



## Revisión

### Correlation and comparison between different measurement sites of waist circumference and cardiovascular risk in children: a systematic review and meta-analysis

*Correlación y comparación entre diferentes lugares de medición de la circunferencia de la cintura y el riesgo cardiovascular en niños: revisión sistemática y metaanálisis*

Angélica María Ballén-Torres<sup>1</sup>, María Lola Evia-Viscarra<sup>1</sup>, Rodolfo Guardado-Mendoza<sup>2</sup>, Daniela Beatriz Muñoz-López<sup>3</sup>, Edgar Efrén Lozada-Hernández<sup>4</sup>, Luis Fernando Meneses-Rojas<sup>5</sup>

<sup>1</sup>Department of Pediatric Endocrinology. Servicios de Salud del Instituto Mexicano del Seguro Social para el Bienestar (IMSS-BIENESTAR). Hospital Regional de Alta Especialidad del Bajío; <sup>2</sup>Metabolic Research Laboratory. Department of Medicine and Nutrition. Universidad de Guanajuato; <sup>3</sup>Department of Medicine and Nutrition. Universidad de Guanajuato. <sup>4</sup>Department of Research. Hospital Regional de Alta Especialidad del Bajío; <sup>5</sup>Department of Gastroenterology. Hospital General de Zona 21. Instituto Mexicano del Seguro Social. León, Guanajuato. Mexico

### Abstract

**Background:** waist circumference (WC) is a component of metabolic syndrome (MetS) and an excellent marker for the risk of cardiovascular disease (CVD) in children. This study aimed to provide information on the anatomical measurement sites of WC and their comparative correlation with MetS and its components in children.

**Methods:** the literature search included papers published between January 2005 and September 2023 that met the following criteria: pediatric patients (2-18 years), WC measurement at different anatomical sites ( $\geq 2$ ), and CVD risk by MetS. The quality of each study was determined using the STROBE and modified GRADE scales. The meta-analysis evaluated the WC<sub>iliac-crest</sub> and WC<sub>middle</sub>.

**Results:** five observational studies (total population: 1,224) were included. WC was measured at 2-4 anatomical sites. In all studies, the correlations between different WC measurement sites and CVD risk were similar. The STROBE assessment ranged from 12-20/22 and the GRADE was A for all the articles. The meta-analysis showed that the heterogeneity ( $I^2$  test) of the WC<sub>iliac-crest</sub> and WC<sub>middle</sub> with CVD variables was substantial.

**Conclusion:** All WC measurement sites showed adequate correlation with CVD risk, with some small individual differences. WC<sub>narrow</sub> and WC<sub>umbilicus</sub> have adequate consistency and could be excellent alternatives in daily clinical practice because of their ease of measurement. Further studies are needed to evaluate the correlation between different WC measurement sites and CVD risk in children stratified according to pubertal stage and sex.

#### Keywords:

Waist circumference.  
Pediatric obesity.  
Cardiovascular risk.  
Metabolic syndrome.

Received: 18/01/2024 • Accepted: 21/04/2024

*Author contributions:* MLEV and AMBT conceived of and designed the study. AMBT collected the data for this study. RGM, DBML, and LFMR evaluated the studies included in this review. MLEV and EELH performed statistical analyses. AMBT, MLEV and LFMR wrote the manuscript.

*Funding statement:* the authors received no specific funding for this study.

*Conflicts of interest:* the authors declare no conflicts of interest.

*Artificial intelligence:* the authors declare not to have used artificial intelligence (AI) or any AI-assisted technologies in the elaboration of the article.

Ballén-Torres AM, Evia-Viscarra ML, Guardado-Mendoza R, Muñoz-López DB, Lozada-Hernández EE, Meneses-Rojas LF. Correlation and comparison between different measurement sites of waist circumference and cardiovascular risk in children: a systematic review and meta-analysis. *Nutr Hosp* 2024;41(5):1105-1115  
DOI: <http://dx.doi.org/10.20960/nh.05144>

#### Correspondence:

María Lola Evia-Viscarra. Department of Pediatric Endocrinology. Servicios de Salud del Instituto Mexicano del Seguro Social para el Bienestar (IMSS-BIENESTAR). Hospital Regional de Alta Especialidad del Bajío. Blvd. Milenio, 130; San Carlos la Roncha. 37544 León, Guanajuato. Mexico  
e-mail: [evialola@hotmail.com](mailto:evialola@hotmail.com)

## Resumen

**Antecedentes:** la circunferencia de la cintura (CC) es un componente del síndrome metabólico (SM) y un excelente marcador de riesgo cardiovascular (RCV). El objetivo de este estudio fue proporcionar información sobre los sitios de medición anatómica de la CC en niños y su correlación comparativa con el SM y sus componentes.

**Métodos:** búsqueda bibliográfica incluyó artículos entre enero 2005 y septiembre 2023 con los siguientes criterios: niños (2-18 años), CC medida en  $\geq 2$  sitios anatómicos y SM. La calidad de cada estudio se evaluó con las escalas STROBE y GRADE modificada. El metaanálisis evaluó la CC cresta iliaca y CC media.

**Resultados:** se incluyó cinco estudios observacionales (población total: 1224). Todos los estudios mostraron similares correlaciones entre los diferentes sitios de medición de CC y el RCV. La evaluación STROBE fue de 12-20/22 y GRADE fue A en todos los artículos. El metaanálisis mostró que la heterogeneidad (prueba  $I^2$ ) de la CC cresta iliaca y la CC media con las variables de RCV fue significativa.

**Conclusión:** todos los sitios de medición de la CC mostraron una correlación adecuada con el RCV, con algunas pequeñas diferencias. CC estrecha y CC umbilical tienen una consistencia adecuada y podrían ser excelentes alternativas en la práctica clínica diaria debido a la facilidad de medición. Se necesitan más estudios para evaluar la correlación entre diferentes sitios de medición de la CC y el riesgo de RCV en niños estratificados según la etapa puberal y el sexo.

### Palabras clave:

Circunferencia de la cintura. Obesidad pediátrica. Riesgo cardiovascular. Síndrome metabólico.

## INTRODUCTION

Waist circumference (WC) is the main indicator of abdominal adiposity and reflects the amount of visceral adipose tissue (VAT). Therefore, it is considered the best measurement for detecting patients at risk of cardiovascular disease (CVD) (1). CVD risk assessment in children with obesity has gained relevance because it may predict increased mortality in adulthood owing to coronary heart disease and stroke (2). Obesity in children is associated with type 2 diabetes *mellitus* (T2DM), dyslipidemia, arterial hypertension, non-alcoholic fatty liver disease, genu valgum, and obstructive sleep apnea (3). The pathophysiological mechanisms that lead to an increased risk of developing CVD with early atherosclerosis in patients with obesity (4) are related to increased insulin resistance (IR) (5) and activation of chronic inflammation (6). The CVD risk in children is assessed based on the presence of metabolic syndrome (MetS). The “Third Report of the National Cholesterol Education Program (NCEP) in Adults Panel of Treatment III” (7) defined MetS when a patient has three of the following components: abdominal obesity measured by WC, increased triglyceride (TG), decreased high-density cholesterol (HDL), hypertension, and fasting hyperglycemia or T2DM (8).

The detection of CVD risk factors is performed with WC measurement in the routine physical exam. Although this evaluation is practical and simple; several recommendations should be made for each anatomical measurement site (Fig. 1). The WC at the narrowest visual abdominal part ( $WC_{\text{narrow}}$ ) was initially described by Lohman et al. (9) and later by the International Society for the Advancement of Kinanthropometry (ISAK) (10), the WC at the midpoint between the lower rib and the top of the iliac crest ( $WC_{\text{midline}}$ ) by the World Health Organization (WHO) (11), and the WC above the border of the iliac crest ( $WC_{\text{iliac-crest}}$ ) by the National Health and Nutrition Examination Survey (NHANES) (12). WC at the level of the umbilicus ( $WC_{\text{umbilicus}}$ ) was described by Croft et al. (13) and Eisenmann et al. (14).  $WC_4$  was unusual and described by Rudolf et al. (15).

Interestingly, the anatomical WC measurement site was not based on comparative correlation studies between different WC measurement sites and cardiovascular risk. The objective of this review is to evaluate anatomical WC measurement sites and their comparative correlation with other MetS components in children.

## METHODS

The study protocol was registered in the Prospero ID CRD42023454847.

### SEARCH STRATEGY

The literature search was conducted by AMBT, without language restriction in Lilacs, MEDLINE/PubMed, Web of Science, and Scopus databases on October 2023. Papers published between January 2005 and August 2023 were included. The PICO framework was used to develop search strategies and ensure comprehensive and bias-free searches: Population: Patients between 2-18 years. Intervention: WC measurement. Comparison: WC measurement at different anatomical sites ( $\geq 2$ ). Diagnostic outcomes: CVD risk or other MetS components.

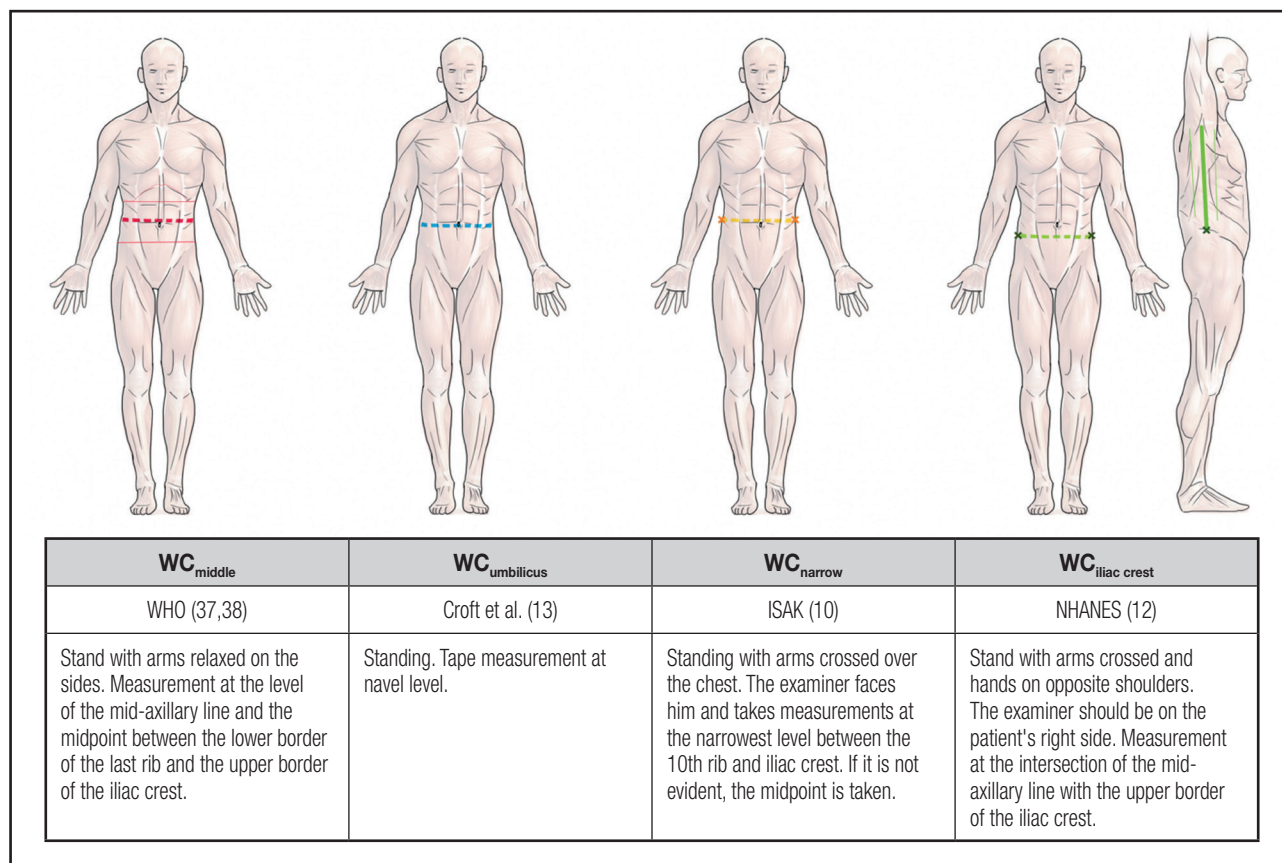
The combination of the boolean descriptors were: “Waist circumference” AND (“measurement anatomical sites” OR “Waist circumference AND pediatric OR cardiometabolic risk factor”). “Waist circumference” AND (“measurement anatomical sites” OR “Waist circumference AND pediatric OR metabolic syndrome”). “Waist circumference measurement sites” and “metabolic syndrome.”

### ELIGIBILITY CRITERIA

Observational studies that had  $\geq 2$  anatomical sites of WC and their correlation with MetS or other components of MetS.

### STUDY QUALITY AND RISK OF BIAS ASSESSMENT

Eligible studies were assessed by two investigators independently (RGM and DBML). Any divergence was resolved by a third evaluator (LFMR). The quality of each study was determined using the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) checklist (22 items) (16). The modified



**Figure 1.**

WC measurement recommendations (WHO: World Health Organization; ISAK: International Standards for Anthropometric Assessment, United Kingdom; NHANES: National Health and Nutrition Examination Survey).

Grading of Recommendations Assessment, Development, and Evaluation (GRADE) scale (Table I) was applied, considering A (good), B (moderate), and C (low) when the paper characteristics were complete, partial, or non-specific, respectively.

**THE STRATEGY OF DATA SYNTHESIS**

The methodology of the systematic review followed the Cochrane Manual guidelines (17), and was adjusted according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) 2020 guidelines (18).

**STATISTICAL ANALYSES**

Meta-analysis was performed with Jamovi 2.2.5 version (by EELH and MLEV) using WC coefficients of correlation with components of MetS included in 4/5 articles. Bosy-Westphal et al. (19) were excluded because they included both children and adults. The WC measurements included in the meta-analysis were the WC<sub>iliac-crest</sub> and WC<sub>middle</sub>. Other WC measurements were

**Table I. GRADE evaluation of the of the scientific articles**

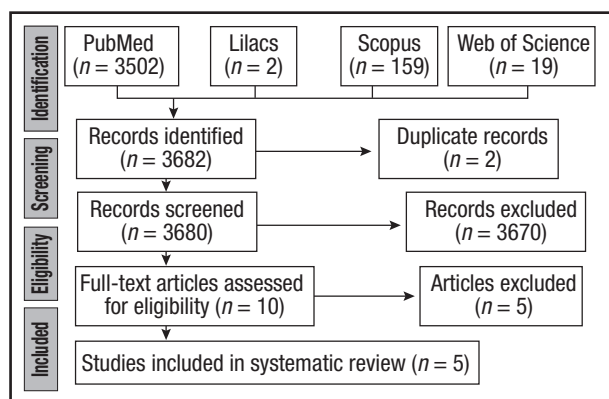
| GRADE score for the characteristics of the paper:<br>A. Completely described<br>B. Partially described<br>C. Not described |
|--|
| Characteristics described in the paper:  |
| Inclusion and exclusion criteria   |
| Methods of sample selection  |
| Stratified by sex, social group, or lifestyle  |
| Baseline valued described  |
| WC measurement method description  |
| Definition of cardiovascular risk or MetS and its components   |
| Bias or confounders taken in account   |
| Statistical analysis applied   |

GRADE: Grading of Recommendations, Assessment, Development and Evaluation; WC: waist circumference; MetS: metabolic syndrome.

excluded because there were only two publications of each one. A random effects model was used to fit the data. Analysis was performed using the Fisher  $r$ -to  $Z$ -transformed correlation coefficient. Heterogeneity was estimated using the Cochran  $Q$  test and  $I^2$  statistic. If  $I^2 > 50\%$  or  $p < 0.1$ , heterogeneity of the results was considered. Studentized residuals and Cook distances were used to determine whether the studies were outliers or influential in the model context. Studies with a studentized residual larger than the  $100 \times (1 - 0.05 / [2 \times k])$  th percentile of a standard normal distribution were considered potential outliers. Studies with Cook's distance greater than the median and six times the interquartile range of Cook's distance were considered influential. The Begg and Muzumbar rank correlation test and Egge's regression test, using the standard error of the observed outcomes as a predictor, were used to verify the funnel plot asymmetry.

## RESULTS

From 3,680 non-duplicate records, ten studies were selected because they evaluated  $\geq 2$  WC anatomical sites. After the full texts were reviewed, five studies were excluded because they did not include MetS components (Fig. 2). Finally, we included five studies with 1,224 children (5-18 years old) of both sexes: Hitze et al. (20) and Bosy-Westphal et al. (19) (both in the German group), Johnson et al. (21), Harrington et al. (22), and López et al. (23). Table II presents the characteristics of the studies. Bosy-Westphal et al. studied prepubertal and pubertal children and adults. Except for the study by López et al. (23), all other studies divided their results by sex. Harrington et al. (22) assessed children according to ethnicities(21). Each article measured WC at to 2-4 anatomical sites, which showed adequate reproducibility. WC of the inferior margin of the ribs ( $WC_{rib}$ ) was measured in the German group (19,20). The WC at the level of the umbilicus ( $WC_{umbilicus}$ ) and  $WC_{narrow}$  was measured by Johnson et al. (21) and Harrington et al. (22). WC was measured four centimeters (cm) above the umbilical scar ( $WC_u$ ) and was evaluated by Hitze et al. (20). The  $WC_{iliac-crest}$  and  $WC_{middle}$  were measured in all studies included in this review.



**Figure 2.**

Flow diagram of search results.

All studies evaluated WC while standing but with different arm positions: hanging freely (19,20) or crossing over the chest (21).

## STATISTICAL EVALUATION IN THE INCLUDED STUDIES

Pearson's correlation was performed between each WC measurement site and MetS components in all articles. In some cases, adjustments for age (19,22), ethnicity (21), or logarithmic transformation of variables are necessary (19-23). The prevalence of obesity and MetS differed between these studies (21-23). Johnson et al. (21) described a different prevalence of MetS according to each definition criterion: the modified National Cholesterol Education Program (NCEP) (24), International Diabetes Federation (IDF) (25), and Cook et al. (26). López et al. (23) described MetS according to the IDF (Table II).

## DIFFERENCES BETWEEN WC MEASUREMENT SITES

The studies showed that the magnitude of WC (cm) was different in measurement sites and that there were inherent to sex, ethnicity and puberal stage (20). The correlation between the magnitude of all WCs was strong ( $r = 0.93$ - $0.99$ ).

## WC MEASUREMENT SITES AND METS

In Hitze et al. (20), in the female (F) group, all WC measurements showed positive correlations with systolic blood pressure (SBP), diastolic blood pressure (DBP), TG, glucose, and Homeostatic Model Assessment for Insulin Resistance (HOMA-IR), but they did not show any correlation with total cholesterol (TC) and low-density cholesterol (LDL). In the male (M) group, all WC measurements were positively correlated with DBP, LDL, and HOMA-IR. They did not show any correlation with the SBP, TG, TC, or glucose levels.

Bosy-Westphal et al. (19) showed that the WC measurement correlation at all anatomical measurement sites was adequate with abdominal fat, but it was better with subcutaneous adipose tissue (SAT) than with VAT. In prepubertal M, the relationship between the  $WC_{iliac-crest}$  with VAT and HOMA-IR was lower than that in the other WC.

In Johnson et al. (21) the correlation coefficient showed a significant positive association between all WC measurement sites and the SBP, DBP, and HOMA-IR. The TG levels were positively correlated with  $WC_{narrow}$  and  $WC_{middle}$ . Correlation between CVD risk variables and  $WC_{narrow}$  or  $WC_{middle}$  was slightly higher. The correlation between MetS and CVD risk was similar for all WC measurements, with only differences. According to the IDF definition of MetS,  $WC_{narrow}$  and  $WC_{middle}$  were significantly associated with MetS, and the number of MetS components. According to Cook's



definition of MetS, there was no association between MetS and all WC; however,  $WC_{\text{narrow}}$  and  $WC_{\text{middle}}$  showed significant odds ratios (OR) with the number of MetS components. For the definition of MetS according to the NCEP, an association was observed between MetS and  $WC_{\text{narrow}}$  or  $WC_{\text{middle}}$  but not with  $WC_{\text{iliac-crest}}$  or  $WC_{\text{umbilicus}}$ .

Harrington et al. (22) evaluated the age-controlled correlation between WC and logVAT was significant in all the groups. The correlation between all anatomical measurement sites and MetS components was good, except for the glucose levels in the white-M and AA-F groups. There was no correlation between any WC measurement site and DBP in the white-M.

López et al. (23) reported a statistically significant correlation between  $WC_{\text{middle}}$  and  $WC_{\text{iliac-crest}}$  and SBP, DBP, TG, and HDL levels. Glucose levels showed a low correlation with all WCs measurements.

The STROBE scale yielded a result between 12 and 20 points for 22 items. The modified GRADE scale results for all included articles were A (good evidence) (Table II).

## META-ANALYSIS

Figure 3 shows the results of the meta-analysis. The random-effects model showed that the correlation of  $WC_{\text{iliac-crest}}$  and  $WC_{\text{middle}}$  with the average variables of MetS differed significantly from 0. The  $WC_{\text{iliac-crest}}$  and  $WC_{\text{middle}}$   $I^2$  test in all evaluations with the CVD risk variables showed heterogeneity: HDL (66.4 %,  $p = 0.028$  and 69.12 %,  $p = 0.019$ ), TG (77.96 %,  $p = 0.006$  and 68.78 %,  $p = 0.028$ ), SBP (94.68 %,  $p < 0.001$  and 95.7 %,  $p < 0.001$ ), and DBP (90.09 %,  $p < 0.001$  and 90.56 %,  $p < 0.001$ ). Glucose presented low heterogeneity in both WC (0 %,  $p < 0.634$ ; 0 %,  $p < 0.722$ ). In the studentized analysis, López et al. (23) presented atypical values for HDL, SBP, and DBP for  $WC_{\text{middle}}$  and  $WC_{\text{iliac-crest}}$ . Glucose analysis in both WC and TG in  $WC_{\text{middle}}$  showed no outliers. Hitze et al. (20) showed the possibility of an outlier in the correlation between  $WC_{\text{middle}}$  and TG levels. Cook's evaluation showed that none of the studies could be considered influential on any of the variables studied. Egge's regression analysis and Begg's rank correlation tests did not indicate asymmetry in the components of MetS evaluated; therefore, the construction of the funnel graph (Fig. 3) with a few studies generated the possibility of bias.

## DISCUSSION

This review shows that the studies included had adequate comparative correlation between all WC measurement sites and other MetS components except glucose which shows a low correlation. It provides an advantage because measurements are easier at certain anatomical points depending on the characteristics of each patient.

The correlation coefficient evaluation in this meta-analysis shows no difference in the correlation between  $WC_{\text{middle}}$  or

$WC_{\text{iliac-crest}}$  and other MetS components. Studies carried out in different countries have shown that measuring  $WC_{\text{middle}}$  can predict the presence of MetS in pediatric patients (27,28). WC is considered a good predictor of MetS because it is positively correlated with MetS (29). A review in adult patients reported that there was no substantial difference in the WC measurement site protocols in terms of cardiovascular morbidity, and mortality (30).

We were not able to perform stratified analyses by sex, age, pubertal stage or ethnicity, because data were insufficient. Children are in constant development: this modifies the correlation between WC measurements at different anatomical sites and other MetS components. Fat distribution is very similar between girls and boys in their first childhood years, and then changes at puberty, the beginning of sexual development (31,32). Other studies have already shown these differences by sex in abdominal fat distribution, and some indicate that the presence of obesity does not seem to modify this distribution; that is, the  $WC_{\text{narrow}}$  is the smallest and the  $WC_{\text{umbilicus}}$  is the largest (33). Furthermore, it is possible that the fat deposits distribution by ethnic group may contribute to different cardio-metabolic risk (34).

Some studies included in this review defined the overwaist at all anatomical sites using percentiles created for  $WC_{\text{middle}}$ . This may have modified their results because each WC anatomical measurement site should have specific percentile value. This is important when evaluating a clinical measurement that constitutes a diagnostic tool for MetS, and to avoid bias. We did not find percentile values for  $WC_{\text{rib}}$  in the literature, and there are only few population studies for the  $WC_{\text{umbilicus}}$  (14) and  $WC_{\text{narrow}}$  (35,36).

In children and adolescents, all definitions of MetS (IDF, Cook, and ATP III) consider the same components, but there is great variability due to lack of standardization of the cut-off points. Similarly, the frequency of  $WC \geq 90^{\text{th}}$  percentile varies according to the WC measurement site and the MetS definition used, modifying therapeutic decisions. The lack of consensus on the WC measurement site in children underestimates or overestimates the CVD risk, as reported in research studies on MetS.

All studies reviewed followed appropriate measurement techniques and standardization, although each author differed in the arms position. The subject's position standing upright with arms relaxed on both sides was described by Lohman et al. (9) to measure  $WC_{\text{narrow}}$  and by the WHO to measure  $WC_{\text{middle}}$  (37,38). Patients standing with arms crossed over their chest and hands on their shoulders were described by NHANES (12) to measure the  $WC_{\text{iliac-crest}}$  and by ISAK (10) to measure  $WC_{\text{narrow}}$  (20-22). Thus, Lennie et al. (39) found significant differences in WC measurements performed at different positions in adults. There have been no studies on this topic in the pediatric population. In the smallest or very restless children, arms crossed on the chest and hands on the shoulders are more comfortable and provide more stability.

Technical difficulties in locating bony anatomical references in children with obesity are uncomfortable for patients (15); therefore,  $WC_{\text{narrow}}$  and  $WC_{\text{umbilicus}}$  facilitate waist measurement. Some children with significant abdominal subcutaneous tissue have a hanging position at the umbilicus, which modifies the umbilical scar position.

**Table II. Structured summary of the results of the included studies**

| Author, year                   | Hitze et al., 2008  | Bosy-Westphal et al., 2009  | Johnson et al., 2010   | Harrington et al., 2013  | López et al., 2016  |
|--------------------------------|---|---|--|--|---|
| <b>Journal name (JRI)</b>      | <i>Obes Facts</i> (3.9)   | <i>J Nutr</i> (4.2)   | <i>J. Pediatr</i> (3.7)  | <i>Pediatr Obes</i> (3.42)   | <i>Endoc Pract</i> (3.86)   |
| <b>Country-City</b>            | Germany-Kiel  | Germany-Kiel  | Canada-Edmonton  | USA-Louisiana  | Mexico-CDMX   |
| <b>n</b>                       | 180   | 234 (prepuberal 74, puberal 160)  | 73   | 371 (White 178, AA 193)  | 366   |
| <b>Inclusion criteria</b>      | Age (median or range)<br>13.2 ± 3.7 years<br>% Female<br>50.5   | 11.97 years (prepuberal, 9.05 years; puberal, 14.9 years)<br>50.5   | 8 to 17 years<br>56.2  | 5 to 18 years<br>52.83   | 10 to 18 years<br>48.9  |
| <b>Exclusion criteria</b>      | Use of drugs that influence the results   | Use of drugs that influence the results   | Not indicated  | Race other than white or AA  | Genetic or endocrine obesity and drugs that influence the results   |
| <b>Measurement sites of WC</b> | WC <sub>fb</sub>  | ✓   | ✓  |  |   |
|                                | WC <sub>middle</sub>  | ✓   | ✓  | ✓  | ✓   |
|                                | WC <sub>iliac crest</sub>   | ✓   | ✓  | ✓  | ✓   |
|                                | WC <sub>narrow</sub>  |   |  | ✓  | ✓   |
|                                | WC <sub>umbilicus</sub>   |   |  | ✓  | ✓   |
| WC <sub>4</sub>                | ✓   |   |  |  |   |
| <b>Anthropometric tape</b>     | Nonelastic (no brand indicated)   | Nonelastic (no brand indicated)   | Spring-loaded (FitSystems) (Calgary, Alberta, Canada)  | Not indicated  | Nonstretchable fiberglass (no brand indicated)  |
| <b>Position of the patient</b> | Stand with arms hanging freely  | Standing  | Standing with arms crossed over the chest. Mirror to see that the tape measure did not slip down the back                      | Standing with arms relaxed at their sides                                | WHO and NHANES measuring method   |
| <b>WC measurement</b>          | 4 trained observers. Each observer simultaneously performed the measurements of the 4 sites. Intraobserver CV: WC <sub>narrow</sub> 0.6 %, WC <sub>4</sub> 1.5 %, WC <sub>middle</sub> 1.1 %, WC <sub>iliac crest</sub> 0.7 %. Inter-observer CV: WC <sub>narrow</sub> 1.9 %, WC <sub>middle</sub> 1.9 %, WC <sub>iliac crest</sub> 3.1 % | 4 trained nutritionists. CV intraobserver e interobserver: WC <sub>narrow</sub> 0.59 y 1.29 %; WC <sub>iliac crest</sub> 1.43 y 2.64 % y WC <sub>middle</sub> 1.19 y 2.52 % | The same clinical performed all measurements of WC sequentially 2 times. If they differed > 0.5 cm, a third sequence was taken | 3 trained technicians. Interobserver and intraobserver CV: 0.98 and 0.99 | Measurements were performed by a pediatric obesity specialist and a pediatric endocrinologist. Interobserver CV ±0.41 cm with kappa 0.95-0.98 at the different WC sites |

(Continues on next page)

Table II (Cont.). Structured summary of the results of the included studies

| Author, year                              | Hitze et al., 2008                             | Bosy-Westphal et al., 2009  | Johnson et al., 2010  | Harrington et al., 2013  | López et al., 2016   |
|---|--|---|---|--|--|
| <b>Definition WC &gt; 90<sup>th</sup></b> | WC <sub>midle</sub> (31)                       | Not considered  | Fernández et al. (32) (WC <sub>lic-crest</sub> )  | Fernández et al. (32) (WC <sub>lic-crest</sub> )   | Fernández et al. (32) (WC <sub>lic-crest</sub> ) <sup>a</sup><br>Klinder et al. (33) (W <sub>midle</sub> )   |
| <b>Definition of Mets</b>                 | Not assessed                                   | Not assessed  | Mets and its components were evaluated without WC with the definitions of IDF, NCEP and by Cook et al.  | Mets components: HDL ≤ 45 mg/dL, TG ≥ 75 mg/dL (5-9 years) or ≥ 90 mg/dL (10-18 years). Fasting hyperglycemia ≥ 100 mmol/L. Hypertension = SBP or DBP ≥ 90th for age, sex, and height. Defined Mets when they had ≥ 2 components except WC | Mets according to IDF if they had 3 components (except WC): TG ≥ 150 mg/dL, HDL < 40 mg/dL (M and F) and in ≥ 16 years in M < 40 mg/dL and in F < 50 mg/dL, glucose ≥ 100 mg/dL, arterial hypertension if SBP and/or DBP ≥ 130/85 mmHg |
| <b>Study design</b>                       | Cross-sectional, observational                 | Cross-sectional observational. Children and adults analyzed comparatively and separately  | Cross-sectional observational in children with obesity (BMI ≥ 85 <sup>th</sup> )  | Cross-sectional, observational   | Cross-sectional, observational in children with and without obesity  |
| <b>Sample size calculation</b>            | Not indicated                                  | Not indicated   | Not indicated   | 400  | Not indicated  |
| <b>Statistical analysis</b>               | Correlation of each WC with components of Mets | The strength of the correlation coefficients was compared with Meng's method. For the correlation between WC and CVD risk, the data were adjusted for age | Pearson correlation. Evaluated the association between BMI and Mets components. Both controlled for age, sex, and ethnicity                               | Pearson correlation, controlled for age  | Pearson correlation  |
|   | Others   | TG, HOMA-IR levels were normalized by logarithmic transformation  | Logistic regression to calculate the OR of Mets and increase in its components related to WC and BMI Z. All variables were adjusted for age and ethnicity | The difference between WC in cm was compared by repeated-measures ANOVA and Turkey post-hoc tests. VAT was adjusted  | We determined the ROC curve to see the WC ≥ P 90 <sup>th</sup> of both measurements and the waist-to-height ratio > 0.5 with the CVD risk variables  |
| <b>Prevalence of overweight/obesity</b>   | 12.2 % with overweight (F:11 %, M:13.5 %)      | Puberty: F: 26.1 %, M: 26.3 %. Prepuberty: F: 0 %, M: 7.7 %   | All with BMI > 85 <sup>th</sup> , 95 % of children with BMI > 95 <sup>th</sup> )  | Obesity according: IOTF: 28 %, CDC: 33.4 %   | Overweight: 24 %. Obesity: 55 %  |
| <b>Prevalence of Mets</b>                 | Not evaluated                                  | Not evaluated   | NCEP: F: 39 %, M: 34 %. IDF: F: 43.9 %, M: 50 %. Cook: F: 39 %, M: 56.8 %   | 13.8 % (M White: 19 %, M AA: 10 %, F White: 16.9 % a F AA: 9.7 %)  | IDF: 32.80 %   |

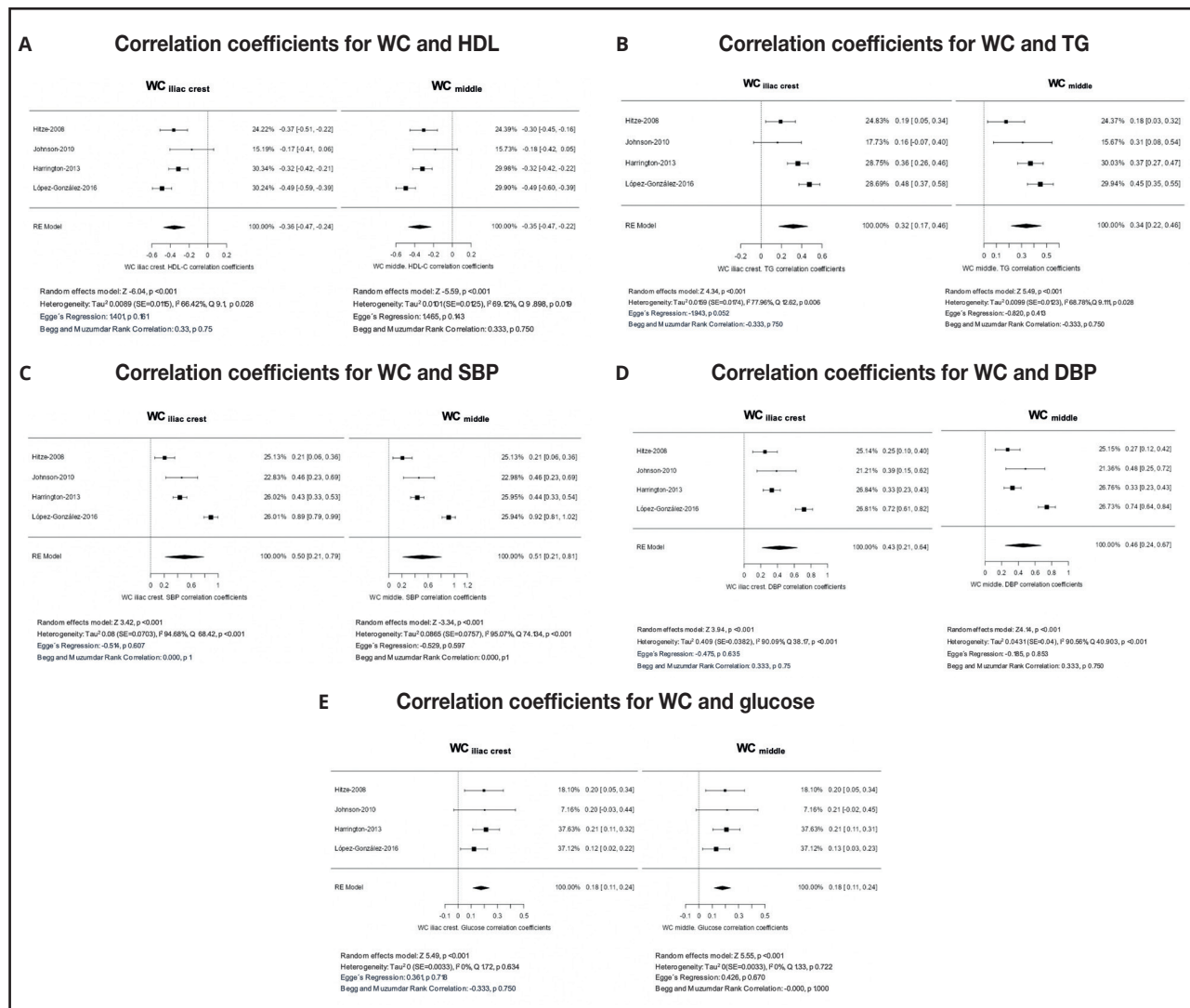
(Continues on next page)

Table II (Cont.). Structured summary of the results of the included studies

| Author, year                                       | Hitze et al., 2008   | Bosy-Westphal et al., 2009  | Johnson et al., 2010  | Harrington et al., 2013  | López et al., 2016  |
|--|--|---|---|--|---|
| <b>Differences between WC measurement sites</b>    | <p>WC<sub>fb</sub> &lt; WC<sub>s</sub> &lt; WC<sub>middle</sub> &lt; WC<sub>lax-crest</sub> (r &gt; 0.93, p &lt; 0.01). WC<sub>fb</sub> and WC<sub>s</sub> (cm) were lower in F.</p> <p>WC<sub>middle</sub> and WC<sub>lax-crest</sub> did not show differences between sexes.</p> <p>Prevalence WC ≥ 90<sup>th</sup> F and M was: WC<sub>fb</sub>: 13.2 % and 15.7 %; WC<sub>s</sub>: 14.3 % and 19.1 %; WC<sub>middle</sub>: 18.7 % and 22.5 % and WC<sub>lax-crest</sub>: 37.4 % and 30.3 %</p>   | <p>WC (cm) in prepuberty and puberty was: WC<sub>fb</sub> &gt; WC<sub>middle</sub> &gt; WC<sub>lax-crest</sub> (p &lt; 0.001)</p>   | <p>WC (cm) by sex:<br/>M: WC<sub>umbilicus</sub> &gt; WC<sub>lax-crest</sub> &gt; WC<sub>middle</sub> &gt; WC<sub>narrow</sub></p> <p>WC<sub>umbilicus</sub> vs WC<sub>lax-crest</sub> (p &lt; 0.01), or vs WC<sub>umbilicus</sub> (p &lt; 0.08) and not significant vs WC<sub>middle</sub>.</p> <p>F: WC<sub>umbilicus</sub> &gt; WC<sub>lax-crest</sub> &gt; WC<sub>middle</sub> &gt; WC<sub>narrow</sub>. WC<sub>umbilicus</sub> was significantly different from all WC.</p> <p>Correlation between all WC: r = 0.93-1, p &lt; 0.0001</p>   | <p>WC (cm): White F WC<sub>umbilicus</sub> &gt; WC<sub>lax-crest</sub> &gt; WC<sub>middle</sub> &gt; WC<sub>narrow</sub> (p &lt; 0.05).</p> <p>White M and AA F no significant difference between WC<sub>lax-crest</sub> and WC<sub>umbilicus</sub></p> <p>AA M WC<sub>umbilicus</sub> was statistically smaller. WC ≥ 90<sup>th</sup> = WC<sub>umbilicus</sub>: 20.7 %, WC<sub>middle</sub>: 29.6 %, WC<sub>lax-crest</sub>: 31.5 %, and WC<sub>narrow</sub>: 31.1 % (evaluated with percentile values of WC<sub>lax-crest</sub>)</p> | <p>All group: High SBP 7.7 %, fasting hyperglycemia 10.7 %, high TG 37.4 %, low HDL 56.4 %</p>  |
| <b>Correlation of each WC with MetS components</b> | <p>F: All WC had positive correlation with SBP, DBP, TG, glucose, HOMA-IR and negative correlation with HDL. Not correlated: TC, y LDL.</p> <p>M: All WC had positive correlation with DBP, LDL, HDL and HOMA-IR. Not correlation: SBP, TG, TC and glucose.</p> <p>The difference between correlation coefficients was significant (p &lt; 0.05) in:<br/>F: WC<sub>lax-crest</sub> vs WC<sub>fb</sub> with the variable TG.<br/>M: WC<sub>lax-crest</sub> vs WC<sub>fb</sub>, WC<sub>s</sub>, and WC<sub>middle</sub> with LDL showed positive correlation (p &lt; 0.01)</p> | <p>In all groups the correlation between all WC with SAT (r = 0.65-0.76) showed better correlation than VAT (r = 0.75 to 0.89).</p> <p>Correlation in prepubertal and pubertal between all WC with VAT was 0.73-0.87, with SBP 0.73-0.93 and similar with cardiovascular risk (numerical data not shown in the paper).</p> <p>Prepubertal M: correlation between WC<sub>lax-crest</sub> with VAT (r = 0.65) and HOMA-IR (r = 0.13, p &lt; 0.05); were lower than the other WC (WC<sub>middle</sub> with VAT r = 0.76, and WC<sub>umbilicus</sub> with HOMA-IR r = 0.45, WC<sub>fb</sub> and VAT r = 0.76 and WC<sub>fb</sub> with HOMA-IR r = 0.33)</p> | <p>Correlation between all WC with: SBP (r = 0.40-0.48), DBP (r = 0.37-0.46), Insulin (r = 0.43-0.63), HOMA-IR (r = 0.43-0.62). TG had positive correlation with WC<sub>umbilicus</sub> and WC<sub>lax-crest</sub>. Correlation among WC<sub>umbilicus</sub>, WC<sub>lax-crest</sub> and WC<sub>middle</sub> was slightly stronger with CVD risk variables. Association of MetS for IDF: WC<sub>umbilicus</sub> (OR 2.18; IC 1.23-3.85; p = 0.007) and WC<sub>middle</sub> (OR 1.81; IC 1.06-3.09; p = 0.03); showed higher risk of MetS and number of MetS components. MetS - Cook definition: MetS was not associated with any WC; but it was associated with the number of components in WC<sub>umbilicus</sub> (OR 2.10; IC 1.28-3.44; p = 0.003), WC<sub>middle</sub> (OR 1.83; IC 1.13-2.98; p = 0.01), MetS-NCEP: was associated in WC<sub>umbilicus</sub> (OR 3.80; CI 1.28-11.33, p = 0.02), and WC<sub>middle</sub> (OR 3.24; CI 1.08-9.72; p = 0.04)</p> | <p>Correlation between all WC and log VAT was significant in all groups: white and black M = 0.81-0.86, white F and AA = 0.87-0.89</p> <p>Correlation of all WC with MetS components was strong except for white M and AA-F in AA-M the correlation between glucose with WC<sub>umbilicus</sub> was lower but not with other WC. There was no correlation between all WC with DBP in white-M</p>   | <p>Correlation between WC<sub>middle</sub> was significant with SBP 0.72, DBP 0.63, glucose 0.13, TG 0.42 and HDL -0.46. Correlation between WC<sub>lax-crest</sub> with: SBP 0.71, DBP 0.61, glucose 0.12, HDL -0.455. Sensitivity WC &gt; 90<sup>th</sup> for WC<sub>umbilicus</sub> and WC<sub>lax-crest</sub>: Hypertension 88.9 and 61.5, hyperglycemia 64.1 and 59, hypertriglyceridemia 64.4 and 64.4, low HDL 63.1 and 65.1 and to identify ≥ 2 components of MetS was 67.9 and 68.8, respectively. The AUC for WC<sub>umbilicus</sub> &gt; 90<sup>th</sup> and WC<sub>lax-crest</sub> &gt; 90<sup>th</sup>: hypertension (0.69 and 0.64), low HDL (0.62 and 0.62) and ≥ 2 components of MetS (0.617 and 0.608)</p> |
| <b>Evaluations of the quality of evidence</b>      | STROBE   | 16  | From 12 to 18   | 20   | From 16 to 19   |
|  | GRADE  | A   | A   | A  | A   |

JRI: journal rank indicator; AA: African American; WHO: World Health Organization; NCHS: National Center of Health Statistics; CV: coefficient of variation; MetS: metabolic syndrome; IDF: International Diabetes Federation; NCEP: National Cholesterol Education Program; HDL: high density lipoprotein cholesterol; TG: triglycerides; SBP: systolic blood pressure; DBP: diastolic blood pressure; M: male(s); F: female(s); BMI: body mass index; CVD: cardiovascular disease; HOMA-IR: Homeostatic Model Assessment for Insulin Resistance; OR: odds ratio; VAT: visceral adipose tissue; NPV: negative predictive value; PPV: positive predictive value; BMI: body mass index; IDF: International Obesity Task Force; CDC: Centers for Disease Control and Prevention; NHANES: National Health and Nutrition Examination Survey; TC: total cholesterol; LDL: low-density lipoprotein cholesterol; SAT: subcutaneous adipose tissue; CI: confidence interval; AUC: area under the curve; STROBE: Strengthening the Reporting of Observational Studies in Epidemiology; GRADE: Grading of Recommendations, Assessment, Development and Evaluation.





**Figure 3.**

Meta-analysis of correlation coefficients for WC<sub>iliac-crest</sub>, WC<sub>middle</sub>, and MetS components (WC: waist circumference; WC<sub>iliac-crest</sub>: waist circumference measured above the iliac crest; WC<sub>middle</sub>: waist circumference measured between the floating rib and the iliac crest; MetS: metabolic syndrome; HDL: high-density lipoprotein cholesterol; TG: triglycerides; SBP: systolic blood pressure; DBP: diastolic blood pressure) (Supplementary material: <https://www.nutricionhospitalaria.org/anexos/05144-01.pdf>).

However, this measurement may still be significant due to the prominence of the abdomen. In daily pediatric practice, WC is measured by physicians, nutritionists, pediatricians, and nurses. Work team training is essential to ensure precision in the technique and the anatomical site of measurement.

In table III, we summarize the strengths and weaknesses of each WC measurement site in children that can be better adapted to daily clinical practice according to training in measurement techniques and the characteristics of children.

A major limitation of our results is related to the fact that only few studies have assessed the correlation between different anatomical WC measurement sites and MetS in children and adolescents. This draws attention, considering that WC is the most relevant point for CVD risk exploration

in clinical practice. Therefore, this systematic review should be the starting point for future studies on the specific characteristics of CVD risk according to age and sex at different measurement sites. It is essential to create specific percentiles for each WC measurement site for each ethnic group or population.

**CONCLUSIONS**

There is similar and adequate correlation between all WC measurement sites and other MetS components in the included studies, regardless the anatomical site of measurement. However, there are differences by age, pubertal development, and

**Table III. Strengths and weaknesses of WC measurement sites**

| Measurement site          | Strengths  | Weaknesses   |
|---------------------------|--|--|
| WC <sub>middle</sub>      | Established measurement protocol.<br>Percentile tables in different ethnic populations. Good correlation with MetS | Difficulty in locating the anatomical point.<br>Measurement can be uncomfortable and requires more time  |
| WC <sub>iliac-crest</sub> | Established measurement protocol.<br>Percentile tables in different ethnic populations. Good correlation with MetS | Difficulty in locating the anatomical point.<br>Difficult to stabilize the tape measure on a curved skin surface   |
| WC <sub>rib</sub>         | Anatomical site location can be easy to locate and measure   | No established measurement protocol.<br>Not commonly used.<br>May underestimate WC measurement   |
| WC <sub>4</sub>           | Less comfortable and easier to locate in overweight or obese patients  | No established measurement protocol.<br>Not commonly used.<br>The point of measurement can be at different sites on the abdomen, which can give a lot of variability |
| WC <sub>umbilicus</sub>   | Description of measurement protocol. Tables of percentiles in different ethnic population                          | Modification of umbilical scar location by adipose tissue descent in patients with obesity   |
| WC <sub>narrow</sub>      | Established measurement protocol.<br>Percentile tables in different ethnic populations. Good correlation with MetS | It is an anatomical site that may be difficult to visualize in some patients   |

ethnicity that have not yet been clearly defined.

**REFERENCES**

- Al-Domi H, Al-Shorman A. Increased waist circumference is associated with subclinical atherosclerosis in schoolchildren. *Diabetes & Metabolic Syndrome: Clinical Research & Reviews* 2019;13(1):264-9. DOI: 10.1016/j.dsx.2018.09.004
- Twig G, Yaniv G, Levine H, Leiba A, Goldberger N, Derazne E, et al. Body-Mass Index in 2.3 Million Adolescents and Cardiovascular Death in Adulthood. *N Engl J Med* 2016;374(25):2430-40. DOI: 10.1056/NEJMoa1503840
- Ness-Abramof R, Apovian CM. Waist circumference measurement in clinical practice. *Nutrition in Clinical Practice* 2008;23(4):397-404.
- Skilton MR, Celermajor DS, Cosmi E, Crispi F, Gidding SS, Raitakari OT, et al. Natural history of atherosclerosis and abdominal aortic intima-media thickness: Rationale, evidence, and best practice for detection of atherosclerosis in the young. *J Clin Med* 2019;8(8):1201. DOI: 10.3390/jcm8081201
- Tagi VM, Samvelyan S, Chiarelli F. An update of the consensus statement on insulin resistance in children 2010. *Front Endocrinol (Lausanne)* 2022;13:1061524. DOI: 10.3389/fendo.2022.1061524
- Henning RJ. Obesity and obesity-induced inflammatory disease contribute to atherosclerosis: a review of the pathophysiology and treatment of obesity. *Am J Cardiovasc Dis* 2021;11(4):504-29.
- National Cholesterol Education Program (NCEP) Expert Panel on Detection Evaluation and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III). Third Report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III) final report. *Circulation* 2002;106(25):3143-421
- Alberti KG, Zimmet P, Shaw J. Metabolic syndrome-a new world-wide definition. A Consensus Statement from the International Diabetes Federation. *Diabetic Medicine* 2006;23(5):469-80. DOI: 10.1111/j.1464-5491.2006.01858.x
- Lohman TG, Roche AF, Martorell R. Anthropometric Standardization Reference Manual. Kinetics, Kinetics, Human: Champaign, IL.; 1988.
- Stewart A, Marfell-Jones M, Olds T, de Ridder JH. International Standards for Anthropometric Assessment. ISAK: United Kingdom; 2011.
- World Health Organization. The WHO STEPwise Approach to Surveillance. 2017.
- NHANES. Anthropometry Procedures Manual. National Health and Nutrition Examination Survey. (CDC. ed) Atlanta, US; 2000.
- Croft JB, Keenan NL, Sheridan DP, Wheeler FC, Speers MA. Waist-to-hip ratio in a biracial population: measurement, implications, and cautions for using guidelines to define high risk for cardiovascular disease. *J Am Diet Assoc* 1995;95(1):60-4. DOI: 10.1016/S0002-8223(95)00014-3
- Eisenmann JC. Waist circumference percentiles for 7- to 15-year-old Australian children. *Acta Paediatr* 2005;94(9):1182-5. DOI: 10.1111/j.1651-2227.2005.tb02071.x
- Rudolf MC, Walker J, Cole TJ. What is the best way to measure waist circumference? *Int J Pediatr Obes* 2007;2(1):58-61. DOI: 10.1080/17477160601095177
- von Elm E, Altman DG, Egger M, Pocock SJ, Gøtzsche PC, Vandenbroucke JP; et al. The strengthening the reporting of observational studies in epidemiology (STROBE) statement: Guidelines for reporting observational studies. *International Journal of Surgery* 2014;12(12):1495-9. DOI: 10.1016/j.ijsu.2014.07.013
- Higgins J, Thomas J. *Cochrane Handbook for Systematic Reviews of Interventions*. 2nd Edition. 2022nd ed. (Higgins J, Thomas J, Chandler J. eds). John Wiley & Sons: Chichester (UK); 2022.
- Page MJ, McKenzie JE, Bossuyt PM, Boutron I, Hoffmann TC, Mulrow CD, et al. The PRISMA 2020 statement: an updated guideline for reporting systematic reviews. *Syst Rev* 2021;10(1):89. DOI: 10.1186/s13643-021-01626-4
- Bosy-Westphal A, Booke CA, Blöcker T, Kossel E, Goele K, Later W, et al. Measurement site for waist circumference affects its accuracy as an index of visceral and abdominal subcutaneous fat in a Caucasian population. *J Nutr* 2010;140(5):954-61. DOI: 10.3945/jn.109.118737
- Hitze B, Bosy-Westphal A, Bielfeldt F, Settler U, Mönig H, Müller MJ. Measurement of waist circumference at four different sites in children, adolescents, and young adults: concordance and correlation with nutritional status as well as cardiometabolic risk factors. *Obes Facts* 2008;1(5):243-9. DOI: 10.1159/000157248

21. Johnson ST, Kuk JL, Mackenzie KA, Huang TT, Rosychuk RJ, Ball GD. Metabolic Risk Varies According to Waist Circumference Measurement Site in Overweight Boys and Girls. *Journal of Pediatrics* 2010;156(2):247-52.e1. DOI: 10.1016/j.jpeds.2009.08.010
22. Harrington DM, Staiano AE, Broyles ST, Gupta AK, Katzmarzyk PT. Waist circumference measurement site does not affect relationships with visceral adiposity and cardiometabolic risk factors in children. *Pediatr Obes* 2013;8(3):199-206. DOI: 10.1111/j.2047-6310.2012.00106.x
23. López-González D, Miranda-Lora A, Klünder-Klünder D, Queipo-García G, Bustos-Esquivel M, Paez-Villa M, et al. Diagnostic performance of waist circumference measurements for predicting cardiometabolic risk in Mexican children. *Endocr Pract* 2016;22(10):1170-6. DOI: 10.4158/EP161291.OR
24. Huang TTK. Finding Thresholds of Risk for Components of the Pediatric Metabolic Syndrome. *Journal of Pediatrics* 2008;152(2). DOI: 10.1016/j.jpeds.2007.09.032
25. Zimmet P, Alberti KG, Kaufman F, Tajima N, Silink M, Arslanian S, et al. The metabolic syndrome in children and adolescents – an IDF consensus report. *Pediatr Diabetes* 2007;8(5):299-306. DOI: 10.1111/j.1399-5448.2007.00271.x
26. Cook S, Weitzman M, Auinger P, Nguyen M, Dietz WH. Prevalence of a Metabolic Syndrome Phenotype in Adolescents Findings From the Third National Health and Nutrition Examination Survey, 1988-1994. *Arch Pediatr Adolesc Med* 2003;158(8):821-7. DOI: 10.1001/archpedi.157.8.821
27. Li YM, Zou ZY, Ma YH, Luo JY, Jing J, Zhang X, et al. Predicting Metabolic Syndrome Using Anthropometric Indices among Chinese Adolescents with Different Nutritional Status: A Multicenter Cross-sectional Study. *Biomedical and Environmental Sciences* 2021;34(9). DOI: 10.3967/bes2021.095
28. Perona JS, Schmidt-RioValle J, Fernández-Aparicio Á, Correa-Rodríguez M, Ramírez-Vélez R, González-Jiménez E. Waist Circumference and Abdominal Volume Index Can Predict Metabolic Syndrome in Adolescents, but only When the Criteria of the International Diabetes Federation are Employed for the Diagnosis. *Nutrients* 2019;11(6). DOI: 10.3390/nu11061370
29. Masquio DCL, Ganen A de P, Campos RM da S, Sanches P de L, Corgosinho FC, Caranti D, et al. Los valores de corte de circunferencia de cintura para predecir el síndrome metabólico en adolescentes obesos. *Nutr Hosp* 2015;31(4). DOI: 10.3305/nh.2015.31.4.8442
30. Ross R, Berentzen T, Bradshaw AJ, Janssen I, Kahn HS, Katzmarzyk PT, et al. Does the relationship between waist circumference, morbidity and mortality depend on measurement protocol for waist circumference? *Obesity Reviews* 2008;9(4):312-25. DOI: 10.1111/j.1467-789X.2007.00411.x
31. Palmer BF, Clegg DJ. The sexual dimorphism of obesity. *Mol Cell Endocrinol* 2015;402:113-9. DOI: 10.1016/j.mce.2014.11.029
32. Denzer C, Thiere D, Muche R, Koenig W, Mayer H, Kratzer W, et al. Gender-specific prevalences of fatty liver in obese children and adolescents: roles of body fat distribution, sex steroids, and insulin resistance. *Journal of Clinical Endocrinology and Metabolism* 2009;94(10):3872-81. DOI: 10.1210/jc.2009-1125
33. Wang J, Thornton JC, Bari S, Williamson B, Gallagher D, Heymsfield SB. Comparisons of waist circumferences measured at 4 sites. *Am J Clin Nutr* 2003;77(2):379-84. DOI: 10.1093/ajcn/77.2.379
34. Pfeiffer C, Willmer T, Dias S, Abrahams Y, Louw J, Goedecke JH. Ethnic and Adipose Depot Specific Associations Between DNA Methylation and Metabolic Risk. *Front Genet* 2020;11. DOI: 10.3389/fgene.2020.00967
35. Katzmarzyk P. Waist circumference percentiles for Canadian youth 11–18 y of age. *Eur J Clin Nutr* 2004;58(7):1011-5. DOI: 10.1038/sj.ejcn.1601924
36. McCarthy H, Jarrett K, Crawley H. The development of waist circumference percentiles in British children aged 5.0–16.9 y. *Eur J Clin Nutr* 2001;55(10):902-7. DOI: 10.1038/sj.ejcn.1601240
37. World Health Organization. Measuring Obesity. Classification and Description of Anthropometric Data. Report on a WHO Consultation of the Epidemiology of Obesity. Warsaw 21-23 October 1987. Copenhagen: WHO, 1989. Nutrition Unit Document, EUR/ICP/NUT 123. 1987.
38. World Health Organisation (WHO). WHO I Waist Circumference and Waist-Hip Ratio. Report of a WHO Expert Consultation. Geneva, 8-11 December 2008. 2008;(December):8-11.
39. Lennie S, Diatuo A, Evill T, Stewart AD. Protocol variations in arm position influence the magnitude of waist girth. *J Sports Sci* 2013;31(12):1353-58. DOI: 10.1080/02640414.2013.781664