

International Journal of Biochemistry Research & Review

24(2): 1-8, 2018; Article no.IJBCRR.45107

ISSN: 2231-086X, NLM ID: 101654445

The Association between Micronutrients Levels and Gestational Diabetes: A Cross Sectional Study in Ashanti Region

Daniel Abera Ataanya^{1*} and Christopher Larbie²

¹KNUST Senior High Secondary School, Kumasi-Ashanti Region, Ghana. ²Department of Biochemistry and Biotechnology, Faculty of Biosciences, College of Science, Kwame Nkrumah University of Science and Technology, Kumasi, Ghana.

Authors' contributions

This work was carried out in collaboration between both authors. Authors DAA and CL designed the study and performed the statistical analysis. Author DAA wrote the protocol, the first draft of the manuscript, managed the literature searches and managed the analyses of the study. Author CL supervised the study. Both authors read and approved the final manuscript.

Article Information

DOI: 10.9734/IJBCRR/2018/45107

Editor(s):

(1) Dr. KV Ramanath, Professor, Department of Pharmacy Practice, SAC College of Pharmacy, B.G.Nagar, Karnataka,

Reviewers:

(1) Mahmood Hassan Dalhat, Usmanu Danfodiyo University Sokoto, Nigeria.
(2) Arthur N. Chuemere, University of Port Harcourt, Nigeria.

Complete Peer review History: http://www.sciencedomain.org/review-history/27412

Original Research Article

Received 11 September 2018 Accepted 16 November 2018 Published 26 November 2018

ABSTRACT

Aims: Micronutrients such as chromium, iron, magnesium and zinc can serve as good therapeutic and preventive agents for several diabetes complications including Gestational Diabetes Mellitus (GDM). The aim of this study was to compare the serum levels of chromium, iron, zinc and magnesium in pregnant women with and without GDM and to assess the association between the levels of these minerals and GDM.

Study Design: A cross-sectional study.

Place and Duration of Study: Antenatal clinics in five selected hospitals in Ashanti region of Ghana between August 2016 to August 2017.

Methodology: A total of 50 pregnant women with GDM and 50 controls of the same gestational age participated in the study. A structured questionnaire was used to collect information on demographic and clinical characteristics. Standard oral glucose tolerance test was used to

*Corresponding author: E-mail: aberadaniel@rocketmail.com;

measure glucose profile. Serum levels of chromium, iron, zinc and magnesium were measured using the atomic absorption spectrophotometer (AAS).

Results: The results indicated that 14% of the respondents with GDM had a previous history of gestational diabetes. However, there was no previous history among the control group. Serum zinc was not significantly different in the two groups $(0.052 \pm 0.01 \text{ mg/dl}; 0.093 \pm 0.03 \text{ mg/dl}, p=0.276)$. Iron levels in the GDM group were high $(0.956 \pm 0.35 \text{ mg/dl})$ compared with the control group $(0.635 \pm 0.41 \text{ mg/dl})$. There was no significant difference of the serum magnesium level in the two groups (p=0.967). Chromium was higher in the GDM group (0.051 ± 0.05) than the control group (0.012 ± 0.06) . There was a significant positive correlation between iron (r = 0.303, p = 0.000) and chromium (r = 0.302, p = 0.002) with the markers of GDM.

Conclusion: The study showed that high serum concentration of iron is associated with hyperglycemia. Serum magnesium and zinc levels did not exhibit any significant differences between GDM women and nondiabetic women. It is recommended that micronutrients supplementation during pregnancy needs to be carefully examined and commence only when significant deficiencies are observed.

Keywords: Diabetes; gestational; insulin; iron overload; hypertension.

ABBREVIATIONS

AAS : Atomic Absorption Spectrophotometry

GDM : Gestational Diabetes Mellitus
GTF : Glucose Tolerance Factor
IGT : Impaired Glucose Tolerance
NDSS : National Diabetes Service Scheme
NGDM : Non-Gestational Diabetes Mellitus
OGTT : Oral Glucose Tolerance Test

1. INTRODUCTION

Gestational diabetes mellitus (GDM) is a type of diabetes that manifests during pregnancy and normally disappears after delivery. About 8% of pregnant women between the gestational ages of 24 to 28 weeks develop gestational diabetes [1]. In pregnancy, the placenta (the blood source for the baby) synthesizes useful hormones that helps in the baby's growth and develop. Some of these hormones (estrogen, cortisol and human placental lactogen) block the action of the mother's insulin leading to insulin resistances. During pregnancy, to keep normal blood glucose levels, the mothers pancreatic cells need to produce up to 3 times the normal amount of insulin because of these hormones. If the body is not able to achieve this, it might result in the development of GDM [1]. Micronutrients have shown to be good therapeutic and preventive agents for several diabetes complications including GDM. Abnormalities in the metabolism of magnesium, chromium, zinc and copper have been shown to relate to diabetes [2].

Chromium, a trace element, is a member of the compounds called 'glucose tolerance factor' (GTF) which are required for the appropriate

metabolism of glucose and lipid sensitivity. It increases insulin sensitivity by inducing the binding of insulin to its receptors in cells and also increasing receptor numbers. Chromium enhances an increased beta-cell sensitivity and glucose utilisation. A research conducted by Scott et al. [3], has revealed that chromium concentration in serum of pregnant women were considerably lower than normal.

Iron supplements and elevated iron stores is associated with increasing oxidative stress and gestational diabetes [4]. Consequently, while iron supplementation may lead to improvement in pregnancy outcomes when maternal iron levels are low, it is also possible that supplementation intended to prevent deficiency problems may have consequences when the woman's iron level is not low or she does not have iron deficiency [4]. There are so many molecular mechanisms that seek to explain these effects. These mechanisms are not clear but are said to include modulation of adipokines, oxidative stress and intracellular signal transduction pathways [4].

Zinc as a mineral impact a lot on pancreatic function as well as insulin secretion. It is believed that iron and folic acid supplementation during pregnancy has a negative influence on zinc absorption. Zinc deficiency impact negatively on carbohydrate metabolism. Studies have revealed how zinc deficiency associate with diabetes mellitus, glucose intolerance, cardiovascular disease and insulin resistance. Thus, Zinc supplementation in pregnancy is very important because it has been suggested to reduce the risk of some pregnancy complications [5].

In this current study we investigated the association between levels of micronutrients in serum and gestational diabetes among pregnant women.

2. PARTICIPANTS AND METHODS

2.1 Sample Collection

This cross-sectional case-controlled study was restricted to pregnant women attending antenatal clinic at the St. Michael Hospital, Ejisu Government Hospital, Kumasi South Hospital and the KNUST Hospital all in Ashanti region of Ghana. These pregnant women were within the gestational age of 24 to 28 weeks. Convenient sampling technique was used till the required number was obtained. All those who qualified to participate in the study signed a consent form. Pregnant women who had already been diagnosed with gestational diabetes for the current pregnancy were included.

Oral Glucose Tolerance test was performed for the participants. They were asked to have an unrestricted diet rich in carbohydrate for three days and then fast overnight before tests were conducted. First, they were tested for fasting blood glucose after the 8-10 hours overnight fasting using a glucometer. The participants were made to drink 100 g/ml of glucose solution. They were then tested for glucose levels using glucometer at 1, 2 and 3 hours afterward. The following were considered as abnormal values of glucose according to the American Diabetes Association [6].

- Fasting blood glucose levels ≥ 95 mg/dL (5.33 mmol/L).
- II. 1 hour post-prandial blood glucose level ≥ 180 mg/dL (10 mmol/L).
- III. 2 hour post-prandial blood glucose level ≥ 155 mg/dL (8.6 mmol/L).
- IV. 3 hour post-prandial blood glucose level ≥ 140 mg/dL (7.8 mmol/L).

For the participants who recorded abnormal values of glucose, the test was repeated after one week to rule out false positives. Those who had abnormal OGTT values were chosen for the study as those with GDM. Those that were normal and matched with those with diabetes were chosen as the control group. A total of 50 pregnant women with GDM and 50 controls of the same gestational age were recruited for the study. A structured, pre-tested questionnaire was used to solicited information such as age,

occupation, educational background, tobacco use, alcohol intake, previous history of GDM, family history of diabetes and nutritional supplementation of the participants. Ethical approval for this project was sought from the Committee on Human Research Publication and Ethics of Kwame Nkrumah University of Science and Technology (CHRPE/AP/015/17).

2.2 Micronutrients Analysis

Four (4) ml of blood was taken from each participant in to vacutainer tube by trained hospital phlebotomist. The serum was separated by centrifuging at 3500 g for 10 min and stored at -20°C prior to micronutrient analysis. After the total number of participants was reached, the samples were digested by mixing 0.5 ml of serum with deionised water in a digestion flask. Five milliliters of Nitric-perchloric acid (5 ml, 1:1 v/v) and 5 ml of sulphuric acid were added to each sample in a fume chamber. The samples were then heated on a heat plate at 2000°C±5 for 30 minutes and allowed to cool. Measurement for chromium, iron, magnesium and zinc was usina the atomic absorption spectrophotometer. The sensitivity of methods for measuring the macronutrients was 1 ppm [7].

2.3 Statistical Analysis

The data was analysed using the Statistical package for social science (SPSS) version 20 (IBM, USA). Continuous variables were expressed as means and standard deviation while categorical variables were expressed as percentages. The significance of the differences in the mean values between study group and controls for normally distributed parameters were determined using the independent t-test for continuous variables and Chi-square test for categorical variables at 95% confidence level. Pearson correlation coefficients were calculated to signify the association between different quantitative variables. P-values <0.05 were considered statistically significant.

3. RESULTS AND DISCUSSION

3.1 Results

A total of 100 pregnant women participated in the study of which 50 had GDM as the study group and 50 were without gestational diabetes mellitus (NGDM) as the control group. Table 1 summaries the demography and clinical characteristics of the participants. With the study

group, age, family history of GDM, previous history of diabetes, the use of alcohol and tobacco, systolic and diastolic blood pressures compared to the control group were not significantly different. The mean values for the standard oral glucose tolerance test (OGTT), showed that the control group recorded 8.98 mmol/L, 7.00 mmol/L and 5.02 mmol/L blood glucose levels for the 1, 2 and 3 hours blood glucose measurements respectively while that for the study group was 14.40 mmol/L, 12.51 mmol/L and 10.42 mmol/L for the same measurement (Table 2). The mean waist circumference for women with GDM was 109.0 cm and that for the women without GDM was 97.2 cm. The mean fasting blood glucose level for the women with and without GDM was 7.90 mmol/L and 4.39 mmol/L respectively. Waist circumference and fasting blood glucose levels were significantly different between the women with and without GDM (P=0.001). The urine glucose level of the participant presented in Table 3 showed that the control group had no trace of glucose in their urine. In the GDM group, 30% had no trace while 38% of them have 2+ of urine glucose. About 2%, 18% and 12% had trace, 3+ and 4+ respectively of urine glucose. The urine glucose levels were significantly different in the two groups (p=0.000). The urine protein of the participants is presented in Table 4. The findings revealed that 80% of the test group and the entire control group had no proteins in their urine. It was also shown that 12%, 6% and 2% of the participant with GDM had trace, 2+ and 3+ of urine protein respectively. There was statistically significant difference between the two groups (p=0.001).

Table 4 shows the serum micronutrients including chromium, iron, magnesium and zinc levels in the study group and control group. There was no significant difference in the levels of zinc and magnesium in the two groups (p=0.278; p=0.967). However, there was significant difference in the iron and chromium levels for the two groups (p=0.000; p=0.002). Table 5 shows the correlation of the measured minerals with the markers of GDM. The results show a weak and an inverse relationship between FBG and serum zinc level (r = -0.041, p=0.276). There was a weak and insignificant correlation with zinc. Magnesium also showed a weak and an inverse or negative relationship (r = -043, p= 0.967) with FBG. There was positive correlation between serum chromium and fasting blood glucose (r = 0.302, p= 0.002). Serum iron also correlated positively with fasting blood glucose (r = 0.303, p = 0.002).

3.2 Discussion

Gestational diabetes is a very complex metabolic disorder that can be associated with several factors. The causes of GDM are multifactorial which include environmental factors affecting insulin sensitivity and genetic factors [8].

Table 1. Socio-demographic characteristics of participants

Subjects	GDM group	Normal group	P-value
-	N (%)	N (%)	
Age	30.98 ± 4.89	30.66 ± 5.14	0.682
Family history of GDM			
Yes	7 (14.0)	1 (2.0)	0.059
No	43 (86.0)	49 (98.0)	
Previous history of GDM			
Yes	5 (10.0)	0 (0.0)	0.056
No	45 (90.0)	50(100.0)	
Alcohol use			
Yes	2 (4.0)	5 (10.0)	0.436
No	48 (96.0)	45 (90.0)	
Tobacco use	. ,	. ,	
Yes	0 (0.0)	0 (0.0)	
No	50 (100.0)	50 (100.0)	
SBP	121.3 ±18.6	122.0 ± 15.1	0.823
DBP	68.2±17.7	68.9 ± 13.8	0.816

P-value is significant at p<0.05

Table 2. Markers of GDM in the participants

Classification	Women with GDM n (%)	Women without GDM n (%)	P-value
WC	109.0 ± 6.93	97.18 ± 5.13	0.001
OGTT			
1-hr	14.40 ± 2.26	8.98 ± 0.54	0.001
2-hr	12.51 ± 2.01	7.00 ± 0.67	
3-hr	10.42 ± 1.74	5.02 ± 0.61	
FBG	7.90 ±1.8	4.39 ± 0.41	0.001

Table 3. Clinical characteristics of study participants

Classification	Women with GDM n (%)	Women without GDM n (%)	P-value
Glucose	11 (70)	11 (70)	
Normal	15 (30)	50 (100)	0.001
Trace	1 (2)	0 (0)	
2+	19 (38)	0 (0)	
3+	9 (18)	0 (0)	
4+	6 (12)	0 (0)	
Protein	,	,	
Normal	40 (80)	50 (100)	0.001
Trace	6 (12)	0 (Ô)	
2+	3 (6)	0 (0)	
3+	1 (2)	0 (0)	

Table 4. Serum micronutrients of study the participants

Macronutrients	GDM group	Control group	P-value
Chromium	0.0506	0.0124	0.002
Iron	0.956	0.635	0.001
Magnesium	1.979	1.974	0.967
Zinc	0.0515	0.0930	0.276

Table 5. Pearson correlation of the measured minerals and gestational diabetes

	Zinc	Magnesium	Chromium	Iron
FBS	041	043	.302**	.303**
p-value	0.276	0.967	0.002	0.001

^{**-} correlation is significant at p<0.01 level (2-tailed) control variables: FBS fasting blood sugar, extent of diabetes

The mean age for pregnant women with gestational diabetes was 30.98 years while that of the controls was 30.66 years. These findings were consistent with two previous studies [5,8], which revealed higher mean ages for the people with diabetes compared to the controls. Family and previous history of diabetes was high among the study group compared with the control group (Table 1). Risk factors such as previous macrosomic neonates, family history of GDM and previous medical history of GDM increase the rate of developing diabetes and hence pregnant women with these factors should be screened early for gestational diabetes mellitus [8].

A greater proportion of the study participants (60%) were mildly diabetic while 40% of them had severe diabetes. All the participants used as controls recorded normal blood glucose levels. These findings conform to the report of Anqiang et al. [9] who reported increasing OGTT values in pregnant women with GDM. The difference in the severity of diabetes may be attributed to the different conditions such as age, BMI, previous history of diabetes and many more confronting the individual pregnant women including elevated blood pressure [10].

Chromium is a micronutrient that is required in small quantity in the body and plays a very important role in human nutrition. Regulation of blood glucose levels is among the several importance of chromium in the body. In this study, it was revealed that there was a significant decrease in the levels of chromium in the pregnant women without GDM (p=0.002). This was different from the findings of Ghosh et al. [11], which show that chromium levels in nondiabetic Indians where higher compared to those with diabetes. Scientist believe that chromium is a co-factor that assist insulin to get to the cell membrane and aid in the transport of glucose in to the cell [12]. Long before now, scientist used to believe that to achieve this, the body will have to convert chromium to glucose tolerance factor which is a large complex. However current research has shown that there is no glucose tolerance factor but chromium acts with a protein known as low molecular weight chromium binding substance to aid in the action of insulin. This means that chromium's role is to assist insulin to bind to its receptors and its deficiency in pregnancy can result in abnormal glucose tolerance and insulin resistance leading to GDM. The findings of this current study may indicate that the test group had GDM not because of chromium deficiency but as a result of other factors.

The study revealed that the serum iron level in pregnant women with GDM was significantly higher than the pregnant women without GDM (p=0.001). These findings were in line with the study conducted by Muhammad et al. [13] that showed that serum iron levels were significantly higher in pregnant women with abnormal glucose tolerance when compared with those with normal glucose tolerance. A study by Ford and Cogswell [14], has shown that elevated iron stores are associated with high frequency of diabetes. Lao et al. [15], has shown in an observational study consisting of 762 Chinese women without diabetes and with singleton pregnancies, selected between 28 to 30 weeks, that the group with a higher haemoglobin level had a significantly elevated incidence of gestational diabetes as well as greater iron concentration. These findings confirm the observation of our study of higher iron levels in the women with GDM than the control group. The increase in the iron levels in the study may be attributed to iron supplementation during pregnancy consumption of high iron rich diets. Pearson correlation analysis for serum iron and fasting blood glucose revealed a statistically significant positive relationship between serum iron and FBS (r= 0.303, p<0.001) which is consistent with

the results of Sarker et al. [16]. This positive association shows that iron store may contribute to the development of gestational diabetes mellitus by inducing oxidative stress on the pancreatic cell.

Magnesium is an important metal that is involved in several levels of insulin production, its binding and activity. Magnesium is also a critical cofactor of several enzymes involved carbohydrate metabolism. In this study, there was no significant difference in the serum magnesium levels for the pregnant women with and without diabetes (p=0.967). These findings are similar to the finding of Walter et al. [17], which indicated no significant difference in the plasma magnesium levels of people with diabetes and the control group. Previous studies have shown low levels of magnesium in pregnant women with GDM than control group [18,19,20]. In the current study, there was also a negative correlation between the serum magnesium levels with fasting blood glucose (r=-0.043, p=0.967). Karim et al. [21] and Mishra et al. [22] also found correlation between negative magnesium levels and fasting blood glucose in diabetes. This association indicates the role magnesium plays in the development and progression of diabetes.

This study showed a lower serum zinc levels in the study group compared with the control group. These findings agree with that of Al-Maroof et al. [23] who reported lower levels of serum zinc levels in GDM. The decreasing levels of zinc in diabetes may be associated with hyperzincuria and also lower gestational absorption of zinc. Pearson correlation analysis also revealed a statistically insignificant negative correlation between serum zinc and FBS (r=-0.41, p=0.276).

4. CONCLUSION

Family history of diabetes, previous history of GDM, urine glucose, urine proteins and alcohol usage were high in the study group compared to the control group. These factors are therefore to be considered as risk factors to the development of gestational diabetes mellitus. Relation between micronutrient including magnesium, zinc and gestational diabetes were not significantly different in the two groups except iron and chromium. Iron level was significantly high in the study group and correlate with the development of GDM. Many reports have shown that chromium have beneficial properties for individuals with GDM. However, results from this

study conducted show lack of any favourable impact of chromium on GDM.

CONSENT

All those who qualified to participate in the study signed a consent form.

ETHICAL APPROVAL

Ethical approval for this project was sought from the Committee on Human Research Publication and Ethics of Kwame Nkrumah University of Science and Technology (CHRPE/AP/015/17).

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- National Diabetes Service Scheme. Gestational diabetes, caring for yourself and your baby. Diabetes Australia. 2010; 1:3-25.
- Hamid HG, Parween AI, Nazk AM. Evaluation of serum chromium levels in patients with type 1 and 2 diabetes mellitus and insulin resistance. International Journal of Basic and Applied Science. 2012;12:69-72.
- Woods SE, Ghodsi V, Engel A, Miller J, James S. Serum chromium and gestational diabetes. J Am Board Fam Med. 2008; 21(2):153-7.
- 4. Prasad DKV, Sheela P, Kumar AN, Kumar NL, Deedi MK, Madhulatha D. Iron levels increased in serum from gestational diabetes mellitus mothers in coastal area of Andhra Pradesh. J Diabetes Metab. 2013;4:269.
- El-Said NM, Sadik NA, Mohammed NAG. Magnesium in type 2 diabetes mellitus and its correlation with glycemic control. International Journal of Research in Medical Sciences. 2015;3(8):1958-1963.
- 6. American Diabetes Association. Gestational diabetes mellitus. Diabetes Care. 2004;27:s88-s89.
- Zheng G, Wang L, Guo Z, Sun L, Wang L, Wang C, Zuo Z, Qui H. Association of serum heavy metals and trace element concentrations with reproductive hormone levels and polycystic ovary syndrome in a

- Chinese population. Biol Trace Elem Res. 2015;167(a):1-10.
- Akhlaghi F, Bagheri, SM, Rajabi O. A comparative study of relationship between micronutrients and gestational diabetes. ISRN Obstetrics and Gynecology. 2012; 2012:1-4.
- 9. Yang A, Zhao J, Lu M, Gu Y, Zhu Y, Chen D, Fu J. Expression of Hepcidin and Ferroportin in the Placenta, and Ferritin and Transferrin Receptor 1 Level in maternal and umblical cord blood in pregnant women without gestational diabetes. Int J Environ Res Public Health. 2016;13(8). pii: E766.
- 10. Hedderson MM, Ferrara A. High blood pressure before and during early pregnancy is associated with an increased risk of gestational diabetes. Diabetes Care. 2008;31(12):2362-2367.
- 11. Ghosh D, Bhattacharya B, Mukherjee B, Manna B, Sinha M, Chowdhury J, Chowdhury S. Role of chromium supplementation in Indians with type 2 diabetes mellitus. J Nutr Biochem. 2002; 13(11):690-697.
- 12. Abbas B, Thabaan AJ. Relationship between late pregnancy and serum chromium concentration in patients with diabetes. Al-Nahrain Journal of Science. 2007;10(1):25-29.
- 13. Afkhami-Ardekani M, Rashidi M. Iron status in women with and without gestational diabetes mellitus. J Diabetes Complications. 2009;23(3):194-8.
- Ford ES, Cogswell ME. Diabetes and serum ferritin concentration among U S adults. Diabetes Care. 1999;22(12):1978-83
- Lao TT, Chan LY, Tam FK, Ho FL. Maternal hemoglobin and risk of gestational diabetes mellitus in Chinese women. Obstetrics and Gynecology. 2002; 99:807-812.
- Sarker MR, Jebunnesa F, Khatun T, Helal R, Ali L, Rahim ATM. Role of maternal iron status in the pathogenesis of gestational diabetes mellitus. Bangladesh Medical Journal. 2011;40(3):56-60.
- Walter RM Jr, Uriu-Hare JY, Olin KL, Oster MH, Anawalt BD, Critchfield JW, Keen CL Copper, zinc, magnesium and manganese status and complications of diabetes mellitus. Diabetes Care. 1991;14:1050-1056.

- Diwan A, Pradhan A, Lingojwar D, Krishna K, Singh P, Almelkar S. Serum zinc, chromium and magnesium levels in type 2 diabetes. Int J Diabet Dev Countries. 2006; 26(3):121-124.
- Tripathy S, Sumathi S, Raj G. Minerals nutritional status of type 2 diabetic subjects. Int J Diab Dev Countries. 2004; 24:27-28.
- Sharma A, Dabla S, Agrawal R, Barijatya H, Kothari R, Kochar D. Serum magnesium: an early predictor of course complications of diabetes mellitus. J Indian Med Assoc. 2007;105(1):16-20.
- Karim R, Nargis W, Begum KA, Subhan S, Uddin M. Serum lipid profile, serum

- magnesium and fasting serum glucose in newly diagnosed type 2 diabetic subjects. Bangladesh J Medical Biochem. 2014; 7(1):3-10.
- 22. Mishra S, Padmanaban P, Deepti G, Sarkar G, Sumathi S, Toora, B. Serum magnesium and dyslipidemia in type 2 diabetes mellitus. Biomedical Research. 2012;23(2):290-299.
- 23. Al-Maroof A, Al-Sharbatti S. Serum zinc levels in diabetic patients and effect of zinc supplementation on glycemic control of type 2 diabetes. Saudi Medical J. 2006; 27(3):340-350.

© 2018 Ataanya and Larbie; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:
The peer review history for this paper can be accessed here:
http://www.sciencedomain.org/review-history/27412