

Original

Sagittal abdominal diameter, but not waist circumference is strongly associated with glycemia, triacylglycerols and HDL-c levels in overweight adults

G. D. Pimentel^{1,2}, F. Moreto², M. M. Takahashi², K. C. Portero-McLellan² and R. C. Burini²

¹Department of Internal Medicine. FCM. State University of Campinas (UNICAMP). Campinas. SP. Brazil. ²Botucatu Medical School. Center for Nutritional and Exercise Metabolism (CeMENutri). Department of Public Health. Sao Paulo State University (UNESP). Botucatu. SP. Brazil.

Abstract

Aim: To correlate the sagittal abdominal diameter (SAD) and waist circumference (WC) with metabolic syndrome-associated abnormalities in adults.

Methods: This cross-sectional study included one-hundred twelve adults (M = 27, F = 85) aging 54.0 ± 11.2 yrs and average body mass index (BMI) of 30.5 ± 9.0 kg/m². The assessment included blood pressure, plasma and anthropometric measurements.

Results: In both men and female, SAD and WC were associated positively with body fat% ($r = 0.53$ vs $r = 0.55$), uric acid ($r = 0.45$ vs $r = 0.45$), us-PCR ($r = 0.50$ vs $r = 0.44$), insulin ($r = 0.89$ vs $r = 0.75$), insulin resistance HOMA-IR ($r = 0.86$ vs $r = 0.65$), LDL-ox ($r = 0.51$ vs $r = 0.28$), GGT ($r = 0.70$ vs $r = 0.61$), and diastolic blood pressure ($r = 0.35$ vs $r = 0.33$), and negatively with insulin sensitivity QUICKI ($r = -0.89$ vs $r = -0.82$) and total cholesterol/TG ratio ($r = -0.40$ vs $r = -0.22$). Glycemia, TG, and HDL-c were associated significantly only with SAD ($r = 0.31$; $r = 0.39$, $r = -0.43$, respectively).

Conclusion: Though the SAD and WC were associated with numerous metabolic abnormalities, only SAD correlated with dyslipidemia (TG and HDL-c) and hyperglycemia (glycemia).

(Nutr Hosp. 2011;26:1125-1129)

DOI:10.3305/nh.2011.26.5.5241

Key words: Sagittal abdominal diameter. Waist circumference. Hyperglycemia. Dyslipidemia. Inflammation. Anthropometric measurements.

DIÁMETRO ABDOMINAL SAGITAL, PERO NO LA CIRCUNFERENCIA DE LA CINTURA SE ASOCIA FUERTEMENTE CON LA GLUCEMIA, TRIACILGLYCEROLS Y HDL-C EN ADULTOS CON SOBREPESO

Resumen

Objetivo: Correlacionar el diámetro abdominal sagital (DAS) y la circunferencia de la cintura (CC) con las anomalías asociadas al síndrome metabólico en adultos.

Métodos: Este estudio transversal incluyó a 112 adultos (H = 27, M = 85) con edad de $54,0 \pm 11,2$ años y un promedio de índice de masa corporal (IMC) de $30,5 \pm 9,0$ kg/m². La evaluación incluía la presión sanguínea y medidas plasmáticas y antropométricas.

Resultados: Tanto en hombres como mujeres, DAS y CC se asociaban positivamente con el % grasa corporal ($r = 0,53$ vs $r = 0,55$), el ácido úrico ($r = 0,45$ vs $r = 0,45$), la us-PCR ($r = 0,50$ vs $r = 0,44$), la insulina ($r = 0,89$ vs $r = 0,75$), la resistencia a la insulina HOMA-IR ($r = 0,86$ vs $r = 0,65$), la LDL-ox ($r = 0,51$ vs $r = 0,28$), GGT ($r = 0,70$ vs $r = 0,61$), y la presión sanguínea diastólica ($r = 0,35$ vs $r = 0,33$), y negativamente con la sensibilidad a la insulina QUICKI ($r = -0,89$ vs $r = -0,82$) y el cociente colesterol total/TG ($r = -0,40$ vs $r = -0,22$). La glucemia, los TG, y la HDL-c se asociaban significativamente sólo con DAS ($r = 0,31$; $r = 0,39$, $r = -0,43$, respectivamente).

Conclusión: Aunque DAS y CC se asociaban con numerosas anomalías metabólicas, sólo DAS se correlacionaba con la dislipemia (TG y HDL-c) y la hiperglucemia (glucemia).

(Nutr Hosp. 2011;26:1125-1129)

DOI:10.3305/nh.2011.26.5.5241

Palabras clave: Diámetro abdominal sagital. Circunferencia de la cintura. Hiperglucemia. Dislipemia. Inflamación. Medidas antropométricas.

Correspondence: Gustavo Duarte Pimentel.
Faculdade de Ciências Médicas - FCM.
Cidade Universitária.
Tessália Vieira de Camargo, 126.
13084-971 Campinas/SP. Brazil.
E-mail: gupimentel@yahoo.com.br

Recibido: 2-II-2011.
1.ª Revisión: 6-III-2011.
Aceptado: 9-III-2011.

Introduction

The “gold standard” measurements of visceral or intraabdominal obesity are obtained by computed tomography, dual-energy-X ray absorptiometry, or magnetic resonance imaging. However, they are expensive and dual-energy-X ray absorptiometry and computed tomography involves radiation exposure. Though, it is impractical for epidemiologic purposes, in the context of primary care, or in clinic routine.^{1,2}

Indirect anthropometric estimates of body composition have proven usefulness for clinical practice and epidemiologic surveys because they are simple, noninvasive, and cheap. Several studies indicate that measures of abdominal fat are better predictors of metabolic syndrome (MS) and inflammation than total body adiposity as assessed by body mass index (BMI) in adults.³⁻⁷

In adult populations, the waist circumference (WC) is the most commonly used indicator of abdominal adiposity and is the main pathological finding in MS.^{8,9} However, the reliability of this measure in people with subcutaneous fat has been questioned¹⁰ because these individuals appear “tummy apron”.

Sagittal abdominal diameter (SAD) is highly correlated with visceral adipose tissue assessment by computed tomography.¹¹ Methodologically the SAD would be better measurement than the WC, because the sliding of subcutaneous fat to the waist sides when the measurement is taken orthostatically.^{3,5,12}

The aim of this study was to correlate two anthropometric measurements (WC and SAD) with MS-associated abnormalities in adults.

Materials and methods

Subjects and methods

This descriptive and cross-sectional study was conducted in patients clinically selected for lifestyle modification program “Mexa-se Pro-Saúde” (2006-2008). One hundred twelve patients (85 female and 27 male) participated of study. The criterion for exclusion was only subjects with liver, kidney, heart, or peripheral vascular disease, as well as chronic alcoholic.

This study was conducted according to the guidelines laid down in the Declaration of Helsinki and all procedures involving human subjects were approved by the Ethics Committee of Sao Paulo State University (UNESP, Brazil) n° 170/2005. Written informed consent was obtained from all subjects.

Anthropometric measurements

Height was measured to nearest 0.5 cm. Body weight was measured to the nearest 0.1 kg without shoes in light indoor clothing. BMI was calculated as the ratio of body weight (kg) divided by height (m)

squared. WC was measured in a supine position between the lowest rib and the iliac crest.¹³

Body fat percentage (%BF) was calculated from the resistance value (ohms) informed by bioelectric impedance analysis (BIA) (Biodynamics® 450 model) and subsequent application in the equation recommended by Segal et al.¹⁴ In order to reduce possible changes in water status, the participants were demanded to follow the recommendations: of avoiding drinking alcoholic beverages as well as caffeine for 24 hours before the test, fasting for 4 hours before the test, avoid intense exercising for at least 12 hours before the test, and let know about the use of medicine based on diuretics (in this case, the participants were not submitted to the test).¹³

The SAD was measured with a portable, sliding beam, abdominal caliper (Holtain, Ltd.; Dyfed, Wales, UK). The caliper’s upper arm was brought down to just above an abdominal mark made midway between the iliac crests, a location that approximates to the L4-L5 interspace. The subject was asked to inhale and exhale gently, and the arm of the caliper was brought down to touch the abdominal mark without compression.³

Clinical and biochemical measurements

Blood pressure was measured in the participant’s right arm after a 5 minutes rest by using an indirect auscultation with a mercury sphygmomanometer. Systolic and diastolic blood pressure was defined as Korotkoff phases 1 and 5, respectively.

Blood samples were drawn from an antecubital vein, and all serum and plasma samples were immediately chilled, kept on ice, centrifuged, and stored at -80°C until analyzed. Fasting glucose, total cholesterol, high-density lipoprotein (HDL-c), triacylglycerols (TG), γ -glutamyl transferase (GGT) and uric acid were quantified by commercial kits by enzymatic colorimetry assay (Labtest Diagnostica, MG, Brazil) in a semi-automatic spectrophotometry. Low-density lipoprotein (LDL-c) was calculated by the equation $LDL-c = total\ cholesterol - (HDL-c + TG/5)$ ¹⁵ and LDL-c subclass by the equation $(TG/HDL-c\ ratio)^{16}$ which is a good predictive factor for oxidized-LDL-c. Fasting insulin was assayed by immunochemical luminescence using commercial kits (DPC Medlab) in automated equipment (Immulite 2000R; DPC Medlab). The insulin resistance was calculated by the Homeostasis Model Assessment of insulin resistance (HOMA-IR) and for insulin sensitivity the QUICKI formula.¹⁷

Plasma by ultra-sensitivity C-reactive protein (us-CRP) was measured using Immulite Kit (DPCâ Medlab-Diagnostic Products Corporation, Los Angeles, CA).

Statistical analysis

Data are presented as means and standard deviations. The normality of the distribution within each sex

Table I
Characteristics of the study population

Variables	Men (n = 27)	Female (n = 85)	p value
Age (years)	53.2 ± 9.6	53.8 ± 10.4	0.82
Body mass index (kg/m ²)	31.3 ± 5.9	29.5 ± 6.0	0.28
Waist circumference (cm)	100.9 ± 14.6	95.7 ± 13.8	0.17
Sagittal abdominal diameter (cm)	23.7 ± 3.0	22.0 ± 4.0	0.14
Body fat (%)	33.5 ± 6.6	34.0 ± 7.4	0.85
Fasting glycemia (mg/dL)	111.0 ± 39.6	97.6 ± 37.2	0.25
Total cholesterol (mg)	194.5 ± 33.7	198.5 ± 36.3	0.83
HDL-cholesterol (mg)	49.2 ± 9.7	52.3 ± 11.5	0.39
LDL-cholesterol (mg)	119.6 ± 30.7	125.7 ± 34.0	0.59
Oxidized-LDL-c	3.7 ± 2.0	2.9 ± 2.4	0.30
Triacylglycerols (mg)	176.2 ± 65.5	147.8 ± 69.8	0.21
Total cholesterol/triglycerides ratio	1.4 ± 0.7	1.8 ± 0.9	0.22
Uric acid (mg/dL)	6.1 ± 1.8	4.5 ± 1.2	0.002*
C-reactive protein (mg/L)	0.45 ± 0.20	0.65 ± 0.84	0.47
Fasting insulin (ng/mL)	9.1 ± 4.6	9.5 ± 8.8	0.96
HOMA-IR	1.3 (0.3-6.8)	4.5 (1.4-20.0)	0.83
QUICKI	0.15 ± 0.02	0.15 ± 0.03	0.48
γ-glutamyl transferase (U/l)	31.5 ± 20.5	26.7 ± 13.5	0.64
Systolic blood pressure (mmHg)	122.6 ± 19.3	122.7 ± 14.5	0.97
Diastolic blood pressure (mmHg)	73.3 ± 10.5	77.0 ± 10.6	0.39

Data are presented as means and standard deviations. HOMA-IR: Homeostasis model assessment for insulin resistance.
*p < 0.05 vs men.

group was tested for all the variables using the Kolmogorov-Smirnov test. Pearson's correlations were used to verify a possible association between WC and SAD with metabolic abnormalities and comparisons between sexes were performed with Student's unpaired t test. All statistical analyses were performed by using SPSS for Windows (version 12.0; SPSS Inc, Chicago, IL). P < 0.05 was considered as statistically significant.

Results

Participants showed an age averaging (mean±Sd) 54.0 ± 11.2 years, BMI of 30.5 ± 9.0 kg/m², SAD of 22.4 ± 3.9 cm (23.7 ± 3.0 cm for male and 22.0 ± 4.0 cm for female, p = 0.14), and WC of 96.3 ± 13.7 cm (101.0 ± 14.8 cm for male and 95.4 ± 13.4 cm for female, p = 0.17) (table I). Uric acid was the only measure that differed between genders. Other general characteristics of subjects studied are shown in table I.

SAD and WC were associated positively (fig. 1) with body fat% (r = 0.53 vs r = 0.55), uric acid (r = 0.45 vs r = 0.45), us-PCR (r = 0.50 vs r = 0.44), insulin (r = 0.89 vs r = 0.75), HOMA-IR (r = 0.86 vs r = 0.65), oxidized-LDL-c (r = 0.51 vs r = 0.28), GGT (r = 0.70 vs r = 0.61), and diastolic blood pressure (r = 0.35 vs r = 0.33), and negatively (fig. 1) with insulin sensibility

QUICKI (r = -0.89 vs r = -0.82) and total cholesterol/TG ratio (r = -0.40 vs r = -0.22). Glycemia, TG, and HDL-c were associated significantly only with SAD (r = 0.31; r = 0.39, r = -0.43, respectively). Total cholesterol and LDL-c were not associated with neither adiposity indicators.

Discussion

The relative utility of several estimates of fat distribution has been controversial. Some investigators have proposed that WC is a better indicator of abdominal fat distribution than is waist-hip ratio, because it requires only one measurement and is more highly correlated with visceral fat^{9,18} and yet is suggest that predictive equation for evaluation of abdominal obesity based on bioelectrical impedance may be very useful in the clinical practice.¹⁹ SAD has been proposed to be even better than WC. However, no large or consistent difference between SAD and WC has been found in relation to visceral fat.¹⁸ The present study and others^{3,5,13} suggest that the SAD may be also a strong predictor of blood metabolic abnormalities.

Petersson et al.⁵ demonstrated that every one-centimeter increase in SAD was associated with an increase of CRP by 0.41 mg/l, corresponding to an increased mean CRP level by 16%. This estimation is

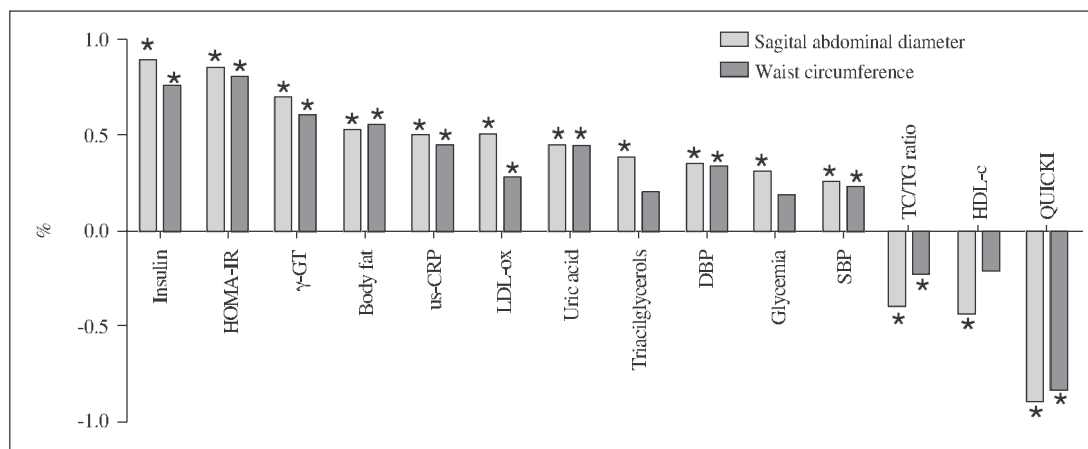


Fig. 1.—Correlation coefficients between anthropometric measurements (SAD and WC) and metabolic components. HOMA-IR: Homeostasis model assessment for insulin resistance, γ -GT: γ -glutamyl transferase, us-CRP: ultra-sensitivity C-reactive protein, LDL-ox: low-density lipoprotein cholesterol oxidized, DBP: diastolic blood pressure, TC/TG ratio: total cholesterol/triglycerides, HDL-c: high-density lipoprotein cholesterol, QUICKI: quantitative insulin sensitivity check index. * $p < 0.05$.

of clinical importance once elevated levels of serum CRP are associated with the MS and cardiovascular diseases.^{20,21} Other studies have obtained a stronger association for SAD to insulin resistance,²² cardiovascular risk⁴ and MS²² than WC, BMI and waist-hip ratio.

In a previous study²³ we established the cut-off points for SAD that corresponded to altered WC (WC > 102 cm for men and > 88 cm for women). The established points were 23.1 cm for men and 20.1 cm for women. Here we showed that either WC or SAD correlated well with plasma markers of metabolic abnormalities.

Recently, López de la Torre²⁴ demonstrated that both female and male and adults and elderly with high WC values are associated with diabetes. However, in the present study only SAD correlated with the MS components (TG, glycemia and HDL-c). Thus, the SAD could be seen as an appropriate method to be used for MS diagnosis purpose.

In summary, this study shown that both SAD and WC associated with numerous metabolic abnormalities. However, only SAD correlated with glycemia, TG and HDL-c, indicating that the SAD is a strong indicator of dyslipidemia and hyperglycemia. Thus, we suggesting that SAD measurement should be adopt in clinical practice and epidemiological studies.

Author contributions

The work presented here was carried out in collaboration between all authors. GDP and RCB defined the research theme. GDP, KCPM, MMT and RCB designed methods and experiments. GDP wrote the manuscript, analyzed the data and interpreted the results. FM performed the plasma biochemistry analysis. All authors have given final approval of submitted version.

Acknowledgements

This work was funded by grants from the Sao Paulo State Research Foundation (FAPESP, Brazil) and CAPES, Brazil. We declare that there are no conflicts of interest.

References

- Fuller NJ, Jebb SA, Laskey MA, Coward WA, Elia M. Four-component model for the assessment of body composition in humans: comparison with alternative methods, and evaluation of the density and hydration of fat-free mass. *Clin Sci* 1992; 82 (6): 687-93.
- Biaggi RR, Vollman MW, Nies MA, Brenner CE, Flakoll PJ, Levenhagen DK, et al. Comparison of air-displacement plethysmography with hydrostatic weighing and bioelectrical impedance analysis for the assessment of body composition in healthy adults. *Am J Clin Nutr* 1999; 69 (5): 898-903.
- Risérus U, Arnlov J, Brismar K, Zethelius B, Berglund L, Vessby B. Sagittal abdominal diameter is a strong anthropometric marker of insulin resistance and hyperproinsulinemia in obese men. *Diabetes Care* 2004; 27 (8): 2041-6.
- Öhrvall M, Berglund L, Vessby B. Sagittal abdominal diameter compared with other anthropometric measurements in relation to cardiovascular risk. *Int J Obes* 2000; 24 (4): 497-501.
- Petersson H, Daryani A, Risérus U. Sagittal abdominal diameter as a marker of inflammation and insulin resistance among immigrant women from the Middle East and native Swedish women: a cross-sectional study. *Card Diabetol* 2007; 6: 1-7.
- Brekke HK, Lenner RA, Taskinen MR, Mansson JE, Funahashi T, Matsuzawa et al. Lifestyle modification improves risk factors in type 2 diabetes relatives. *Diabetes Res Clin Prac* 2005; 68 (1): 18-28.
- Pimentel GD, Portero-McLellan KC, Oliveira ER, Spada AP, Oshiiwa M, Zemdeg JC et al. Long-term nutrition education reduces several risk factors for type 2 diabetes mellitus in Brazilians with impaired glucose tolerance. *Nutr Res* 2010; 30 (3): 186-90.
- Alberti KG, Zimmet P, Shaw J. The metabolic syndrome—a new worldwide definition. *Lancet* 2005; 366 (9491): 1059-62.

9. Grundy SM. Hypertriglyceridemia, insulin resistance, and the metabolic syndrome. *Am J Cardiol* 1992; 83 (9B): 25-9.
10. Nordhamn K, Södergren E, Olsson E, Karlström B, Vessby B, Berglund L et al. Reliability of anthropometric measurements in overweight and lean subjects: consequences for correlations between anthropometric and other variables. *Int J Obes* 2000; 24 (5): 652-7.
11. Schoen RE, Thaete FL, Sankey SS, Weissfeld JL, Kuller LH. Sagittal diameter in comparison with single slice CT as a predictor of total visceral adipose tissue volume. *Int J Obes Relat Metab Disord* 1998; 22 (4): 338-42.
12. Frenhani PB, Pimentel GD, Portero-McLellan KC, Burini RC. Sagittal abdominal diameter as a predictor of visceral abdominal fat, insulin resistance, dyslipidemia and inflammation in overweight Brazilian adults. *Clin Nutr Suppl* 2008; 3 (Suppl. 1): 126 [Abstract].
13. Heyward VH, Stolarczyk LM. Avaliação da composição corporal aplicada. 1.ed. Barueri: São Paulo, 2000.
14. Segal KR, Van Loan M, Fitzgerald PI, Hodgdon JA, Van Itallie TB. Lean body mass estimation by bioelectrical impedance analysis: a four-site cross-validation study. *Am J Clin Nutr* 1988; 47 (1): 7-14.
15. Friedewald T, Levy RI, Frederickson DS. Estimation of the concentration of low-density lipoprotein cholesterol in plasma, without use of the preparative ultracentrifuge. *Clin Chem* 1972; 18 (6): 499-502.
16. Tsimihodimos V, Gazi I, Kostara C, Tselepis AD, Elisaf M. Plasma lipoproteins and triacylglycerol are predictors of small, dense LDL particles. *Lipids* 2007; 42 (5): 403-9.
17. Levy J, Matthews DR, Hermans MP. Correct homeostasis model assessment (HOMA) evaluation uses the computer program. *Diabetes Care* 1998; 21 (12): 2191-2.
18. Clasey JL, Bouchard C, Teates CD, Riblett JE, Thorner MO, Hartman ML et al. The use of anthropometric and dual-energy X-ray absorptiometry (DXA) measures to estimate total abdominal and abdominal visceral fat in men and women. *Obes Res* 1999; 7 (3): 256-64.
19. Piernas Sánchez CM, Morales Falo EM, Zamora Navarro S, Garaulet Aza M. Study and classification of the abdominal adiposity throughout the application of the two-dimensional predictive equation Garaulet et al., in the clinical practice. *Nutr Hosp* 2010; 25 (2): 270-4.
20. Koenig W, Sund M, Frohlich M, Fischer HG, D Löwel H, Döring A et al. C-Reactive protein, a sensitive marker of inflammation, predicts future risk of coronary heart disease in initially healthy middle-aged men: results from the MONICA (Monitoring Trends and Determinants in Cardiovascular Disease) Augsburg Cohort Study, 1984 to 1992. *Circulation* 1999; 99 (2): 237-42.
21. Tracy RP, Lemaitre RN, Psaty BM, Ives DG, Evans RW, Cushman M et al. Relationship of C-reactive protein to risk of cardiovascular disease in the elderly. Results from the Cardiovascular Health Study and the Rural Health Promotion Project. *Arterioscler Thromb Vasc Biol* 1997; 17 (6): 1121-7.
22. Valsamakis G, Chetty R, Anwar A, Banerjee AK, Barnett A, Kumar S. Association of simple anthropometric measures of obesity with visceral fat and the metabolic syndrome in male Caucasian and Indo-Asian subjects. *Diabet Med* 2004; 21 (12): 1339-45.
23. Pimentel GD, Portero-McLellan KC, Maestá N, Corrente JE, Burini RC. Accuracy of sagittal abdominal diameter as predictor of abdominal fat among Brazilian adults: a comparison with waist circumference. *Nutr Hosp* 2010; 25 (4): 656-61.
24. López de la Torre M, Bellido D, Soto A, Carreira J, Hernández Mijares A. Standardisation of the waist circumference (WC) for each range of body mass index (BMI) in adult outpatients attended to in Endocrinology and Nutrition departments. *Nutr Hosp* 2010; 25 (2): 262-9.