Effect of Borage on hippocampal TNF-α protein and gene in the Amyloid β-Peptide(25–35)-Induced of Alzheimer model in rat

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Alzheimer’s disease (AD) is a neurodegenerative disorder and most common form of dementia that leads to memory impairment. In the present study we have examined the investigating effect of Borago officinale on hippocampal TNF-α protein in the Beta Amyloid peptid induced of inflammation in the rat. Wistar male rats received intrahippocampal (IHP) injection of the Aβββββ(25–35) and borage extract orally (100 mg/kg). Inflammation was confirmed by pathologist. In order to measure the expression of the TNF-α gene, RT-PCR method and for extraction of the TNF-α protein western blotting method was used. Results showed that using Borage lead to reduce TNF-α gene and protein. This data suggests that borage could improve the reduction of inflammation factors such as TNF-α and borage consumption may lead to an improvement of AD-induced cognitive dysfunction.

Key words: TNF-α, Alzheimer, RT-PCR, Western Blot, Amyloid β-Peptide

Alzheimer’s disease (AD) and inflammation of brain immune system happen simutaneously. Based on Walsh and et al 2012 investigations, increase of complement proteins, cytokines, microglia cells and astrocytes are observed (Walsh et al., 2012). These protein groups can do intermediation and set all aspects of natural and acquired immunes (Choi and et al, 2014). Some of these cytokines can lead to inflammation in the body. TNF-α can be probably considered as the most inflammatory cytokines. Based on Rinne and et al investigations, Monocytes and Macrophage are two important features lead to TNF-α producing cells (Rinne and et al, 2003).

In Alzheimer disease around the nerve cells setting Beta Amyloid can lead to microglia activity, free radical producing and inflammatory processing and hurt the healthy neurons (Sarter and et al., 2004).

In order to measure the effect of anti-inflammatory Borage, a large number of studies were carried out by several investigators such as Kast and et al. They found that oil of Borage causes the reduction of inflammation because of the existence of GLA and increasing of the synthesis of Prostaglandin E and cAMP (Mattson and et al., 2004).

Cameron and et al studied the Herbal treatment of arthritis disease and emphasized that Borage reduces the rate of inflammatory cytokines because of the existence of GLA (Cameron and et al., 2006).
Although a large number of investigations were carried out in Alzheimer disease, it seems that there is no sufficient studies in the current field. In the present study, effects of Borage in decreasing inflammation and decreasing of inflammatory factors such as TNF-α were studied. In order to examine these effects, peptid beta amyloid and Alzheimer model in rat were used.

MATERIALS AND METHODS

The Aβ(25–35) was purchased from sigma-Aldrich company (St Louis, MO, USA). Borage leaves were obtained in dried condition from Research Institute for Islamic and Complementary Medicine (Tehran, Iran). Aβ 25–35 was solubilized in sterile water at 1 µg/µL concentration and stored at -20°C(Allen 2002).

Animals

28 250-300 gram weight male wistar rats were chosen for experimental procedure. all animals were housed in group and remained 21 degree centigrade along all the experimental procedure. Humidity of the house condition was 50%. 12 hr light/12 hr dark cycle was used for light housing condition.(Shankar and et al 2008).

The rats randomly were classified in to the following groups: the control or intact group that was left undisrupted; the sham-operated group; the Aβ 25–35 model group which received single bilateral intrahippocampal (IHP) injections of Aβ25–35; the borage-treated group that received borage extract (orally, 100 mg/kg) following IHP injection of Aβ25–35 for 14 days (Kast and et al 2001).

Preparation of Borage Extract

Dried borage leaves were cleaned carefully, then ground into coarse powder by electrically driven device. The powdered material was soaked into aqueous ethanol (80%) for one week with occasional shaking. The extract was filtered through a Whatman filter paper and evaporated to dryness under reduced pressure at a maximum of 40°C using a rotary evaporator. Borago officinalis produced 10.9% dried extract. The extract was completely dissolved in purified water and kept at 4°C(La Ferla and et al 1995).

Intrahippocampal Injection of Aβ25–35

The animals were anesthetized with the ketamine (100 mg/kg) and xylazine (10 mg/kg) and were transferred to a stereotaxic apparatus (Stoelting, Wood Dale, IL, USA). The injection was made using a 10 µL microsyringe (Hamilton-Reno, NV, USA). Relative to the bregma and with the stereotaxic arm at 0°, the coordinates for the toothed gyrus were posterior -3.6; lateral α2.3; and dorsal 3 mm (Conforti and et al 2008).

Aβ solution (6 µL) was bilaterally injected into the area over 1 µL/2 min. the cannula was left in place for 2 min after each injection to allow for spread. Sham operated rats received vehicle solution(10). The skin was then stitched and the animals were left to recover in a warm box before returning to their home cages. The injection place was checked by injection trypan blue instead of peptide in preliminary experiments (Figure 1).

Histological Verification

For verification of the injection place a light microscope (Olympus, Japan) was used, the trypan blue injected rats were perfused with 4% paraformaldehyde in 0.1 M phosphate buffer (pH = 7.3 ) and the hippocampi were serially sectioned into 10 µm coronal parts by a microtome (Leica Instruments, Germany). After deparaffinization and rehydration, sections were stained in 0.1% cresyl violet for 3 minutes. Finally, the parts were photographed with a digital camera (Olympus, DP 11, Japan) attached to a microscope (Olympus Provis, Ax70, Japan) and the stained pieces were qualitatively analyzed for the injection site(Klementiev and et al 2007).

RT-PCR Experiment

The day after the last treatment, the animals were euthanized by cervical dislocation. The brains were removed; hippocampi were immediately dissected out on ice, and then frozen in liquid nitrogen and kept at -80 °C until use.

Total mRNA was extracted from the frozen hippocampi by using phenol-chloroform method. Tissue samples were homogenized in 1000 il RNATM (Cinnagen, Teh-ran, Iran) then 200 il ice-cold chloroform was added. The homogenates were centrifuged (Eppendrof, Ham-burg, Germany) at 12000 g for 20 min in 4°C. The RNA of the supernatant was precipitated with isopropanol, and washed with ethanol 75%. The air-dried RNA pellet was dissolved in RNase free water. cDNA first-strand synthesis was performed using a cDNA synthesis Kit (Quigaen, Hilden, Germany). First strand cDNA was used as template for subsequent
PCR with a PCR master kit (Cinnagen, Tehran, Iran), and primers (Cinnagen, Tehran-Iran) as follow:

\[ \text{β-Actin Forward } 5' - \text{GTG GGC CGC CCT AGG CAC CAG-3'} \]
\[ \text{Reverse 5'} - \text{GGC CTT AGG GTT CAG AGG GG-3'} \]
\[ \text{TNF-α Forward } 5' - \text{AAC TTC GGG GTG ATT GGT CC-3'} \]
\[ \text{Reverse 5'} - \text{CTG AGT GTG AGG GTC TGG GC-3'} \]

The PCR reactions included initial denaturation at 95 °C for 3 min, followed by 35 cycles with 95 °C for 50 s, 58 °C for 45 s and 72 °C for 50 s for TNF-α.

The reactions were terminated by 72 °C for 7 min of elongation period. The same annealing temperature was used for β-actin. PCR products were separated by electrophoresis in 1.5% agarose gel at 100V. Our semi-quantitative analysis were made using a digital imaging system (UVIdoc, Houston, Texas, USA).

**Western blot analysis**

Hippocampus tissue collected for Western blot analyses were snap frozen in liquid nitrogen and stored at -70°C until use. Total protein was extracted by homogenization in RIPA buffer (150 mM NaCl, 50 mM Tris-HCl, pH7.5, 1% Triton X, 0.5% Na deoxycholate, 1 mM PMSF) as described previously.

Protein concentration of homogenates was measured and 30 ug resolved by 12% SDS-PAGE and transferred to nitrocellulose membranes (MSI, Westboro, MA, USA). Membranes were blocked in 5% non-fat milk for 1 h at room temperature and probed with polyclonal rabbit anti-mouse StAR antibody diluted 1:5000 in Tris-buffered saline/0.1% Tween-20 (TBST). Following this membranes were incubated with biotinylated goat anti-rabbit IgG for 1 h at room temperature (diluted 1:10,000 in TBST). Membranes were then incubated with Avidin Biotin Enzyme Reagent for 1 h at room temperature and protein signals detected by enhanced chemiluminescence (Supersignal West Pico ECL substrate, Pierce, Rockford, IL, USA) and quantitated by densitometry. A common tissue sample was included on each gel to allow for standardization of chemiluminescence levels and exposure times. Staining of each gel (post transfer) and membrane with Coomassie Brilliant Blue (Sigma Chemical Co.) assessed the accuracy of sample loading and the efficiency of protein transfer. This procedure was repeated for each animal/experimental group.

**Statistical analyses**

Data were presented as mean ± S.E.M and analyzed by SPSS version 21 software. The data of the IL-1 Beta protein and gene were analyzed using two-way analysis of variance (ANOVA), Tukey multiple comparisons tests were used to analyze the significance of the differences between the groups, when appropriate value of p<0.05 was considered significant.

Briefly, the Homogenous test was used to prove the variance homogeneity among the experiment groups.

**RESULTS**

Histological analysis showed that injection of Aβ was in their desired location, according to the atlas of Paxinos and Watson (Figure 1) (Conforti and et al 2008).

**The study of the expression of TNF-α gene**

The obtained results from RT-PCR show that TNF-α gene has a significant reduction among the intact groups in comparison with Beta amyloid and Borage (p<0.05 for both groups of Beta amyloid), that show the increase for the expression of this gene among the Beta amyloid and Borage groups.

Prescribing the Borage caused a significant reduction in TNF-α gene after creating Alzheimer model.

The result obtained from the statistic analyzing TNF-α in attendance groups by ANOVA method showed that there is a significant difference among the experimenting groups.

The Tukey test was used because of the significant difference. the results of this test showed that a significant difference was not observed under the conditions of experiment among the Sham and Intact groups from the view point of TNF-α. Also a significant difference was observed between the Sham and Intact group in comparison with Beta amyloid and Borage groups.

In this test, a significant difference was observed between the Beta amyloid and Borage groups from the view point of TNF-α.

Briefly the Homogenous test was used to prove the variance homogeneity among the attendance groups that the result of this test was
not significant among the attendance groups under the conditions of experiment, that it showed the variance homogeneity among the test groups.

**The study of the rate of the expression of protein in TNF-α**

Dansitometry, the nitro selolose membrane of the protein in Borage and Beta amyloid showed that the rate of this protein in Borage and Beta amyloid groups had a significant increase in comparison with intact and Sham groups. The prscription of the Borage after creating Alzheimer model caused a significant reduction of the rate of the expression of the TNF-α protein in comparison with the Beta amyloid group.

![Fig. 1. Schematic photograph representing the microinjection site of Amyloid β into the hippocampus (black arrow) (85bp, 260bp)](image)

**DISCUSSION**

The results obtained from the present study showed that the prescription of Beta amyloid leads to the increase of the expression of gene and protein TNF-α in hippocampus area. Also the prescription of Borage juice after creating Alzheimer model reduced the injected inflammation caused by Beta amyloid.

![Fig. 2. Western Blot analysis of the TNF-α expression in sham and treatment groups (52 KD, 42 KD)](image)

In present study the mutual injection in hippocampus Beta amyloid25-35 leads to creating inflammation and increasing inflammatory proteins. In agreement with our study Yamaguchi and et al. reported that prescription of Beta amyloid causes Alzheimer(Zussi and et al 2011).

The various experimental studies reported the role of rich feeding from anti-inflammations and anti-oxidants in recovery of neuro toxicity and
recovery of the act of the brain (Cappai and et al 2008).

Great numbers of the medicinal plants were used in neurotic disease as anti-inflammation. Among these plants, the Borago Officinalis has a great importance because of having much Gama Linolenic Acid (GLA) (Yamaguchi and Kawashima 2001).

Gundgrade and et al, in a research that they did about Borago Officinale, got this result that Borage reduces the danger of catching diseases such as cancer, diabet and Alzheimer (Gundgrade and et al 2002).

In a research that Kapoor and et al did about the effect of GLA a natural anti-inflammation, got this result that GLA prevents producing of the IL-1 Beta and its releasing by monocyte and in this way causes the reduction of inflammation. Also GLA causes secretion of prostaglandin E and this factor finally causes the reduction of producing TNF-α and in this way the inflammation finally reduces (Kappor and et al 1999).

Sadiya and et al in a research that they did about Borago Officinale declared that this plant has more than 20% GLA and traditionally it is used in the treatment of Romatoid and has positive and useful effect on the brain (Sadiya and et al 2000).

Kast and et al in a research that they did about the anti-inflammation effect of the oil of Borago Officinale got this result that it causes the reduction of inflammation because of the existence of GLA and increasing of the synthesis of prostaglandin E and CAMP (Kast and et al 1998).

In a research that Cameron and et al did about the Herbal treatment of arthritis romatoid disease, emphasized that Borago Officinale reduces the rate of inflammatory cytokins because of the existence of GLA (Cameron and et al 2006).

The result of this study showed that using the Borage juice after the injection of Beta amyloid lead to the reduction of inflammation. In the present study after analyzing the results of the experiment by SPSS software version 21, it was confirmed that the surgical operation has had no effect on the results of the experiment. By the injection of Beta amyloid in hippocampus, the rate of the gene expression and protein TNF-± in the Borage and Beta amyloid group get a significant increase in comparison with the Sham and Intact groups which this is a confirmation on the created inflammation after the injection of Beta amyloid. After the consumption of the alcoholic juice of Borago Officinale the rate of the gene expression and hippocampic TNF-± protein get a significant reduction in comparison with Beta amyloid group which it is the confirmation in the hypothesis of the present study which declares that the consumption of Borago Officinale casues the reduction of the created inflammatory factors in the Alzheimer model which is created by peptid Beta amyloid in hippocampus.

Generally it seems that the Borago Officinale reduces the effect of inflammatory proteins and in this way it reduces the effects of inflammatory Beta amyloid.

Brifly, the results of the present study declare that the prescription of the Borago Officinale reduces the injected inflammation by Beta amyloid (25-35) and therefore it may be useful in the treatment of some disease as Alzheimer.

REFERENCES


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