Neuroprotective role of curcumin on the hippocampus against the structural and serological alterations of streptozotocin-induced diabetes in Sprague Dawley rats

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ARTICLE INFO

Article type: Original article

Article history:
Received: Sep 6, 2016
Accepted: Apr 13, 2017

Keywords:
Curcumin
Diabetes
Hippocampus
Neuroprotective Rats

ABSTRACT

Objective(s): Diabetes mellitus causes impaired memory and cognitive functions. The hippocampus plays a key role in memory and learning. Curcumin attenuates diabetic nephropathy in vivo. Curcumin has shown a neurogenic effect and cognition-enhancing potential in aged rats. The aim of this study is to evaluate the possible protective role of curcumin on the histological and serological changes of the hippocampus in diabetic rats.

Materials and Methods: Forty albino rats were divided into four groups, ten rats each. Group 1 control rats, group 2 rats received curcumin orally (200 mg/kg/day for six weeks), group 3 rats were injected intraperitoneally with streptozotocin (STZ) (100 mg/kg, single dose), group 4 received a single injection of STZ and received curcumin orally for six weeks. Parafin sections of hippocampus were prepared and stained with hematoxylin and eosin stain, and immunohistochemical staining for GFAP and caspase-3. Morphometrical and statistical analyses were performed. Glycemic status and parameters of oxidative stress was measured.

Results: Examination of hippocampus of diabetic rats showed disorganization of small pyramidal cells in CA1, many cellular losses in the pyramidal cells of CA3, many degenerated granule cells in the dentate gyrus. GFAP positive astrocyte and caspase-3 positive neuron counts were significantly increased. There were significant serum glucose elevation and significant lowered levels of oxidative stress parameters as compared to control rats. Curcumin administration improved the structural and serological alterations of the hippocampus with significant reduction in serum glucose level.

Conclusion: Curcumin ameliorates the deteriorus effect of diabetes on the hippocampus through its antioxidant, antiapoptotic and anti-inflammatory efficacies.

Introduction

Diabetes mellitus is a metabolic disorder characterized by the presence of high blood sugar (hyperglycaemia) resulting from either deficient levels of the hormone insulin or from defective insulin action or both (1).

The rapidly rising prevalence of diabetes mellitus is highlighted all over the world. It was 2.8% in 2000 and will rise up to 4.4% in 2030. The total number of people suffering from diabetes was 171 million in 2000 and may rise as many as 366 million in 2030 (2).

Uncontrolled hyperglycaemia can cause severe damage to almost all of the body’s major organs including vasculopathy, nephropathy, neuropathy and retinopathy (1). Peripheral neuropathy was known as the only complication involving the nervous system. In the last decade, it has become clear that central nervous system is also involved in both human and animal (3).

The central nervous system alterations include abnormal expression of hypothalamic neuropeptides, astrogliosis in the cerebellum, dysfunction of the glutamate transporter and neuronal degeneration (4).

Cognitive and behavioral impairments are being the most reported central nervous system dysfunction (5). Human subjects with type 1 or type 2 diabetes mellitus exhibited lower cognition compared to non-diabetic subjects (6). Impaired cognitive functions were reported in rodent models of diabetes. Rats with streptozotocin (STZ) induced diabetes showed impaired learning ability (7).

The hippocampal formation is a complex part of the limbic system. It is formed of the hippocampus proper, the dentate gyrus and the subiculum. It plays a key role in cognitive functions like memory and learning (8).

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Oxidative stress plays an important role in the pathogenesis of diabetic complications. The most important free radicals are reactive oxygen species (ROS) and reactive nitrogen species. ROS include free radicals such as superoxide and hydroxyl (9). A body of evidence is stating that apoptosis is implicated in several diabetic complications. These include neuronal cells apoptosis in diabetic neuropathy, myocardial apoptosis in cardiac pathogenesis and mesangial cell apoptosis in diabetic nephropathy (10).

Diabetes is a great problem to public health. Effective management of diabetes is a pressing goal of the research worldwide. Current treatments, like hypoglycemic drugs or insulin, causes many adverse effects and faces significant challenges as insulin resistance or hypoglycaemic attacks. The recent trend is to use medicinal natural products which may offer the same efficacy without so many side effects (11).

One of these natural compounds is curcumin which has considered a promising natural therapeutic agent. It is the most active component found in turmeric (Curcuma longa Linn.), a popular Indian spice. Curcumin, a nontoxic polyphenol, has been shown to contain antioxidant, antiapoptotic, anti-infection, hypcholesterolemic and anti-inflammatory activities (12).

Curcumin attenuated podocyte apoptosis caused by high glucose in vitro and diabetic nephropathy in vivo (13). Curcumin had shown a neurogenic effect and cognition-enhancing potential in aged rats (14). Curcumin ameliorated the harmful effects of glutamate neurotoxicity on the hippocampus (15).

So, the aim of the present study was to evaluate the possible protective role of curcumin on the histological and serological changes of the hippocampus in diabetic rat.

Materials and Methods

Experimental animals

Forty healthy male rats of the Sprague Dawely strain aged four to six months were locally bred at the animal house of Research Center and Bilharzial Research Unit, Faculty of Medicine, Ain Shams University. The rats were housed in stainless steel cages, five rats per cage. The rats were exposed to dark/light cycle and allowed for free access to diet and water with suitable environmental conditions and good ventilation and at a temperature of 25 °C.

Induction of diabetes

After 12 hr of fasting: the animals of model groups received a single 100 mg/kg intraperitoneal injection of STZ dissolved in 0.1 mol/l cold citrate, pH 4.5. The animals were allowed to drink 5% glucose solution to overcome STZ-induced hypoglycemia. After 72 hr, animals with blood glucose levels greater than 200 mg/dl were considered diabetic (16).

Curcumin administration

Curcumin, available in the form of a powder, was administrated at a dose of 200 mg/kg/day dissolved in 1 ml/kg corn oil by daily gavage. Corn oil, available in the form of an oily solution, was used as a solvent for curcumin (Sigma Chemical Co., Cairo, Egypt).

Experimental design

The rats were divided into four groups, ten rats each.

Group 1: control group: rats received single intraperitoneal injection of citrate buffer alone.

Group 2: non diabetic received curcumin: rats received curcumin orally in a dose of 200 mg/kg/day for six weeks (17).

Group 3: diabetic group: rats were injected intraperitoneally with a single dose of 100 mg/kg of STZ (16).

Group 4: diabetic received curcumin: rats received a single intraperitoneal injection of STZ (100 mg/kg) then received curcumin orally in a dose of 200 mg/kg/day for six weeks.

Sampling

The separated blood: samples were attained from animals’ tail vein, and then centrifuged for 15 min. The separated serum was used for the detection of glucose and insulin.

At the end of the experiment the animals were anesthetized and decapitated. The skulls were opened and the brains were collected immediately and carefully. Coronal sections were obtained and left immersed in the 10% formalin to continue fixation two more days. Then, the specimens were processed for preparation of paraffin blocks. Paraffin sections (5 µm) were prepared and stained with hematoxylin and eosin stain (18).

Immunohistochemical study for glial fibrillary acidic protein (GFAP) and caspase-3

Sections were taken on positive slides and immunostained using avidin-biotin technique. Paraffin sections (5 µm) were deparaffinized in xylene, rehydrated and pretreated with 0.01% hydrogen peroxide (H₂O₂) for blocking endogenous peroxidase activity and unmasking of the antigenic site was carried out by transmitting sections into 0.01 M citrate buffer (pH 6) for 10 min and in ethanol for 10 min. Microwave-assisted antigen retrieval was then performed for 20 min. Sections were incubated overnight at 4 °C with the diluted primary antibody at dilution 1/100 monoclonal mouse antibodies for GFAP and caspase-3. Sections were incubated with the avidin-biotin complex (ABC) reagent for 60 min then incubated in peroxidase substrate solution for 6-10 min. Finally, haematoxylin was used as a counter stain, dehydration in absolute alcohol, clearing and
mounting were done. Immunoreactivity was visualized as dark brown cytoplasmic staining for GFAP and caspase-3 (19).

Quantitative morphometric study: Ten fields per section were taken from high-power (400x) using Olympus digital camera installed on Olympus microscope. The number of immunopositive cells for GFAP and caspase-3 was counted in sections of each animal of the different studied groups, using ImageJ software program and averaged per field for each animal.

**Biochemical analysis**

**Catalase (CAT) assay**
Catalase (CAT) activity was assayed in tissue homogenates of hippocampus using a kit provided by Abcam according to manufacturer’s instructions, based on the elimination of H₂O₂ from a solution containing 30 mmol/l H₂O₂ in 10 mmol/l potassium phosphate buffer (pH 7) at 240 nm. Results were explained as mmol of H₂O₂ consumed/min/mg protein (20).

**Glutathione peroxidase (GPx) assay**
Glutathione peroxidase (GPx) activity in homogenates of hippocampus tissue of rats was measured by a Glutathione Peroxidase Assay Kit provided by Abcam according to manufacturer’s instructions. The results were expressed as units/mg protein (21).

**Superoxide dismutase (SOD) assay**
Superoxide dismutase activity in homogenates of hippocampus tissue of rats was measured using a Superoxide Dismutase Activity Colorimetric Assay Kit provided by Abcam, according to manufacturer’s instructions. Assay method was based on a competitive inhibition assay using xanthine-xanthine oxidase system to reduce nitroblue tetrazolium. Results were explained as U/mg protein (22).

**Reduced glutathione (GSH) assay**
Reduced glutathione (GSH) was assayed using the method of Ellman, by treating 0.5 ml of the supernatant with 0.5 ml of Ellman’s reagent consisted of (19.8 mg of 5,5-dithiobisnitrobenzoic acid in 100 ml of 0.1% sodium nitrate) and 3 ml of phosphate buffer (0.2 M, pH 8). This method depends on reduction of Ellman’s reagent by SH group of (GSH) to form 5, 5'-dithio-2-nitrobenzoic acid in phosphate buffer. The produced yellow color was measured spectrophotometrically at 412 nm, using a biochemical analyzer manufactured by (ERMA INC. Tokyo, JAPAN, model: AE-600N) (23).

**Statistical analysis**
The values were represented as mean±standard deviation (±SD). The data were analyzed by one-way ANOVA with post-hoc test for multiple comparisons between groups using SPSS software (SPSS Inc., Chicago, Illinois, USA).

**Results**

**Histological results**

*Hematoxylin and eosin stained sections*
Light microscopic examination of H&E-stained paraffin sections of the rats of control group (group 1) showed C-shaped hippocampus. It is divided into four regions CA1, CA2, CA3 and CA4 and dentate gyrus. Areas in between compact zones of cells comprise the molecular layer. The core of the dentate gyrus is forming the hilus (Figure 1).

_Cornu Ammonis_ CA1 is formed of 5–6 compact layers of small pyramidal cells arranged with characteristic palisade appearance. The pyramidal cells showed vesicular nuclei, prominent nucleoli and pale basophilic cytoplasm (Figure 2). CA3 is formed of zone of large pyramidal cells (Figure 3). The molecular layer consists of interacting axons and dendrites, glial cells, and capillaries (Figures 2, 3).
The dentate gyrus is formed of compact layers of small granule cells. The dentate hilar region contains axons, dendrites, and interneurons (Figure 4).

The non-diabetic received curcumin (group 2) showed the same histological appearance like control (group 1).

Examination of the diabetic group (group 3) revealed disorganization and dispersion of pyramidal cells in CA1 region, most of them having eosinophilic cytoplasm and pyknotic nucleus. Some having halo around them (Figure 5). The pyramidal cells of CA3 region were disorganized with many areas of cell loss. The cells appeared contracted. Many degenerated with eosinophilic cytoplasm were observed. Their nuclei were shrunken dark pyknotic or karyolytic or hyperchromatic (Figure 6). The molecular layer showed increased glial cells and widened congested capillaries (Figures 5, 6).

Figure 3. A photomicrograph of a coronal section of the hippocampus of an adult control rat showing layers of large pyramidal cells in CA3 region (arrows) and molecular layer with many glial cells (g) and capillaries (c). H & E; X 400

Figure 4. A photomicrograph of a coronal section of the hippocampus of an adult control rat showing dentate gyrus formed of layers of compact granular cells (red arrows). H & E; X 400

Figure 5. A photomicrograph of a coronal section of the hippocampus of an adult diabetic rat showing dispersed degenerated pyramidal cells with eosinophilic cytoplasm and pyknotic nucleus (arrow) with some having halo around them (arrow head). Notice widened congested capillaries (c) and glial cells (g). H & E; X 400

Figure 6. A photomicrograph of a coronal section of the hippocampus of an adult diabetic rat showing disorganization and areas of cell loss (CL) of contracted pyramidal cells (P) of CA3 region; some kariolytic nuclei (K), pyknotic nuclei (double arrows) and hyperchromatic nuclei (arrow head). Notice the dilated capillaries (C) and increased glial cells (g) in molecular layer. H & E; X 400

Figure 7. A photomicrograph of a coronal section of the hippocampus of an adult diabetic rat showed many degenerated granule cells in the dentate gyrus with vacuolated cytoplasm (black arrows) and pyknotic nucleus (red arrow). Notice the dentate hilus with increased glial cells (g). H & E; X 400
The dentate gyrus showed marked disorganization. Many degenerated granule cells with vacuolated cytoplasm and pyknotic nucleus were also detected. The dentate hilus showed many glial cells (Figure 7). Areas of hemorrhage and areas of cell loss were detected in the dentate gyrus (Figure 8).

H&E stained paraffin sections of the diabetic group 4 showed preservation of most of pyramidal cells of CA1 region. Most of them showed vesicular nuclei. Few nuclei were pyknotic (Figure 9). Large pyramidal cells of CA3 region were preserved. Most of them had centrally located vesicular nuclei and apparent nucleoli (Figure 10).

The dentate gyrus was preserved. The granule cells were compactly arranged with rounded pale vesicular nuclei the dentate gyrus. The dentate hilus showed nearly normal neurons (Figure 11).

**Figure 8.** A photomicrograph of a coronal section of the hippocampus of an adult diabetic rat showed wide area of hemorrhage (h) in the dentate gyrus. Notice area of cell loss (CL). H & E; X 400

**Figure 9.** A photomicrograph of a coronal section of the hippocampus of an adult rat of the diabetic received curcumin group showing preservation of most of pyramidal cells of CA1 region (arrow head). Notice few cells with pyknotic nuclei (arrows). H & E; X 400

**Figure 10.** A photomicrograph of a coronal section of the hippocampus of an adult rat of the diabetic treated with curcumin group showing preservation of large pyramidal cells of CA3 region (arrows). H & E; X 400

**Figure 11.** A photomicrograph of a coronal section of the hippocampus of an adult rat of the diabetic received curcumin group showing preservation of the dentate gyrus (DG). H & E; X 400

**Figure 12.** Immunohistochemical staining for the demonstration of GFAP in the hippocampus of an adult control rat showing GFAP-positive staining in cytoplasm and processes of scattered astrocytes; they appear small, with few processes (arrows). GFAP; × 400
Table 1. Effect of curcumin on GFAP positive astrocytes number and caspase-3 positive neurons in all studied groups

|                          | Number of GFAP positive astrocytes / field | Number of caspase-3 positive neurons / field | Significance of GFAP positive astrocytes number and caspase-3 positive neurons |
|--------------------------|--------------------------------------------|----------------------------------------------|--------------------------------------------------------------------------------|
| Control group            | 11.75 ± 1.46                               | 1.26 ± 0.5                                   | Versus any of the other studied groups (P<0.001)                                |
| Non diabetic received curcumin | 10.62 ± 1.46                               | 1.07 ± 0.6                                   | Versus control group (P<0.01)                                                  |
| Diabetic group           | 35.5 ± 1.83                                | 10.61 ± 1.2                                  |                                                                                |
| Diabetic received curcumin | 16.1 ± 1.73                                | 3.72 ± 0.8                                   |                                                                                |

Data expressed as mean ± standard deviation. * highly statistically significant difference; ** statistically significant difference

**Immunohistochemical stains**

**GFAP immunostaining**

The control group showed GFAP positive reaction in cytoplasm and in the few processes of scattered small astrocytes of molecular layer (Figure 12). The diabetic group showed many intensely stained GFAP-positive astrocytes with increased size of their cell body and increased number and length of their processes in the molecular layer and pyramidal layer (Figure 13). The diabetic received curcumin group showed relatively fewer astrocytes with thin processes in the molecular layer (Figure 14).

**Caspase-3 immunostaining**

The control group showed negative immunostaining reaction for caspase-3 in the pyramidal cells (Figure 15). The diabetic group showed marked cytoplasmic expression of the caspase-3 in most of the pyramidal cells (Figure 16). The diabetic received curcumin group showed minimal brownish immunoreaction for caspase-3 in some pyramidal cells less than those of diabetic group (Figure 17).

**Figure 13.** Immunohistochemical staining for the demonstration of GFAP in the hippocampus of an adult diabetic rat showing many GFAP-positive astrocytes with thick, long and intensely stained processes in the molecular layer and pyramidal layer (arrows). GFAP, ×400

**Figure 14.** Immunohistochemical staining for the demonstration of GFAP in the hippocampus of an adult rat of the diabetic received curcumin group showing relatively fewer astrocytes with thin processes in the molecular layers (arrows). GFAP, ×400

**Figure 15.** Immunohistochemical staining for the demonstration of caspase-3 in the hippocampus of an adult control rat showing negative immunoreaction in pyramidal cells. Caspase-3, ×400
Curcumin role on hippocampus in diabetic rats

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Table 2. Effect of curcumin on glycemic status in all studied groups

| Parameters          | Group 1 (control) | Group 2 (non-diabetic received curcumin) | Group 3 (diabetic group) | Group 4 (diabetic received curcumin) |
|---------------------|-------------------|----------------------------------------|--------------------------|-------------------------------------|
| FBG (mg/dl)         | 68.40±23.3        | 64.7±19.5                              | 214±74.2 a, b            | 69.65±19.86 c                      |
| Postprandial BG (mg/dl) | 103.2±18.45    | 98±15.42                              | 230±82.1a, b             | 92.16±20.1 c                       |
| Insulin (ng/ml)     | 2.7±0.8           | 2.9±0.7                                | 0.73±0.54 a, b           | 2.5±0.45 c                         |
| HOMA-IR             | 1.46±0.36         | 1.23±0.52                              | 4.68±1.5 a, b            | 1.34±0.470 c                       |

Data are expressed as mean ± SD, FBG: fasting blood glucose, Postprandial BG: 2-hours blood glucose after oral glucose loading (3 g/kg b. w.), HOMA-IR: homeostasis model assessment of insulin resistance. a Significantly different from control group, b significantly different from non-diabetic group received curcumin, c significantly different from diabetic group.

Figure 16. Immunohistochemical staining for the demonstration of caspase-3 in the hippocampus of an adult diabetic rat showing marked cytoplasmic expression of the caspase-3 in most of the pyramidal cells (arrow). Caspase-3, × 400

Figure 17. Immunohistochemical staining for the demonstration of caspase-3 in the hippocampus an adult rat of the diabetic received curcumin group showing weak immunoreactivity of caspase-3 in the cytoplasm of the pyramidal cells with brown cytoplasmic deposits (arrow). Caspase-3, × 400

Quantitative immunohistochemical assessment and statistical analysis

As revealed in (Table 1, Histogram 1) there was a highly statistically significant increase (P<0.001) in the number of GFAP positive astrocytes and caspase-3 positive neurons of the diabetic group as compared to control group. However, the diabetic received curcumin group revealed a dramatic decrease in the number of GFAP positive astrocytes and caspase-3 positive neurons as compared to the diabetic group (P<0.001).

Biochemical results

Effect of curcumin on blood glycemic status

Table 2 shows the mean levels of fasting blood glucose, post prandial blood glucose, insulin and HOMA-IR for all studied groups. The diabetic rats had significantly (P<0.01) elevated blood glucose concentrations compared with non-diabetic rats. Feeding rats with curcumin had significantly (P<0.05) reduced blood glucose concentrations and HOMA-IR, however, significantly elevated mean insulin levels in comparison to control. Also, the treatment with curcumin significantly improved the glycemic status in diabetic rats when compared with untreated diabetic rats.

Effect of curcumin on parameters of oxidative stress

Table 3 shows significantly (P<0.05) lower levels of GSH, SOD, CAT and GPx in diabetic rats when compared with the control rats. Treatment with curcumin significantly (P<0.05) increased their levels.

Table 3. Effect of curcumin on oxidative stress parameters in all studied groups

| Groups                      | Group 1 (control) | Group 2 (non-diabetic received curcumin) | Group 3 (diabetic group) | Group 4 (diabetic received curcumin) |
|-----------------------------|-------------------|----------------------------------------|--------------------------|-------------------------------------|
| GPx (μgGSHconsumed/min/mg protein) | 25.36 ± 2.3       | 27.24 ± 1.97                           | 12.35 ± 2.14 a, b       | 22.45 ± 1.75 c                      |
| GSH (μg/mg protein)         | 9.32±2.8          | 10.6±3.4                               | 3.19±0.76 a, b          | 8.6±3.9 c                           |
| SOD (units/mg protein)      | 4.8±0.38          | 5.32±0.47                              | 2.23±0.31 a, b          | 4.6±0.51 c                          |
| CAT (μmoles of H2O2)        | 15.60 ± 0.62      | 16.52 ± 1.19                           | 6.31±0.73 a, b          | 14.30±0.46 c                        |

Data are expressed as mean±SD, a significantly different from control group, b significantly different from non-diabetic group received curcumin, c significantly different from diabetic group.
Discussion

Diabetes mellitus is associated with lower cognitive performance and increased risk for dementia. The incidence of cognitive function was found to be declined by 40% in people with diabetes. It was reported that there is an overall 50-100% increase in the incidence of dementia in diabetic subjects (24).

Examination of diabetic hippocampus sections revealed that diabetes caused marked histological alterations. Many degenerative changes were seen in the pyramidal cells. Disorganization of pyramidal cells detected in CA1 region with absence of characteristic palisade arrangement. Many areas of cell loss leaving empty spaces were detected in CA3 region. The dentate gyrus showed marked disorganization, vacuolation and hemorrhage and decreased granule cells. Similar findings were found by other researchers (25). However, Zhou et al. (26) stated that STZ-induced diabetic rats showed neuronal cell death at the CA3 and dentate gyrus regions of the hippocampus.

Moreover, Pamidi and Satheeshaya Nayak, (27) reported that untreated diabetes mellitus can induce the damage in the hippocampus within two weeks of diabetic period or earlier and prolonged untreated diabetes leads to severe hippocampal neurodegeneration.

The current study revealed marked cytoplasmic expression for the caspase-3, an indicator of apoptosis, in most of the pyramidal cells. There was a highly statistically significant increase in caspase-3 positive neuron count of the diabetic group as compared to other studied groups.

Vincent et al. (28) reported increased caspase-3 activity indicating neuronal and Schwann cells death in uncontrolled hyperglycemia.

Zhao et al. (29) proved that diabetes increase bax and caspase-3 expression leading to apoptosis of the pyramidal neurons in STZ-induced diabetic rats. Hyperglycemia elevates ROS in hippocampal neuronal cells resulted in decrease in ATP activity and increase in caspases-3 and 9 leading to caspase-dependent apoptosis (30).

Several possible mechanisms have been explained regarding diabetes induced neurodegeneration. Diabetes mellitus and the accompanying hyperglycemia is a chronic endogenous stressor that is accompanied with increased oxidative stress in brain, particularly the hippocampus, by accelerating free radical generation. These radicals contribute to increased neuronal degeneration by inducing oxidation to proteins, DNA alterations and lipids oxidation in cell membranes (31). The present study indicated statistical high significant reduction in the levels of the parameters of oxidative stress in the diabetic rats.

Another mechanism is the activated polyol pathway during hyperglycaemia consuming NADPH which is the essential cofactor for regenerating reduced glutathione. Depletion of glutathione lowers the threshold for intracellular oxidative damage (32). This mechanism also explains the significantly (P<0.05) lower levels of reduced GSH in the diabetic rats of the current study.

Also, another important mechanism expected to lead to neuronal loss in the diabetic rats of this study could be due to lower insulin and HOMA-IR which has a neuroprotective anti-apoptotic effect (33).

In the present study, GFAP positive astrocytes were significantly increased in the diabetic group as compared to control group supported by the findings of Baydas et al (34). Similar findings were found by Saravia et al. (35) who stated that GFAP positive astrocytes increased 3-fold in the hippocampus of the diabetic mice compared to the control group.

In contrast, other studies have shown a decrease in the GFAP immunoreactivity in the astrocytes of diabetic tissues. GFAP immunoreactivity was reduced in the olfactory bulb of 8-week STZ-diabetic animals (36). A decrease in the number of GFAP-immunoreactive astrocytes was also detected in the gray matter of the spinal cords of STZ-induced diabetic rats (37). These conflicting reports could result from discrepancy in immunohistochemical methods, diabetic models, doses of STZ, and antibody sources used by different researchers (38, 39).

Primary function of glial cells is to surround neurons and hold them in place, to nurture the neurons, to insulate one neuron from another and to destroy and remove dead neurons, a process essential to neural plasticity and stability. However, damaged neurons may activate glia, which may then kill neurons (40).

Treatment with curcumin significantly improved the glycemic status in diabetic rats. The results of our study showed significant decrease in both fasting and postprandial blood glucose levels as well as HOMA-IR after treatment with curcumin. Mahesh et al. (41), also reported a significant decrease in blood glucose levels in STZ-induced diabetes model after treatment with curcumin.

In the present study, the diabetic group received curcumin showed preserved histological structure of the hippocampus and the dentate gyrus. There was high significant reduction in the number of caspase-3 positive neurons as compared to the diabetic group. These finding may be explained on the basis that curcumin reduce the alterations in lysosomal enzymes activities, thus minimizing the injury due to diabetes (42). In line with the current results, it was reported that treatment of diabetic rats with curcumin cause reduction of pancreatic caspase-3 content (43) and decrease of cardiac caspase-3 immunoreactivity expression (44).
Curcumin induces proliferation and migration of neural progenitor cells and enhances neural plasticity and repair in vitro and in vivo in the adult hippocampus (45).

The present study showed decrease in GFAP immunohistochemical staining and in the number of GFAP positive astrocytes of the diabetic group received curcumin compared with the diabetic group.

Curcumin caused significant decrease in GFAP immunohistochemistry staining in retinal Müller glia in diabetic rat (46).

Tissue concentration of antioxidant parameters showed significant increase in the diabetic group received curcumin and that may suggest a significant protecting role of curcumin against harmful effects of diabetes. Results of Borra et al. (47) showed a significant protective effect of curcumin that may be due to modulating lipid peroxidation and augmenting antioxidant defense system.

El-Bahr, (48) attributed the curcumin induced anti-diabetic effect to the fact that it maintains normal glucose metabolism via amelioration of the diabetes induced oxidative stress via increasing the activities of antioxidant enzymes including superoxide dismutase, glutathione peroxidase and increasing the reduced glutathione concentration.

**Conclusion**

The results of the current work suggest that curcumin has a neuroprotective beneficial effect as it could protect rat hippocampus from STZ induced altered histological and serological changes owing to its antioxidant, anti-inflammatory, and antiapoptotic effects. Therefore, diabetic patients and high risk people for diabetes are recommended to consume daily doses of curcumin in their meals.

**Acknowledgment**

The authors wish to thank Dr Amgad Gaber Elsaid (Lecturer of Anatomy and Embryology, Faculty of Medicine, Ain Shams University, Egypt) for his efforts in conducting and reviewing this paper.

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