

ORIGINAL PAPER

ORYGINALNY ARTYKUŁ NAUKOWY

**EFFECT OF OBESITY ON THE RELATIONSHIP BETWEEN GLYCOSYLATED
HEMOGLOBIN AND INSULIN RESISTANCE IN YOUNG ADULTS**

**WPLYW OTYŁOŚCI NA ZWIĄZEK MIĘDZY HEMOGLOBINĄ
GLIKOZYLOWANĄ A INSULINOOPORNOŚCIĄ U MŁODYCH OSÓB
DOROSŁYCH**

**Sümeyra Topal^{1(A,B,C,D,E,F)}, Sinem Yalınzoğlu Çaka^{2(A,B,C,D,E)}, Nursan Çınar^{3(A,D,E,F)},
Sevin Altınkaynak^{3(A,D,E,G)}**

¹Department of Pediatric Nursing, Faculty of Health Science, Kahramanmaraş İstiklal University,
Kahramanmaraş, Türkiye

²Department of Pediatric Nursing, Faculty of Health Science, Kocaeli University, Kocaeli, Türkiye

³Department of Pediatric Nursing, Faculty of Health Science, Sakarya University, Türkiye

Topal S, Yalınzoğlu Çaka S, Çınar N, Altınkaynak S. Effect of obesity on the relationship between glycosylated hemoglobin and insulin resistance in young adults. Health Prob Civil. <https://doi.org/10.5114/hpc.2024.139159>

Tables: 4

Figures: 1

References: 36

Submitted: 2024 Jan 24

Accepted: 2024 Apr 22

Address for correspondence / Adres korespondencyjny: Sinem Yalınzoğlu Çaka, Department of Pediatric Nursing, Faculty of Health Science, Kocaeli University, Kabaoğlu, Baki Komsuoğlu

Boulevard No. 515, Umuttepe, 41001 Izmit/Kocaeli, Türkiye, e-mail: sinemyalnizoglu@gmail.com,
phone: +90 (262) 303 10 00

ORCID: Sümeyra Topal <https://orcid.org/0000-0001-6316-4043>, Sinem Yalınzoğlu Çaka
<https://orcid.org/0000-0002-1572-7013>, Nursan Çınar <https://orcid.org/0000-0003-3151-9975>, Sevin
Altınkaynak <https://orcid.org/0000-0001-6417-0402>

Copyright: © John Paul II University in Białá Podlaska, Sümeyra Topal, Sinem Yalınzoğlu Çaka, Nursan Çınar, Sevin Altınkaynak. This is an Open Access journal, all articles are distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 4.0 International (CC BY-NC-SA 4.0) License (<http://creativecommons.org/licenses/by-ncsa/4.0/>), allowing third parties to copy and redistribute the material in any medium or format and to remix, transform, and build upon the material, provided the original work is properly cited and states its license.

Summary

Background. The purpose of this prospective case-control research was to compare insulin resistance in obese and non-obese students with regard to Glycosylated Hemoglobin (HbA1c) and the Homeostatic Model Assessment Insulin Resistance (HOMA-IR).

Material and methods. This study was conducted with a total of 122 students who were determined to be obese (n=35), overweight (n=43) and normal weight (n=44), among students at a university in Türkiye (n=891). Students were first classified according to their Body Mass Index (BMI) and blood samples were taken for fasting blood glucose, HgA1c, and insulin levels.

Results. Of the 891 students participating in the study, 4.83% (n=43) were overweight and 3.93% (n=35) were obese. HbA1c levels were found below 5.7 for 72.7% of normal weight students and HbA1c levels were found between 5.7-6.4 for 65.7% of obese students. There was a significant statistical difference between them ($p=0.001$). When the HOMA-IR value

was >3.16 , there was a significant difference between the obese group and the other groups ($p=0.042$). There was a significant positive correlation between BMI and both HbA1c ($r=0.299, p=0.001$) and HOMA-IR ($r=0.432, p=0.000$).

Conclusions. It was found that HOMA-IR was more common in obese students, and early diagnosis of insulin resistance might be HbA1c. HbA1c can be used since it is inexpensive and simple to evaluate insulin resistance in obese young adults.

Keywords: HbA1c, insulin resistance, young adults, nursing care, obesity

Streszczenie

Wprowadzenie. Celem tego prospektywnego badania kliniczno-kontrolnego było porównanie młodych osób dorosłych, którzy nie mieli nadwagi z tymi, którzy byli otyli pod względem związku między hemoglobina glikozylowaną (HbA1c) a oceną modelu homeostazy insulinooporności (HOMA-IR).

Material i metody. Badania przeprowadzono z udziałem 122 studentów, u których stwierdzono otyłość ($n=35$), nadwagę ($n=43$) i prawidłową masę ciała ($n=44$), wśród studentów uniwersytetu w Turcji ($n=891$). Studenci zostali najpierw sklasyfikowani według wskaźnika masy ciała (BMI), a następnie pobrano od nich próbki krwi w celu oznaczenia poziomu glukozy na czczo, HgA1c i insuliny.

Wyniki. Spośród 891 studentów biorących udział w badaniu, 4,83% ($n=43$) miało nadwagę, a 3,93% ($n=35$) było otyłych. Poziom HbA1c wynosił poniżej 5,7 u 72,7% studentów z prawidłową masą ciała, a poziom HbA1c między 5,7 a 6,4 u 65,7% studentów otyłych. Istniała między nimi istotna różnica statystyczna ($p=0,001$). Gdy wartość HOMA-IR wynosiła $>3,16$, występowała istotna różnica między grupą otyłych studentów a pozostałymi grupami ($p=0,042$). Stwierdzono istotną dodatnią korelację między BMI a HbA1c ($r=0,299, p=0,001$), jak i HOMA-IR ($r=0,432, p=0,000$).

Wnioski. Stwierdzono, że HOMA-IR występował częściej u otyłych studentów, a wczesną diagnozą insulinooporności może być HbA1c. Badanie HbA1c jest niedrogi i proste dlatego może być stosowane w ocenie insulinooporności u otyłej młodzieży.

Słowa kluczowe: HbA1c, insulinooporność, młodzi dorośli, opieka pielęgniarska, otyłość

Introduction

Obesity is defined as a high level of fat caused by an imbalance between energy intake and consumption to a level where it harms health [1]. Obesity is a problem of energy metabolism that is becoming more common in both developed and developing nations. Obesity has become one of the leading causes of disability and death, affecting not only adults but also children and adolescents worldwide [2]. The World Health Organization (WHO) study states that obesity is becoming more commonplace each year. According to the WHO data, it is reported that adult obesity has more than doubled worldwide since 1990, and obesity in adolescence has quadrupled. It is emphasized that in 2022, 43% of adults aged 18 and over will be overweight and 16% will live with obesity [3]. Like in other nations across the world, the prevalence of obesity is rising daily in our own country. According to the results of the Türkiye Health Survey (2022), when calculated according to body mass index, 20.2% of people aged 15 and over were obese and 35.6% were overweight. When examined by gender, it is seen that 23.6% of women were obese and 30.9% were overweight, while 16.8% of men were obese and 40.4% were overweight [4].

Obesity causes negative metabolic changes on blood pressure, triglyceride, cholesterol, and insulin resistance. Insulin resistance is linked to the obesity-diabetes relationship. The primary factors contributing to the development of type 2 diabetes mellitus

(T2DM) are insulin resistance and hyperinsulinemia. Insulin resistance and severe hyperinsulinemia combine with central obesity. Specifically, research has shown that the height of Body Mass Index (BMI) closely correlates with an increased risk of diabetes [5-9]. In research conducted by Kılıç et al., fasting blood glucose and HbA1c levels of individuals aged 18 and older with different BMI were analyzed, and it was determined that the prevalence of diabetes increased with weight gain [4]. In the study conducted by Lee et al. [9] on 126 obese and overweight children in the 7–14 age range, it is recommended that obesity is linked to decreased glucose tolerance and that the HbA1c level should be checked in addition to the oral glucose tolerance test [9].

Obesity is a pathophysiologic component of T2DM and is frequently associated with insulin resistance [10]. Hyperglycemia, or elevated blood sugar, is a symptom of diabetes mellitus (DM), a multifaceted chronic condition caused by insufficient insulin production, action, or both [11]. Although the exact mechanism is unknown, obesity may contribute to long-term, low-grade systemic and local inflammation that develops into insulin resistance-associated DM [3,6]. Furthermore, hyperinsulinemia and insulin resistance may play a role in the development of obesity [1].

Aim of the work

Several researchers suggested the connection between obesity and T2DM. However, from what we have observed, little is known about the combined effects of HbA1c level and overweight/obesity on the increased risk of insulin resistance, especially in the young Turkish population. The appearance of obesity from an early age and the regular increase in insulin resistance may explain the earlier onset of impaired glucose tolerance in children and adolescents.

The purpose of this prospective case control research was to compare insulin resistance in obese and non-obese students with regard to HbA1c.

Material and methods

Study design

The study involves two stages.

The 1st stage of the study

The population of the study comprised 1121 students enrolled in the Sakarya University, Faculty of Health Sciences for the 2016-2017 academic year, while the sample was made up of 891 students who willingly agreed to take part in the research.

The 2nd stage of the study

The G*Power (v3.1.7) tool was used for power analysis to calculate the sample size. The effect size was calculated using HbA1c mean (5.68 and 5.92) and standard deviation (0.26 and 0.42) values obtained from the publication entitled “HbA1c and glucose intolerance in obese children and adolescents” by Lee et al. [9]. In accordance with the calculation, it was shown that a minimum of 35 students needed to participate in each group. 35 obese students, 43 overweight students, and 44 normal weight students constituted the control group. All students who were obese and met the sampling criteria were included in the study. Overweight and normal weight students were selected by a simple random sampling method.

The following individuals were excluded from the study: those who did not consent to participate, those with systemic infections or genetic and chronic disorders, as well as anyone taking medication (metformin, corticosteroid, etc.) that may alter the test findings.

Instruments

Height and weight measurements were performed with the DESIS M301 brand stadiometer. The device was installed and calibrated before using. Weight measurement was taken with a digital scale sensitive to 100 grams. Care was taken to ensure that the individual wore as little clothing as possible and no shoes. During the measurement, individuals could not receive any support and had to step on the scale evenly. BMI were calculated by Weight (kg) / Height² (m) formula. According to BMI values, 18.5-24.9 was normal weight, 25.0-29.9 was pre-obesity, 30.0-39.9 was obesity and above 40 was considered morbid obesity [3].

Biochemical analyses

Blood samples were obtained by two researchers (ST and SYÇ) from the students in the case and control groups for HgA1c, fasting blood glucose and insulin levels. Peripheral venous blood samples were collected after a 12-hour fasting period. In the study, HgA1c and fasting blood glucose tests were measured with the Architect® ci4100 Analyzer (Abbott Diagnostics Ltd.). Serum insulin was measured with a Roche Cobas E601 Immunology Analyzer. Blood collection, anthropometric and biochemical measurements were documented with ISO 9001 and CE standards. Blood samples were transferred to the hospital where the samples were tested. Venous blood samples were collected in EDTA tubes and stored in the refrigerator (+4 °C) until sent to the laboratory. The insulin resistance was calculated (Fasting

Glucose (mg/dl)/Fasting Insulin (MIU/ml) /405) using the Homeostasis Model Assessment-Insulin Resistance (HOMA-IR) index. If HOMA-IR value was above 3.16, it was evaluated as insulin resistance [7,8,12]. Those with HbA1c \geq 6.5% were considered DM, and those with HbA1c values of between 5.7 and 6.4 were considered to have a prediabetes risk [13].

Ethical dimension of the study

The study received ethical approval (Approval number: 16214662/050.01.04/146) from the Medical Faculty Ethics Committee at Sakarya University. Written and verbal approvals were obtained by reading the informed consent to the students participating in the study. The Declaration of Helsinki was followed in the conduct of the study.

Statistical analysis

Statistical analyses were performed using the SPSS 25.0 software. Frequency (f) and percentage (%) of descriptive statistics were used as the statistical methods in the study. An independent group t-test and one-way ANOVA test was used for those with normal distribution, and Kruskal Wallis-H test was used because the distribution was non-standard. Bonferroni correction was used to adjust the significance level of multiple comparisons to control the type I error rate. A Chi square test was used to determine the difference between the expected frequencies. The Spearman correlation analysis was used to investigate the relationship between the values. In addition, $p < 0.05$ was taken as the significance level.

Results

A total of 122 young adults consisting of 35 obese, 43 overweight and 44 normal weight individuals according to BMI values were included in the study (Figure 1). The mean BMI value was 33.08 ± 2.57 in the obese group, 26.70 ± 1.25 in the overweight group and 21.94 ± 1.68 in the normal weight group. The ratios of normal weight, overweight and obese females were respectively 77.3%, 65.1%, and 91.4%.

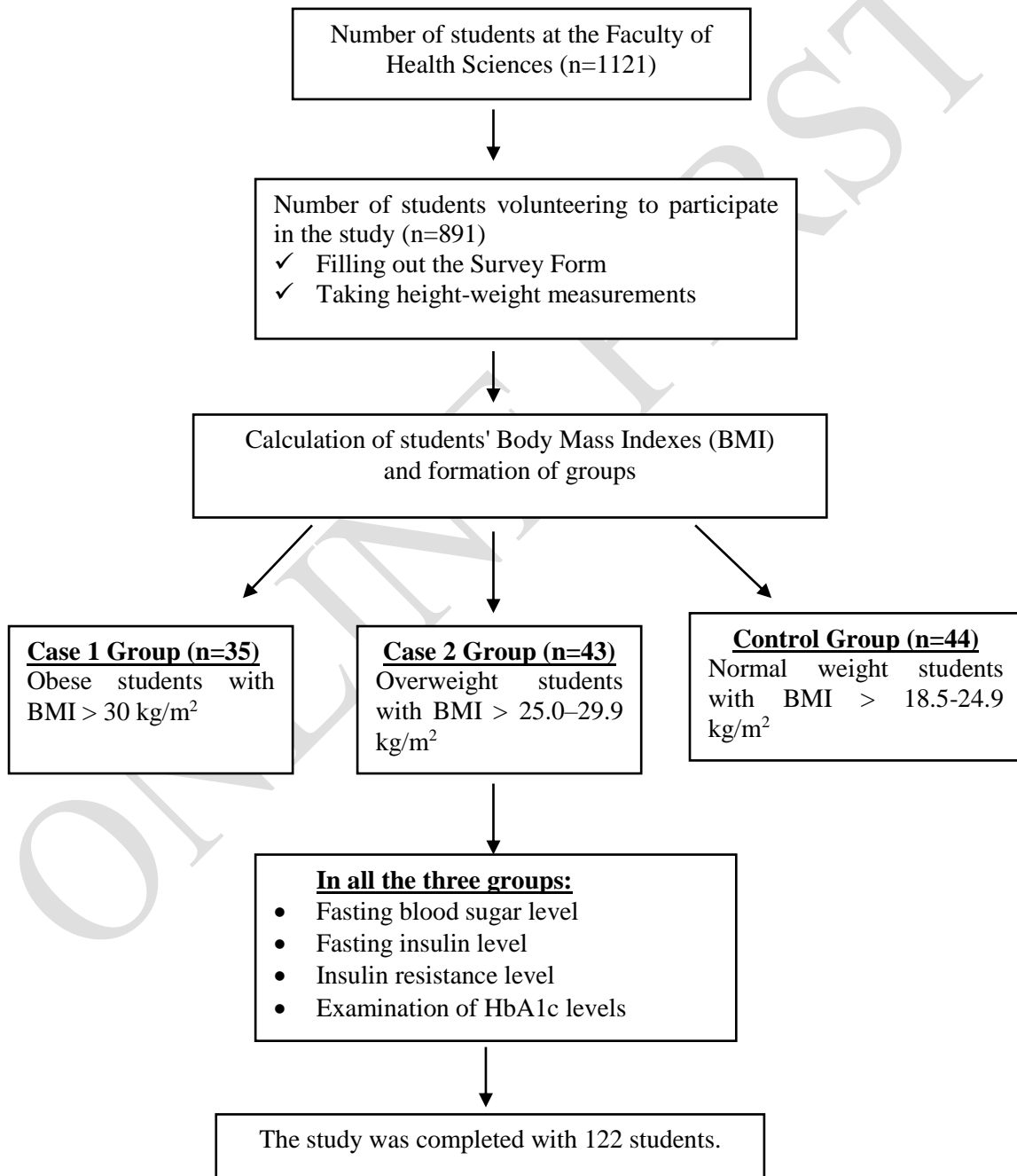


Figure 1. Flow chart of the study

Among the students who participated in the research, the average age of those with normal weight was 20.43±0.9. The mean age was 20.39±0.97 for the overweight group, and 20.05±1.05 for the obese, and there was no statistically significant difference between the groups ($p=0.176$). It was determined that 77.3% of the participants in the normal weight group were female, 84.1% were middle income, 36.4% lived with their friends, 75% did not smoke, and 77.3% preferred public transportation. It was determined that 65.1% of the overweight participants were women, 79.1% were middle income, 30.2% lived with friends, 86.0% did not smoke, and 67.4% preferred public transportation. Of those in the obese group, 91.4% were women, 88.6% were middle income, 31.4% lived with family members, 82.9% did not smoke, and 57.1% preferred public transportation. It was determined that they preferred public transportation. While there was no significant difference between the groups in terms of age, working status, place of residence, smoking/alcohol use, and regular exercise status, a significant difference was performed with the mode of transportation ($p=0.028$) (Table 1).

Table 1. Comparison of demographic characteristics of adolescents according to BMI

Variables	BMI						Test	
	Normal weight		Pre-obesity		Obesity		KW	p
	Mean±SD	Mean±SD	Mean±SD	Mean±SD	Mean±SD	Mean±SD		
Age (years)	20.43±0.94		20.39±0.97		20.05±1.05		3.479	0.176
Variables	n	%	n	%	n	%	χ^2	p
Gender								
Female	34	77.3	28	65.1	32	91.4	7.556	0.023
Male	10	22.7	15	34.9	3	8.6		

Working status								
Yes	6	13.6	3	7.0	1	2.9	2.824	0.239
No	38	86.4	40	93.0	34	97.1		
Perceived economic level								
Income higher than expenses	7	15.9	6	14.0	4	11.4	4.220	0.339
Income equivalent to expenses	37	84.1	34	79.1	31	88.6		
Income lower than expenses	0	0.0	3	7.0	0	0.0		
Family type								
Nuclear family	33	75.0	32	74.4	28	80.0	2.039	0.766
Extended family	10	22.7	10	23.3	5	14.3		
Fragmented family	1	2.3	1	2.3	2	5.7		
Place of residence								
Dormitory	12	27.3	11	25.6	10	28.6	9.643	0.431
Private dormitory	8	18.2	9	20.9	7	20.7		
Alone at home	0	0.0	0	0.0	1	2.9		
At home with family	8	18.2	10	23.3	11	31.4		
At home with a relative/friend	16	36.4	13	30.2	6	17.2		
Smoking status								
Smoking	11	25.0	6	14.0	6	17.1	1.828	0.419
Not smoking	33	75.0	37	86.0	29	82.9		
Alcohol use								
Yes	3	6.8	4	9.3	4	11.4	0.623	0.797
No	41	93.2	39	90.7	31	88.6		
Regular exercise								
Yes	4	9.1	6	14.0	2	5.7	1.417	0.438
No	40	90.9	37	86.0	33	94.3		

Mode of transportation								
Usually on foot	10	22.7	10	23.3	15	42.9	9.514	0.028
Public transport	34	77.3	29	67.4	20	57.1		
Private car	0	0.0	4	9.3	0	0.0		

Notes: BMI – Body Mass Index; KW – Kruskal-Wallis Test; χ^2 – Chi-square test; * $p < 0.05$.

HOMA-IR value was higher in obese young adults (3.27 ± 2.12) compared to normal weight (2.21 ± 2.85) and overweight (3.10 ± 3.84) young adults. In normal weight, overweight and obese students, HOMA-IR > 3.16 was 11.4%, 27.9% and 34.3%, respectively (Table 2). Insulin resistance was higher in the obese group and there was a significant statistical difference between them ($p < 0.001$). There was no statistically significant difference between glucose ($p = 0.355$) and insulin ($p = 0.152$) values of normal weight, overweight and obese students. HbA1c levels were below 5.7 in 72.7% of normal weight young adults; HbA1c levels were between 5.7 and 6.4 in 55.8% of overweight young adults, and HbA1c levels were between 5.7 and 6.4 in 65.7% of obese young adults, and there was a significant statistical difference between them ($p = 0.001$). HbA1c was higher in the obese group (5.74 ± 0.25) compared to the normal weight (5.47 ± 0.33) and overweight groups (5.69 ± 0.32), and there was a significant statistical difference between them ($p < 0.001$) (Table 2).

Table 2. HOMA-IR, glucose, insulin and HbA1c values of normal weight, overweight and obese young adults

Variables	BMI			Test	<i>p</i>
	Normal weight (1) Mean \pm SD	Pre-obesity (2) Mean \pm SD	Obesity (3) Mean \pm SD		
HOMA-IR	2.21 \pm 2.85	3.10 \pm 3.84	3.27 \pm 2.12	KW: 17.902	0.000** 1<2; 1<3

Glucose (mg/dl)	84.18±8.66		85.81±9.93		87.17±8.86		F: 1.045	0.355
Insulin (mIU/ml)	10.25±10.95		13.87±13.63		14.98±8.78		F: 1.913	0.152
HbA1c (%)	5.47±0.33		5.69±0.32		5.74±0.25		F: 8.879	0.000 **1<2; 1<3
Variables	n	%	n	%	n	%	χ^2	p
HOMA-IR <3.16	39	88.6	31	72.1	23	65.7	6.280	0.042*
HOMA-IR >3.16	5	11.4	12	27.9	12	34.3		
HbA1c<5.7	32	72.7	19	44.2	12	34.3	13.012	0.001**
HbA1c ≥ 5.7-6.4	12	27.3	24	55.8	23	65.7		

Notes: KW – Kruskal Wallis H Test; F – One Way ANOVA Test; χ^2 – Chi-squared test; * p <0.05;

** p <0.001

The glucose and insulin levels were significantly higher (p <0.001) in obese individuals with a HOMA-IR value of >3.16 compared to the obese group with a HOMA-IR value of <3.16. HbA1c and BMI did not show a statistically significant difference (p =0.558, 0.411 respectively). Also, HbA1c was significantly higher in normal weight individuals with a HOMA-IR value of >3.16 compared to the normal weight group with a HOMA-IR value of <3.16 (p =0.038). BMI, glucose and insulin values did not show a statistically significant difference (p =0.532, 0.463, 0.070 respectively) (Table 3).

Table 3. Comparison of glucose, insulin and HbA1c levels in the normal and obese weight groups according to HOMA-IR value

Obese group	HOMA-IR (<3.16) Mean±SD	HOMA-IR (>3.16) Mean±SD	t	p
Glucose (mg/dl)	85.04±8.20	91.25±8.98	-2.061	0.047*
Insulin (MIU/ML)	10.33±3.09	23.90±9.34	-4.897	0.000**

HbA1c (%)	5.73±0.26	5.78±0.25	-0.547	0.588
BMI (kg/m²)	32.81±2.50	33.58±2.75	-0.833	0.411
Normal weight	Mean±SD	Mean±SD	t	p
Glucose (mg/d)	83.38±6.34	90.40±19.27	-0.809	0.463
Insulin (MIU/ML)	7.37±2.06	32.72±23.14	-2.448	0.070
HbA1c (%)	5.45±0.34	5.64±0.14	-2.328	0.038*
BMI (kg/m²)	21.88±1.66	22.39±1.94	-0.629	0.532

Notes: t – Independent Sample t test; * $p < 0.05$, ** $p < 0.001$.

While a positive correlation was observed between BMI and insulin ($r=0.429$, $p < 0.001$), HbA1c ($r=0.299$, $p < 0.001$) and HOMA-IR ($r=0.432$, $p < 0.001$), there was no significant relationship with glucose ($r=0.107$, $p=0.204$) (Figure 2, Table 4).

Table 4. Relationship Between BMI, glucose, insulin, HbA1c and HOMA-IR values

Variables		BMI	Glucose	Insulin	HbA1c	HOMA-IR
BMI	r	-	0.107	0.429	0.299	0.432
	p	-	0.240	0.000**	0.001**	0.000**
Glucose	r	-	-	0.307	0.131	0.447
	p	-	-	0.001	0.151	0.000**
Insulin	r	-	-	-	0.235	0.983
	p	-	-	-	0.009**	0.000**
HbA1c	r	-	-	-	-	0.255
	p	-	-	-	-	0.005**

Notes: r – Spearman correlation analysis; * $p < 0.05$; ** $p < 0.001$.

Discussion

In this study, which was carried out to compare insulin resistance in obese and non-obese students with regard to HbA1c, some sociodemographic characteristics of case and control groups (mean age, economic level, family type, place of residence etc.) were determined. There was no statistically significant difference between the groups and both groups showed a homogeneous distribution in terms of the mentioned characteristics. The fact that similar results were obtained in both groups is one of the strengths of the study. However, there was a significant difference between the groups according to gender and mode of transportation (Table 1).

According to estimates from the World Obesity Federation, 254 million children between the ages of 5--19 and 206 million in 2025 will be obese [14]. Although there is no study reflecting the country-wide in our country, the frequency of childhood obesity is between 1.6% and 7.8% in the studies [14-19]. Among 891 young adults included in the study, 4.83% of them were overweight and 3.93% of them were obese, which is consistent with the findings of the studies carried out in our country. One significant risk factor for the onset of obesity is gender. The likelihood of becoming overweight or obese is twice as high in females [14,15,18,19]. Comprehending the disparities between genders when it comes to obesity may significantly influence how the disease is treated, as well as how community and individual initiatives for the pandemic's prevention and control are planned. In our study similar to the literature ratios of normal weight, overweight, and obese young females were higher compared to young males, and there was a statistically significant relationship between them. This significant difference may be due to the fact that there were more female students than males in the study. However, new literature is emerging on the importance of considering gender-related issues such as phases of the menstrual cycle, use of hormonal

contraception, and the impact of menopause when evaluating a patient with obesity [20-21]. In future studies, it would be useful to investigate potential reasons behind this difference for the female gender, such as differences in social norms, cultural influences or biological factors. Another important difference is the relationship between students' BMI value and mode of transportation. The number of students using public transportation is observably higher in the group with normal weight. The use of public transportation vehicles such as buses and trains actually requires students to be on the move during transportation (transfer between vehicles, walking distance, etc.). When the literature was examined, studies found that commuting to school by car was associated with obesity in students [22-24]. Wen et al. [22], in their study on the relationship between school travel patterns and obesity in children aged 10-13, reported that there was a positive relationship between driving to school every day and obesity [22]. Therefore, to prevent overweight/obesity, it may be recommended to encourage children to walk or cycle instead of traveling by car.

Chronic diseases such as DM, metabolic syndrome and hypertension caused by obesity and the prevalence of obesity are parallel. The key mechanism in the relationship between obesity and T2DM is insulin resistance. Insulin resistance causes hyperinsulinism, and when the beta cells start to secrete insulin at the expected level after a period of euglycemic state, hyperglycemia and then T2DM develop [1]. The BMI is known to have a considerable impact on the HOMA-IR score. It should be noted that as obesity becomes more common, HOMA-IR values might also vary. Obese people had higher HOMA-IR scores than teenagers who were overweight or of normal weight [25]. Teens who were normal weight, overweight, or obese had HOMA-IR rates of 11.4%, 27.9%, and 34.3%, respectively. Between these groups, there was a statistically difference, and HOMA-IR and BMI showed a positive link. According to a Japanese study, children who were overweight had significantly higher levels of fasting insulin and HOMA-IR, and there was no association found between

their BMI, HbA1c, and fasting glucose [26]. The interaction between HOMA-IR and BMI before adolescence may predict the likelihood of experiencing metabolic syndrome during puberty [27]. Therefore, reducing insulin resistance in childhood may contribute to the prevention of metabolic syndrome in adolescence and adulthood. Additionally, it is predicted that both metabolic syndrome and T2DM in adulthood may result from the presence of pediatric metabolic syndrome [28]. Additionally, T2DM in adulthood can also be predicted using the HOMA-IR index for insulin resistance at early ages [27,28]. However, the variability of HOMA-IR threshold levels used to define insulin resistance greatly hinders the interpretation of the results [29].

HbA1c is associated with a 6-8-week average glucose level formed by the irreversible bond established by glucose with hemoglobin. HbA1c between the range of 5.7-6.4 is the diagnostic criterion of prediabetes risk, and HbA1c \geq %6.5 is the diagnostic criterion of DM [13]. In this study, HbA1c was higher in the obese group compared to normal weight and overweight group. In the study carried out by Onal et al. [25] in the 3-15 age group, it was found that HbA1c in obese and normal weight children did not show a statistical significance. Upon reviewing the literature, no study has looked at the connection between HbA1c and BMI in youths older than 19.

Glucose and insulin levels were significantly higher in obese individuals with a HOMA-IR value of >3.16 compared to the obese group with a HOMA-IR value of <3.16 . Glucose and insulin levels did not show any statistically significant difference in normal weight individuals with a HOMA-IR value of >3.16 compared to the normal weight group with a HOMA-IR value of <3.16 . HOMA-IR, HbA1c. Insulin levels increased as the BMI increased, in other words, there was a positive correlation between them. Studies in overweight and obese –young people have found a statistically significant weak to moderate linear correlation between BMI and insulin, glucose, and HOMA-IR [19,30-32]. Research

suggests that obesity is strongly linked to a higher risk of metabolic disorders such as dyslipidemia, insulin resistance, and T2DM [33]. Insulin resistance is an integral feature of the metabolic syndrome and is an important determinant of the development of T2DM [34]. The main basis for obesity's link to T2DM is its ability to trigger insulin resistance. An imbalance between energy expenditure and food intake results in an excessive build-up of adipose tissue, which is the cause of obesity. In addition to being the primary location for storing extra energy from food consumption, adipose tissue is now understood to be an endocrine organ [1]. In order to take precautions against the risk factors of metabolic syndrome, healthcare professionals should plan personalized interventions in different dimensions and according to need. Today, it is known that healthcare personnel who receive special training specific to the subject to be trained are effective in reducing the risks of diseases and reducing costs [35,36].

HbA1c levels were below 5.7 in 72.7% of normal weight students, between 5.7 and 6.4 in 55.8% of overweight students, and between 5.7 and 6.4 in 65.7% of obese students, and there was a statistically significant difference among these groups. In our study, there was no student with a HbA1c level of >6.5 . However, as BMI increases, HbA1c also increases, meaning there was a positive correlation between them. These results indicate that obesity-associated insulin resistance has a major role in the development of DM. In our research, there was no statistically significant difference in HOMA-IR between those with HbA1c $>5.7-6.4$ and those with HbA1c <5.7 . A positive correlation was observed in the research carried out by Önal et al. [25]. We believe that the HOMA-IR cut-off point of 2.5 is connected to this outcome.

Study limitations

There are some limitations of our study that should be noted. Of the 891 students who participated in the study, 35 were obese, 43 were overweight and 813 were normal weight. In the normal weight group, 44 students were selected by simple random sampling method. The number of obese and overweight students in the group was low. HbA1c (cheap, simple and fasting is not required) can be used as a diagnostic criterion in the early diagnosis of insulin resistance. In the future, it may be recommended to conduct longitudinal studies with larger populations examining modifiable risk factors for obesity-related diseases, including interventions targeting the long-term effects of obesity on insulin resistance, and addressing its relationship with HbA1c.

Another limitation of the study is that although BMI remains a useful tool in many studies for assessing general trends in weight status and identifying individuals at high risk for certain obesity-related health conditions, other bioelectrical analytical measurements (skeletal muscle, body fat, etc.) are not used. The fact that the study was conducted in a cross-sectional design and the presence of potential confounding variables that were not taken into account also limits the study. There is a need for randomized controlled long-term studies on the subject.

Conclusions

In conclusion, increasing obesity in young people augers for metabolic syndrome in the future. Insulin resistance is one of the diagnostic criteria of metabolic syndrome. For T2DM to be diagnosed early, insulin resistance must be identified. This purpose, HbA1c can be used since it is cheap and simple (fasting is not required) to evaluate insulin resistance in

obese young people. Therefore, more frequent monitoring of overweight and obese young people to protect their health will make significant contributions to public health.

Disclosures and acknowledgements

The authors would like to sincerely thank all the study participants for their participation. The authors declare no conflicts of interest with respect to the research, authorship, and/or publication of this article. This study was supported by Sakarya University Department of Scientific Research Projects Unit (Project No: 2017-11-00-007). Artificial intelligence (AI) was not used in the creation of the manuscript.

References:

1. Wondmkun YT. Obesity, insulin resistance, and type 2 diabetes: associations and therapeutic implications. *Diabetes, Metabolic Syndrome and Obesity*. 2020; 13: 3611-3616. <https://doi.org/10.2147/DMSO.S275898>
2. Ünlü TN, Deniz D. Obesity policies applied in Turkey; Germany application comparison. *Journal of Selcuk Health*. 2022; 3(1): 62-78.
3. World Health Organization (WHO). Obesity and overweight [Internet]. Geneva: WHO; 2021 [access 2024 Apr 5]. Available from: <https://www.who.int/news-room/fact-sheets/detail/obesity-and-overweight#:~:text=The%20diagnosis%20of%20overweight%20and,help%20the%20diagnosis%20of%20obesity.>

4. Türkiye İstatistik Kurumu (TÜİK). [Women in statistics] [Internet]. Ankara: Türkiye İstatistik Kurumu; 2023 [access 2024 Apr 8]. Available from: <https://data.tuik.gov.tr/Bulten/Index?p=Istatistiklerle-Kadin-2023-53675> (in Turkish).
5. Kılıç AF, Bayrak M, Çadircı K. Body Mass Index and hemoglobin A1c levels in diabetic adults. *Pharmata*. 2023; 3(4): 84-86. <https://doi.org/10.5152/Pharmata.2023.23015>
6. Ghobrial BB, ElKabarity RH, Hassan AMM. Association of glycosylated hemoglobin levels with insulin resistance in adolescent among medical students Ain Shams University. *QJM: An International Journal of Medicine*. 2023; 116(Suppl. 1): hcad069-025. <https://doi.org/10.1093/qjmed/hcad069.025>
7. Keskin M, Kurtoglu S, Kendirci M, Atabek ME, Yazici C. Homeostasis model assessment is more reliable than the fasting glucose/insulin ratio and quantitative insulin sensitivity check index for assessing insulin resistance among obese children and adolescents. *Pediatrics*. 2005; 115(4): 500-503. <https://doi.org/10.1542/peds.2004-1921>
8. Kurtoğlu S, Hatipoğlu N, Mazıcıoğlu M, Kendirci M, Keskin M, Kondolot M. Insulin resistance in obese children and adolescents: HOMA-IR cut-off levels in the prepubertal and pubertal periods. *J Clin Res Pediatr Endocrinol*. 2010; 2(3): 100-106. <https://doi.org/10.4274/jcrpe.v2i3.100>
9. Lee HS, Park HK, Hwang JS. HbA1c and glucose intolerance in obese children and adolescents. *Diabetic Medicine*. 2012; 29(7): 102-105. <https://doi.org/10.1111/j.1464-5491.2012.03596.x>
10. Hardy OT, Czech MP, Corvera S. What causes the insulin resistance underlying obesity?. *Curr Opin Endocrinol Diabetes Obes*. 2012; 19(2): 81. <https://doi.org/10.1097/MED.0b013e3283514e13>

11. Christopher B, Newgard CB. Molecular and metabolic mechanisms of insulin resistance and β -cell failure in type 2 diabetes. *Nat Rev Mol Cell Biol.* 2008; 9: 193-205.
<https://doi.org/10.1038/nrm2327>
12. Sahin NM, Kinik ST, Tekindal MA. OGTT results in obese adolescents with normal HOMA-IR values. *J Pediatr Endocrinol Metab.* 2013; 26: 285-291.
<https://doi.org/10.1515/jpem-2012-0206>
13. American Diabetes Association. Standards of medical care in diabetes-2016 abridged for primary care providers. *Clinical Diabetes.* 2016; 34: 3-21.
<https://doi.org/10.2337/diaclin.34.1.3>
14. Jebeile H, Kelly AS, O'Malley G, Baur LA. Obesity in children and adolescents: epidemiology, causes, assessment, and management. *Lancet Diabetes Endocrinol.* 2022; 10(5): 351-365. [https://doi.org/10.1016/S2213-8587\(22\)00047-X](https://doi.org/10.1016/S2213-8587(22)00047-X)
15. Yilmaz FK, Yazar O. The trends of prevalence of obesity in Turkey from 2008-2016. *Iranian Journal of Diabetes and Obesity.* 2021; 13(2): 65-74. <https://doi.org/10.18502/ijdo.v13i2.6474>
16. Bereket A, Atay Z. Current status of childhood obesity and its associated morbidities in Turkey. *J Clin Res Pediatr Endocrinol.* 2012; 4(1): 1-7. <https://doi.org/10.4274/jcrpe.506>
17. Batur T, Akbay Hİ, Çokluk E, Esendemir A. Evaluation of the hemoglobin A1c test in detecting pediatric prediabetes. *Journal of Contemporary Medicine.* 2023; 13(1): 1-5.
<https://doi.org/10.16899/jcm.1095786>
18. Ameye H, Swinnen, J. Obesity, income and gender: the changing global relationship. *Global Food Security.* 2019; 23: 267-281. <https://doi.org/10.1016/j.gfs.2019.09.003>
19. Kim S, Song K, Lee M, Suh J, Chae HW, Kim HS, et al. Trends in HOMA-IR values among South Korean adolescents from 2007-2010 to 2019-2020: a sex-, age-, and weight

- status-specific analysis. *International Journal of Obesity*. 2023; 47(9): 865-872.
<https://doi.org/10.1038/s41366-023-01340-2>
20. Kalra S, Kapoor N, Kota S, Das S. Person-centred obesity care – techniques, thresholds, tools and targets. *Eur Endocrinol*. 2020; 16: 11-3.
<https://doi.org/10.17925/EE.2020.16.1.11>
21. Kapoor N, Arora S, Kalra S. Gender disparities in people living with obesity – an uncharted territory. *Journal of Mid-Life Health*. 2021; 12(2): 103-107.
https://doi.org/10.4103/jmh.jmh_48_21
22. Wen LM, Merom D, Rissel C, Simpson JM. Weight status, modes of travel to school and screen time: A cross-sectional survey of children aged 10-13 years in Sydney. *Health Promot. J. Austr.* 2010; 21: 57-63. <https://doi.org/10.1071/HE10057>
23. Fulton JE, Shisler JL, Yore MM, Caspersen CJ. Active transportation to school: findings from a national survey. *Res. Q. Exerc. Sport*. 2005; 76: 352-357.
<https://doi.org/10.1080/02701367.2005.10599306>
24. Tudor-Locke C, Ainsworth BE, Popkin BM. Active commuting to school: an overlooked source of childrens' physical activity?. *Sports Med*. 2001; 31: 309-313.
<https://doi.org/10.2165/00007256-200131050-00001>
25. Onal ZE, Atasayan V, Gürbüz T, Hepkaya E, Nuhoglu C. Association of glycosylated hemoglobin (HbA1c) levels with insulin resistance in obese children. *Afr Health Sci*. 2014; 14: 533-538. <https://doi.org/10.4314/ahs.v14i3.6>
26. Fujii C, Sakakibara H. Association between insulin resistance, cardiovascular risk factors and overweight in Japanese schoolchildren. *Obesity Research & Clinical Practice*. 2012; 6: 1-8. <https://doi.org/10.1016/j.orcp.2011.04.002>

27. Kostovski M, Simeonovski V, Mironska K, Tasic V, Gucev Z. Metabolic profiles in obese children and adolescents with insulin resistance. *Open Access Macedonian Journal of Medical Sciences*. 2018; 6(3): 511-518. <https://doi.org/10.3889/oamjms.2018.097>
28. Morrison JA, Glueck CJ, Horn PS, Schreiber GB, Wang P. Pre-teen insulin resistance predicts weight gain, impaired fasting glucose, and type 2 diabetes at age 18-19 y: a 10-y prospective study of black and white girls. *Am J Clin Nutr*. 2008; 88(3): 778-88. <https://doi.org/10.1093/ajcn/88.3.778>
29. Gayoso-Diz P, Otero-González A, Rodríguez-Alvarez MX, Gude F, García F, De Francisco A et al. Insulin resistance (HOMA-IR) cut-off values and the metabolic syndrome in a general adult population: effect of gender and age: EPIRCE cross-sectional study. *BMC Endocr Disord*. 2013; 13: 47. <https://doi.org/10.1186/1472-6823-13-47>
30. Lee CT, Lin CY, Strong C, Lin YF, Chou YY, Tsai MC. Metabolic correlates of health-related quality of life among overweight and obese adolescents. *BMC Pediatrics*. 2018; 18: 1-8. <https://doi.org/10.1186/s12887-018-1044-8>
31. Tornquist D, Tornquist L, Reuter CP, Horta JA, Renner JDP, Burgos MS. Effects of an interdisciplinary intervention on insulin resistance indicators in overweight and obese adolescents. *Journal of Human Growth and Development*. 2020; 30(2): 274-282. <https://doi.org/10.7322/jhgd.v30.10379>
32. Ling JCY, Mohamed MNA, Jalaludin MY, Rampal S, Zaharan NL, Mohamed Z. Determinants of high fasting insulin and insulin resistance among overweight/obese adolescents. *Scientific Reports*. 2016; 6(1): 36270. <https://doi.org/10.1038/srep36270>
33. Jung UJ, Choi MS. Obesity and its metabolic complications: the role of adipokines and the relationship between obesity, inflammation, insulin resistance, dyslipidemia and nonalcoholic fatty liver disease. *International Journal of Molecular Sciences*. 2014; 15(4): 6184-6223. <https://doi.org/10.3390/ijms15046184>

34. Lillioja S, Mott DM, Spraul M, Ferraro R, Foley JE, Ravussin E, et al. Insulin resistance and insulin secretory dysfunction as precursors of non-insulin-dependent diabetes mellitus. *N. Engl. J. Med.* 1993; 329: 1988-1992.
<https://doi.org/10.1056/NEJM199312303292703>
35. Sundell J. Resistance training is an effective tool against metabolic and frailty syndromes. *Advances in Preventive Medicine.* 2011: 984683. <https://doi.org/10.4061/2011/984683>
36. Said MA, Abdelmoneim MA, Alibrahim MS, Kotb AAH. Aerobic training, resistance training, or their combination as a means to fight against excess weight and metabolic syndrome in obese students – which is the most effective modality?. A randomized controlled trial. *Applied Physiology, Nutrition, and Metabolism.* 2021; 46(8): 952-963.
<https://doi.org/10.1139/apnm-2020-0972>