

Effects of metformin on middle cerebral artery flow velocity in newly diagnosed type 2 Diabetic patients

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Abstract

Type 2 diabetes mellitus is a major cause of cerebrovascular disease. Risk of ischemic stroke is higher in diabetic patients and it is about 2-6 times that in normal population. Atherosclerotic lesions happen early in the disease and patients with type 2 diabetes mellitus. Transcranial Doppler can be used for examination of cerebral arteries and it can detect early atherosclerotic changes. Metformin an old oral anti-diabetic drug but still cornerstone in the treatment guidelines of type 2 diabetes mellitus. It improves endothelial function with preferable cardiovascular effects. Twenty patients were recruited for the study. Transcranial Doppler examination done for all patients before initiation of treatment. Patients received metformin for twelve weeks and they were re-examined with transcranial Doppler. The data of present study showed that a highly significant increase in diastolic velocity and mean flow velocity after treatment with significant reduction in pulsatility index and resistive index. In conclusion, metformin treatment increases cerebral diastolic flow velocity with reduction in vascular resistance in patients with type 2 diabetes.

Keywords: Metformin; Transcranial Doppler; Middle cerebral artery (MCA); Type 2 diabetes mellitus


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Introduction

Stroke is major global health problem. It is considered the 6th cause of death in United States but may be considered the second cause of death in developing countries [1]. There is little data available about risk of stroke in Iraq and even in Middle East area [2]. A study

about prevalence of silent stroke in Iraqi Kurdish population was found and it was about 19% in normal asymptomatic individuals [3]. However, few and old reports from KSA and other Gulf countries recorded annual incidence from 29 to about 60 per 100000 people [4]. The incidence of stroke is increasing in these countries and the risk is expanding, therefore, urgent strategy for

prevention and better clinical care of stroke patients is needed [2]. Diabetes mellitus is a wellrecognized risk factor for cerebrovascular diseases. The association between stroke risk and diabetic patients is well known and many studies concerned with this concept. Previous studies have been found that the risk of stroke and cerebrovascular complication was greater in diabetic patients than those without the disease. The risk was 3 folds greater in patients with type 2 diabetes mellitus than in non-diabetic population. Furthermore, morbidity and mortality that happen with stroke is greater and more severe in diabetic patients than those who are nondiabetic [5]. Also patients with metabolic syndrome and pre-diabetic have one and half fold increase in the risk of stroke than other population [6]. This is due to multiple risk factors like hyperlipidemia, hypertension and insulin resistance which manifested as glucose intolerance.

Early microangiopathic changes are also attributed to cerebrovascular complication and stroke in diabetic patients as macrovascular complications [7]. Transcranial Doppler is useful tool for detection of cerebral vasculopathy. It can detect systolic and diastolic velocities, resistive index and pulsatility index [8-10]. Diastolic flow in cerebral arteries is important because the perfusion of tissues occurs during diastole, therefore it is important to maintain adequate diastolic flow to keep adequate cerebral perfusion and

overcome any changes in general circulation [11]. Resistive index and pulsatility index represent vascular resistance [12]. Increase in resistive index means decrease in compliance of artery and atherosclerosis [13]. Increase in pulsatility index means reduction in perfusion of cerebral tissue [14-16]. Metformin is the only member of the biguanide group of oral anti-diabetic drugs available for use today. Metformin is considered the drug of choice for the treatment of patients with type 2 diabetes mellitus. American academy of endocrinology and diabetology guidelines in 2016 mentioned the use of metformin in all stages of diabetes and combined it with other agents in high HbA1c [17, 18]. Metformin is regarded as an oral antidiabetic drug because it lowers blood glucose concentrations in T2D without causing overt hypoglycemia.

Metformin is also frequently described as an insulin sensitizer leading to reduction in insulin resistance and significant reduction of plasma fasting insulin level. The improvement in insulin sensitivity by metformin could be ascribed to its positive effects on insulin receptor expression and tyrosine kinase activity. Metformin is the only biguanide that can be used in clinical practice [18]. In addition to anti-hyperglycemic effect of metformin, it also improves endothelial function. In this regard, it does so by several mechanisms; first: it decreases free fatty acids and triglycerides. Moreover, it inhibits hepatic lipogenesis and increase

peripheral fatty acids utilization [19]. Second: metformin mitigates oxidized LDL mediated endothelial injury [20]. Third: it activates SIRT as well as increases its expression in endothelial cells. Previous studies showed that increase SIRT activity and expression is protective against endothelial injury and considered as anti-atherosclerotic factor [21]. Fourth: metformin may increase endothelial NO via AMPK activation pathway. Recent studies demonstrated that metformin increases the bioavailability of nitric oxide and eNOS through activation of AMPK pathway [22]. Furthermore, AMPK pathway activation increases the activity of peroxisome proliferator activated receptor gamma which is a transcription factor increases lipid oxidation and utilization in skeletal muscles as well as has important role in regulation of glucose metabolism [23]. Activation of AMPK pathway also reduces endoplasmic reticulum stress which is associated with endothelial dysfunction and accelerated atherosclerotic changes as well as oxidative stress [24]. Fifth mechanism: metformin reduces oxidative stress [25] and finally metformin significantly reduces inflammatory mediators of atherosclerosis like adhesion molecules such as VCAM and ICAM [23].

In previous clinical trial where patients with polycystic ovarian syndrome used metformin, there was significant reduction in vascular oxidative stress, hyperinsulinemia,

dyslipidemia and endothelial dysfunction [26]. The net result of above review is that metformin improves endothelial injury and may preserve endothelial function. It has been found that short-duration metformin treatment mitigates arterial stiffness and endothelial dysfunction in young women with polycystic ovary disease [27]. In the United Kingdom Prospective Diabetes Study, metformin was unique among other oral hypoglycemic agents in reduction of cardiovascular morbidity and mortality that is associated with diabetes mellitus [28]. Besides that, metformin reduces risk of retinopathy, neuropathy and other microvascular complications [29]. Metformin also prevent cerebrovascular disease and remodeling in diabetic patient and reduces risk of ischemic stroke as well as it may reverse established diabetes-related vascular structural changes [30]. Our study is conducted to confirm the beneficial effect of early treatment of metformin on cerebral hemodynamics in patients with type 2 diabetes mellitus as examined by transcranial Doppler.

Subjects and methods

Twenty patients with newly diagnosed type 2 diabetes mellitus were included in our study. The study started from November 2014 and end in October 2016. A verbal consent was taken from each participant. The study was approved by Kufa Medical College Ethical Committee for clinical trials. The diabetic patients were selected from center of diabetes and endocrinology in

Al Sadr medical City in Najaf City/Iraq. They were diagnosed with diabetes mellitus type 2 according to World Health

Organization definition of diabetes as fasting plasma glucose ≥ 7.0 mmol/L (126 mg/dL), 2-hour post-load plasma glucose ≥ 11.1 mmol/L (200 mg/dL) or HbA1C $\geq 6.5\%$ [31]. Patients were randomly selected by endocrinologist after diagnosis and all of them have newly diagnosed type 2 diabetes. They underwent investigation regarding fasting blood sugar and glycated hemoglobin. Then transcranial Doppler examination and measurements were under taken at TCD department/ Middle Euphrates Neuroscience center/ Al Sadr Medical city/Najaf. Studies were conducted in the morning (9:00 am) after overnight fasting and subjects refrained from products containing caffeine 24-hours before study sessions. The time required for every subject to complete study sessions was approximately 30 minutes. Digital transcranial Doppler with M-mode WAKIe with continuous monitoring and physiological test software (Atysmedical, France) and EZ-Dop DWL, COMPUMEDICS, GmbH, Germany. These instruments are provided with head-band as probe holder for pulse-wave 2 MHz phase array transducer. To study the MCA blood flow velocities, the subject was asked to tilt his head to a side and to breath normal quiet breathing, the MCA was first identified by Doppler probe

through trans-temporal window, usually M1 segment. For identification of middle cerebral artery, spectral wave form window was started at 20mm and then decreased gradually to 12mm to get rid of ultrasound noise and obtain accurate measure. The depth of Doppler beam that was adjusted at 45-55mm, utilizing a trans-temporal window above zygomatic arch in the preauricular area. To obtain MCA spectral wave form, probe is directed backward and slightly upward. The pulse wave Doppler sample volume was adjusted to maximum level in the device and the power 75 MW for identification and localization of middle cerebral artery spectral waveform [32, 33]. To recognize the waveform of middle cerebral artery, the waves are above zero line in spectral wave display screen. After localization of middle cerebral artery waveform and getting the optimal signal of middle cerebral artery. The means of peak systolic velocity, diastolic velocity, resistive index and pulsatility index of 10 cardiac cycles were automatically recorded by specific TCD software of the device. Then, patients were treated with metformin (Glucophage) 1000mg daily for twelve weeks. After twelve weeks, glycated hemoglobin level with transcranial Doppler examination were repeated. Version 20 SPSS statistical program was used for analysis of data. Paired t-test was used for comparison between means of before and after treatment analysis for the same patients.

Results

Mean age of patients was 44.55 ± 1.18 years. Patients consist of seven women with a ratio of 35% and 13 men with a ratio of 65%. Metformin group also showed a significant decrease in FBS from 159.25 ± 6.24 mg/dL to 129.1 ± 5.37 mg/dL after twelve weeks of treatment with metformin 1 g/day ($P < 0.000$). For metformin group HbA1c decreased from $7.76 \pm 0.26\%$ to $6.86 \pm 0.16\%$ ($P < 0.000$). There was no statistical difference in peak systolic velocity between baseline reading (71.35 ± 1.84 cm/sec) and post-treatment one (72.1 ± 1.1 cm/sec) ($P = 0.517$). On the hand, there was statistical significant Effect of metformin on glycemic control and Doppler indices

increase in diastolic velocity and mean flow velocity from (25.35 ± 0.96 cm/sec, 40.68 ± 1.02 cm/sec) to (28.6 ± 0.5 cm/sec, 43.11 ± 0.51 cm/sec) respectively and p values as follow ($P < 0.000$ and 0.001). Metformin group showed significant decrease in

Table 1. pulsatility index and resistive index after treatment. Pulsatility index decreased from 1.14 ± 0.04 to 1.01 ± 0.025 ($P = 0.001$) and resistive index also decreased from 0.64 ± 0.0134 to 0.61 ± 0.009 ($P = 0.007$).

Parameter	Metformin group		
	Mean \pm SEM		
	Before	After	P value
FBS (mg/dL)	159.25 ± 6.24	129.1 ± 5.37	0.000 (H.S)
HbA1c %	7.76 ± 0.26	6.86 ± 0.16	0.000 (H.S)
Peak systolic velocity (cm/sec)	71.35 ± 1.84	72.1 ± 1.1	0.517 (N.S)
Diastolic velocity (cm/sec)	25.35 ± 0.96	28.6 ± 0.5	0.000 (H.S)
Mean velocity (cm/sec)	40.68 ± 1.02	43.11 ± 0.51	0.001 (S)
Pulsatility Index	1.14 ± 0.04	1.01 ± 0.025	0.001 (S)
Resistive Index	0.64 ± 0.0134	0.61 ± 0.009	0.007 (S)

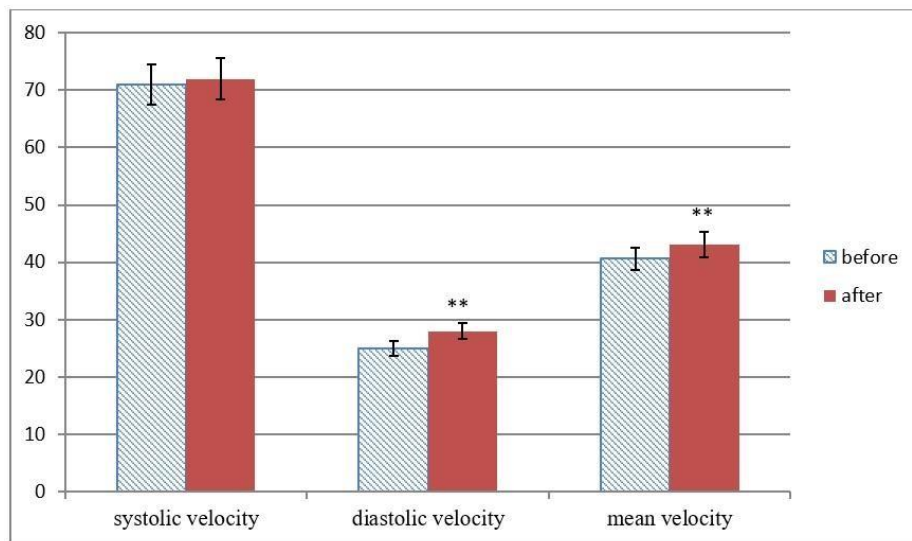


Figure 1.

Bar charts explain the effect of metformin on systolic, diastolic and mean velocities of middle cerebral artery by TCD examination in newly diagnosed patients with type 2 DM. ** means $P < 0.05$.

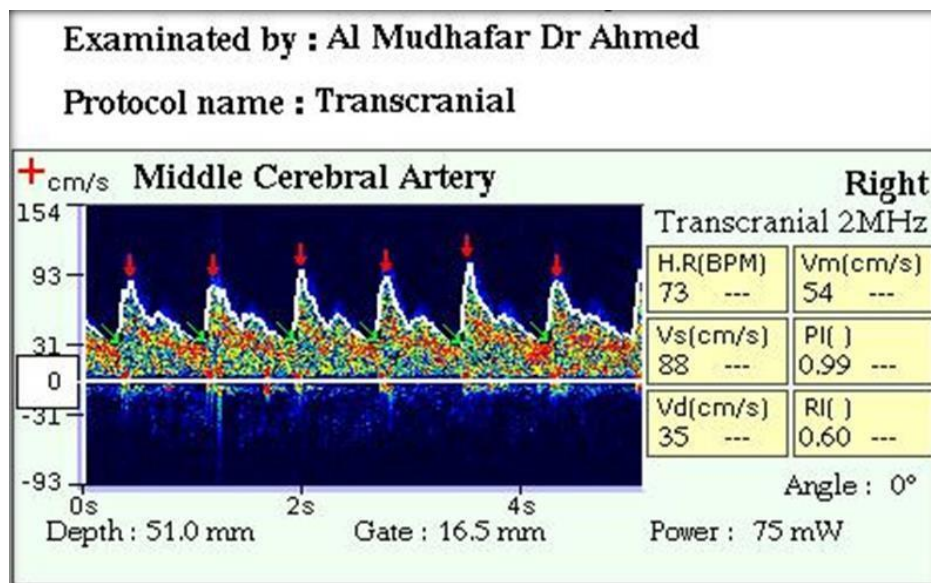


Figure 2.

Spectral wave form of right middle cerebral artery of patient with newly diagnosed type 2 DM before treatment. The yellow arrow referred to diastolic wave.

Examined by : Al Mudhafar Dr Ahmed

Protocol name : Transcranial

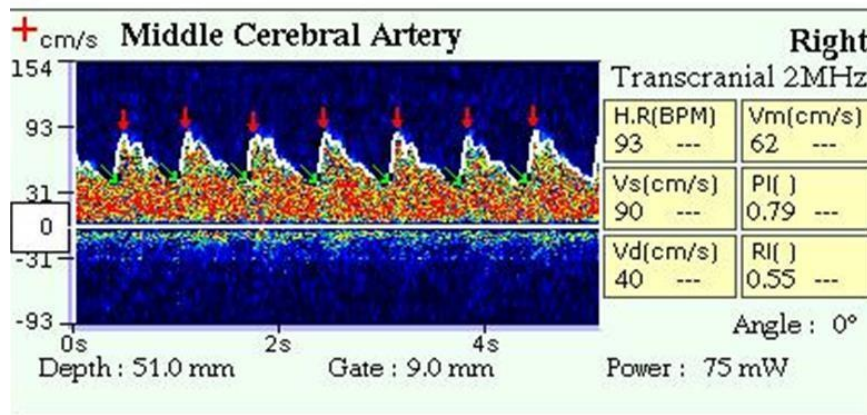


Figure 3.

Spectral waveform of right middle cerebral artery of the same patient above in Figure 2 after treatment with metformin for 12 weeks. The yellow arrow referred to diastolic velocity. There was increase in both systolic and diastolic but the latter is more with decrease in both RI and PI.

Discussion

Nowadays, there is clear and sufficient evidence that type 2 diabetes mellitus is associated with accelerated atherosclerosis, therefore an increased risk of stroke and other vascular disease exists. These vascular complications are associated with high morbidity and mortality. From this point, it is of paramount importance to predict any evidence of vasculopathy and manage it earlier [34].

In our study, metformin exerted significant reduction in blood glucose and HbA1c. It is well known that metformin decreases insulin resistance in all tissues especially the liver [35]. It also decreases hepatic output of glucose

[36]. Moreover, metformin decreases β cells apoptosis and slows diabetes disease progression [37-39]. Many of these effects are mediated through AMP-kinase pathway [22, 23, 30]. Nowadays, metformin use is recommended in the onset of diabetes type 2 [17]. Furthermore, it is highly recommended in pre-diabetic patients and metabolic syndrome [40].

Localized baseline increase in systolic velocity may indicate stenosis of cerebral vessel [41]. In our study, we did not record abnormally high systolic velocity and patients with high systolic velocity that indicate variable degree of stenosis were excluded from our study. On the other hand, diastolic velocity is

important as diastolic flow is vital for brain perfusion as cerebral perfusion occurs during diastole [11, 12]. Patients with impaired diastolic flow have high risk of developing cerebrovascular accident [13, 14].

Therefore, improvement in diastolic velocity may protect the patient from ischemic stroke as it keeps adequate blood flow during diastole. Pulsatility index and resistive index may reflect the compliance of the vessel and therefore, high PI and/or RI may indicate increase vascular resistance and increase in arterial stiffness [15, 16]. Previous studies showed that diabetic patients have higher resistive index than normal population and this may increase micro- and macro-vascular complications [32].

Furthermore, it may predict the risk of cerebrovascular disease. It is clear fact that early diagnosis and treatment of vascular complications in diabetic patients is prompt and it is highly recommended. In our study, metformin not only improve diastolic blood flow velocity but it also lower resistive and pulsatility indices. No previous study was found regarding the effect of metformin on cerebrovascular hemodynamics by transcranial Doppler. This finding further ensures the positive cardiovascular effects of metformin. Early use of metformin with regular exercises and dietary control delays the onset of cardiovascular complications of diabetes [42]. In a large randomized clinical trial, metformin decreases the risk of stroke in diabetic patients [43].

Moreover, chronic metformin treatment significantly reduces infarct size in animal model [44]. The mechanism beyond that is chronic metformin therapy down-regulate AMPK in cerebral vessels and decreases lactate accumulation during ischemia [42-45]. However, acute metformin treatment during ischemic stroke may cause activation of AMPK which has harmful effects on neurons during ischemia with accumulation of lactate [44].

We concluded, that chronic metformin therapy has beneficial effect on cerebrovascular hemodynamics and may lower the risk of stroke in patients with type 2 diabetes mellitus.

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