never smoked

(76%). In all, 41% of the women reported at least 1 prior pregnancy. The most common initial primary infertility diagnosis was unexplained infertility (46%). WC was positively related to BMI (*r* = 0.69, *P* < .0001), but not to other personal or reproductive characteristics (Table 1).

Waist circumference was inversely associated with estradiol (E2) trigger levels. The adjusted mean (95% confidence interval [CI]) E2 trigger levels for women in increasing tertiles of WC were 2319 (2116–2521), 2118 (1936–2301), and 2011 (1838–2185) pmol/L (*P* trend = 0.01) (Table 2). However, we did not observe associations of WC with endometrial thickness, day 3 FSH levels, oocyte yield, or fertilization rate (Table 2).

Waist circumference was inversely related to the probabilities of implantation, clinical pregnancy, and live birth per initiated treatment cycle (Figure 1). The probabilities of implantation, clinical pregnancy, and live birth for women in the highest category of WC were, respectively, 14%, 13%, and 15% lower than those of women in the lowest category of WC (Figure 1). Results did not change after excluding height from all of the models. A similar pattern was observed when the analyses were conducted using the World Health Organization suggested cut-off values²⁷ for WC (Supplementary Table 1), when WC was modeled as a continuous variable

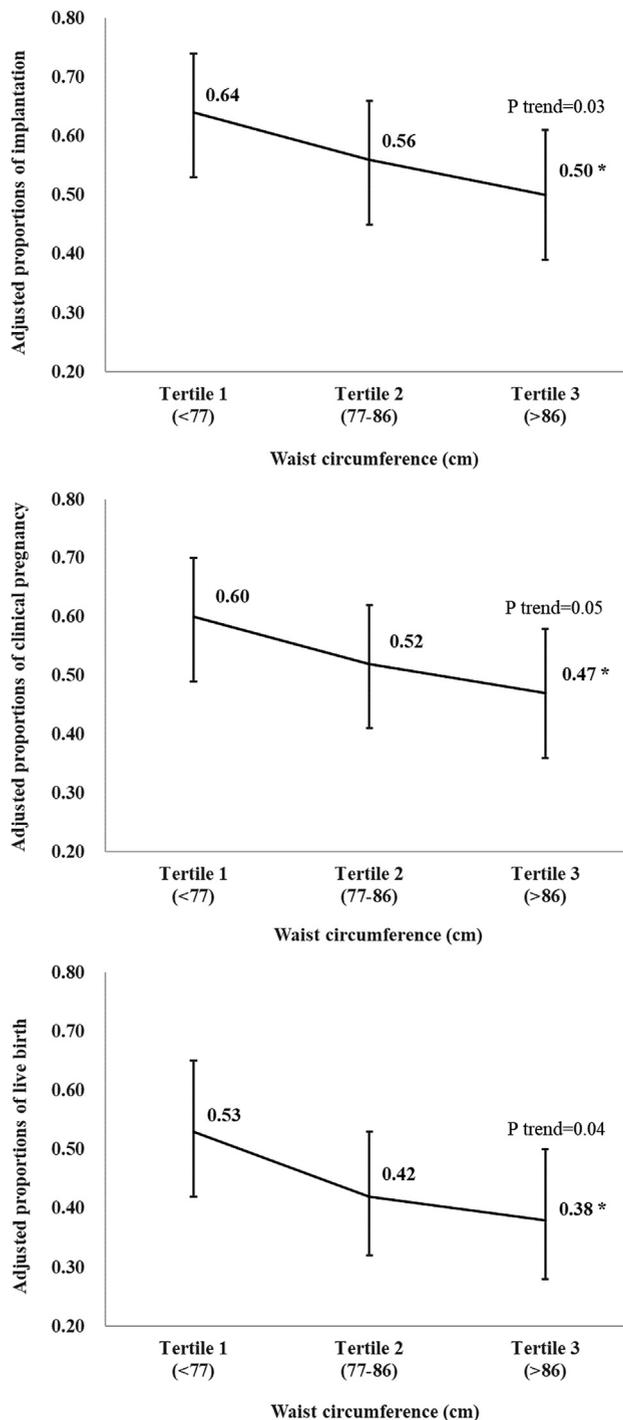
allowing for nonlinear relations (Supplementary Figures 1–3), and in sensitivity analyses using different approaches to account for the collinearity between WC and BMI (Supplementary Table 2).

We then classified women in joint categories of WC and BMI. Women with BMI ≥25 kg/m² and WC ≥77 cm had the lowest probability (95% CI) of live birth per cycle started (38% [27–50%]), whereas women with a BMI between 18.5 and 25 kg/m² and WC <77 cm had the highest probability of live birth (54% [42–66%]) (Figure 2). Overweight women with higher WC always had the lowest probability of live birth (Supplementary Table 3). In contrast, overweight women with lower WC had chances of treatment success similar to women with normal BMI. The results were similar using different WC cut-off values (Supplementary Table 3).

Sensitivity analyses showed that the relation between WC and ART outcomes did not differ between fresh and cryo cycles or by primary infertility diagnosis. Further adjusting for male partner BMI and WC in the subset of women for whom these data were available did not change the findings either (data not shown). Moreover, although graphic evaluation of the relation between WC and ART outcomes suggested nonlinear relations (Supplementary Figures 1–3), formal tests for nonlinearity showed that nonlinear models not are superior to

FIGURE 1

Adjusted probabilities (95% confidence intervals) of implantation, clinical pregnancy, and live birth per initiated treatment cycle across tertiles of waist circumference among women undergoing infertility treatment with assisted reproductive technology. All models were adjusted for age, race/ethnicity (white vs. nonwhite), smoking (ever vs. never), infertility diagnosis (male factor, female factor, unexplained), day 3 follicle-stimulating hormone, body mass index, and height. * $P < .05$ compared with tertile 1



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linear models in describing the relation between WC and ART outcomes in this study population.

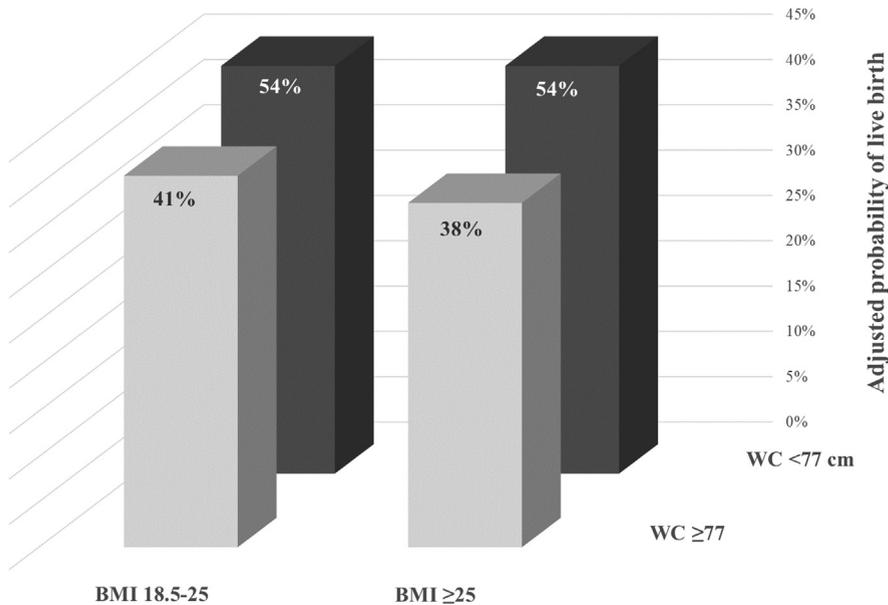
Comment

We examined the associations of baseline WC with embryological and clinical outcomes in a cohort of women undergoing ART. Waist circumference was associated with lower E2 trigger levels and, more importantly, lower probability of implantation, clinical pregnancy, and live birth per treatment cycle independently of BMI and other potential confounders. When women were jointly classified in categories of BMI and WC, overweight women with higher WC had the lowest probabilities of live birth. In contrast, overweight women with lower WC had probabilities of live birth similar to those of women with normal BMI. These data suggest that WC provides information regarding a woman's chances of treatment success with ART above and beyond her BMI.

To our knowledge, only 2 previous studies have evaluated the relation between WC and outcomes of infertility treatments. Hansen et al conducted a secondary data analysis from a randomized, multicenter clinical trial in women undergoing intrauterine insemination (IUI), which found that neither BMI nor WC was associated with the probability of live birth.²⁸ In addition, Moran et al evaluated the association of WC and ART outcomes in a pilot weight loss study among obese women presenting for infertility treatment. They found that decreases in WC were a better predictor of higher probability of live birth than changes in BMI, which did not predict ART outcomes.²⁹ Waist circumference has also been related to lower fecundability independently of BMI in cohorts of pregnancy planners without a history of infertility.^{30,31} Moreover, visceral fat area was related to response to ovulation induction with clomiphene citrate among women with polycystic ovary syndrome (PCOS) independently of BMI.³² Furthermore, emerging evidence suggests that WC may be a risk factor for pregnancy loss.^{33,34} Collectively, these data suggest that abdominal

FIGURE 2

Adjusted probability of live birth in joint categories of body mass index (18.5–25 or ≥ 25 kg/m²) and waist circumference (<77 or ≥ 77 cm) for women undergoing infertility treatment with assisted reproductive technology. All models were adjusted for age, race/ethnicity (white vs. nonwhite), smoking, infertility diagnosis (male factor, female factor, unexplained), day 3 follicle-stimulating hormone, and height



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obesity may have an impact on fertility above and beyond global measures of adiposity, which has been the main focus of the literature on the relation between adiposity and fertility.

The observed association between WC and ART outcomes may reflect a true biological effect. Waist circumference has been linked to insulin resistance and chronic inflammation. Insulin resistance is an important pathogenic factor in common metabolic disorders and plays a key pathogenic role in infertility.³⁵ Waist circumference has been reported as an independent predictor of insulin resistance after controlling for BMI,³⁶ as well as a great predictor of insulin sensitivity.³⁷ Moreover, WC was related to chronic inflammation,^{38,39} which can further affect ovulation and steroidogenesis.^{40–42} Whether the observed relation is mediated through these or other mechanisms deserves further evaluation.

Strengths of our study include its prospective design, with no loss to

follow-up, and robust assessment of intermediate and clinical outcomes, all of which aid in the interpretation of the findings. Both BMI and WC were measured by trained study staff, which minimized variability of the measurements. The sample size of the study population allowed us to evaluate clinically relevant outcomes with sufficient statistical power, including live birth. The main limitation of the study is the relatively small size of groups of women jointly classified according to their BMI and WC, which resulted in stratum sizes as small as 13 treatment cycles in some analyses. Although the results were robust regardless of cutoff values used in these joint models, additional larger cohort studies are warranted. In addition, WC was measured only once at enrollment, and thus any changes in WC during ART treatment were not captured by our study.

In conclusion, we found an association of high WC with lower probability

of live birth among women undergoing infertility treatment with ART. This association was independent of BMI, was consistent across different cut-off points of WC used in the analysis, and, in stratified analyses, was more pronounced for women with high WC and high BMI. These results suggest that central adiposity may be important for human fertility, above and beyond overall adiposity.

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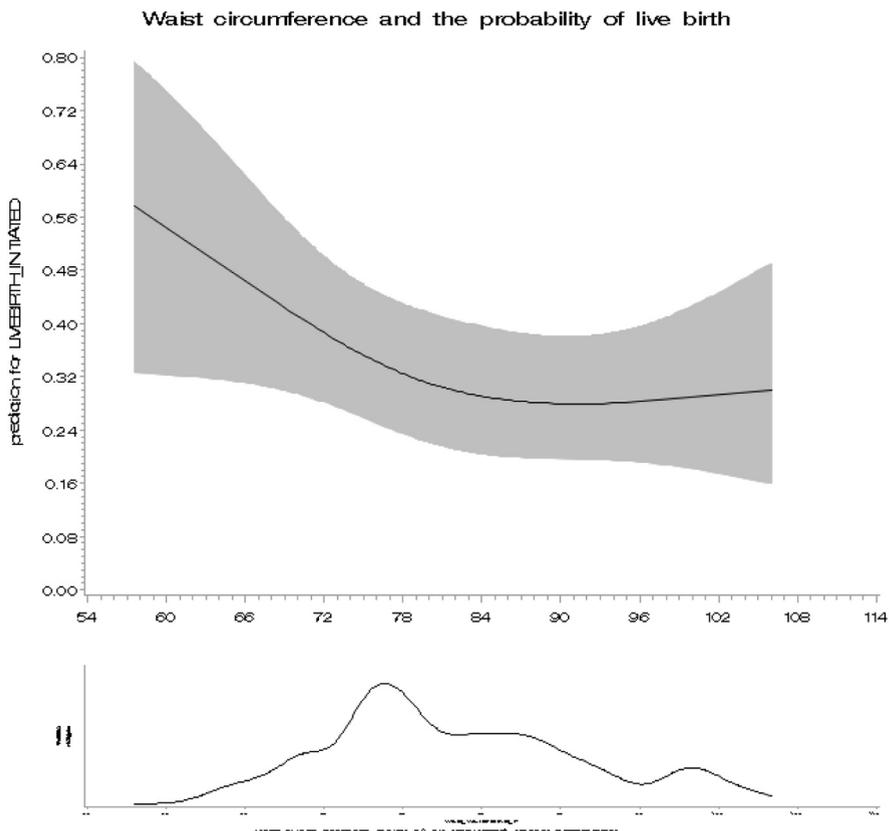
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FIGURE S1
Restricted cubic spline plot of the association between waist circumference and probability of live birth. P-value for non-linear relation: 0.20

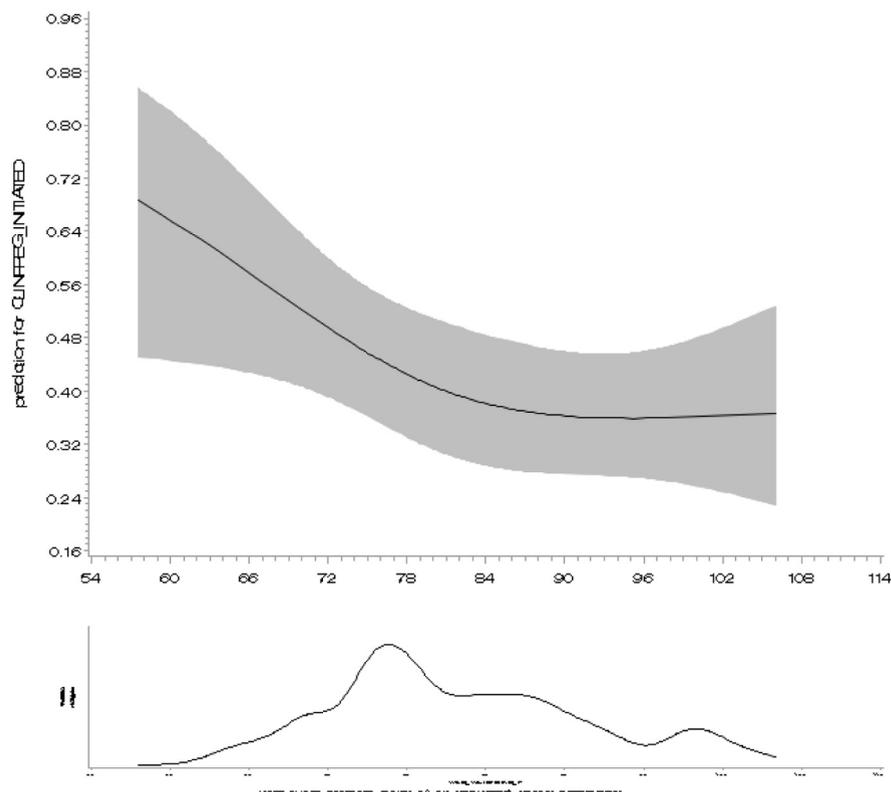


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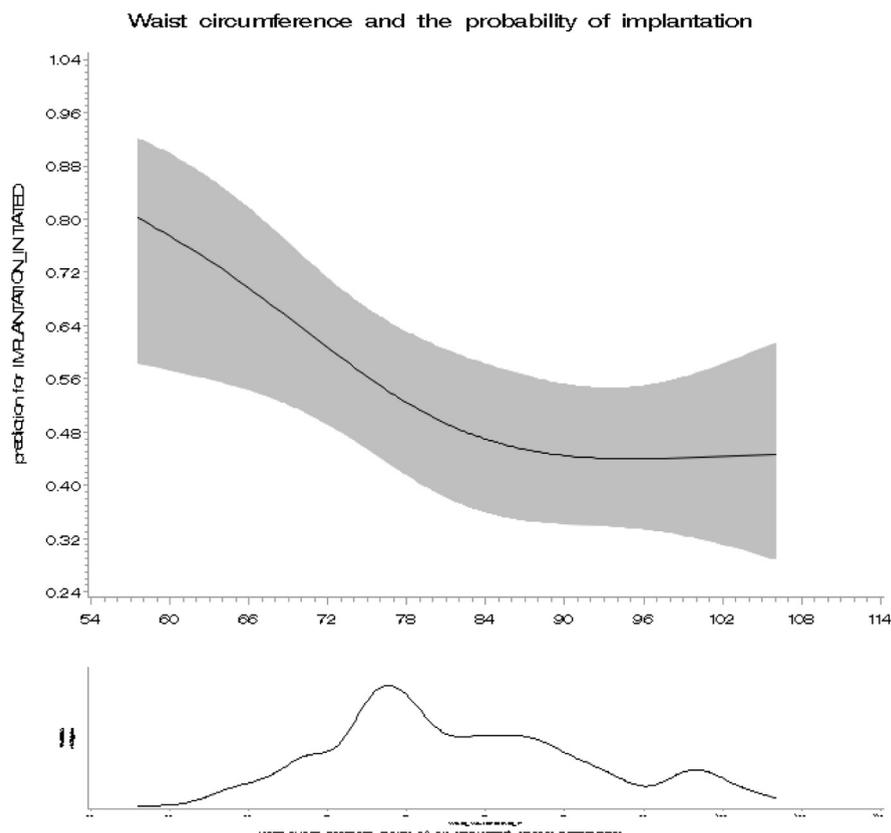
FIGURE S2

Restricted cubic spline plot of the association between waist circumference and probability of clinical pregnancy. P-value for non-linear relation: 0.19

Waist circumference and the probability of clinical pregnancy



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FIGURE S3**Restricted cubic spline plot of the association between waist circumference and probability of implantation. P-value for non-linear relation: 0.13**

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SUPPLEMENTARY TABLE 1

Association between World Health Organization—suggested waist circumference cut-off values and clinical assisted reproductive technology outcomes (N = 264 women, cycles = 445)

Adjusted proportions (mean, 95% CI) ^a	WC ≤80		WC >80		P _{trend}		
	Cycles = 215		Cycles = 230				
Implantation	0.62	(0.53–0.71)	0.51 ^b	(0.42–0.61)	.04		
Clinical pregnancy	0.57	(0.48–0.66)	0.48	(0.39–0.58)	.09		
Live birth	0.48	(0.38–0.58)	0.41	(0.31–0.51)	.19		
Adjusted proportions (mean, 95% CI) ^a	WC ≤80		WC >80		P _{trend}		
	Cycles = 322		Cycles = 123				
Implantation	0.60	(0.51–0.67)	0.49	(0.37–0.61)	.08		
Clinical pregnancy	0.55	(0.47–0.63)	0.46	(0.34–0.58)	.12		
Live birth	0.47	(0.38–0.55)	0.38	(0.27–0.51)	.20		
Adjusted proportions (mean, 95% CI) ^a	Waist circumference, cm						
	WC ≤80		80 < WC ≤ 88		WC >88.0		P _{trend}
Cycles = 215		Cycles = 107		Cycles = 123			
Implantation	0.62	(0.53–0.71)	0.54	(0.42–0.65)	0.49 ^b	(0.38–0.61)	.04
Clinical pregnancy	0.57	(0.48–0.66)	0.51	(0.39–0.62)	0.46	(0.34–0.58)	.08
Live birth	0.48	(0.38–0.58)	0.43	(0.31–0.56)	0.38	(0.27–0.51)	.15

CI, confidence interval; WC, waist circumference.

^a All models were adjusted for age, race (white vs. nonwhite), smoking (ever vs. never), infertility diagnosis (male factor, female factor, unexplained), day 3 follicle-stimulating hormone, body mass index, and height; ^b $P < .05$ compared with the first category.

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SUPPLEMENTARY TABLE 2

Association between waist circumference and probability of clinical assisted reproductive technology outcomes (N = 264 women, cycles = 445)

Adjusted proportions (95% CI) ^a	Implantation						P _{trend}
	Measured waist circumference, cm						
	Tertile 1 (<77.0)		Tertile 2 (77.0–86.0)		Tertile 3 (>86.0)		
Model 1	0.64	(0.54–0.74)	0.56	(0.46–0.66)	0.50 ^a	(0.39–0.61)	.03
Model 2	0.64	(0.53–0.74)	0.56	(0.45–0.66)	0.50 ^a	(0.39–0.61)	.03
Model 3	0.63	(0.52–0.73)	0.56	(0.45–0.66)	0.52	(0.40–0.63)	.14
Adjusted proportions (95% CI) ^a	BMI-adjusted waist circumference						P _{trend}
	Tertile 1 (<80.9)		Tertile 2 (80.9–86.7)		Tertile 3 (>86.7)		
	Model 4	0.66	(0.56–0.75)	0.51 ^a	(0.40–0.61)	0.52 ^a	
Model 5	0.66	(0.55–0.75)	0.50 ^a	(0.40–0.61)	0.53	(0.42–0.64)	.03
Adjusted proportions (95% CI) ^a	Clinical pregnancy						P _{trend}
	Measured waist circumference, cm						
	Tertile 1 (<77.0)		Tertile 2 (77.0–86.0)		Tertile 3 (>86.0)		
Model 1	0.60	(0.49–0.70)	0.51	(0.41–0.62)	0.47 ^a	(0.37–0.58)	.06
Model 2	0.60	(0.49–0.70)	0.52	(0.41–0.62)	0.47 ^a	(0.36–0.58)	.05
Model 3	0.61	(0.50–0.71)	0.52	(0.41–0.62)	0.46	(0.34–0.58)	.06
Adjusted proportions (95% CI) ^a	BMI-adjusted waist circumference						P _{trend}
	Tertile 1 (<80.9)		Tertile 2 (80.9–86.7)		Tertile 3 (>86.7)		
	Model 4	0.62	(0.52–0.72)	0.48 ^a	(0.37–0.58)	0.47 ^a	
Model 5	0.62	(0.52–0.72)	0.48 ^a	(0.37–0.58)	0.47 ^a	(0.36–0.58)	.01
Adjusted proportions (95% CI) ^a	Live birth						P _{trend}
	Measured waist circumference, cm						
	Tertile 1 (<77.0)		Tertile 2 (77.0–86.0)		Tertile 3 (>86.0)		
Model 1	0.53	(0.42–0.65)	0.42	(0.32–0.53)	0.38 ^a	(0.28–0.50)	.04
Model 2	0.53	(0.42–0.65)	0.42	(0.32–0.53)	0.38 ^a	(0.28–0.50)	.04
Model 3	0.53	(0.41–0.65)	0.42	(0.31–0.53)	0.39	(0.27–0.52)	.10
Adjusted proportions (95% CI) ^a	BMI-adjusted waist circumference						P _{trend}
	Tertile 1 (<80.9)		Tertile 2 (80.9–86.7)		Tertile 3 (>86.7)		
	Model 4	0.53	(0.39–0.62)	0.41	(0.27–0.49)	0.38 ^a	
Model 5	0.52	(0.41–0.63)	0.41	(0.31–0.52)	0.38	(0.28–0.50)	.04

BMI, body mass index; CI, confidence interval.

Model 1: Adjusted for age, race (white vs. nonwhite), smoking (ever vs. never), infertility diagnosis (male factor, female factor, unexplained), day 3 follicle-stimulating hormone, and height.

Model 2: Model 1 adjusted for BMI residual (waist circumference—adjusted BMI).

Model 3: Model 1 adjusted for measured BMI.

Model 4: Adjusted for age, race (white vs. nonwhite), smoking (ever vs. never), infertility diagnosis (male factor, female factor, unexplained), day 3 follicle-stimulating hormone, and height.

Model 5: Model 4 adjusted for measured BMI.

^a $P < .05$ compared with the first category.

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SUPPLEMENTARY TABLE 3

Adjusted probabilities (95% confidence intervals) of implantation, clinical pregnancy and live birth among women undergoing infertility treatment with assisted reproductive technology, jointly stratified by body mass index (BMI) and waist circumference (WC) categories (N = 257 women, cycles = 435)

	121		162		13		139	
Number of cycles	18.5 ≤ BMI <25				BMI ≥25			
Adjusted proportions (95% CI) ^a	WC <77.0		WC ≥77.0		WC <77.0		WC ≥77.0	
Implantation	0.64	(0.52–0.74)	0.55	(0.45–0.65)	0.73	(0.39–0.92)	0.49 ^b	(0.38–0.61)
Clinical pregnancy	0.61	(0.49–0.72)	0.50	(0.40–0.60)	0.66	(0.35–0.88)	0.48	(0.37–0.60)
Live birth	0.54	(0.42–0.66)	0.41	(0.32–0.52)	0.54	(0.24–0.81)	0.38 ^b	(0.27–0.50)
	191		92		17		135	
Number of cycles	18.5 ≤ BMI <25				BMI ≥25			
Adjusted proportions (95% CI) ^a	WC ≤80		WC >80		WC ≤80		WC >80	
Implantation	0.62	(0.52–0.71)	0.53	(0.40–0.66)	0.76	(0.48–0.91)	0.48 ^b	(0.37–0.60)
Clinical pregnancy	0.58	(0.48–0.67)	0.48	(0.35–0.60)	0.67	(0.40–0.86)	0.47	(0.36–0.59)
Live birth	0.49	(0.39–0.59)	0.43	(0.30–0.56)	0.45	(0.21–0.71)	0.39	(0.28–0.51)
	250		33		34		118	
Number of cycles	18.5 ≤ BMI <25				BMI ≥25			
Adjusted proportions (95% CI) ^a	WC ≤86		WC >86		WC ≤86		WC >86	
Implantation	0.59	(0.50–0.67)	0.60	(0.41–0.76)	0.71	(0.52–0.85)	0.45 ^b	(0.33–0.56)
Clinical pregnancy	0.54	(0.45–0.63)	0.53	(0.35–0.70)	0.68	(0.49–0.82)	0.43	(0.32–0.55)
Live birth	0.46	(0.37–0.55)	0.49	(0.31–0.68)	0.59	(0.40–0.76)	0.32 ^b	(0.22–0.45)
	263		20		49		103	
Number of cycles	18.5 ≤ BMI <25				BMI ≥25			
Adjusted proportions (95% CI) ^a	WC ≤88		WC >88		WC ≤88		WC >88	
Implantation	0.59	(0.50–0.67)	0.64	(0.38–0.83)	0.66	(0.49–0.79)	0.45 ^b	(0.33–0.57)
Clinical pregnancy	0.54	(0.46–0.63)	0.55	(0.31–0.76)	0.62	(0.46–0.76)	0.43	(0.32–0.56)
Live birth	0.46	(0.37–0.55)	0.56	(0.32–0.77)	0.52	(0.35–0.69)	0.33	(0.23–0.46)

CI, confidence interval.

^a All models were adjusted for age, race (white vs. nonwhite), smoking, infertility diagnosis (male factor, female factor, unexplained), day 3 follicle-stimulating hormone, and height; ^b $P < .05$ compared with the first category.

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