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JUDUL: Abdominal Diameter Profiles have Relationship with Insulin Resistance in Obese Female Adolescents

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1.	Submission Artikel	23 November 2019	2-5
2.	Proses Review dan Revisi	25 November 2019	6-28
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==Paper Information==

Paper Title: Abdominal diameter profiles have relationship with insulin resistance in obese female adolescents

Abstract:

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Method: This study was conducted in June-August 2019 at Diponegoro University Semarang. this study used cross sectional design with a total of 120 female students aged 17-21 years and selected using purposive sampling method. Study variables included SAD, SAD/Height, ADI and insulin resistance level. SAD was measured by abdominal caliper, SAD/Height was ratio of SAD divided by height. ADI was ratio of SAD divided by thigh circumference. The value of insulin resistance was obtained from the calculation of homeostasis model assessment insulin resistance (HOMA-IR). Data normality was analyzed by Kolmogorov-smirnov test, all data were not normal distribution so the bivariate test analyzed by Rank spearman test.

Results: As many as 42,5% subjects had SAD which included at risk

and 83,3% subjects had suffering insulin resistance. There were significant relationship between SAD, SAD/Height, ADI with insulin resistance (SAD ($p=0,001$; $r=0,297$), SAD/Height ($p<0,001$; $r=0,345$) and ADI ($p=0,003$; $r=0,273$)). In addition, SAD, SAD/Height, ADI were significant relationship with fasting blood glucose (SAD ($p=0,048$; $r=0,181$), SAD/Height ($p=0,022$; $r=0,209$) and ADI ($p=0,003$; $r=0,270$)).

Conclusion: There were significant relationship between SAD, SAD/Height, ADI with insulin resistance and fasting blood glucose in obese adolescent female.

Keywords: Sagittal Abdominal Diameter (SAD), SAD/Height, Abdominal Diameter Index (ADI), insulin resistance.

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Abdominal diameter profiles have relationship with insulin resistance in obese female adolescents

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ABSTRACT

Purpose: This study aimed to determine the relationship of abdominal diameter consist of Sagittal Abdominal Diameter (SAD), Ratio SAD/Height and Abdominal Diameter Index (ADI) with insulin resistance in obese female adolescents.

Method: This study was conducted in June-August 2019 at Diponegoro University Semarang. this study used cross sectional design with a total of 120 female students aged 17-21 years and selected using purposive sampling method. Study variables included SAD, SAD/Height, ADI and insulin resistance level. SAD was measured by abdominal caliper, SAD/Height was ratio of SAD divided by height. ADI was ratio of SAD divided by thigh circumference. The value of insulin resistance was obtained from the calculation of homeostasis model assessment insulin resistance (HOMA-IR). Data normality was analyzed by Kolmogorov-smirnov test, all data were not normal distribution so the bivariate test analyzed by Rank spearman test.

Results: As many as 42,5% subjects had SAD which included at risk and 83,3% subjects had suffering insulin resistance. There were significant relationship between SAD, SAD/Height, ADI with insulin resistance (SAD ($p=0,001$; $r=0,297$), SAD/Height ($p<0,001$; $r=0,345$) and ADI ($p=0,003$; $r=0,273$)). In addition, SAD, SAD/Height, ADI were significant relationship with fasting blood glucose (SAD ($p=0,048$; $r=0,181$), SAD/Height ($p=0,022$; $r=0,209$) and ADI ($p=0,003$; $r=0,270$)).

Conclusion: There were significant relationship between SAD, SAD/Height, ADI with insulin resistance and fasting blood glucose in obese adolescent female.

Keywords: Sagittal Abdominal Diameter (SAD), SAD/Height, Abdominal Diameter Index (ADI), insulin resistance.

INTRODUCTION

Obesity is one factor of that cause various diseases such as cardiovascular disease, hypertension, cancer, and diabetes mellitus. Most countries in Asia have increasing of overweight and obesity prevalence in this last decade. Increased of obesity prevalence does not only occurred in adult group but also occurred in adolescent group. Indonesia is the one of the countries that have high prevalence of abdominal obesity in the Southeast Asian region (Ramachandran & Snehalatha, 2010). The prevalence of visceral/abdominal obesity at age ≥ 15 years based on RISKESDAS 2013 as many as 26,6% and increase to 31% in 2018 (Kementrian Kesehatan RI, 2013) (Kementrian Kesehatan RI, 2018). A study of adolescent in Semarang showed that 46 of 566 adolescents suffering abdominal obesity (Hanifah & Dieny, 2016). The prevalence of abdominal obesity in female higher than male, the result data of RISKESDAS 2013 showed that female group who suffering abdominal obesity is 56,3% while the male group is 43,7% (Sudikno, Syarief, Dwiriani, & Riyadi, 2016).

Abdominal obesity can reflect excess of visceral fat tissue or subcutaneous or both. Visceral fat contributes to the pathogenesis of diabetes mellitus, glucose intolerance and insulin resistance. Insulin resistance is a impaired condition of glucose absorption that induced by insulin (Ye, 2013). Fat tissue distribution in intraabdominal area caused increasing metabolic such as lipogenesis and lipolysis activity. Lipolysis produce *free fatty acids* (FFA), FFA from visceral fat brought to the portal circulation and liver. Excessive of FFA secretion from visceral fat through the liver cause toxic effect for example gluconeogenesis and insulin resistance. (Ye, 2013) (Westphal, 2008) Study of 220 obese children and adolescents in Brazil shows that 33,2% subjects suffering insulin resistance (Romualdo, Nobrega, & Escrivao, 2014). Study of 52 obese

adolescents in Semarang also shows that 91,6% subjects suffering insulin resistance (Nuraini, Sulchan, & Dieny, 2017).

Excessive of visceral fat tissue in abdominal obesity can measure by *Computed Tomography* (CT) as gold standard, but measurement of Waist Circumference (WC), and abdominal diameter also can be used as alternative. Abdominal diameter measurement consist of three components, namely Sagittal Abdominal Diameter (SAD), Sagittal Abdominal Diameter/Height (SAD/Height), Abdominal Diameter Index (ADI) (Carneiro Roriz et al., 2016). SAD can describe the visceral fat size, moreover it can also predicting metabolic syndrome from visceral fat area. SAD is closely related with visceral fat mass than other anthropometric measurements. SAD measured by supine position when subcutaneous fat move to side of waist. SAD measurement at this position reflect size of intraabdominal fat in antero-posterior like in CT image measurement. SAD strongly related with glucose intolerance, cardiovascular disease and good indicator for predicting insulin resistance (Anjana et al., 2004; Carneiro Roriz et al., 2016; Firouzi, Tucker, LeCheminant, & Bailey, 2018; Gletsu-Miller et al., 2013).

SAD/Height is derivative of SAD measurement, which is use height as divisor of SAD. SAD/Height is anthropometric indicator that significantly related to visceral fat tissue (Carneiro Roriz et al., 2016)(Khan, Rissanen, Bullard, & Knekt, 2014). ADI is derivative of SAD measurement divided by thigh circumference. ADI is significantly related to several indicators of metabolic syndrome, and maybe has significant relationship with insulin resistance.(Ehrlich & Smith, 2011)A study in Brazil also showed that SAD/Height and ADI is good indicator for measurement visceral fat tissue in women (Roriz et al., 2014). Study on abdominal diameter (SAD, SAD/Height and ADI) as insulin resistance indicator has not been much done especially in Indonesia. Therefore, based on this background the researchers interest to identifying the relationship of abdominal diameter (SAD, SAD/Height and ADI) with insulin resistance in obese adolescent female.

METHODS

This study was conducted in June-August 2019 with female student subject at Diponegoro University Semarang. This study was a observational analytic with cross sectional design. The subjects were selected by screening process which obtained 1260 subjects, after that 215 subjects were included to inclusion criteria and selected as many as 120 subjects who involved in this study using purposive sampling method. The inclusion criteria were female student in Diponegoro University Semarang, aged 17-21 years, having waist circumference >80 cm, not pregnant, not consuming alcohol, not consuming drug that can related to glucose and insulin level, willing to fast minimum 8 hours before taking blood sample. The exclusion criteria was the subject resign during the study.

The independent variables in this study were abdominal diameter consisting of SAD, SAD/Height, and ADI. SAD measured by abdominal caliper which is manually measurement. Measurement were conducted with supine position, subject relax on the table. One of caliper's arm was placed under the subject (*iliac crest*) and the other caliper's arm was slowly moving down on subject abdomen with normal breathing (Khan et al., 2014). SAD categorized as normal if <19,3 cm and at risk if $\geq 19,3$ cm (Sampaio, Simões, Assis, & Ramos, 2007). SAD/Height was derivative of SAD, which is use height (in centimeter) as SAD devisor. ADI was derivative of SAD measurement divided by thigh circumference (in centimeter). The thigh circumference was measured on the right side of the body, at the midpoint between inguinal crease and proximal border on the patella (Roriz et al., 2014).

The dependent variable in this study was insulin resistance. Insulin resistance is a diruption of the biological response to insulin which cause body insulin requirement increase and then resulting hyperinsulinemia to maintain normal glucose level. Insulin resistance level obtained from calculating *Homeostasis Model Assessment Insulin Resistance* (HOMA-IR), with formula:

$$\frac{\text{fasting insulin} \left(\frac{mU}{L} \right) \times \text{fasting glucose} \left(\frac{mmol}{L} \right)}{22,5}$$

Cut off the normal value of HOMA-IR in adolescent was <1,65 (Nuraini et al., 2017). Blood sample of subject was taking by laboratory officer. Before the blood was taken, the subject fasting for 8-12 hours and then blood was taken 5 cc.

Univariate data analyze was used to describe the characteristic of each variable in this study. Data normality test analyzed by Kolmogorov-smirnov test. Bivariate data analyzed by Rank spearman test for abnormal data distribution. The purpose of bivariate analysis were to determine relationship of each abdominal diameter variable with insulin resistance.

RESULT

Subject Characteristic

Table of subject characteristic shows that age range of subjects were 17-21 years old. The highest fasting blood glucose of subject was 206 mg/dL while the lowest fasting blood glucose was 68 mg/dL. The median insulin level of subject was 0,82 μ U/mL. The highest HOMA-IR value of subject was 8,32 with median value 2,33. SAD range of subject was 14,90-26,10 cm.

Table 1. Subject characteristic

Variable	Min	Max	Median \pm SD
Age (year)	17	21	19 \pm 0,8
Weight (kg)	47,80	107,40	66,60 \pm 11,38
Height (cm)	141,20	171,40	157,40 \pm 5,44
Sagittal Abdominal Diameter(cm)	14,90	26,10	18,68 \pm 2,12
Sagittal Abdominal Diameter/Height (ratio)	0,09	0,17	0,12 \pm 0,014
Abdominal Diameter Index/ADI	0,03	0,56	0,32 \pm 0,41
Fasting blood glucose(mg/dL)	68,00	206,00	87,00 \pm 13,15
Insulin (μ U/mL)	2,71	52,60	10,82 \pm 6,92
HOMA-IR	0,54	18,32	2,33 \pm 2,09

This result study shows that almost half of the subjects (42,5%) had SAD that categorized at risk. Majority of subjects (89,2%) had normal fasting insulin level. Almost of subjects (97,5%) had normal fasting blood glucose, whereas conversely most of subjects (83,3%) had suffering insulin resistance.

Table 2. Frequency distribution of sagittal abdominal diameter, fasting blood glucose , fasting insulin and HOMA-IR

Characteristic	n	%
SagittalAbdominal Diameter		
No risk	69	57,5
At risk	51	42,5
Fasting Blood Glucose		
Normal	117	97,5
High	3	2,5
Fasting Insulin		
Normal	107	89,2
High	13	10,8
HOMA-IR		
Normal	20	16,7
Resistance	100	83,3

Relationship of Abdominal Diameter with Insulin Resistance

Bivariate analysis in this study used Rank Spearman test. Table 3 shows that SAD,SAD/Height and ADI had significant relationship with insulin resistance and fasting blood glucose ($p < 0,05$), with positive correlation.

Table 3. Relationship of abdominal diameter with insulin resistance and fasting blood glucose

Variable	Fasting blood glucose		HOMA-IR	
	r	p	r	p
Sagittal abdominal diameter	0,181	0,048	0,297	0,001
SAD/Height	0,209	0,022	0,345	0,000
Abdominal diameter index	0,270	0,003	0,273	0,003

DISCUSSION

The result of this study shows that SAD median was $18,68 \pm 2,12$ cm with minimum value 14,90 cm and maximum value 26,10 cm. SAD is indicator that can describe visceral fat size (Stokic, Srdic, Brtko, & Tomic-Nagalic, 2012). SAD measured by supine position, visceral abdominal fat tend to lift the abdominal wall toward sagittal, abdominal subcutaneous fat pressure the stomach and subcutaneous fat moves to the waist, so SAD can reflect the visceral abdominal fat volume (Vasques et al., 2015). As many as 42,5% subjects had SAD which categorized at risk. SAD in women closely related with hyperlipidemia, insulin resistance and cardiovascular risk, moreover it can also be used as a predictor of cardiometabolic risk score (Riserus, Faire, Berglund, & Hellenius, 2010). The study of children and adolescent in Arab shows that SAD significantly related with the severity of metabolic syndrome components (Al-attas et al., 2012).

This study showed that the HOMA-IR most of subjects (83,3%) included in insulin resistance category, however, almost of subjects had normal fasting blood glucose and normal fasting insulin. The study of 38 obese adolescent in Semarang also shows that same result with this study. This condition can occur because hyperglycemia is the last part from insulin resistance condition, which is long enough condition of impaired glucose intolerance and or impaired fasting blood glucose. Carbohydrate metabolism in adolescence also still good, so the body can maintaining normal glucose level (Ramadhan & Wijayanti, 2017). The high incidence of insulin resistance who suffering by subjects increased the risk of metabolic syndrome, because insulin resistance is a risk factor for metabolic syndrome that first arise (Nuraini et al., 2017). The highest HOMA-IR value in the subject was 18,32. Adolescent group with metabolic syndrome significantly had higher of HOMA-IR. HOMA-IR is indicator that strongly correlated with total and basal insulin. The HOMA-IR mean progressively increase with following sex maturation increase and weight from normal to obese. The study in India shows that the increase of HOMA-IR significantly related with body mass index, moreover there was significant different between HOMA-IR in adolescent with normal nutritional status, overweight and obesity (Singh, Garg, Tandon, & Marwaha, 2013).

Bivariate analysis in this study showed that SAD, SAD/Height, and ADI significant related to insulin resistance with positive correlation. A study of 75 overweight and obese adolescent showed that SAD and insulin resistance had significant relationship, with correlation that not much different from this study ($r=0,485$). The study also showed that SAD described a better correlation with insulin resistance than other anthropometric measurements. SAD can predict insulin resistance because it measures abdominal visceral fat than subcutaneous fat. Abdominal visceral fat also had strongly related with cardio metabolic disease (Krishnappa, Yashoda, Boraiah, & Vishwa, 2015). The study of women subject in Brazil comparing SAD with body mass index (BMI), waist circumference (WC), waist to hip ratio (WHR) as resistance insulin indicator, the result show that SAD and BMI was strongest indicator to indicate the occurrence of insulin resistance. BMI was widely used to identify metabolic risk but it can not be used alone, because it can not distinguish muscle adipose tissue or body fat distribution, whereas SAD had reported as a good marker for visceral adipose tissue in various ethnic group (Vasques et al., 2015).

In contrast with SAD, study about relationship between SAD/Height ratio and ADI with insulin resistance still limited, but study of 194 subjects in Brazil shows that SAD/Height and ADI had significant related and positive correlation with visceral adipose tissue (Roriz et al., 2014). A study of 1347 subjects in Korea showed that high visceral fat mass and low adiponectin level were correlated with increased risk of insulin resistance and β cell dysfunction. Obesity is well known as risk factor for insulin resistance. Abdominal obesity can cause insulin resistance by stimulating metabolic product formation which derived from fat, hormone and cytokines. Insulin resistance can cause endothelial dysfunction and insulin signaling pathway changes. Several previous studies show that visceral adipose tissue can inhibit adiponectin secretion. Adiponectin mediates the insulin sensitivity system and glucose homeostasis (Moon, Ha, Han, Kim, & Kim, 2019). Insulin resistance is adaptive response of *Free Fatty Acid* (FFA) increase, and related with visceral adipose tissue level. FFA increase can change the liver, muscle and other tissue metabolism toward lipid deposition and oxidation. Insulin secretion increase offset the glucose capacity decrease. Subcutaneous adipose tissue taking FFA and storing excess calories in visceral adipose tissue (Donohoe, Doyle, & Reynolds, 2011). Excess FFA secretion from visceral fat will be

oxidized, stored (lipids droplets) or metabolized to toxic derivatives (DAG and ceramides). Toxic derivatives cause insuline resistance, impaired cell function (lipotoxicity) or cause apoptosis. Apoptosis in the pancreas cause decrease and impairment of β cell capacities to secrete insulin which increase the risk of developing type 2 diabetes mellitus. The consequence of insulin resistance increase is endogenous glucose production by liver and glucose utilization decrease by peripheral tissue. As result are glycemia and insulin pancreas secretion increase, beside that occure hepatic insulin clearance impaired, and causing hyperinsulinemia (Castro, Koika, Kim, & Bergman, 2015).

This study also showed that significant relationship between abdominal diameter (SAD, SAD/Height, and ADI) with fasting blood glucose level. Abdominal diameter measurement can reflect visceral fat. Visceral/abdominal fat stores are considered as implication of insulin resistance, cardiovascular and other metabolic condition which related to type 2 diabetes mellitus. Excessive visceral fat related with impaired of insulin sensivity. Visceral fat related with impaired glucose regulation, because it is closely related to insulin resistance (Kantartzis et al., 2010). Lypolysis of visceral fat can cause free circulated of FFA in obese individual. This cause FFA enter to portal vein which can accumulated in the liver and cause impaired of glucose tolerance and insulin homeostasis. Visceral fat tissue and liver fat tissue related with insulin resistance in diabetic patient. A study of 297 subject showed that subject with isolated impaired fasting glucose and isolated impaired glucose tolerance had excess abdominal fat (visceral fat, abdominal subcutaneous fat and liver fat) (Borel et al., 2015).

CONCLUSION

As many as 42,5% subjects had SAD were categorized at risk, while 83,3% subject had suffering resistance insulin. There were significant relationship between abdominal diameter (SAD, SAD/Height, ADI) with insulin resistance and fasting blood glucose in obese adolescent female, with positive correlation.

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- a. explain first about the specific terminology (e.g riskesdas, kementerian kesehatan etc), because these are not a universal terminology
- b. please better present about the step of samples selection by flow chart, so the number of the samples clearly step by step
- c. please revise the writing of the symbol of comma in decimal to the metric form, because in English language the decimal symbol by dot (example 23.5 not 23,5)
- d. please declare explicit about the ethical clearance (the ethic committee institution approval and the number of approval)
- e. please delete the correlation coefficient (r) from the abstract, because it is not usually to write
- f. please avoid repeat to write the content of the table (the percentage or the frequency) in the narrative, you must explain the other information from the table (such as why the result have the big range between minimum and maximum value, and the information about the extreme value of the data)
- g. the conclusion not only about significant result, but also include the suggestion that can be improved by health administration office or the other institution that have the responsibility to make the nutrition improvements.

Abdominal diameter profiles have relationship with insulin resistance in obese female adolescents

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ABSTRACT

Purpose: This study aimed to determine the relationship of abdominal diameter consist of Sagittal Abdominal Diameter (SAD), Ratio SAD/Height and Abdominal Diameter Index (ADI) with insulin resistance in obese female adolescents.

Method: This study was conducted in June-August 2019 at Universitas Diponegoro, Central Java. this study used cross sectional design with a total of 120 female students aged 17-21 years and selected using purposive sampling method. Study variables included SAD, SAD/Height, ADI and insulin resistance level. SAD was measured by abdominal caliper, SAD/Height was ratio of SAD divided by height. ADI was ratio of SAD divided by thigh circumference. The value of insulin resistance was obtained from the calculation of homeostasis model assessment insulin resistance (HOMA-IR). Data normality was analyzed by Kolmogorov-smirnov test, all data were not normal distribution so the bivariate test analyzed by Rank spearman test.

Results: As many as 42,5% subjects had SAD which included at risk and 83,3% subjects had suffering insulin resistance. There were significant relationship between SAD, SAD/Height, ADI with insulin resistance (SAD (p=0.001), SAD/Height (p<0.001) and ADI (p=0.003). In addition, SAD, SAD/Height, ADI were significant relationship with fasting blood glucose (SAD (p=0.048), SAD/Height (p=0.022) and ADI (p=0.003).

Conclusion: There were significant relationship between SAD, SAD/Height, ADI with insulin resistance and fasting blood glucose in obese adolescent female.

Keywords: Sagittal Abdominal Diameter (SAD), SAD/Height, Abdominal Diameter Index (ADI), insulin resistance.

INTRODUCTION

Obesity is one factor of that cause various diseases such as cardiovascular disease, hypertension, cancer, and diabetes mellitus. Most countries in Asia have increasing of overweight and obesity prevalence in this last decade. Increased of obesity prevalence does not only occurred in adult group but also occurred in adolescent group. Indonesia is the one of the countries that have high prevalence of abdominal obesity in the Southeast Asian region (Ramachandran & Snehalatha, 2010). The prevalence of visceral/abdominal obesity at age ≥ 15 years based on Basic Health Research 2013 as many as 26.6% and increase to 31% in 2018 (Ministry of Health Republic of Indonesia, 2013) (Ministry of Health Republic of Indonesia, 2018). A study of adolescent in Semarang showed that 46 of 566 adolescents suffering abdominal obesity (Hanifah & Dieny, 2016). The prevalence of abdominal obesity in female higher than male, the result data of Basic Health Research 2013 showed that female group who suffering abdominal obesity is 56.3% while the male group is 43.7% (Sudikno, Syarief, Dwiriani, & Riyadi, 2016).

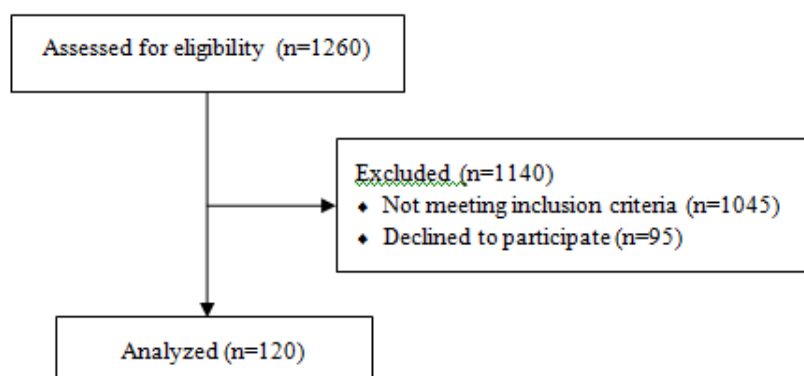
Abdominal obesity can reflect excess of visceral fat tissue or subcutaneous or both. Visceral fat contributes to the pathogenesis of diabetes mellitus, glucose intolerance and insulin resistance. Insulin resistance is a impaired condition of glucose absorption that induced by insulin (Ye, 2013). Fat tissue distribution in intraabdominal area caused increasing metabolic such as lipogenesis and lipolysis activity. Lipolysis produce *free fatty acids* (FFA), FFA from visceral fat brought to the portal circulation and liver. Excessive of FFA secretion from visceral fat through the liver cause toxic effect for example gluconeogenesis and insulin resistance. (Ye, 2013) (Westphal, 2008) Study of 220 obese children and adolescents in Brazil shows that 33.2% subjects suffering insulin resistance (Romualdo, Nobrega, & Escrivao, 2014). Study of 52 obese adolescents in Semarang also shows that 91.6% subjects suffering insulin resistance (Nuraini, Sulchan, & Dieny, 2017).

Excessive of visceral fat tissue in abdominal obesity can measure by *Computed Tomography* (CT) as gold standard, but measurement of Waist Circumference (WC), and abdominal diameter also can be used as alternative. Abdominal diameter measurement consist of three components, namely Sagittal Abdominal Diameter (SAD), Sagittal Abdominal Diameter/Height (SAD/Height), Abdominal Diameter Index (ADI) (Carneiro Roriz et al., 2016). SAD can describe the visceral fat size, moreover it can also predicting metabolic syndrome from visceral fat area. SAD is closely related with visceral fat mass than other anthropometric measurements. SAD measured by supine position when subcutaneous fat move to side of waist. SAD measurement at this position reflect size of intraabdominal fat in antero-posterior like in CT image measurement. SAD strongly related with glucose intolerance, cardiovascular disease and good indicator for predicting insulin resistance (Anjana et al., 2004; Carneiro Roriz et al., 2016; Firouzi, Tucker, LeCheminant, & Bailey, 2018; Gletsu-Miller et al., 2013).

SAD/Height is derivative of SAD measurement, which is use height as divisor of SAD. SAD/Height is anthropometric indicator that significantly related to visceral fat tissue (Carneiro Roriz et al., 2016) (Khan, Rissanen, Bullard, & Knekt, 2014). ADI is derivative of SAD measurement divided by thigh circumference. ADI is significantly related to several indicators of metabolic syndrome, and maybe has significant relationship with insulin resistance. (Ehrlich & Smith, 2011) A study in Brazil also showed that SAD/Height and ADI is good indicator for measurement visceral fat tissue in women (Roriz et al., 2014). Study on abdominal diameter (SAD, SAD/Height and ADI) as insulin resistance indicator has not been much done especially in Indonesia. Therefore, based on this background the researchers interest to identifying the relationship of abdominal diameter (SAD, SAD/Height and ADI) with insulin resistance in obese adolescent female.

METHODS

This study was conducted in June-August 2019 with female student subject at Diponegoro University Semarang. This study was a observational analytic with cross sectional design. The subjects were selected by screening process which obtained 1260 subjects, after that 215 subjects were included to inclusion criteria and selected as many as 120 subjects who involved in this study using purposive sampling method. The inclusion criteria were female student in Diponegoro University Semarang, aged 17-21 years, having waist circumference >80 cm, not pregnant, not consuming alcohol, not consuming drug that can related to glucose and insulin level, willing to fast minimum 8 hours before taking blood sample. The exclusion criteria was the subject resign during the study. This study had received permission from the Health Research Ethics Committee with Number 373/EC/KEPK/FK UNDIP/VIII/2019.



The independent variables in this study were abdominal diameter consisting of SAD, SAD/Height, and ADI. SAD measured by abdominal caliper which is manually measurement. Measurement were conducted with supine position, subject relax on the table. One of caliper's arm was placed under the subject (*iliac crest*) and the other caliper's arm was slowly moving down on subject abdomen with normal breathing (Khan et al., 2014). SAD categorized as normal if <19.3 cm and at risk if ≥ 19.3 cm (Sampaio, Simões, Assis, & Ramos, 2007). SAD/Height was derivative of SAD, which is use height (in centimeter) as SAD divisor. ADI was derivative of SAD measurement

divided by thigh circumference (in centimeter). The thigh circumference was measured on the right side of the body, at the midpoint between inguinal crease and proximal border on the patella (Roriz et al., 2014).

The dependent variable in this study was insulin resistance. Insulin resistance is a disruption of the biological response to insulin which cause body insulin requirement increase and then resulting hyperinsulinemia to maintain normal glucose level. Insulin resistance level obtained from calculating *Homeostasis Model Assessment Insulin Resistance (HOMA-IR)*, with formula:

$$\frac{\text{fasting insulin} \left(\frac{mU}{L}\right) \times \text{fasting glucose} \left(\frac{mmol}{L}\right)}{22.5}$$

Cut off the normal value of HOMA-IR in adolescent was <1.65 (Nuraini et al., 2017). Blood sample of subject was taken by laboratory officer. Before the blood was taken, the subject fasting for 8-12 hours and then blood was taken 5 cc.

Univariate data analyze was used to describe the characteristic of each variable in this study. Data normality test analyzed by Kolmogorov-smirnov test. Bivariate data analyzed by Rank spearman test for abnormal data distribution. The purpose of bivariate analysis were to determine relationship of each abdominal diameter variable with insulin resistance.

RESULT

Subject Characteristic

Table of subject characteristic shows that age range of subjects were 17-21 years old. The highest fasting blood glucose of subject was 206 mg/dL while the lowest fasting blood glucose was 68 mg/dL. The median in insulin level of subject was 0.82 μ U/mL. The highest HOMA-IR value of subject was 18.32 with median value 2.33. The median in SAD measurement was 18.68 cm. The highest SAD measurement of subject was 26.10 cm.

Table 1. Subject characteristic

Variable	Min	Max	Median \pm SD
Age (year)	17	21	19 \pm 0.8
Weight (kg)	47.80	107.40	66.60 \pm 11.38
Height (cm)	141.20	171.40	157.40 \pm 5.44
Sagittal Abdominal Diameter (cm)	14.90	26.10	18.68 \pm 2.12
Sagittal Abdominal Diameter/Height (ratio)	0.09	0.17	0.12 \pm 0.014
Abdominal Diameter Index/ADI	0.03	0.56	0.32 \pm 0.41
Fasting blood glucose (mg/dL)	68.00	206.00	87.00 \pm 13.15
Insulin (μ U/mL)	2.71	52.60	10.82 \pm 6.92
HOMA-IR	0.54	18.32	2.33 \pm 2.09

This result study shows that almost half of the subjects (42.5%) had SAD that categorized at risk. Majority of subjects (89.2%) had normal fasting insulin level. Almost of subjects (97.5%) had normal fasting blood glucose, whereas conversely most of subjects (83.3%) had suffering insulin resistance.

Table 2. Frequency distribution of sagittal abdominal diameter, fasting blood glucose, fasting insulin and HOMA-IR

Characteristic	n	%
Sagittal Abdominal Diameter		
No risk	69	57.5
At risk	51	42.5
Fasting Blood Glucose		
Normal	117	97.5
High	3	2.5
Fasting Insulin		
Normal	107	89.2
High	13	10.8
HOMA-IR		
Normal	20	16.7
Resistance	100	83.3

Relationship of Abdominal Diameter with Insulin Resistance

Bivariate analysis in this study used Rank Spearman test. Table 3 shows that SAD, SAD/Height and ADI had significant relationship with insulin resistance and fasting blood glucose ($p < 0.05$), with positive correlation.

Table 3. Relationship of abdominal diameter with insulin resistance and fasting blood glucose

Variable	Fasting blood glucose		HOMA-IR	
	r	p	r	p
Sagittal abdominal diameter	0.181	0.048	0.297	0.001
SAD/Height	0.209	0.022	0.345	<0.001
Abdominal diameter index	0.270	0.003	0.273	0.003

DISCUSSION

The result of this study shows that SAD median was 18.68 ± 2.12 cm with minimum value 14.90 cm and maximum value 26.10 cm. This result similar to study that conducted in Brazil by Feliciano et al. showed that in overweight adolescent the SAD measurement reached 26.3 cm (Feliciano Pereira P. et al, 2015). SAD is indicator that can describe visceral fat size (Stokic, Srdic, Brtka, & Tomic-Nagalic, 2012). SAD measured by supine position, visceral abdominal fat tend to lift the abdominal wall toward sagittal, abdominal subcutaneous fat pressure the stomach and subcutaneous fat moves to the waist, so SAD can reflect the visceral abdominal fat volume (Vasques et al., 2015). As many as 42.5% subjects had SAD which categorized at risk. SAD in women closely related with hyperlipidemia, insulin resistance and cardiovascular risk, moreover it can also be used as a predictor of cardiometabolic risk score (Riserus, Faire, Berglund, & Hellenius, 2010). The study of children and adolescent in Arab shows that SAD significantly related with the severity of metabolic syndrome components (Al-attas et al., 2012).

The result of this study shows that HOMA-IR median was 2.33 ± 2.09 cm with minimum value 0.54 cm and maximum value 18.32 cm. The high score of HOMA-IR similar to González-Zavala et al.'s study that found high score of HOMA-IR in overweight adolescent subjects that reached 18.32 (González-Zavala et al, 2015). This study showed that the HOMA-IR most of subjects (83.3%) included in insulin resistance category, however, almost of subjects had normal fasting blood glucose and normal fasting insulin. This result similar to Yi Kung Hye et al.'s study that found high prevalence of insulin resistant in obese adolescent subject was 47.1% (Yi Kung Hye et al. 2014). The study of 38 obese adolescent in Semarang also shows similar result with this study. This condition can occur because hyperglycemia is the last part from insulin resistance condition, which is long enough condition of impaired glucose intolerance and or impaired fasting blood glucose. Carbohydrate metabolism in adolescence also still good, so the body can maintaining normal glucose level (Ramadhan & Wijayanti, 2017). The high incidence of insulin resistance who suffering by subjects increased the risk of metabolic syndrome, because insulin resistance is a risk factor for metabolic syndrome that first arise (Nuraini et al., 2017). The highest HOMA-IR value in the subject was 18.32. Adolescent group with metabolic syndrome significantly had higher of HOMA-IR. HOMA-IR is indicator that strongly correlated with total and basal insulin. The HOMA-IR mean progressively increase with following sex maturation increase and weight from normal to obese. The study in India shows that the increase of HOMA-IR significantly related with body mass index, moreover there was significant different between HOMA-IR in adolescent with normal nutritional status, overweight and obesity (Singh, Garg, Tandon, & Marwaha, 2013).

Bivariate analysis in this study showed that SAD, SAD/Height, and ADI significant related to insulin resistance with positive correlation. A study of 75 overweight and obese adolescent showed that SAD and insulin resistance had significant relationship, with correlation that not much different from this study ($r = 0.485$). The study also showed that SAD described a better correlation with insulin resistance than other anthropometric measurements. SAD can predict insulin resistance because it measures abdominal visceral fat than subcutaneous fat. Abdominal visceral fat also had strongly related with cardio metabolic disease (Krishnappa, Yashoda, Boraiah, & Vishwa, 2015). The study of women subject in Brazil comparing SAD with body mass index (BMI), waist circumference (WC), waist to hip ratio (WHR) as resistance insulin indicator, the result show that SAD and BMI was strongest indicator to indicate the occurrence of insulin resistance. BMI was widely used to identify metabolic risk but it can not be used alone, because it can not distinguish

muscle adipose tissue or body fat distribution, whereas SAD had reported as a good marker for visceral adipose tissue in various ethnic group (Vasques et al., 2015).

In contrast with SAD, study about relationship between SAD/Height ratio and ADI with insulin resistance still limited, but study of 194 subjects in Brazil shows that SAD/Height and ADI had significant related and positive correlation with visceral adipose tissue (Roriz et al., 2014). A study of 1347 subjects in Korea showed that high visceral fat mass and low adiponectin level were correlated with increased risk of insulin resistance and β cell dysfunction. Obesity is well known as risk factor for insulin resistance. Abdominal obesity can cause insulin resistance by stimulating metabolic product formation which derived from fat, hormone and cytokines. Insulin resistance can cause endothelial dysfunction and insulin signaling pathway changes. Several previous studies show that visceral adipose tissue can inhibit adiponectin secretion. Adiponectin mediates the insulin sensitivity system and glucose homeostasis (Moon, Ha, Han, Kim, & Kim, 2019). Insulin resistance is adaptive response of *Free Fatty Acid* (FFA) increase, and related with visceral adipose tissue level. FFA increase can change the liver, muscle and other tissue metabolism toward lipid deposition and oxidation. Insulin secretion increase offset the glucose capacity decrease. Subcutaneous adipose tissue taking FFA and storing excess calories in visceral adipose tissue (Donohoe, Doyle, & Reynolds, 2011). Excess FFA secretion from visceral fat will be oxidized, stored (lipids droplets) or metabolized to toxic derivatives (DAG and ceramides). Toxic derivatives cause insulin resistance, impaired cell function (lipotoxicity) or cause apoptosis. Apoptosis in the pancreas cause decrease and impairment of β cell capacities to secrete insulin which increase the risk of developing type 2 diabetes mellitus. The consequence of insulin resistance increase is endogenous glucose production by liver and glucose utilization decrease by peripheral tissue. As result are glycemia and insulin pancreas secretion increase, beside that occur hepatic insulin clearance impaired, and causing hyperinsulinemia (Castro, Koika, Kim, & Bergman, 2015).

This study also showed that significant relationship between abdominal diameter (SAD, SAD/Height, and ADI) with fasting blood glucose level. Abdominal diameter measurement can reflect visceral fat. Visceral/abdominal fat stores are considered as implication of insulin resistance, cardiovascular and other metabolic condition which related to type 2 diabetes mellitus. Excessive visceral fat related with impaired of insulin sensitivity. Visceral fat related with impaired glucose regulation, because it is closely related to insulin resistance (Kantartzis et al., 2010). Lypolysis of visceral fat can cause free circulated of FFA in obese individual. This cause FFA enter to portal vein which can accumulated in the liver and cause impaired of glucose tolerance and insulin homeostasis. Visceral fat tissue and liver fat tissue related with insulin resistance in diabetic patient. A study of 297 subject showed that subject with isolated impaired fasting glucose and isolated impaired glucose tolerance had excess abdominal fat (visceral fat, abdominal subcutaneous fat and liver fat) (Borel et al., 2015).

CONCLUSION

As many as 42,5% subjects had SAD were categorized at risk, while 83,3% subject had suffering resistance insulin. There were significant relationship between abdominal diameter (SAD, SAD/Height, ADI) with insulin resistance and fasting blood glucose in obese adolescent female, with positive correlation. As it is important to monitor the growth and development of adolescents over time, it is advisable to standardize the use of one measure of body fat location. Considering that abdominal fat, more than total fat, has been associated with cardiometabolic risk, it is recommended the use abdominal diameter measurement (SAD, SAD/Height, ADI) as a measure that reflects the adipose tissue in this region, in the assessment of the nutritional status of adolescents.

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Abdominal Diameter Profiles have Relationship with Insulin Resistance in Obese Female Adolescents

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ABSTRACT

Purpose: This study aimed to determine the relationship of abdominal diameter consist of Sagittal Abdominal Diameter (SAD), Ratio SAD/Height and Abdominal Diameter Index (ADI) with insulin resistance in obese female adolescents.

Method: This study was conducted in June-August 2019 at Universitas Diponegoro, Central Java. this study used cross sectional design with a total of 120 female students aged 17-21 years and selected using purposive sampling method. Study variables included SAD, SAD/Height, ADI and insulin resistance level. SAD was measured by abdominal caliper, SAD/Height was ratio of SAD divided by height. ADI was ratio of SAD divided by thigh circumference. The value of insulin resistance was obtained from the calculation of homeostasis model assessment insulin resistance (HOMA-IR). Data normality was analyzed by Kolmogorov-smirnov test, all data were not normal distribution so the bivariate test analyzed by Rank spearman test.

Results: As many as 42.5% subjects had SAD which included at risk and 83.3% subjects had suffering insulin resistance. There was significant relationship between SAD, SAD/Height, ADI with insulin resistance (SAD (p=0.001), SAD/Height (p<0.001) and ADI (p=0.003). In addition, SAD, SAD/Height, ADI were significant relationship with fasting blood glucose (SAD (p=0.048), SAD/Height (p=0.022) and ADI (p=0.003).

Conclusion: There were significant relationship between SAD, SAD/Height, ADI with insulin resistance and fasting blood glucose in obese adolescent female.

Keywords: Sagittal Abdominal Diameter (SAD), SAD/Height, Abdominal Diameter Index (ADI), insulin resistance

INTRODUCTION

Obesity is one factor of that cause various diseases such as cardiovascular disease, hypertension, cancer, and diabetes mellitus. Most countries in Asia have increasing of overweight and obesity prevalence in this last decade. Increased of obesity prevalence does not only occurred in adult group but also occurred in adolescent group. Indonesia is the one of the countries that have high prevalence of abdominal obesity in the Southeast Asian region (1). The prevalence of visceral/abdominal obesity at age ≥ 15 years based on Basic Health Research 2013 as many as 26.6% and increase to 31% in 2018 (2,3). A study of adolescent in Semarang showed that 46 of 566 adolescents suffering abdominal obesity (4). The prevalence of abdominal obesity in female higher than male, the result data of Basic Health Research 2013 showed that female group who suffering abdominal obesity is 56.3% while the male group is 43.7% (5).

Abdominal obesity can reflect excess of visceral fat tissue or subcutaneous or both. Visceral fat contributes to the pathogenesis of diabetes mellitus, glucose intolerance and

insulin resistance. Insulin resistance is a impaired condition of glucose absorption that induced by insulin (6). Fat tissue distribution in intraabdominal area caused increasing metabolic such as lipogenesis and lipolysis activity. Lipolysis produce *free fatty acids* (FFA), FFA from visceral fat brought to the portal circulation and liver. Excessive of FFA secretion from visceral fat through the liver cause toxic effect for example gluconeogenesis and insulin resistance (6,7). Study of 220 obese children and adolescents in Brazil shows that 33.2% subjects suffering insulin resistance (8). Study of 52 obese adolescents in Semarang also shows that 91.6% subjects suffering insulin resistance (9).

Excessive of visceral fat tissue in abdominal obesity can measure by *Computed Tomography*(CT) as gold standard, but measurement of Waist Circumference (WC), and abdominal diameter also can be used as alternative. Abdominal diameter measurement consist of three components, namely Sagittal Abdominal Diameter (SAD), Sagittal Abdominal Diameter/Height (SAD/Height), Abdominal Diameter Index (ADI) (10). SAD can describe the visceral fat size, moreover it can also predicting metabolic syndrome from visceral fat area. SAD is closely related with visceral fat mass than other

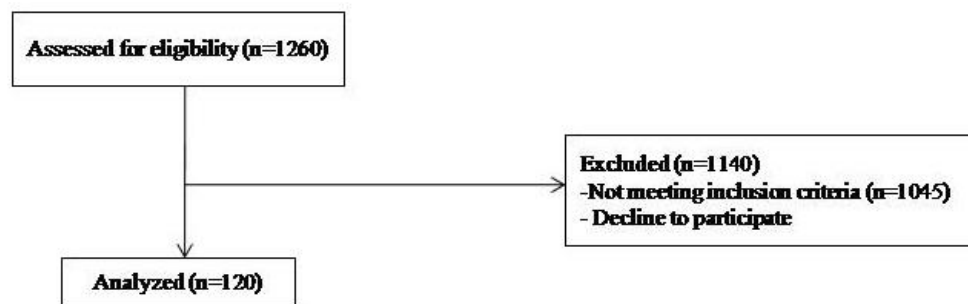


Figure 1. The flow diagram of subject inclusion and exclusion in this research

anthropometric measurements. SAD measured by supine position when subcutaneous fat move to side of waist. SAD measurement at this position reflect size of intraabdominal fat in antero-posterior like in CT image measurement. SAD strongly related with glucose intolerance, cardiovascular disease and good indicator for predicting insulin resistance (10-13).

SAD/Height is derivative of SAD measurement, which is use height as divisor of SAD. SAD/Height is anthropometric indicator that significantly related to visceral fat tissue (10,14). ADI is derivative of SAD measurement divided by thigh circumference. ADI is significantly related to several indicators of metabolic syndrome, and maybe has significant relationship with insulin resistance (15). A study in Brazil also showed that SAD/Height and ADI is good indicator for measurement visceral fat tissue in women (16). Study on abdominal diameter (SAD, SAD/Height and ADI) as insulin resistance indicator has not been much done especially in Indonesia. Therefore, based on this background the researchers interest to identifying the relationship of abdominal diameter (SAD, SAD/Height and ADI) with insulin resistance in obese adolescent female.

METHODS

This study was conducted in June-August 2019 with female student subject at Diponegoro University Semarang. This study was a observational analytic with cross sectional design. As described in **Figure 1**, of 1260 participants from this study were selected by screening process which obtained 1260 subjects, after that 215 subjects were included to inclusion criteria and selected as many as 120 subjects who involved in this study using purposive sampling method. The inclusion criteria were female student in Diponegoro University Semarang, aged 17-21 years, having waist circumference >80 cm, not pregnant, not consuming alcohol, not consuming drug that can related to glucose and insulin level, willing to fast minimum 8 hours before taking blood sample. The exclusion criteria was the subject resign during the study. This study had received permission from the Health Research Ethics Committee with Number 373/EC/KEPK/FK UNDIP/VIII/2019.

The independent variables in this study were abdominal diameter consisting of SAD, SAD/Height, and ADI. SAD measured by abdominal caliper which is manually measurement. Measurement were conducted with supine position, subject relax on the table. One of caliper's arm was placed under the subject (*iliac crest*) and the other caliper's arm was slowly moving down on subject abdomen with normal breathing (14). SAD categorized as normal if <19.3 cm and at

Table 1. Subject characteristic

Variable	Min	Max	Median±SD
Age (year)	17	21	19± 0.8
Weight (kg)	47.80	107.40	66.60±11.38
Height (cm)	141.20	171.40	157.40±5.44
Sagittal Abdominal Diameter(cm)	14.90	26.10	18.68±2.12
Sagittal Abdominal Diameter/Height (ratio)	0.09	0.17	0.12±0.014
Abdominal Diameter Index/ADI	0.03	0.56	0.32±0.41
Fasting blood glucose(mg/dL)	68.00	206.00	87.00±13.15
Insulin (μU/mL)	2.71	52.60	10.82±6.92
HOMA-IR	0.54	18.32	2.33±2.09

risk if ≥19.3 cm (17). SAD/Height was derivative of SAD, which is use height (in centimeter) as SAD divisor. ADI was derivative of SAD measurement divided by thigh circumference (in centimeter). The thigh circumference was measured on the right side of the body, at the midpoint between inguinal crease and proximal border on the patella (16).

The dependent variable in this study was insulin resistance. Insulin resistance is a disruption of the biological response to insulin which cause body insulin requirement increase and then resulting hyperinsulinemia to maintain normal glucose level. Insulin resistance level obtained from calculating *Homeostasis Model Assessment Insulin Resistance* (HOMA-IR), with formula (18):

$$HOMA\ IR = \frac{fasting\ insulin\ \left(\frac{mU}{L}\right) \times fasting\ glucose\ \left(\frac{mmol}{L}\right)}{22.5} \quad (1)$$

Cut off the normal value of HOMA-IR in adolescent was <1.65 (9). Blood sample of subject was taking by laboratory officer. Before the blood was taken, the subject fasting for 8-12 hours and then blood was taken 5 cc.

Univariate data analyze was used to describe the characteristic of each variable in this study. Data normality test analyzed by Kolmogorov-smirnov test. Bivariate data analyzed by Rank spearman test for abnormal data distribution. The purpose of bivariate analysis were to determine relationship of each abdominal diameter variable with insulin resistance.

RESULT

Subject Characteristic

Table 1 shows subject characteristic of this study. Age range of subjects were 17-21 years old. The highest fasting blood glucose of subject was 206 mg/dL while the lowest fasting blood glucose was 68 mg/dL. The median in insulin level

Table 2. Frequency distribution of sagittal abdominal diameter, fasting blood glucose, fasting insulin and HOMA-IR

Characteristic	n	%
Sagittal Abdominal Diameter		
No risk	69	57.5
At risk	51	42.5
Fasting Blood Glucose		
Normal	117	97.5
High	3	2.5
Fasting Insulin		
Normal	107	89.2
High	13	10.8
HOMA-IR		
Normal	20	16.7
Resistance	100	83.3

Table 3. Relationship of abdominal diameter with insulin resistance and fasting blood glucose

Variable	Fasting blood glucose		HOMA-IR	
	r	p	r	p
Sagittal abdominal diameter	0.181	0.048	0.297	0.001
SAD/Height	0.209	0.022	0.345	<0.001
Abdominal diameter index	0.270	0.003	0.273	0.003

of subject was 0.82 μ U/mL. The highest HOMA-IR value of subject was 18.32 with median value 2.33. The median in SAD measurement was 18.68 cm. The highest SAD measurement of subject was 26.10 cm.

Frequency distribution of sagittal abdominal diameter, fasting blood glucose, fasting insulin and HOMA-IR were presented in **Table 2**. Almost half of the subjects (42.5%) had SAD that categorized at risk. Majority of subjects (89.2%) had normal fasting insulin level. Almost of subjects (97.5%) had normal fasting blood glucose, whereas conversely most of subjects (83.3%) had suffering insulin resistance.

Relationship of Abdominal Diameter with Insulin Resistance

Bivariate analysis in this study used Rank Spearman test. Relationship of abdominal diameter with insulin resistant and fasting blood glucose were presented in **Table 3**. In this study, we found that SAD, SAD/Height and ADI had significant relationship with insulin resistance and fasting blood glucose ($p < 0.05$), with positive correlation.

DISCUSSION

The result of this study shows that SAD median was 18.68 \pm 2.12cm with minimum value 14.90 cm and maximum value 26.10 cm. This result similar to study that conducted in Brazil by Feliciano et al. showed that in overweight adolescent the SAD measurement reached 26.3 cm (19). SAD is indicator that can describe visceral fat size (20). SAD measured by supine position, visceral abdominal fat tend to lift the abdominal wall toward sagittal, abdominal subcutaneous fat pressure the stomach and subcutaneous fat moves to the waist, so SAD can reflect the visceral abdominal fat volume (21). As many as 42.5% subjects had SAD which categorized at risk. SAD in women closely related with hyperlipidemia, insulin resistance and cardiovascular risk, moreover it can also be used as a predictor of cardiometabolic risk score (22). The study of

children and adolescent in Arab shows that SAD significantly related with the severity of metabolic syndrome components (23).

The result of this study shows that HOMA-IR median was 2.33 \pm 2.09cm with minimum value 0.54 cm and maximum value 18.32 cm. The high score of HOMA-IR similar to González-Zavala et al.'s study that found high score of HOMA-IR in overweight adolescent subjects that reached 18.32 (24). This study showed that the HOMA-IR most of subjects (83.3%) included in insulin resistance category, however, almost of subjects had normal fasting blood glucose and normal fasting insulin. This result similar to Yi Kung Hye et al.'s study that found high prevalence of insulin resistant in obese adolescent subject was 47.1% (25). The study of 38 obese adolescent in Semarang also shows similar result with this study. This condition can occur because hyperglycemia is the last part from insulin resistance condition, which is long enough condition of impaired glucose intolerance and or impaired fasting blood glucose. Carbohydrate metabolism in adolescence also still good, so the body can maintaining normal glucose level (26). The high incidence of insulin resistance who suffering by subjects increased the risk of metabolic syndrome, because insulin resistance is a risk factor for metabolic syndrome that first arise (9). The highest HOMA-IR value in the subject was 18.32. Adolescent group with metabolic syndrome significantly had higher of HOMA-IR. HOMA-IR is indicator that strongly correlated with total and basal insulin. The HOMA-IR mean progressively increase with following sex maturation increase and weight from normal to obese. The study in India shows that the increase of HOMA-IR significantly related with body mass index, moreover there was significant different between HOMA-IR in adolescent with normal nutritional status, overweight and obesity (27).

Bivariate analysis in this study showed that SAD, SAD/Height, and ADI significant related to insulin resistance with positive correlation. A study of 75 overweight and obese adolescent showed that SAD and insulin resistance had significant relationship, with correlation that not much different from this study ($r=0.485$). The study also showed that SAD described a better correlation with insulin resistance than other anthropometric measurements. SAD can predict insulin resistance because it measures abdominal visceral fat than subcutaneous fat. Abdominal visceral fat also had strongly related with cardio metabolic disease (28). The study of women subject in Brazil comparing SAD with body mass index (BMI), waist circumference (WC), waist to hip ratio (WHR) as resistance insulin indicator, the result show that SAD and BMI was strongest indicator to indicate the occurrence of insulin resistance. BMI was widely used to identify metabolic risk but it can not be used alone, because it can not distinguish muscle adipose tissue or body fat distribution, whereas SAD had reported as a good marker for visceral adipose tissue in various ethnic group (21).

In contrast with SAD, study about relationship between SAD/Height ratio and ADI with insulin resistance still limited, but study of 194 subjects in Brazil shows that SAD/Height and ADI had significant related and positive correlation with visceral adipose tissue (16). A study of 1347 subjects in Korea showed that high visceral fat mass and low adiponectin level were correlated with increased risk of insulin resistance and β cell dysfunction. Obesity is well known as risk factor for insulin resistance. Abdominal obesity can cause insulin resistance by stimulating metabolic product formation which derived from

fat, hormone and cytokines. Insulin resistance can cause endothelial dysfunction and insulin signaling pathway changes. Several previous studies show that visceral adipose tissue can inhibit adiponectin secretion. Adiponectin mediates the insulin sensitivity system and glucose homeostasis (29). Insulin resistance is adaptive response of *Free Fatty Acid* (FFA) increase, and related with visceral adipose tissue level. FFA increase can change the liver, muscle and other tissue metabolism toward lipid deposition and oxidation. Insulin secretion increase offset the glucose capacity decrease. Subcutaneous adipose tissue taking FFA and storing excess calories in visceral adipose tissue (30). Excess FFA secretion from visceral fat will be oxidized, stored (lipids droplets) or metabolized to toxic derivatives (DAG and ceramides). Toxic derivatives cause insuline resistance, impaired cell function (lipotoxicity) or cause apoptosis. Apoptosis in the pancreas cause decrease and impairment of β cell capacities to secrete insulin which increase the risk of developing type 2 diabetes mellitus. The consequence of insulin resistance increase is endogenous glucose production by liver and glucose utilization decrease by peripheral tissue. As result are glycemia and insulin pancreas secretion increase, beside that occur hepatic insulin clearance impaired, and causing hyperinsulinemia (31).

This study also showed that significant relationship between abdominal diameter (SAD, SAD/Height, and ADI) with fasting blood glucose level. Abdominal diameter measurement can reflect visceral fat. Visceral/abdominal fat stores are considered as implication of insulin resistance, cardiovascular and other metabolic condition which related to type 2 diabetes mellitus. Excessive visceral fat related with impaired of insulin sensitivity. Visceral fat related with impaired glucose regulation, because it is closely related to insulin resistance (32). Lypolysis of visceral fat can cause free circulated of FFA in obese individual. This cause FFA enter to portal vein which can accumulated in the liver and cause impaired of glucose tolerance and insulin homeostasis. Visceral fat tissue and liver fat tissue related with insulin resistance in diabetic patient. A study of 297 subject showed that subject with isolated impaired fasting glucose and isolated impaired glucose tolerance had excess abdominal fat (visceral fat, abdominal subcutaneous fat and liver fat) (33).

CONCLUSION

As many as 42,5% subjects had SAD were categorized at risk, while 83,3% subject had suffering resistance insulin. There were significant relationship between abdominal diameter (SAD, SAD/Height, ADI) with insulin resistance and fasting blood glucose in obese adolescent female, with positive correlation. As it is important to monitor the growth and development of adolescents over time, it is advisable to standardize the use of one measure of body fat location. Considering that abdominal fat, more than total fat, has been associated with cardiometabolic risk, it is recommended the use abdominal diameter measurement (SAD, SAD/Height, ADI) as a measure that reflects the adipose tissue in this region, in the assessment of the nutritional status of adolescents.

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Fillah Dieny <fillahdieny@gmail.com>

Your article has been published in EJGM

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Balas Ke: editor@ejgm.co.uk

Kepada: Fillah Fithra Dieny <fillahdieny@gmail.com>, A Fahmy Arif Tsani <fahmi_tsani@yahoo.com>

Dear Dr. Dieny,

We would like to inform you that your article entitled **Abdominal Diameter Profiles have Relationship with Insulin Resistance in Obese Female Adolescents** has been published in **Electronic Journal of General Medicine**:

URL: <http://www.ejgm.co.uk/article/abdominal-diameter-profiles-have-relationship-with-insulin-resistance-in-obese-female-adolescents-7882>DOI: <https://doi.org/10.29333/ejgm/7882>

Thank you publishing with us. We are looking forward to your future publications.

Best regards,

Electronic Journal of General Medicine

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ABDOMINAL DIAMETER PROFILES HAVE RELATIONSHIP WITH INSULIN RESISTANCE IN OBESE FEMALE ADOLESCENTS
(Fillah Fithra Dieny, Ria F Setyaningsih, Deny Yudi Fitrianti, Firdananda Fikri Jauharany, Yehezkiel D Putra, A Fahmy Arif Tsani)

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Paper No: 14-793

Dear Prof. /Dr. /Mr. /Mrs. /Miss/ Ms.

NOTIFICATION OF ACCEPTANCE FOR PRESENTATION IN INTERNATIONAL CONFERENCE ON HEALTH AND WELL BEING (ICHWB2019)

Thank you for your participation in the International Conference on Helth and Well Being (ICHWB2019).
The ICHWB2019 committee has completed the review for your paper and suggested the following recommendations.

Presentation Status

- ✓ Accepted for ICHWB2019 presentation

Publication

- ✓ Current status: Accepted

Publication in journal indexed by SCOPUS. You might be requested to revise your paper (will be notified, if requested).

Payment: International USD 400 / Indonesia IDR 5.000.000

Payment Arrangement (before 27 November 2019)

Please proceed the payment arrangement.

The payment information and the account address can be found in <http://iseth.ums.ac.id/2019/reg19>

For presentation schedule purpose, proceed with the online registration and upload the payment proof:

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Looking forward to meeting you in ICHWB2019 Conference.

Sincerely Yours,



Setyaningrum Rahmawaty, Ph.D.

Conference Chair ICHWB2019

International Conference on Health and Well Being (ICHWB2019)

setyaningrum.rahmawaty@ums.ac.id
