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## The Effects of Viscosity on Blood Glucose Level of Some People in Mosul City

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Article Information	Abstract				
Article history: Received: Abril21,2024 Reviewer: June,4,2024 Accepted: June,4,2024 Available online	Background: Blood viscosity reflects blood viscosity. It is a measure of the blood's ability to flow through blood vessels. Excessive viscosity is a strong indicator of cardiovascular diseases as it can impede blood flow and/or oxygen supply to body tissues. Objectives: This study was conducted to investigate the effect of viscosity on blood sugar levels.				
<i>Keywords:</i> Glucose, , GLUT, GLUT -2, plasma proteins ,P.C.V, viscosity.	Materials and Methods: Seventy-two random blood samples were collected from healthy individuals without diabetes from qualified medical laboratories in the city of Mosul. The blood samples were divided into three groups based on packed cell volume (PCV) results. The first group had a PCV reading below the normal range. The second				
<i>Correspondence:</i>	group had a PCV reading within the normal range. The third group had a PCV reading above the normal range and included both male and female participants aged between 22 and 60 years. Results: The study results showed that an elevated PCV level in the third group led to a decrease in glucose levels compared to the first and second groups. In males, blood sugar levels in the second group were significantly lower than those in the first group. In females, the second group also showed a significant decrease in glucose levels compared to the first group. Conclusion: Increased viscosity due to proteins affects the function of glucose-carrying proteins and results in lower blood glucose levels.				

# تأثير اللزوجة على مستوى سكر الدم لدى بعض سكان مدينة الموصل خلاصة

الخلفية: لزوجة الدم تعكس لزوجة الدم. وهو مقياس لقدرة الدم على المرور عبر الأوعية الدموية. تعتبر اللزوجة المفرطة مؤشرا قويا لأمراض القلب والأوعية الدموية لأنها يمكن أن تعيق تدفق الدم و/أو إمدادات الأكسجين إلى أنسجة الجسم. الأهداف: أجريت هذه الدراسة لمعرفة تأثير اللزوجة على مستويات السكر في الدم. مؤهلين ومتخصصين في مدينة الموصل. تم تقسيم عينات الدم إلى ثلاث مجموعات على أساس نتائج حجم الخلايا المعبأة مؤهلين ومتخصصين في مدينة الموصل. تم تقسيم عينات الدم إلى ثلاث مجموعات على أساس نتائج حجم الخلايا المعبأة (PCV). المجموعة الأولى التي كانت فيها قراءة PCV أقل من المعدل الطبيعي. المجموعة الثانية حيث كانت قراءة VCV ضمن المعدل الطبيعي. المجموعة الثالثة حيث كانت قراءة VCV أعلى من المعدل الطبيعي وتضم فئتين ذكور وإناث تتراوح أعمارهم بين (٢٢ – ٦٠) سنة. النتائج: أظهرت نتائج الدراسة أن ارتفاع مستوى VCV في المجموعة الثالثة يؤدي إلى انخفاض مستوى الجلوكوز في التتائج: أظهرت نتائج الدراسة أن ارتفاع مستوى VCV في المجموعة الثالثة يؤدي إلى انخفاض مستوى الجلوكوز في التعاطف مع المجموعة الأولى والثالثة. في المجموعة الثالثة يؤدي إلى انخفاض مستوى الجلوكوز في عن المعدل الطبيعي. المجموعة الثالثة حيث كانت متراءة VCV في الدم في المجموعة الثالثة يؤدي إلى انخفاض مستوى الجلوكوز في الاتائج: أظهرت نتائج الدراسة أن ارتفاع مستوى VCV في المجموعة الثالثة يؤدي إلى انخفاض مستوى الجلوكوز في الموطف مع المجموعتين الأولى والثانية. في الذكور، كانت مستويات السكر في الدم في المجموعة الثانية أقل بشكل محوظ مما كانت عليه في المجموعة الأولى. أما المجموعة الثانية من الإناث فقد انخفض مستوى الجلوكوز بفارق معنوي ما محوظ مما كانت عليه في المجموعة الأولى. أما المجموعة الثانية من الإناث فقد انخفض مستوى الجلوكوز بفارق معنوي عن المجموعة الأولى.

#### Introduction

Blood is a red viscid fluid flowing regularly inside blood vessels of the body. It consists of two parts, cellular part (red blood cells, white blood cells and platelets), and fluid part which contain vitamins particles, hormones, electrolytes, as well as proteins which give the blood its viscous property <sup>[1]</sup>. Viscosity is an important property of the fluid that have internal friction for the terraced fluid layers that slide over each other<sup>[2,3,4]</sup>. Viscosity is working independently from fluid velocity, i.e., it doesn't change with the blood velocity, but it can effect on the speed of the flowing blood <sup>[5]</sup>. Plasma proteins can do their effects on augmenting viscosity either directly under physiologic and pathologic conditions and that what is the globulin do, or indirectly, through aggregation of deformed red blood cells, and that is what fibrinogen do <sup>[6,7]</sup>.

Hypoalbuminemia might lead to increase viscosity due to increase in fibrinogen and finally leads to aggregation of disfigured red blood cells <sup>[8]</sup>. After knowing the most important causes of high blood viscosity, we should know the bad consequences of it as a result of increased percentages of plasma proteins and slowdown of blood flow, with possible cardiovascular problems in the form of heart attack, cardiomegaly, blood vessels fibrosis, and impairment of renal function and function of other systems in the body due to ischemic changes <sup>[9]</sup>. The blood has many functions, the most important of which is transportation of oxygen necessary for the cellular respiration, in addition to that, it will deal with metabolic waste products like Co2 and solid products and excrete them through specialized excretory system, transport hormones from endocrine glands to target organs, in addition to that, the blood keeps the body temperature fixed and defends against microorganism. The blood also transports the food particles which has been absorbed in the intestine and delivered to the cells and tissues. The increasing rate of blood viscosity will be affect directly or indirectly on the function of the blood <sup>[10]</sup>, and this may change the normal rate of blood components<sup>[11]</sup>,.

Blood glucose is regarded as the main source of energy in the body and it is the corner stone in most of the metabolic process inside the cells in order to proceed in their life cycle <sup>[12]</sup>. After absorption of glucose by intestinal mucosa, it will enter the blood and then move to the cells through proteins which is present on the cytoplasmic membrane of cells which includes (GLUT 1 - 4). Each type of cells having a specific type of protein present on its cytoplasmic membrane, for example, GLUT -1 is present on the red blood cell membrane <sup>[13]</sup>, GLUT - 2 is present on the cytoplasmic membrane of epithelial cells of intestine ,liver and pancreas <sup>[14]</sup>. GLUT -3 a protein carrier for sugar is present on the cytoplasmic membrane of nervous cells, while the GLUT - 4 protein is present in the fat cells and striated muscles <sup>[15,16]</sup>. All these proteins help glucose to enter the cells where cytoplasmic membrane separates the intracellular fluid from extracellular fluid. The protein on the plasma membrane binds to the glucose and introduces it into the cell by selective method <sup>[17]</sup>. The function of these proteins is affected by osmotic pressure <sup>[18]</sup>. The levels of glucose and lipids also vary with a decrease in the proportion of albumin due to the high proportion of fibrinogen, which increases blood viscosity. When the blood protein is increased, it may increase

the concentration of glucose transporting proteins and thus directly proportional to the process of entering glucose into the cells, so that the concentration of glucose inside the cell become more than outside it <sup>[19,20]</sup>.

The aim of this experimental study is to find out the effects of high blood viscosity (as a result of increased plasma proteins) on blood glucose levels.

## **Materials and Methods**

The samples used in the study included 72 random samples of people without diabetes. Samples were collected from two specialized laboratories for the period from November 5<sup>th</sup>, 2019 to October 20<sup>th</sup>, 2020 in the city of Mosul /Iraq, with ages ranging between (22-60) years, males and females. The samples were kept in special test tubes after taking the sample and under ideal laboratory conditions, using sterilized equipments.

The samples were divided according to the sex into male and females and as follow :

## 1. Male samples:

**Group** (1) included samples of people whose PCV test results were less than normal ranges (38%).

**Group (2)** included samples of people whose PCV test results were within normal ranges (38-44%).

**Group (3)** included samples of people whose PCV test results were above normal ranges (44%).

## 2. Female samples:

**Group** (1) included samples of women whose PCV test results were less than normal ranges

Group (2) included samples of women whose PCV test results were within normal ranges

**Group (3)** included samples of women whose PCV test results were above normal ranges.

## Laboratory tests

1. Hematology tests:

The PCV of all samples was measured using Micro –Hematocrit centrifuge, a German origin apparatus <sup>[21]</sup>.

2. Biochemistry tests:

The glucose level of the samples was measured by the enzymatic method using Spectrophotometer ,721- 2000, a Chinese –origin device at a wavelength 500nm <sup>.</sup>

<u>The statistical analysis</u>: Analysis of variance ANOVA ONE WAY was used and the factors were tested using Duncan, a multi-range test to show the significant differences.

## Results

The results of the analysis of statistical variance for the mean of the three male samples showed a significant difference in the levels of the PCV test and its effect on glucose level , as shown in table -1

parameters	Groups	means	Sig.
	Group 1	38.51	1.000
		(a)	
P.C.V	Group 2	47.53	1.000
		(b)	
	Group 3	53.07	1.000
		(c)	
	Group 1	90.98	.077
		(d)	
	Group 2	90.41	.077
Glucose		(e)	
	Group 3	86.20	.077
		(f)	

**Table -1-** shows the mean differences between the three groups of the variables for males (PCV, glucose).

Similar letters are not significantly different from each other at the 0.05 probability level .

To clarify the difference between the results of the three groups, a letter code was given for each result, and that the similarity in letters means that there is no significant difference between the groups. We note the results of PCV test of the first group (a), the second group (b) and the third group (c), where there was a significant difference at the significant degree of 1.000, while the glucose results for the first group (d) and the second group (e) and the third group (f) at the significant degree .077, it also gave a significant difference between the groups.

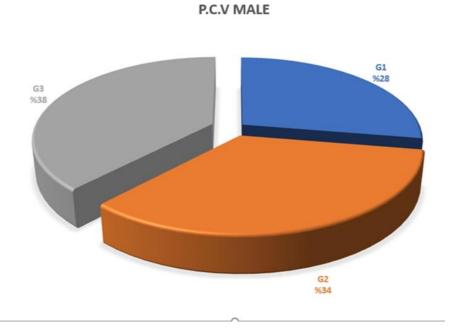
After completing the analysis of variance for the results of (PCV) and glucose for the groups, and from following the tables and figures, it's clear that there is an inverse relation between the levels of PCV and glucose level. When a comparison is made between figure (1) and (2), we note that the first group is the lowest in terms of PCV than the second group and the third group with discrepancy between the means. The first group represented 38.51 while the mean of the second group were 47.53 and the third 53.07. As for the level of glucose for the three groups (first, second and the third), the mean degree of glucose in the first group was 90.98, and for the second and third, they were 90.41 and 86.20 respectively.

groups	means	Sig
Group 1	33.36	1.000
	(a)	
Group 2	40.36	1.000
	(b)	
Group 3	46.75	1.000
Group 1	(c)	
	166.10	.239
	(d)	
	Group 1 Group 2 Group 3	Group 1   33.36     (a)     Group 2   40.36     (b)     Group 3   46.75     (c)     Group 1   166.10

	Group 2	96.70	.239
Glucose		(e)	
	Group 3	90.00	.239
		(f)	

**Table -2-** shows the mean differences between the three groups of variables (PCV, glucose) in female.

Similar letters are not significantly different from each other at the probability level 0.05.



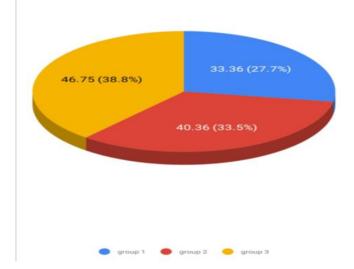
#### Figure (1) shows the comparison between the mean PCV test for males.

Figure (1) represents the significant differences between the three groups of PCV test, and c was the highest compared with b and a.



Figure (2) showing the comparison between the mean glucose test for males

Figure -2- shows that there are differences in blood glucose level for the three groups, as (d) is the highest in glucose level compared with (e) and (f).

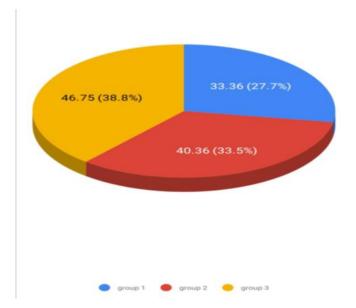


The results of the statistical analysis of the tests of female samples of three groups show significant differences for the PCV test with a significant degree of 1.000 Figure (3), as well as blood glucose test with a significant degree of 239 as in table -2.

The same applies to female samples as well as to the male, as the PCV mean of the first group was 33.36 compared to second group 40.36 and the third 46.75. As for glucose level, the female samples of the first group were the highest with a mean of 166.10, compared with the mean of the second and third groups which were 96.70 and 90.00 respectively.

Fig -3- shows the difference in pcv ratios between the three groups in females, and we notice the difference between group number (1) and the control group (3). While the difference between group (2) and (3) was the smallest ratio

Fig -4- gives the indicators of the difference in the concentration of glucose in the three groups, where the third group was the lowest compared to the first and second groups, and the relationship of the increase in PCV was inverse with concentration of glucose, just as in the male samples. We noticed an intersection between the results of the study with other studies <sup>[10,15,20]</sup>, where these studies showed that high glucose is a major cause of high viscosity and this intersection may be due to several reasons, including the environmental level or nutrition or a difference in some living habits such as daily physical activity.



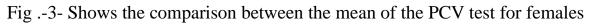


Fig -3- shows the difference between the three groups, as the third group is the highest in PCV compared to the first and second group.

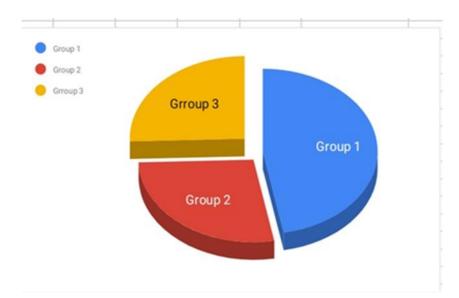


Fig -4- shows the comparison between the mean glucose test for females

#### DISCUSSION

This discrepancy and the inverse relationship between the levels of PCVs and glucose is due to the fact that the viscosity occurred due to increase in blood proteins and that the increase in the blood proteins was the reason for the increase in the level of activity of glucose transporting proteins across the cell membranes **GLUT 1- 4** and this is consistent with the study <sup>[17,22].</sup> in increasing the activity of glucose transporting proteins through the cell membrane. The balanced flow of glucose through the narrow orifice represented by GLUT -2 increases the vestibular tension of the extracellular fluid, which in turn raises the hydrostatic pressure, so that water, glucose and salt enter, and the flowing water prevents the glucose from leaving with ferritin and thus works to restore the activity of vestibular tension again <sup>[23]</sup>.

The inverse relationship between PCV and glucose levels explain that an increase in blood protein increases osmotic pressure and thus an increase in the activity of glucose transporting proteins <sup>[24,25]</sup> and high levels of protein in the blood led to a decrease in glucose levels and this what was stated in the study, although the increase in PCV of the samples was within the normal limits, but with significant differences between groups and the difference in glucose concentration ,which is also within the normal ranges with the differences between groups ,but this decrease in the normal levels of glucose may result in

long term complications for both the circulatory system and the brain ,which affects the individual in doing sports activities or make sleep difficulties <sup>[26,27]</sup>.

## CONCLUSION

The increase in viscosity by the action of proteins (under physiological and biological factors) affects the function of glucose – carrying proteins and decrease in blood glucose, this may result in major health complications.

## ETHICAL DECLARATIONS

#### Acknowledgements

### **Ethics Approval and Consent to Participate**

Before embarking on the research, the approval of the College Council was obtained under approval reference no. 103065, and then it was presented to the Scientific Committee, Finally we have got approval.

I agree with the above statements and declare that this submission as outlined in the Guide for Authors and in the Ethical Statement.

All blood samples were drawn from humans. We obtained their consent to conduct this research.

## Availability of Material

1) This material is the authors' own original work, which has not been previously published elsewhere.

2) The paper is not currently being considered for publication elsewhere.

3) The paper reflects the authors' own research and analysis in a truthful and complete manner.

4) The paper properly credits the meaningful contributions of co-authors and coresearchers.

5) All sources used are properly disclosed (correct citation). Literally copying of text must be indicated as such by using quotation marks and giving proper reference.

#### **Authors' Contributions**

All authors have been personally and actively involved in substantial work leading to the paper, and will take public responsibility for its content.

#### REFERENCES

- 1- Archana M, and Archana N (2016). Glucose transporters physiological and pathological roles. 8(1): 5–9.
- 2- AL Rawi gh (1980). Design and analysis of agricultural experiments. The National Library in Baghdad.
- 3- Roland G (2014). Cellular Regulation of Glucose Uptake by Glucose Transporter GLUT4. Advances in Clinical Chemistry, Chapter Six - 66: 173-240.
- 4- Erika D, and Francisco B (2019). Association between Blood Viscosity and Cardiovascular Risk Factors in Patients with Arterial Hypertension in a High Altitude Setting. Cureus: 11(1):122 -125.
- 5- Rebecca A (2017). Cell Glucose Transport and Glucose Handling During Fetal and Neonatal Development. Fetal and Neonatal Physiology. 428-435.
- **6-** Eunseop Y, and Yang J (2014). Changes in velocity profile according to blood viscosity in a microchannel. Bio microfluidics, 8(3).
- 7- Guandi H (2021). Specifically Targeted Transport of Plasma Membrane Transporters: From Potential Mechanisms for Regulating Cell Health or Disease to Applications. MDPI journals, 11(10), 736.
- 8- Jaap A (2022). Albumin is an interface between blood plasma and cell membrane, and not just a sponge. Clinical Kidney Journal, 15(4).
- Gregory D, and Quirijn D (2020). The Role of Blood Viscosity in Infectious Diseases. Cureus, 12(2).
- **10-** Rajiv A (2001). Add- on angiotensin receptor blocked. Kidney I international,59.

- 11- Tiannan W, and Jing W (2020). Current understanding of glucose transporter 4 expression and functional mechanisms. World J Biol Chem, 11(3): 76–98.
- 12- Kay B, and Jean C (2002). Activation of GLUT1 by metabolic and osmotic stress: potential involvement of AMP-activated protein kinase (AMPK). National Center for Biotechnology Information, (11):115.
- **13-** Mingjun C, and Jing G (2017). Composition and Function of Cell Membranes. Membrane Biophysics, 1–20.
- 14- Richard. J (2008). Osmotic Water Transport with Glucose in GLUT2 and SGLT. Biophys J, 94(10): 3912–3923.
- **15-** Kwaan C (2010). Role of plasma proteins in whole blood viscosity. Clinical Hem rheology and Microcirculation ,44(3):167-76.
- **16-** Rustad H (2010). Correction for trapped plasma in micro Hematocrit Determinations, 2282-2289.
- **17-** Staff Chemists (2012). Synthesize Artificial cell membrane. Science Daily. University of California San Diego.
- **18-** John E (2012). Introduction to Physiology: The Human Body. Textbook Of Medical Physiology. 14-15.
- **19-** Nader M (2013). Whey Protein Hydro Lysate Increases Translocation of GLUT-4 to the Plasma Membrane Independent of Insulin in Westar Rats. PLoS ONE, 8 (8).
- **20-** Teresa E, and Agostino G (2013). Blood viscosity in subjects with normoglycemia and prediabetes. Diabetes Care Publish Ahead of Print, published online September :2-15.
- **21-** Ivana V (2014). Glucose Transporters in the Mammalia blood Article in periodical biologorem.
- 22- Philipp S (2015). Renate S. The distribution of whole blood viscosity, its determinants and relationship with arte. Cardiology and Cardiovascular Medicine.

- **23-** John E (2016). Guyton hall textbook of medical physiology. Scandinavian journal of clinical and laboratory Investigation :677-679.
- **24-** Navale M , Para jape N (2016). Glucose transporters: physiological and pathological roles .NCBI, 8(1): 5–9, 2016.
- 25- Sanjan K (2016). Hypoglycemia. the neglected complication.
- 26- Mazhar M (2018). Hyperglycemia associated blood viscosity1can be nexus stimuli. Clinical Hem rheology and Microcirculation, 71 (4):1-10.Klabunde E (2021). Viscosity of blood. Cardiovascular physiology concepts.

٤- إريكا د، وفرانسيسكو ب (٢٠١٩). العلاقة بين لزوجة الدم وعوامل الخطر القلبية الوعائية لدى
المرضى الذين يعانون من ارتفاع ضغط الدم الشرياني في الأماكن المرتفعة. كيوريوس: ١١(١):١٢٢ ١٢٥.

٥- ريبيكا أ (٢٠١٧). نقل الجلوكوز في الخلايا والتعامل مع الجلوكوز أثناء نمو الجنين وحديثي الولادة.
فسيولوجيا الجنين وحديثي الولادة. ٤٢٨-٤٣٥.

٦- إيونسوب واي، ويانغ جي (٢٠١٤). التغييرات في ملف تعريف السرعة وفقًا لزوجة الدم في القناة الدقيقة. الموائع الدقيقة الحيوية, ٨(٣).

٧- غواندي ح (٢٠٢١). النقل المستهدف على وجه التحديد لناقلات غشاء البلازما: من الآليات المحتملة لتنظيم صحة الخلية أو المرض إلى التطبيقات. مجلات MDPl، (10)، 736.

٨- جاب أ (٢٠٢٢). الألبومين هو واجهة بين بلازما الدم وغشاء الخلية، وليس مجرد إسفنجة. مجلة الكلى السريرية، ١٥(٤).

٩- غريغوري د، وكويرجن د (٢٠٢٠). دور لزوجة الدم في الأمراض المعدية. قوريوس، ١٢ (٢).

١٠- راجيف أ (٢٠٠١). إضافة إلى حجب مستقبلات الأنجيوتنسين. الكلي أنا الدولية، ٥٩.

١١ - تيانان دبليو، وجينغ دبليو (٢٠٢٠). الفهم الحالي للتعبير عن ناقل الجلوكوز ٤ والآليات الوظيفية.
العالم جي بيول كيم، ١١(٣): ٧٦ - ٩٨.

١٢- كاي بي، وجان سي (٢٠٠٢). تنشيط GLUT1 عن طريق الإجهاد الأيضي والأسموزي: احتمال تورط بروتين كيناز المنشط بـ AMP (AMPK). المركز الوطني لمعلومات التكنولوجيا الحيوية،
١١٥:(١١).

١٣– مينغجون سي، وجينغ جي(٢٠١٧). تكوين ووظيفة أغشية الخلايا. غشاء الفيزياء الحيوية، ١–٢٠. ١٤– ريتشارد. ي (٢٠٠٨). نقل المياه الأسموزي مع الجلوكوز في GLUT2 وSGLT. بيوفيس جي، ١٩٤(١٠): ٣٩٢٣–٣٩٢٣.

١٥- كوان سي (٢٠١٠). دور بروتينات البلازما في لزوجة الدم كله. ريولوجيا الهيم السريرية ودوران
الأوعية الدقيقة ,٤٤(٣):٢٧-٧٦.

١٦ رستاد (٢٠١٠). تصحيح البلازما المحاصرة في تحديدات الهيماتوكريت الدقيقة، ٢٢٨٢-٢٢٨٩.
١٧ - طاقم الكيميائيين (٢٠١٢). توليف غشاء الخلية الاصطناعية. علم يوميا. جامعة كاليفورنيا - سان دييغو.

١٨- جون إي (٢٠١٢). مقدمة في علم وظائف الأعضاء: جسم الإنسان. كتاب مدرسي في علم وظائف الأعضاء الطبية. ١٤-١٥.

١٩- نادر م (٢٠١٣). يزيد بروتين مصل اللبن هيدرو ليسات من انتقال 4-GLUT إلى غشاء البلازما المستقل عن الأنسولين في فئران ويستار . بلوس واحد، ٨ (٨).

٢٠ تيريزا إي، وأجوستينو جي (٢٠١٣). لزوجة الدم في الأشخاص الذين يعانون من ارتفاع السكر في الدم ومرض السكري. نشر رعاية مرضى السكري قبل الطباعة، تم نشره على الإنترنت في الفترة من ٢
إلى ١٥ سبتمبر.

٢١- إيفانا الخامس (٢٠١٤). ناقلات الجلوكوز في دم الثدييات مقالة في علم الأحياء الدوري.

٢٢- فيليب س (٢٠١٥). رينات س. توزيع لزوجة الدم الكاملة ومحدداتها وعلاقتها بالفن. أمراض القلب وطب القلب والأوعية الدموية.

٢٣- جون إي (٢٠١٦). كتاب جويتون هول في علم وظائف الأعضاء الطبي. المجلة الاسكندنافية للتحقيقات السربرية والمخبرية:٦٧٧-٦٧٩.

٢٥ - سانجان ك (٢٠١٦). نقص سكر الدم. التعقيد المهمل.

٢٦- مظهر م (٢٠١٨). يمكن أن يكون ارتفاع السكر في الدم المرتبط بلزوجة الدم من المحفزات الرابطة. ريولوجيا الهيم السريرية ودوران الأوعية الدقيقة، ٧١ (٤): ١-١٠.كلابوندي إي (٢٠٢١). لزوجة الدم. مفاهيم فسيولوجيا القلب والأوعية الدموية.