Introduction

Diabetes mellitus is a chronic endocrine disorder characterized by increased blood glucose levels as a consequence of impaired insulin function or secretion. It should be noted that the prevalence of diabetes is increasing worldwide. In addition to hyperglycemia, hypercholesterolemia is also attributed to diabetes. In this respect, atherosclerosis after dyslipidemia is the most prevalent complication of diabetes mellitus. The association between diabetes and age-related dementias like Alzheimer’s disease has been a matter of debate. In this regard, hypercholesterolemia caused by diabetes leads to elevated brain amyloid-beta levels as well as severe learning and memory impairments. So, it is said that type II diabetes is a risk factor for vascular dementia. Since increased cholesterol synthesis concomitant with impaired cholesterol absorption is reported in diabetes, alteration of cholesterol homeostasis in brain can help understand its dyslipidemia mechanism in diabetes. Furthermore, over 25% of the body’s cholesterol pool is located in the Central Nervous System (CNS) and brain cells are segregated from lipoproteins in the systemic circulation by the blood-brain barrier (BBB).
brain barrier. So, cholesterol homeostasis in the brain is unique and dependent on its own de novo synthesis and degradation with a specific extracellular lipid transport system. It is of note that brain cholesterol metabolism is also involved in several physiological function in the CNS including myelin production. Moreover, SREBP-Cleavage Activating Protein (SCAP), which is a chaperone of Sterol Regulatory Element-Binding Protein 2 (SREBP2), inhibits the overload of intracellular cholesterol whenever cells preserve adequate cholesterol. It has been proved that the upregulation of SREBP-2 expression as a result of SCAP over-expression elevates cholesterol uptake, which results in cholesterol accumulation in diabetic rats. So, regarding previous findings about memory loss of hypercholesterolemia caused by diabetes, it should be elucidated that diabetes complications affect other cholesterol metabolism-related gene in the brain and also lead to a more accurate understanding of alterations of cholesterol metabolism in diabetes. It has been shown that the variation of both plasma and B-cell islet cholesterol homeostasis may correlate to the pathogenesis of diabetes. Lack of ATP binding cassette subfamily A member 1 (ABCA1), which is one of the main players of cholesterol metabolism modulating cholesterol efflux, leads to overload of cholesterol in B-cell islet and impaired insulin release.

There is increasing interest in the application of herbal medicines against human disorders worldwide. *Zingiber officinale* (Z. officinale), also known as ginger, is amongst important herbal medicines possessing anti-diabetic properties, the anti-hyperglycemic effects of which have been indicated in several studies. Besides *in vivo* studies, findings from clinical trials showed the protective effect of ginger extract in the reduction of blood glucose levels. However, the effect of hydroalcoholic extract of ginger on the alteration of some enzymes involved in brain cholesterol homeostasis, including 3-hydroxy-3-methylglutaryl-coenzyme A reductase (HM G-CoA reductase) and cytochrome P450 family 46 subfamily A member 1 (CYP46A1), is poorly understood in diabetes.

It is of interest to check the effect of ginger extract on brain cholesterol homeostasis in a streptozotocin-induced diabetic rat model given the increased levels of major components of lipid profile in diabetes mellitus.

**Materials and Methods**

**Materials**

The dried root of ginger (*Z. officinale*) was purchased from Goldaru Isfahan Pharmaceutical Company, Isfahan, Iran. Mouse anti-β-actin antibody was obtained from Sigma Aldrich, PVDF was obtained from Roche Applied Science, and mouse anti-CYP 46A1 was purchased from Santa Cruz. Mouse Anti-HMG-CoA reductase was kindly donated by Professor TY Chang, Geisel School of Medicine at Dartmouth, Hanover, NH, USA. All other chemicals were purchased from Merck chemicals, Germany.

**Animal model**

40 male Wistar rats weighting 200-250 gr (8-10 weeks old), which were obtained from research center and experimental animal house of Jundishapur University of medical sciences, Ahvaz, Iran were selected for experimental tests in this study. Before all analytical tests, animals were acclimatized under standard ad libitum conditions for 3 days.

**Extract preparation**

The dried roots of ginger (*Z. officinale*) were purchased from Goldaru Herbal Pharmaceutical Company, Isfahan, Iran. The voucher samples were pre-served for reference in the Biochemistry Department of Medical School, Ahvaz Jundishapur University of Medical Sciences. Ginger extraction was done by filtration. Briefly, 200 gr of ginger roots was crushed with a blender and then soaked in 1400 ml of 70% methyl alcohol for 3 days. After filtration of homogenized mixture using Whatman filter paper No.40, the filtrate was placed under vacuum at 50°C to evaporate methanol. Finally, 25 gr crystallized extract was obtained.

**Induction of diabetes by streptozotocin**

Induction of diabetes was performed by intravenous administration of 40 mg/kg streptozotocin (STZ) dissolved in cold 0.1 M citrate buffer (pH=4.5). After 3 days of STZ administration, tail vein blood was taken to measure fasting blood glucose with a glucometer. Diabetes was verified according to plasma glucose concentrations higher than 350 mg/dl.

**Experimental animal groups**

40 rats were randomly divided into 4 groups. Group 1 consisted of 10 rats as non-diabetic control, and each rat received 1.5 ml/kg distilled water daily by gavage. Also, two weeks after induction of diabetes, diabetic rats were randomly divided into 3 experimental animal groups (10 rats each) as follows: Group 2 as non-treated diabetic group in which each rat received 1.5 ml/kg distilled water daily by gavage, Group 3 as the diabetic group that received 200 mg/kg hydroalcoholic extract of ginger dissolved in 1.5 ml/kg distilled water daily by gavage, and Group 4 as the diabetic group that received 400 mg/kg hydroalcoholic extract of ginger dissolved in 1.5 ml/kg distilled water daily by gavage. The treatment lasted for eight weeks, and all experiments were done after two weeks of STZ administration.

**Tissue preparation and western blotting**

Mice from experimental groups were anesthetized and their brain homogenates were prepared as follows. After dissection, the brain was washed with PBS and homogenized in ice-cold RIPA buffer with protease inhibitor cocktail using sonication. An equal amount of proteins in brain homogenates was subjected to 8% sodium dodecylsulfate-polyacrylamide gel electropho
resis (SDS-PAGE) and transferred to a polyvinylidene fluoride membrane (Roche). After blocking with blocking buffer containing 5% skim milk in TBST, the membrane was incubated overnight with specific primary antibodies for HMG-CoA reductase (1:5000 dilution, Abcam), CYP46A1 (1:500 dilution, Santacruz), and B-actin (1:5000 dilution, Sigma). After washing the membrane with TBST, it was incubated for 1.5 hr with specific goat anti-mouse (1:4000 dilution, Sigma) secondary antibodies for HMG-CoA reductase, CYP46A1, and B-actin. After chemiluminescence reaction, the bands were visualized using ChemiDoc™ (Bio-Rad).

Statistical analysis
Statistical analysis was performed with SPSS (Version 18) Software. Descriptive statistics presented data as mean±SD. Analysis of Variance (ANOVA) was used to check significant differences between groups in western blotting analysis. To quantify the difference between protein levels, the data were analyzed using ImageJ software. The quantification reflects the relative amounts as a ratio of each protein band relative to the lane’s loading B-actin as control. For all statistical analysis, p<0.05 was considered as the significance level.

Results
Effect of hydroalcoholic Z. officinale extract on HMG-CoA reductase levels in brain
Brain homogenate was prepared using sonication in ice-cold RIPA buffer with protease inhibitor cocktail. The effect of ginger extract on the protein levels of HMG-CoA reductase in brain homogenate was analyzed by SDS-PAGE and western blotting. Although the level of HMG-CoA reductase in diabetic rats treated with 200 mg/kg concentration of ginger extract was approximately equal to the control group, treatment of streptozotocin-induced diabetic rats with hydroalcoholic extract of ginger for eight weeks caused reduced HMG-CoA reductase compared to diabetic group in the brain. This effect was more pronounced with 400 mg/kg of ginger extract (Figure 1), indicating the effect of ginger on rate-controlling enzyme of mevalonate pathway, which is the metabolic pathway producing cholesterol and other isoprenoids.

Effect of hydroalcoholic Z. officinale extract on brain CYP46A1 levels
To check the effect of ginger on brain CYP46A1 content, the brain homogenate was subjected to SDS-PAGE and western blotting. CYP46A1 level in diabetic non-treated rats decreased compared to control group in the brain but the influence of ginger treatment increased CYP46A1 levels compared to diabetic rats since it reached to control level with 400 mg/kg concentration of ginger extract. Among the treated diabetic groups, increased level of CYP46A1 in diabetic group receiving 200 mg/kg of ginger was significant in comparison to the control group and level of CYP46A1 in diabetic ones receiving 400 mg/kg of ginger was almost as normal as the control group.

Discussion
In this study, the effect of hydroalcoholic Z. officinale extract was investigated on brain HMG-CoA reductase and CYP46A1 levels in streptozotocin-induced diabetic rats. Although the changes in levels of enzymes involved in lipid metabolism due to diabetes have been a matter of debate, our data demonstrated the up-regulation of HMG-CoA reductase in diabetes, and the use of ginger extract caused a modulation in HMG-CoA reductase level and could return it back to the basal level of control group. Although CYP46A1 protein level decreased in diabetes, its level is increased by ginger extract treatment; with 400 mg/kg of ginger extract, it reached to control group’s level, which could reduce the cholesterol accumulation and it has beneficial effects on cholesterol hemostasis in the brain.

Dyslipidemia is among the most important complications of diabetes, which predisposes people to cardiovascular diseases. A significant rise of triglyceride and cholesterol along with decrease of HDL cholesterol can be attributed to endogenous alterations of lipid homeostasis caused by diabetes. Since the neural cells in CNS are segregated from lipoproteins in the systemic circulation by the Blood-Brain Barrier (BBB), here the focus was on the level of two enzymes involved in brain cholesterol hemostasis in a diabetic rat...
CYP46A1 is a cholesterol mechanism. First, it may regulate brain cholesterol content through two thesis and cholesterol elimination under steady suggesting a parallel balance between cholesterol syn-
reduced and almost reached ging. ginger extract shown can correct 46A1 level in a steady
24S mainly expressed in the brain, enzymatically produces brain.
It is well documented that CYP46A1, hydroxycholesterol (24S which is increased by ginger extract, it
is increased by ginger extract, it was
found that the administration of ginger extract significantly reduced HMG-CoA reductase protein level in comparison to control or diabetic group. This reduction was more pronounced in diabetic group receiving 400 mg/kg of ginger. By contrast, up-regulation of CYP 46A1 in diabetic rats treated with ginger extract in comparison to diabetes groups verified its role in stimu-
lar oxidase damage, apoptosis, and inflammation in streptozotocin-induced diabetic rats revealed the neuroprotective effect of gin-
ger against structural brain alteration associated with diabetes.

**Conclusion**

Taken together, our findings revealed a possible mechanism for the cholesterol-lowering effect of ginger in a diabetic animal model. Our data regarding the regulatory effect of ginger on players involved in brain cholesterol homeostasis along with other reports showing the reduction of brain oxidative damage, apoptosis, and inflammation in streptozotocin-induced diabetic rats revealed the neuroprotective effect of ginger against structural brain alteration associated with diabetes.

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![Figure 2. Effect of hydro alcoholic extract of ginger on the protein levels of CYP46A1 in the brain. A) Representative immune-blotting showing specific bands for CYP46A1. β-actin is used as an internal control. B) Graphic presentation of data obtained from western blot analysis. Each bar shows mean±SD. From left to right, lanes 1, 2, and 3 are related to control; lanes 4, 5, and 6 are related to diabetic group (DM) without any treatments; lanes 7, 8, and 9 are related to diabetic group receiving 200 mg/kg ginger extract; and lanes 10, 11, and 12 are related to diabetic group receiving 400 mg/kg ginger extract, respectively.](image-url)
Effect of Hydroalcoholic Ginger Extract on Main Proteins Involved in Brain Cholesterol Homeostasis

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