

# Obesity and pregnancy

Cristina Oana Daciana TEODORESCU<sup>a</sup>, Alexandru HERDEA<sup>b</sup>, Adham CHARKAOUI<sup>c</sup>, Andrei TEODORESCU<sup>d</sup>, Andreea-Iuliana MIRON<sup>e</sup>, Amarin Remus POPA<sup>f</sup>

<sup>a</sup>“Alessandrescu-Rusescu” National Institute for Mother and Child Health, Department of Obstetrics and Gynecology, Polizu Hospital, Bucharest, Romania

<sup>b</sup>“Carol Davila” University of Medicine and Pharmacy, Department of Pediatric Orthopedics, “Grigore Alexandrescu” Children’s Emergency Hospital, Bucharest, Romania

<sup>c</sup>Department of Morphological and Functional Sciences, Faculty of Medicine and Pharmacy, “Dunarea de Jos” University of Galati, Romania

<sup>d</sup>University of Oradea, Faculty of Medicine and Pharmacy, Department of Morphological Sciences, Oradea, Romania

<sup>e</sup>“Carol Davila” University of Medicine and Pharmacy; Department of Oncology, “Coltea” Clinic Hospital, Bucharest, Romania

<sup>f</sup>University of Oradea, Faculty of Medicine and Pharmacy, Department of Diabetes, Nutrition, Metabolic and Internal Diseases, Oradea, Romania

## ABSTRACT

**Introduction:** Obesity is the most frequent metabolic disturbance that can target women of reproductive age, among other population groups, and when obese pregnant women become patients, it represents a serious risk factor for both mother and fetus.

**Aim:** The aim of this study is to offer an overview of the effects exerted by this disturbance on pregnancy.

**Material and methods:** The study targets 157 patients admitted to “Alessandrescu-Rusescu” National Institute for Mother and Child Health – Polizu (INSMC), Bucharest, Romania. In order to define the criterion for obesity, WHO classification (body mass index > 30 kg/m<sup>2</sup>) was used. Data was collected retrospectively after acceptance by the Ethics Committee. Also, we gathered anthropometric data (weight, body mass index and analysis regarding the metabolic profile, including total cholesterol, HDL cholesterol, LDL cholesterol, triglyceride, blood sugar, glycosylated haemoglobin) from all subjects. Each analysis was correlated with each patient’s body mass index. Another correlation was made between metabolic profile, antenatal complications and onset of gestational diabetes and premature birth. Statistical analysis was conducted using GraphPad 8 and MedCalc 14.1.

**Results:** Patients had an average body weight of 66.75 kg with a standard deviation of 12.99 kg and a median of 64 kg. Average body mass index was 25.05 kg/m<sup>2</sup>, with a standard deviation of 5.03 kg/m<sup>2</sup> and a median of 24.2 kg/m<sup>2</sup>. There is a directly proportional and statistically significant correlation between the

Address for correspondence:

Andreea Iuliana Miron

Email: [mironandreea01@gmail.com](mailto:mironandreea01@gmail.com)

Article received on the 27<sup>th</sup> of August 2020 and accepted for publication on the 28<sup>th</sup> of September 2020

*values of blood sugar, glycosylated haemoglobin, LDL, TG, uric acid and BMI. Also, there is an inversely proportional and statistically significant correlation between the values of HDL and BMI. The CT/HDL ratio, low HDL level and elevated LDL level are the main risk factors for premature birth, while the CT/HDL ratio, low HDL level and elevated TG are the main risk factors for the onset of gestational diabetes.*

**Conclusion:** *According to the results of this study, the onset of obesity in pregnant woman is rather dependent on each patient's metabolic profile than body weight.*

**Keywords:** obesity, pregnancy, the metabolic syndrome.

## BACKGROUND

Obesity is the most frequent metabolic disturbance which is affecting various groups of people, including female patients of reproductive age. However, when obesity occurs during pregnancy, it becomes a major risk factor for both mother and fetus (1).

Given the alarming growth of obesity, the World Health Organization (WHO) called it one of the most serious global health issues of the XXI<sup>st</sup> century. Obesity has been defined as a condition characterised by the presence of an excessive amount of body fat which threatens patients' health, thus raising morbidity and mortality. According to WHO, the prevalence of obesity in pregnancy is between 1.8 and 25.3%. The etiology is likely multifactorial, the simplest explanation being that obesity is caused by a disturbance in the energy balance of the body, that is, more calories accumulated than spent (2).

Obesity is associated with a wide range of medical complications, including diabetes, cardiovascular disease, dyslipidemia, arterial hypertension, cancer and osteoarthritis. Also, obesity is associated with an increase in patient mortality (3). A body mass index (BMI) above 22.5-25 kg/m<sup>2</sup> is a strong predictor for general patient mortality (4). Obese as well as underweight women are less fertile compared to those with normal body weight. Among obese women, subfertility is due to a reduction in, or absence of, ovulation frequency (5). Also, a predominant cause for oligo- or anovulation is represented by the polycystic ovary syndrome, which is characterized by central visceral obesity in 40% of cases (6). In addition, in the larger context of metabolic disturbances, low levels of HDL cholesterol and elevated levels of triglycerides and LDL chole-

sterol can be found. Metabolic disturbances raise the risk of developing arterial hypertension, coronary artery disease and thrombosis (7). Pregnancy is characterized by a significant elevation of plasma lipid concentrations during gestation. According to the literature, plasma cholesterol and triglyceride concentrations increase by 25-50% and 200-400%, respectively (8).

Metabolic syndrome is defined as an association of signs including abdominal obesity, elevation of serum triglyceride level, lowering of HDL-cholesterol level, arterial hypertension and insulin resistance (9). All the above-mentioned signs are consequences of an unhealthy lifestyle that can also lead to perivisceral accumulation of fat tissue (10). The incidence of metabolic syndrome is perpetually rising; at present, it affects approximately 20% of the world's population. The proportion is increasing with age (12). Although there are no doubts regarding the increased frequency of well defined signs for the metabolic syndrome, its pathogenesis is still controversial (13). Comparative physiology describes metabolic syndrome as being similar to the biological process by which animals store fat in anticipation of calorie deficit periods (14). Obesity in humans is probably a consequence of a genetic predisposition, in conjunction with environmental factors (15).

This study aims to reveal the extent of the problem, offering a general view of the consequences of obesity on pregnancy.

## MATERIAL AND METHOD

We used the definition for obesity corresponding to WHO classification (BMI > 30 kg/m<sup>2</sup>). Also, we considered the onset of obesity before the gestation period.

**Inclusion criteria:** pregnant women aged over 18 who were admitted to INSMC and agreed to be enrolled in the study.

**Exclusion criteria:** patients with obesity caused by endocrine disorders, patients aged 18 and those who did not agree to participate in the study.

Data was collected after obtaining acceptance from the Ethics Committee.

Anthropometric data (body weight, BMI, and routine analysis regarding the evaluation of metabolic profile, including total cholesterol, HDL cholesterol, LDL cholesterol, triglyceride, blood sugar, and glycosylated haemoglobin) were registered for all patients. Each analysis has been correlated with each patient’s BMI. Another correlation was made between metabolic profile, antenatal complications and onset of gestational diabetes and premature birth.

One hundred and fifty seven patients who were admitted to INSMC participated in the present study. Data confidentiality was ensured for each patient. Also, patient enrollment started only after obtaining the Ethics Committee’s approval. Both the principal investigators and the study coordinator attended „Good Clinical Practice” online courses and had a valid diploma for the entire duration of the study.

Statistical analysis was conducted using GraphPad 8 and MedCalc 14.1. In order to compare differences between averages, we used t-student tests for two lots. ANOVA test with Bonfferoni correction was used when we had to analyse three lots or more. We analysed the r correlation coefficient, which can range between -1 and 1. The inversely proportional cor-

relation between the studied parameters can vary between -1 and 0.

Values equal, or close, to 0 show the lack of any correlation, while those between 0 and 1 show a linear directly proportional relationship between variables. For each analysis, we studied the Gaussian data distribution. If this was respected, we used the Pearson coefficient, if not, we used the Spearman coefficient. All analyses with p value < 0.05 were declared valid and statistically significant. These correlations were graphically represented using linear regression analysis methods. For comparing the distribution frequency of different parameters, we used the Chi Square test and where appropriate, hazard ratio (RR) or odds ratio (OR) in order to investigate the chances and risks.

### RESULTS

Patients’ average body weight was 66.75 kg, with a standard deviation (SD) of 12.99 kg and a median of 64 kg (Figure 1).

Average BMI was 25.05 kg/m<sup>2</sup>, with a SD of 5.03 kg/m<sup>2</sup>, and a median of 24.2 kg/m<sup>2</sup> (Figure 2).

The average blood sugar value in the first trimester of pregnancy was 97.04, with a SD of 20.84 and a median of 95 mg/dL (Figure 3).

There was a positive directly proportional correlation between blood sugar and BMI values (Figure 4).

The average glycosylated haemoglobin value in the first trimester of pregnancy was 5.75, with a SD of 0.8 and a median of 5.7 (Figure 5).

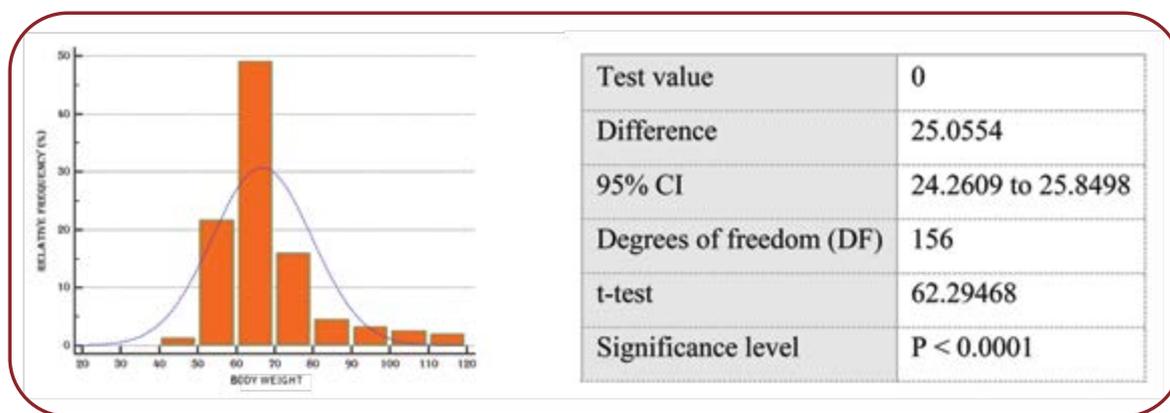


FIGURE 1. Gaussian curve representing the body weight of pregnant women involved in the study

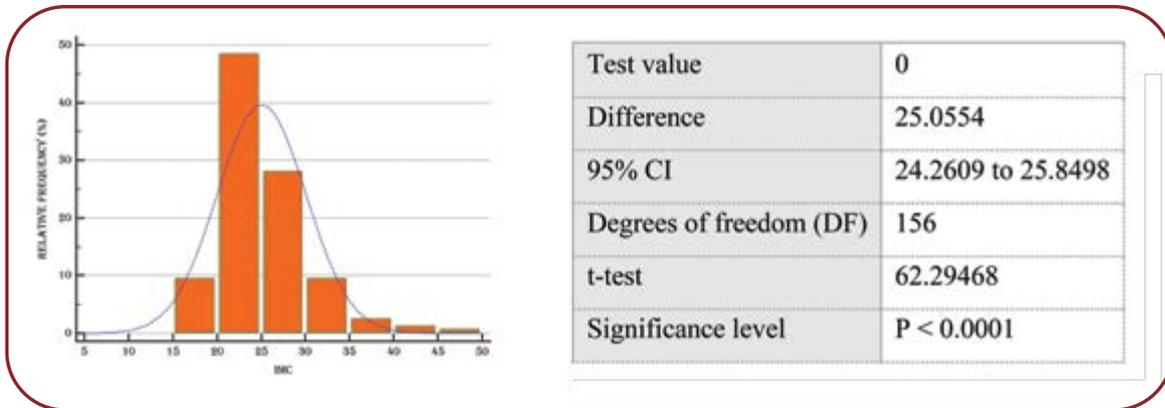


FIGURE 2. Gaussian curve of participants' BMI

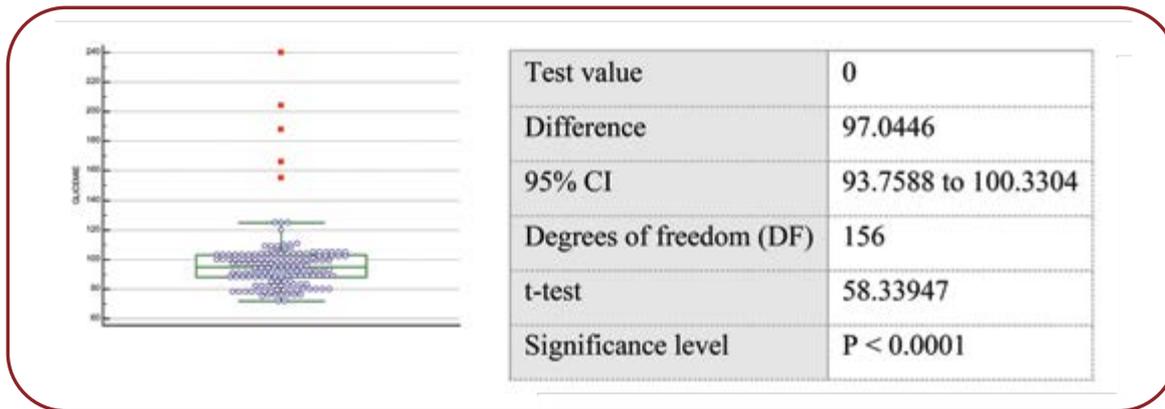


FIGURE 3. Graphical representation of blood sugar values in the first trimester of pregnancy

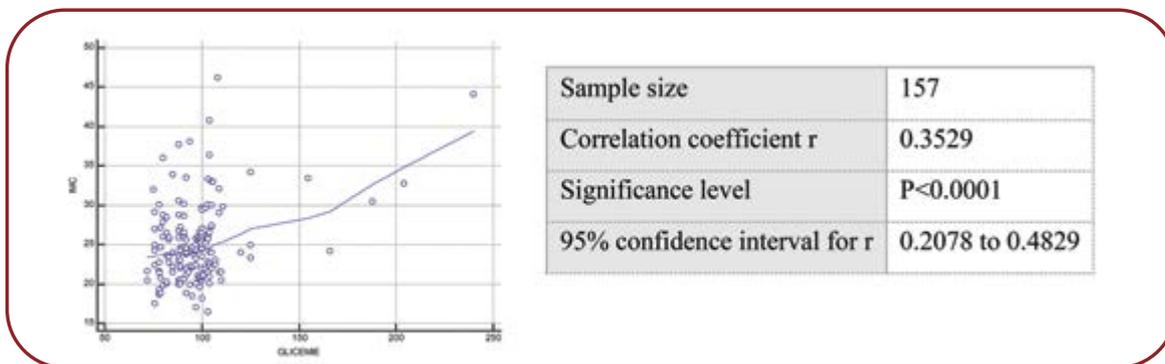


FIGURE 4. Correlation between blood sugar and BMI values

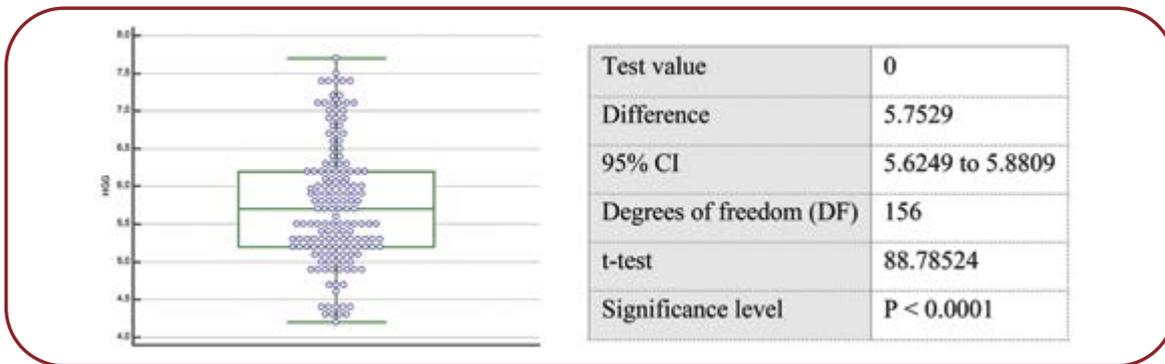


FIGURE 5. Graphical representation of glycosylated haemoglobin value in the first trimester of pregnancy

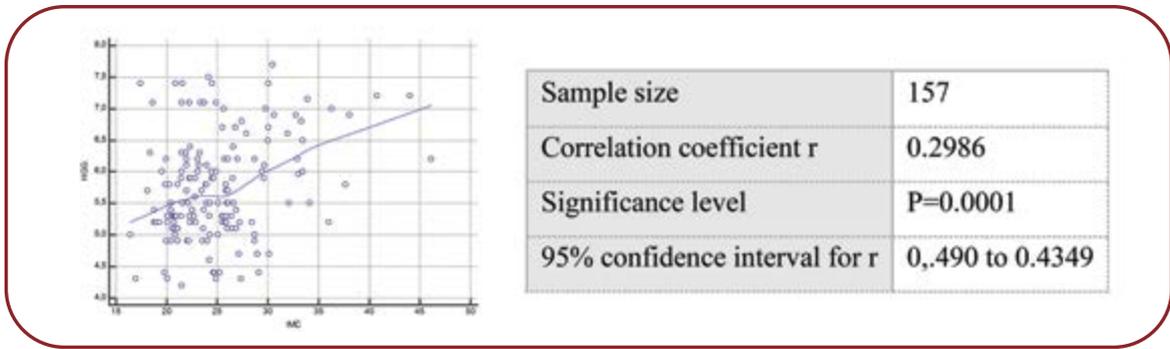


FIGURE 6. Correlation between glycosylated haemoglobin and BMI in the first trimester of pregnancy

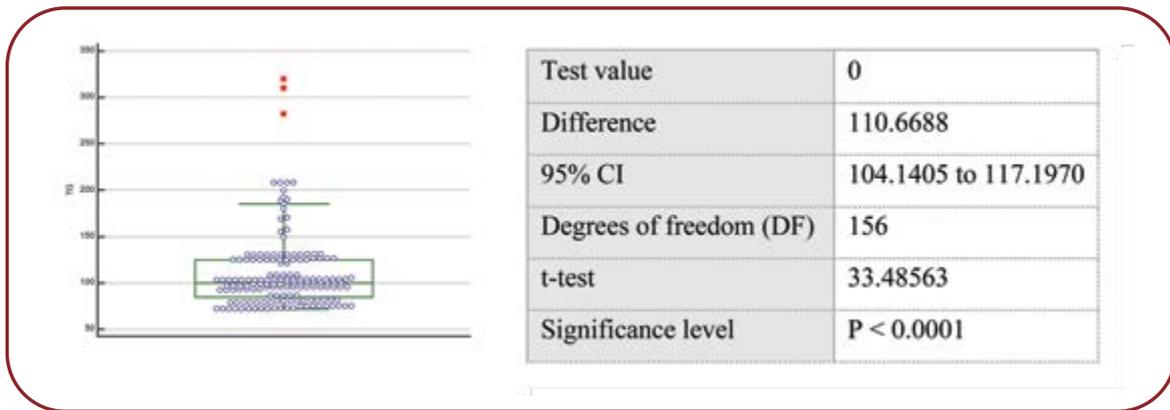


FIGURE 7. Triglyceride values in the first trimester of pregnancy

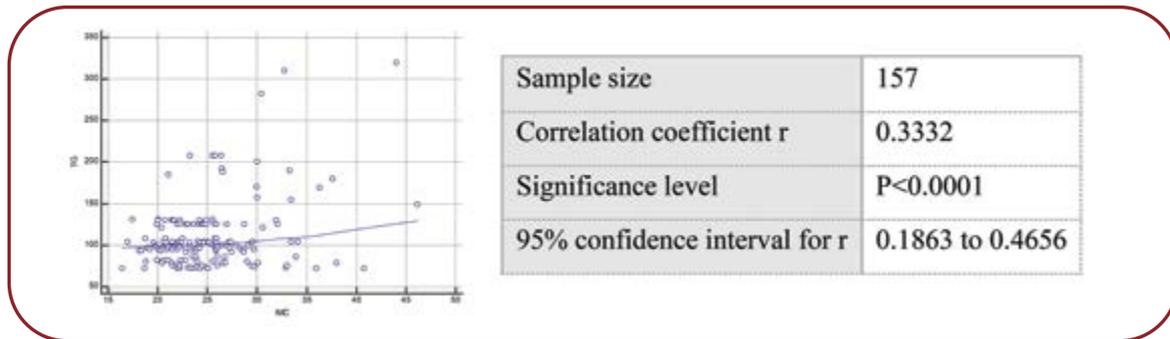


FIGURE 8. Correlation between triglyceride and BMI values in the first trimester of pregnancy

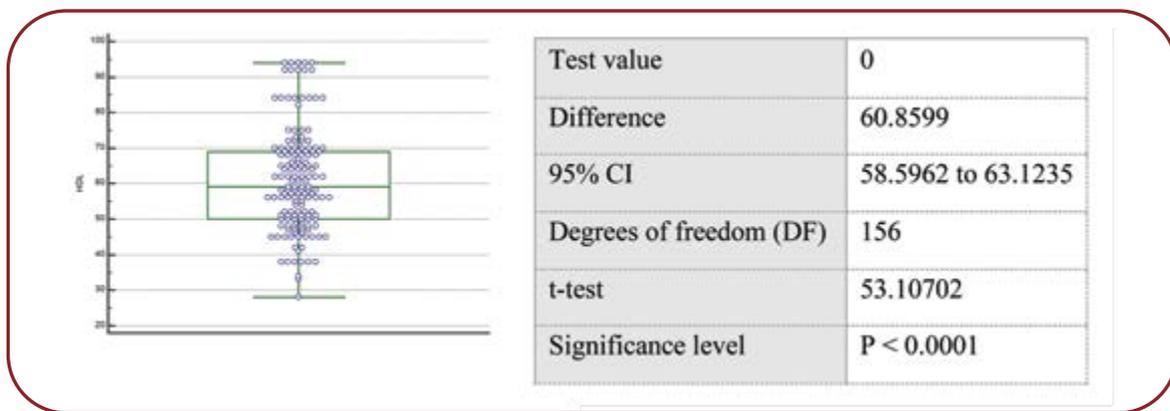


FIGURE 9. The average HDL value in the first trimester of pregnancy

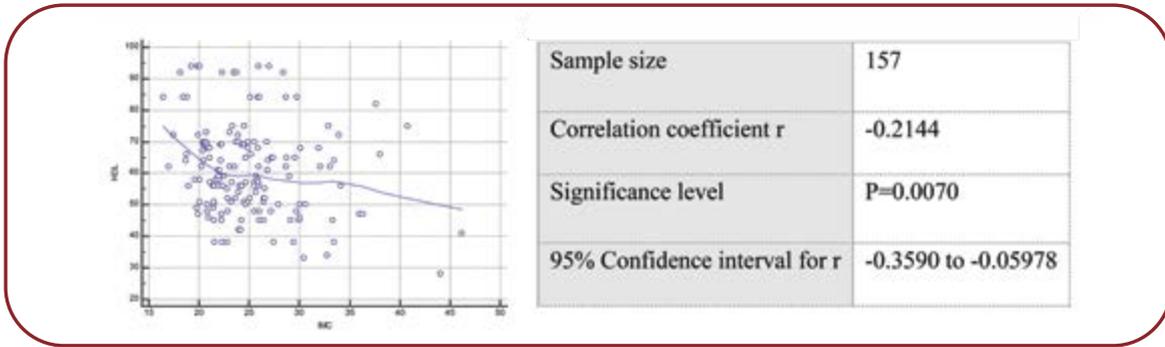


FIGURE 10. Correlation between average HDL and BMI values in the first trimester of pregnancy

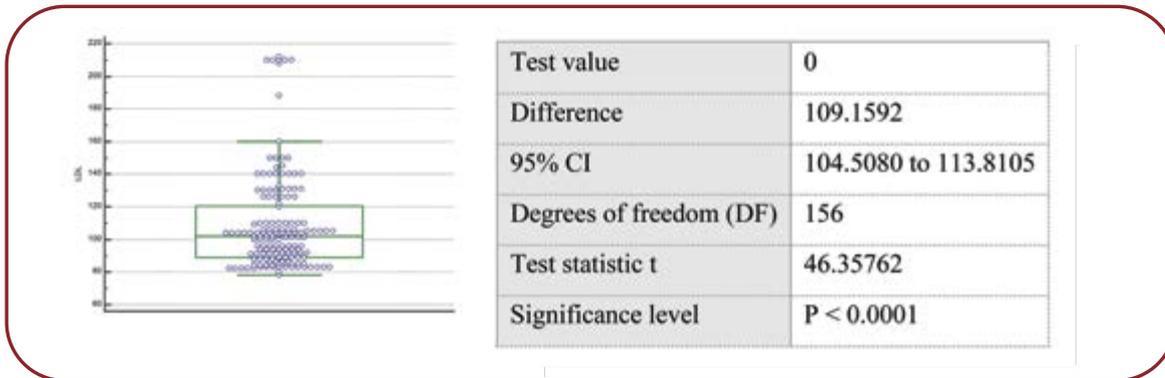


FIGURE 11. The average LDL value in the first trimester of pregnancy

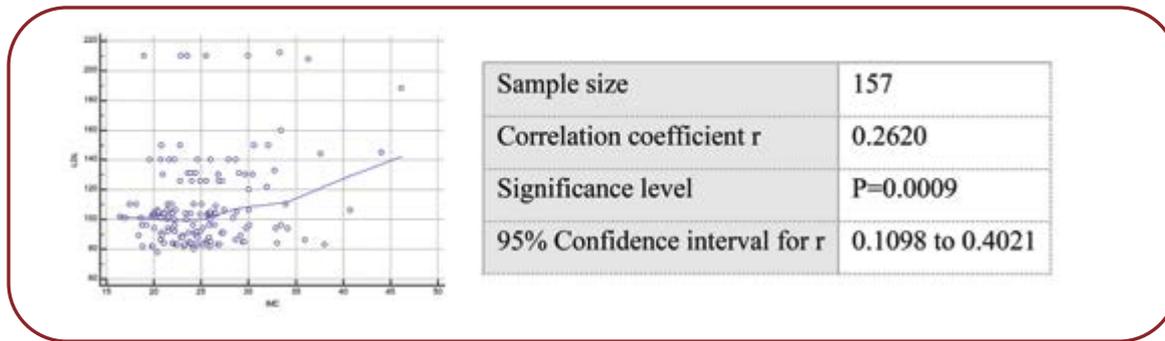


FIGURE 12. Correlation between average LDL and BMI values in the first trimester of pregnancy

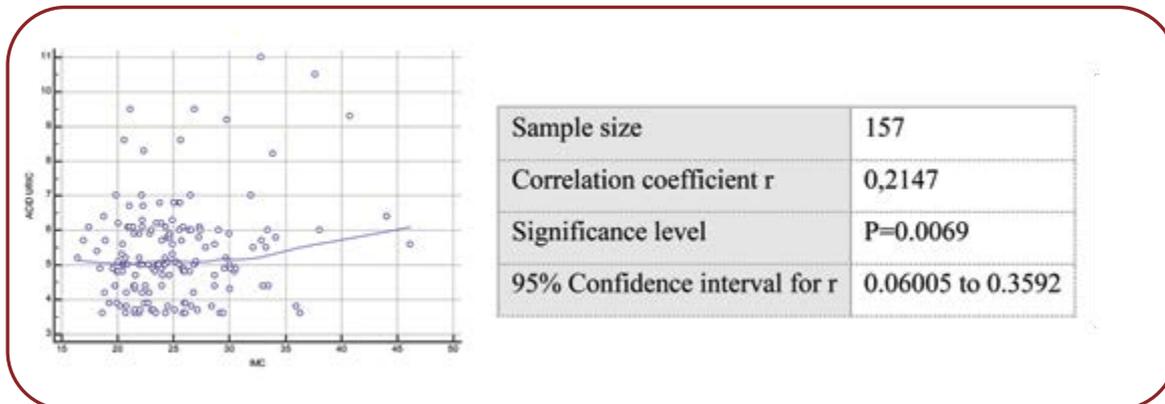
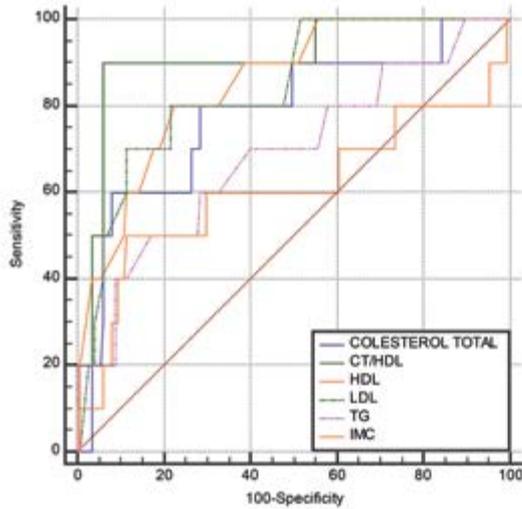


FIGURE 13. Correlation between uric acid and BMI values



Variable 1	Total cholesterol (TC)
Variable 2	TC_HDL TC/HDL
Variable 3	HDL
Variable 4	LDL
Variable 5	TG
Variable 6	IMC
Classification variable	Premature birth Premature birth

Sample size	157
Positive group:	Premature birth = 10
Negative group:	Premature birth = 0

	AUC	SE <sup>a</sup>	95% CI <sup>b</sup>
Total cholesterol (TC)	0.778	0.0821	0.705 to 0.840
TC_HDL	0.910	0.0511	0.854 to 0.950
HDL	0.850	0.0556	0.784 to 0.902
LDL	0.839	0.0591	0.773 to 0.893
TG	0.689	0.0951	0.610 to 0.760
IMC	0.606	0.118	0.525 to 0.683

**FIGURE 14.** Correlation between metabolic profile, antenatal complications and premature birth

There was a directly proportional correlation between the BMI value and the level of glycosylated haemoglobin.

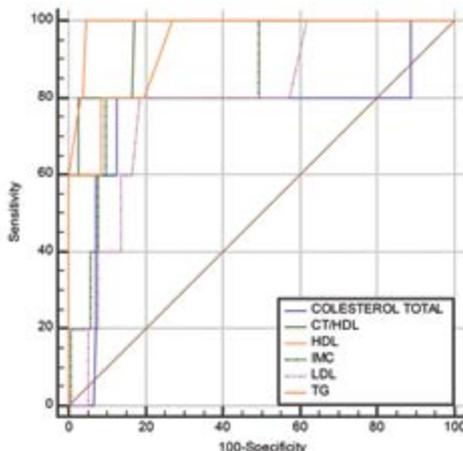
The average triglyceride value in the first trimester was 110.66, with a SD of 41.41 and a median of 100 (Figure 7).

There was a directly proportional and statistically significant correlation between the triglyceride level and the BMI value (Figure 8).

The average HDL value in the first trimester of pregnancy was 60.85, with a SD of 14.35 and a median of 59 (Figure 9).

There was an inversely proportional correlation between HDL and BMI, with P=0.007 (Figure 10).

The average LDL value in the first trimester of pregnancy was 102, with a SD of 29.5 and a median of 102 (Figure 11).



Variable 1	Total cholesterol (TC)		
	TC		
Variable 2	TC_HDL		
	TC/HDL		
Variable 3	HDL		
Variable 4	IMC		
Variable 5	LDL		
Variable 6	TG		
Classification variable	Gestational diabetes		
	Gestational diabetes		
Sample size	157		
Positive group: Gestational diabetes = 1	5		
Negative group: Gestational diabetes = 0	152		
	<b>AUC</b>	<b>SE <sup>a</sup></b>	<b>95% CI <sup>b</sup></b>
Total cholesterol (TC)	0.755	0.145	0.680 to 0.820
TC_HDL	0.961	0.0307	0.918 to 0.985
HDL	0.988	0.00959	0.956 to 0.999
IMC	0.853	0.0808	0.787 to 0.904
LDL	0.792	0.0917	0.720 to 0.853
TG	0.936	0.0427	0.886 to 0.969

**FIGURE 15.** Correlation between metabolic profile, antenatal complications and gestational diabetes

There was a directly proportional correlation between LDL and BMI, with P=0.0009 (Figure 12).

There was a directly proportional and statistically significant correlation between uric acid and BMI values (Figure 13).

## CONCLUSION

There is a directly proportional and statistically significant correlation between the values of blood sugar, glycosylated haemoglobin, LDL, TG, uric acid and BMI. Also, there is an inversely proportional and statistically significant correlation between HDL cholesterol and BMI values. The CT/HDL ratio, low HDL levels and high LDL levels are the main risk factors for premature

birth. The CT/HDL ratio, low HDL levels and high TG levels are the main risk factors for onset of gestational diabetes. In other words, the onset of the pathology in question is rather dependent on a patient's metabolic profile rather than body weight. □

Conflicts of interest: none declared.

Financial support: none declared.

## REFERENCES

1. **Nohr EA, Timpson NJ, Andersen CS, et al.** Severe obesity in young women and reproductive health: the Danish National Birth Cohort. *PLoS One* 2009;4:e8444.
2. **World Health Organization.** Global database on body mass index WHO, Geneva (2009).
3. **Cedergren MI.** Maternal morbid obesity and the risk of adverse pregnancy outcome. *Obstet Gynecol* 2004;103:219-224.
4. **Jensen DM, Damm P, Sorensen B, Molsted-Pedersen L, et al.** Pregnancy outcome and prepregnancy body mass index in 2459 glucose-tolerant danish women. *Am J Obstet Gynecol* 2003;189:239-244.
5. **Catalano M.** Management of obesity in pregnancy. *Obstet Gynecol* 2007;109(2 pt 1):419-433.
6. **Brown K, Apuzzio J, Weiss G.** Maternal obesity and associated reproductive consequences. *Womens Health (Lond Engl)* 2010;6:197-203.
7. **Sewell MF, Huston-Presley L, Super DM, Catalano P.** Increased neonatal fat mass, not lean body mass, is associated with maternal obesity. *Am J Obstet Gynecol* 2006;195:1100-1103.
8. **Usha Kiran TS, Hemmadi S, Bethel J, Evans J.** Outcome of pregnancy in a woman with an increased body mass index. *BJOG* 2005;112:768-772.
9. **Metwally M, Ong KJ, Ledger WL, Li TC.** Does high body mass index increase the risk of miscarriage after spontaneous and assisted conception? A meta-analysis of the evidence. *Fertil Steril* 2008;90:714-726.
10. **Homko CJ, Sivan E, Reece EA, Boden G.** Fuel metabolism during pregnancy. *Semin Reprod Endocrinol* 1999;17:119-125.
11. **Heslehurst N, Simpson H, Ells LJ, et al.** The impact of maternal BMI status on pregnancy outcomes with immediate short-term obstetric resource implications: a meta-analysis. *Obes Rev* 2008;9:635-683.
12. **Asbee SM, Jenkins TR, Butler JR, et al.** Preventing excessive weight gain during pregnancy through dietary and lifestyle counseling: a randomized controlled trial. *Obstet Gynecol* 2009;113(2 Pt 1):305-312.
13. **Claesson IM, Sydsjo G, Brynhildsen J, et al.** Weight gain restriction for obese pregnant women: a case-control intervention study. *Br J Obstet Gynaecol* 2008; 115:44-50.
14. **Kinnunen TI, Pasanen M, Aittasalo M, et al.** Preventing excessive weight gain during pregnancy – a controlled trial in primary health care. *Eur J Clin Nutr* 2007;61:884-891.
15. **Olson CM, Strawderman MS, Reed RG.** Efficacy of an intervention to prevent excessive gestational weight gain. *Am J Obstet Gynecol* 2004;191:530-536.