Research Article

Improvement of Electroacupuncture on APP/PS1 Transgenic Mice in Spatial Learning and Memory Probably due to Expression of Aβ and LRP1 in Hippocampus

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Objectives. To explore the alterations of β-amyloid (Aβ) and low density lipoprotein receptor-related protein-1 (LRP1) in APP/PS1 mice after electroacupuncture (EA) treatment and further to explore the mechanism. Methods. Forty 6-month-old APP/PS1 mice were randomly divided into a model group and an EA group, with twenty wild-type mice used as a normal control group. Mice in the EA group were treated with EA at GV 20 (bài huì) and bilateral KI 1 (yǒng quán) acupoints for 6 weeks. The Morris water maze was applied to assess the spatial memory in behavior. Immunohistochemistry (IHC), ELISA, Western blotting, and so forth were used to observe the expression of LRP1 and Aβ. Results. The Morris water maze test showed that, compared with the normal control group, the model group’s learning and memory capabilities were significantly decreased (P < 0.05; P < 0.01). The EA group was reversed (P < 0.05; P < 0.01). The hippocampal expression of Aβ in the EA group was significantly decreased compared to the model group (P < 0.01). The expression of LRP1 in the model group was significantly lower than that in the normal control group (P < 0.01); the expression in the EA group was significantly higher than that in the model group (P < 0.01). Conclusions. EA therapy can improve the learning and memory capabilities of APP/PS1 mice. The underlying mechanism may lie in the upregulation of an Aβ transport receptor and LRP1.

1. Introduction

Alzheimer’s disease (AD), the most common cause of dementia in the elderly population, is a kind of neurodegenerative diseases mainly indicated by progressive cognition and memory impairment [1, 2].

In recent decades, the pathological mechanisms of AD have been widely studied [3, 4]. The accumulation of amyloid β-peptide (Aβ) plays the most important role in the pathogenesis of AD [5, 6]. A wealth of evidence has indicated that Aβ1–42 deposits participate in the process of neuronal loss which then leads to the occurrence of dementia in AD patients [7]. The high level of Aβ in sporadic AD lies in the imbalance of Aβ production and clearance, especially disordered Aβ clearance [8]. Recent research has focused on Aβ clearance pathways through the cranial microvascular saturable efflux system, namely, transport across the blood–brain barrier (BBB), which has both a fast transport speed and large transport volume. Therefore, the relationship between brain microvessels and the clearance of Aβ is also crucial [9, 10].

A leading hypothesis supports the fact that the main clearing pathway of Aβ in the brain is the transportation of Aβ through the BBB into peripheral blood which has a strong
clearing ability to it [5, 11, 12]. In addition, the BBB model cultured in vitro by the brain microvascular endothelial cell line has also been used to detect whether Aβ can be transported through the BBB, and the finding was affirmative [13, 14]. Aβ is a polar, soluble macromolecular substance [15], and it cannot be freely exchanged between the brain and peripheral blood via free diffusion. Therefore, if Aβ transportation across BBB exists, there must be Aβ specific transporters in the BBB.

Low density lipoprotein receptor-related protein-1 (LRP1) is known to function as a BBB clearance (or efflux) transporter for Aβ. Efflux of Aβ is initiated when it binds directly to LRP1 at the abluminal membrane of the brain endothelial cell [16]. Yamada et al. [17] proved that the brain microvascular endothelial cell uptaking Aβ mainly relies on LRP1 under the BBB-specific cellular context. Bell et al. [18] found in animal experiments that the isotope-labeled Aβ injected into the caudate nucleus of mice would be cleared out from the brain rapidly, and the labeled Aβ was found in the plasma. The clearance of Aβ could be inhibited by LRP1 specific antibodies. Further studies [16, 19, 20] suggest that, in pathological conditions, the abnormality of Aβ levels in the brain might be associated with the altered expression of LRP1 in cranial microvessels.

One survey reported that 55% of patients with AD had tried at least one form of complementary medicine with the hopes that these therapies could improve their overall quality of life and delay further decline in cognitive functioning. Clinical research has shown that acupuncture can improve the mental and behavioral conditions of AD patients, as well as the cognitive function [21–23]. Electroacupuncture (EA) is a simple and effective modern acupuncture method tried at least one form of complementary medicine with the brain might be associated with the altered expression of LRP1 in cranial microvessels.

2. Materials

2.1. Animal Model and Grouping

2.1.1. Animals. 6-month-old APPswe/PSI double transgenic male mice were used as the animal model of AD, and wild-type mice with the same age and sex were used as the normal control group. Animals were purchased from Model Animal Research Center of Nanjing University (animal lot: SCXK (Ning) 2010-000), weighing 34.2 ± 3.98 g. All experimental procedures comply with the guidelines of the "Principles of Laboratory Animal Care" formulated by the National Institute of Health and the legislation of the People's Republic of China for the use and care of laboratory animals. The experimental protocols were approved by the Animal Experimentation Ethics Committee of Beijing University of Chinese Medicine. Efforts were made to minimize the number of animal uses and the suffering of the experimental animals.

2.1.2. Animal Grouping and Intervention. 40 APP/PSI transgenic mice were randomly divided into two groups, a model group (M) (n = 20) and an electroacupuncture group (EA) (n = 20), while 20 wild-type mice were selected as the normal control group (C). All mice were raised with a regular diet in single cages in the barrier system of the Animal Center of Beijing University of Chinese Medicine.

Regarding EA group, EA on GV 20 (băihuí) and bilateral KI 1 (yǒng quán) was given to the mice, with transversely puncturing at a depth of 2-3 mm by disposable sterile acupuncture needles (0.25 mm × 13 mm) (Beijing Zhongyuan Taihe Medicine Company, Ltd.). The anode and cathode of EA were, respectively, connected to the left and right KI 1 (yǒng quán), with a dilatational wave at a frequency of 2/15 Hz, 1 mA, and an intensity that the needle tremors, while animals keep quiet by Han's acupuncture needle stimulator (Beijing Huaxi Industrial Development Company, Han's LH202H type). GV 20 (băihuí) is located at the intersection of the sagittal midline and the line linking two rat ears [26]. KI 1 (yǒng quán) is located on the sole of the foot, at the indentation near the front part, between the second and third metatarsal bones, 1/3 of the distance from the webs of the toes to the heel [28]. The EA treatment was performed with mice restrained in mouse bags, 15 mins per time, once every other day for 6 weeks. Mice in the normal control group and the model group were just restrained in the same mouse bags for 15 min, once every other day, totally for six weeks [29].

3. Methods

3.1. Learning and Memory Behavioral Testing. Morris water maze testing was conducted in a round pool to test the behavior of learning and memory [30]. Water maze training was given on the 2nd–6th days after EA intervention for 6 weeks, with water temperature at 22 ± 2°C. One day before the experiment, mice in each group were placed on the platform for 10 s to adapt to the environment, and each mouse was offered a period of free swimming in the maze for 1 min with the fourth quadrant as the entry point. On the first to fourth days of the experiment, mice in each group were given a place navigation test. Mice were firstly placed on the platform to adapt to the environment for 10 s and then were sequentially put into water facing the wall away from the four quadrants in sequence. The time recording was stopped after 5 s of the mouse on the platform, and the maximum swim time was set at 60 s. On the fifth day, the spatial probe test was performed. The platform was removed, and mice were sequentially placed directly into the water from the four quadrants. A computer connected to an image analyzer (BS-124S Morris water maze video analysis system: provided by the Pharmacology Laboratory of Dongzhimen Hospital affiliated to Beijing University of Chinese Medicine, Shanghai...
Mobile Information Technology Co., Ltd.) monitored the swim pattern.

3.2. Collection of Brain Tissues and Detection of Related Indexes

3.2.1. \(A\beta_{1-42}\) Immunohistochemistry

Sample Preparation. The brains of 2 mice in each group were fixed in paraformaldehyde after cardiac perfusion and then trimmed, dehydrated with ethanol, made transparent with xylene, embedded in paraffin, and sectioned on a coronal plane.

\(A\beta_{1-42}\) Immunohistochemical ABC Method. Sections were firstly dewaxed and hydrated and were put into 0.01 mol/L citrate buffer for antigen thermal remediation for 10 min and 3% methanol hydrogen peroxide at room temperature for 10 min. Then the sections were blocked in 5% normal goat serum at 37°C for 30 min and incubated with primary antibody diluent (USA, Abcam, ab10148, 1:100) for one night at 4°C. After incubation within the primary antibody diluent, the sections were rinsed with phosphate-buffered saline (PBS) and then incubated with secondary antibody diluent (Boster Biological Engineering, goat anti-rabbit HRP-IgG, 1:1000) for 90 minutes at 37°C. The sections were rinsed with PBS before incubation with ABC complex for 90 minutes at 37°C and were placed into diaminobenzidine (DAB) solution for 10 minutes after being rinsed another time with PBS. After being redyed with hematoxylin, they were dehydrated and mounted after transparency. The sections were observed under the microscope (BX53, Olympus Corporation, Japan).

3.2.2. Laser Confocal Imaging of Frozen Section Observation with Laser Scanning Confocal Microscope

Sample Preparation. With 3 mice in each group, the analytes were fixed in paraformaldehyde after cardiac perfusion. 30% sucrose was then added after 24 hours, and brain tissues were frozen sectioned with OTC embedded when sunk into the bottom of the bottle.

Proteinase K was added at 37°C after the brain tissues were washed three times with PBS with 5 mins each. 30 min later, LRPI (USA, Abcam, ab28320, 1: 70) and \(A\beta_{1-42}\) (USA, Abcam, ab10148, 1:100) were added equivalently after washing for three times with PBS as above. With incubation overnight at 4°C, the second antibodies (FITC (USA, Abcam, ab6785, 1: 250) and Alexa Fluor® 647 (USA, Abcam, ab150079, 1: 150)) were added after washing with the same method. With incubation for 1.5 hours at indoor temperature, DAPI (Beijing Zhongshan Golden Bridge, ZLI-9557) was added to fix the sections after washing as before. A laser scanning confocal microscope (FV1000, Olympus Corporation, Japan) was used for sections observation.

3.3. Collection and Preservation of Brain Tissues. After the water maze test, mice in each group were anesthetized with 0.3% sodium pentobarbital (30 mg/kg). The 15 hippocampi were taken with craniotomy and prepared for ELISA and Western blotting samples in time on the left and right sides, respectively, and then preserved in a refrigerator at 80°C.

3.4. Preparation of Samples

3.4.1. For \(A\beta\) ELISA. The 6 right hippocampi were weighed in each group and homogenized with 8 times the volume of mixed liquor of 5 M guanidine hydrochloride, 50 mM Tris hydrochloric acid (pH 8.0), and 1 mM PMSF on ice. Then the hippocampi were centrifuged with 16000 r/min, at 4°C, for 20 min, and the supernatant was obtained. The diluted samples were prepared by separately mixing the supernatant with a standard diluent of 3200 times the volume (KHB3441, Invitrogen, USA) and 800 times the volume (KHB3441, Invitrogen, USA) with the hippocampus of the model and EA groups. The samples in the normal control group were not diluted.

3.4.2. For LRPI Western Blotting. The 6 left hippocampi were added to the RIPA lysate solution containing 1 mM PMSF with the weight ratio of 1: 100 in each group, and then the total protein was extracted from tissues after homogenizing. Then the hippocampi were centrifuged at 2000 r/min, at 4°C, for 10 minutes, after extraction, and the volume of the supernatant was calculated. The samples were then prepared for Western blotting.

3.4.3. For LRPI ELISA. The remaining 9 hippocampi were weighed in each group and homogenized with a dilution of 1: 8 in PBS. The supernatant was extracted with the same extracting method of 6 right hippocampi. The diluted samples to be tested were prepared by adding the supernatant with a standard dilution of 1: 5 (DRE20100, RD, USA) in all groups.

3.5. Double Antibody Sandwich Method for ELISA. The ELISA kit was used to, respectively, detect samples. First, 50 \(\mu\)L of antibody operating solution was added into each plate at room temperature for 120 minutes. Then 100 \(\mu\)L horseradish peroxidase labeled secondary antibody operating solution was added to each plate at room temperature and dark reaction for 30 minutes. Complete plate washing was performed after four attempts. A 100 \(\mu\)L chromogenic substrate operating solution was added to each plate at room temperature and dark reaction for 30 minutes. Plate washing was performed after four completed attempts. Then 100 \(\mu\)L stop solution was added and mixed into each plate. The absorbance was detected at the 450 nm place with a microplate reader within 30 minutes. The standard protein line was drawn with Excel. The sample concentration was converted according to the sample readings, and \(A\beta_{1-42}\), \(A\beta_{1-40}\), and LRPI concentrations were calculated according to the sample diluted concentration.

3.6. Western Blotting. SDS-PAGE electrophoresis was performed with a 10% separating gel and a 5% stacking gel and transferred to a 0.45 \(\mu\)m PVDF membrane. Membrane blocking was performed using 5% nonfat milk in Tris buffered saline supplemented with 0.1% Tween 20 (TBST). The first antibody (USA, Abcam, ab92544, 1:20000; ab8227, 1:2000)
Table 1: Comparison of escape latency time in each group in Morris water maze place navigation test (x ± s, n = 20).

| Groups               | Cases | Day 1     | Day 2     | Day 3     | Day 4     |
|----------------------|-------|-----------|-----------|-----------|-----------|
| Normal control group (C) | 20    | 54.19 ± 6.13 | 45.96 ± 11.64 | 40.33 ± 14.89 | 37.93 ± 13.78 |
| Model group (M)      | 20    | 58.32 ± 4.6 | 60.00 ± 0.00 | 55.39 ± 8.30 | 54.65 ± 9.87 |
| EA group (EA)        | 20    | 58.14 ± 3.2 | 57.05 ± 5.72 | 52.45 ± 9.40 | 48.65 ± 12.71 |

Notes: ** compared with the normal control group, P < 0.01; * compared with the model group, P < 0.05; *** compared with the model group, P < 0.01; **** compared with the first day of the same group, P < 0.01.

Table 2: Comparison of platform crossover number and swimming distance in platform quadrant of each group in Morris water maze spatial probe test (x ± s, time, cm, n = 20).

| Groups               | Cases | Platform crossover number | Swimming distance in platform quadrant |
|----------------------|-------|---------------------------|----------------------------------------|
| Normal control group (C) | 20    | 1.74 ± 1.4                | 367.35 ± 142.89                         |
| Model group (M)      | 20    | 0.50 ± 0.44               | 192.46 ± 72.00                          |
| EA group (EA)        | 20    | 0.90 ± 0.71               | 296.61 ± 105.84                         |

Notes: ** compared with the normal control group, P < 0.01; * compared with the model group, P < 0.05; *** compared with the model group, P < 0.01.

was added prior to incubation for one night at 4°C. The secondary antibody (USA, Abcam, ab6721, 1:2000) was added before shaking and incubating at room temperature for 1.5 h. HRP-ECL luminous liquid was added and the X-ray film exposure was completed in a dark room following the developing and fixing. After calibrating the markers, analysis and scanning were performed, and the relative expression of LRP1 (LRP1/β-actin gray value) was compared in each group.

4. Statistical Analysis

SPSS 17.0 software was used to conduct the statistical analysis. All data were presented as means ± standard deviation (x ± s). Variance analysis of multigroup repeated measurement design date was adopted for the data of the Morris water maze behavioral escape latency. One-way ANOVA was used after the test of normal distribution and homogeneity of variance, and LSD method was used for pairwise comparisons for the ELISA detection and Western blotting. If there was a non-normal distribution or heterogeneity of variance for the data, a nonparametric test would be used. Statistical significance was set to P < 0.05, while highly statistical significance was set to P < 0.01.

5. Results

5.1. Effect of EA on Spatial Learning and Memory. Statistical results showed that, according to variance analysis of repeated measurement and effect between groups, the escape latency time in each group was significantly decreased with the increase of training days, and there were significant differences among groups (P < 0.05). There was a significant difference for the training time (day) (P < 0.05), but there was no significant difference for the interaction between the training time and groups (day × group) (P > 0.05). Pairwise comparison results showed that the escape latency time in the model group was significantly longer than that in the control and EA groups (see Table 1 and Figure 1).

One-way ANOVA was used to determine the number of platform crossovers and the swimming distance in the platform quadrants for the spatial probe test for the three groups, and the differences were statistically significant (P < 0.05). The number of platform crossovers and the swimming distance in the platform quadrant of the model group were significantly lower than those in the normal control group (P < 0.01). Compared with the model group, the number of platform crossovers of the EA group was significantly higher (P < 0.05), and the swimming distance in the platform quadrant in the EA group was significantly longer (P < 0.01) (see Table 2).

The swimming trajectories of mice in the normal control group was mostly concentrated in the original target platform quadrant, and the subjects tended to search mainly. However, the type of searching in the model group was random. Compared to the model group, the swimming trajectories of mice in the EA group were more concentrated in the original target quadrant or adjacent quadrants, and the
5.2.1. Immunohistochemistry Results of Aβ

The expression of LRP1 was more than that of model group. Reduced, and there were only some diffuse plaques, while expressed. In EA group, the senile plaques were relatively dense-core plaques deposited, while less LRP1 was around the vascular endothelial cells. In the model group, there were not found, while LRP1 was expressed mostly inside the cell and negative ones outside the cell. In the model group, there were positive expressions of Aβ1–42 and plaque deposits with compactness outside the cell. Compared with the model group, the expression of Aβ1–42 in EA group was significantly weakened, and there were a few diffuse senile plaques (see Figure 3).

5.2.2. Laser Confocal Imaging of Aβ1–42 and LRP1 in Hippocampus

In the normal control group, plaque deposits of Aβ1–42 were not found, while LRP1 was expressed mostly around the vascular endothelial cells. In the model group, there were dense-core plaques deposited, while less LRP1 was expressed. In EA group, the senile plaques were relatively reduced, and there were only some diffuse plaques, while the expression of LRP1 was more than that of model group. Aβ1–42, LRP1, and cell nucleus were labeled with laser confocal imaging (see Figure 4).

5.2.3. Effect of EA on Aβ1–42 and Aβ1–40 Expression.

The ELISA results were shown in Table 3. LSD test was used for pairwise comparison. Results showed that Aβ1–42 and Aβ1–40 expression levels and Aβ1–42/Aβ1–40 values in the model group were all higher than those of the normal control group (P < 0.01); Aβ1–42 and Aβ1–40 expression levels in the EA group were lower than those in the model group (P < 0.01), while there was no significant difference for Aβ1–42/Aβ1–40 between the EA and model groups.

5.2.4. Effect of EA on Relative Expression Level of LRP1.

Western blotting results of LRP1 were shown in Table 4 and Figure 5. The LRP1/β-actin gray value was the relative expression level of LRP1.

LRPI/Actin. There was a significant difference between the model group and the normal control group (P < 0.01), and the EA group was significantly higher than the model group (P < 0.01) (see Table 4).

Table 3: Effect of EA on Aβ1–42 and Aβ1–40 expression (x ± s, pg/mg, n = 6).

| Groups                  | Cases | Aβ1–42 | Aβ1–40 | Aβ1–42/Aβ1–40 |
|-------------------------|-------|--------|--------|---------------|
| Normal control group (C)| 6     | 0.14 ± 0.38 | 0.077 ± 0.01 | 1.79 ± 0.61   |
| Model group (M)         | 6     | 6119.76 ± 670.13** | 801.05 ± 219.24** | 7.97 ± 1.61** |
| EA group (EA)           | 6     | 1326.58 ± 501.40** | 297.05 ± 112.89** | 7.02 ± 1.73   |

Notes: ** compared with the normal control group, P < 0.01; *** compared with the model group, P < 0.01.

Table 4: Effect of EA on relative expression level of LRP1 (x ± s, n = 6).

| Groups                  | Cases | LRPI gray value (x10^4) | Actin gray value (x10^4) | LRP1/Actin |
|-------------------------|-------|------------------------|--------------------------|------------|
| Normal control group (C)| 6     | 6.77 ± 1.17            | 11.16 ± 0.86             | 0.60 ± 0.08 |
| Model group (M)         | 6     | 3.87 ± 0.76**          | 11.77 ± 1.24             | 0.33 ± 0.07** |
| EA group (EA)           | 6     | 6.34 ± 1.32**          | 11.86 ± 0.74             | 0.53 ± 0.10** |

Notes: ** compared with the normal control group, P < 0.01; *** compared with the model group, P < 0.01.

5.2. Effect of EA on Aβ and LRP1

5.2.1. Immunohistochemistry Results of Aβ1–42 in Hippocampus.

In the normal control group, there were brown positive expressions of Aβ1–42 inside the cell and negative ones outside the cell. In the model group, there were positive expressions of Aβ1–42 and plaque deposits with compactness outside the cell. Compared with the model group, the expression of Aβ1–42 in EA group was significantly weakened, and there were a few diffuse senile plaques (see Figure 3).

5.2.2. Laser Confocal Imaging of Aβ1–42 and LRP1 in Hippocampus.

In the normal control group, plaque deposits of Aβ1–42 were not found, while LRP1 was expressed mostly around the vascular endothelial cells. In the model group, there were dense-core plaques deposited, while less LRP1 was expressed. In EA group, the senile plaques were relatively reduced, and there were only some diffuse plaques, while the expression of LRP1 was more than that of model group. Aβ1–42, LRP1, and cell nucleus were labeled with laser confocal imaging (see Figure 4).

5.2.3. Effect of EA on Aβ1–40 and Aβ1–42 Expression.

The ELISA results were shown in Table 3. LSD test was used for pairwise comparison. Results showed that Aβ1–42 and Aβ1–40 expression levels and Aβ1–42/Aβ1–40 values in the model group were all higher than those of the normal control group (P < 0.01); Aβ1–42 and Aβ1–40 expression levels in the EA group were lower than those in the model group (P < 0.01), while there was no significant difference for Aβ1–42/Aβ1–40 between the EA and model groups.

5.2.4. Effect of EA on Relative Expression Level of LRP1.

Western blotting results of LRP1 were shown in Table 4 and Figure 5. The LRP1/β-actin gray value was the relative expression level of LRP1.

LRPI/Actin. There was a significant difference between the model group and the normal control group (P < 0.01), and the EA group was significantly higher than the model group (P < 0.01) (see Table 4).

Table 5: Effect of EA on LRP1 expression (x ± s, pg/mg, n = 9).

| Groups                  | Cases | LRP1 |
|-------------------------|-------|------|
| Normal control group (C)| 9     | 0.21 ± 0.25 |
| Model group (M)         | 9     | 0.17 ± 0.1** |
| EA group (EA)           | 9     | 0.19 ± 0.11 |

Notes: ** compared with the normal control group, P < 0.01.

6. Discussions

The disordered clearance of Aβ is the main pathogenesis of AD. Aβ is cleared out through the BBB or degraded through the pathway of brain cell fluid drainage [31, 32]. The Aβ level between the brain parenchyma and brain interstitial fluid is inversely proportional to the age of mice. For the older mice, the Aβ level of brain parenchyma increases, while the Aβ level of brain interstitial fluid decreases [33]. The experimental study has found that LRP1 can transport Aβ out from BBB to be involved in the clearance of Aβ vessels [34], and the Aβ level can be decreased by transporting it.

In 1997, Narita et al. [35] discovered that there was a correlation between LRP1 and AD. However, studies on the pathogenesis of AD have shown that the functions of LRP1 are complicated, and some results are contradictory. On one hand, LRP1 can internalize APP and decompose it as Aβ in organelle lumen; on the other hand, LRP1 on neuronal membrane can also perform enzymatic degradation through the endocytosis of Aβ with its ligand a2M and ApoE; meanwhile, LRP1 on brain microvascular endothelial cells may mediate the outflow transport of Aβ across the BBB. Therefore, LRP1 is correlated to the generation and clearance of Aβ in some degree [36–38]. Studies have indicated that, for the AD model, the transportation of Aβ across the BBB
mediated by LRP1 in the brain is damaged [39, 40]. The inhibition of LRP1 expression in brain of healthy mice could decrease the outflow rate of Aβ in brain by 30% [41] and prompt the sedimentation of Aβ in brain.

Studies have shown that, by injecting iodine-labeled Aβ into the brain, the rapid transport process of Aβ across the BBB can be observed [42]. This process can be inhibited by LRP1’s sensitive inhibitor (RAP) and α2 macroglobulin. All these evidences prove that LRP1 on the BBB may be the main carrier of Aβ transported out from the brain; that is, LRP1 is involved in the clearance of Aβ, and it also has a negligible effect on the prevention of AD symptoms.

Experiments have shown that electroacupuncture can improve learning and memory abilities in transgenic mice and decrease the Aβ level in brain. The mechanism may be related to the effect of electricity on the cerebral microvessels and the Aβ transport receptor LRP1 [13, 18, 43].

Our results show that, in the water maze escape latency, mice in the normal control group and the EA group had separately retained the spatial memory in the 2nd and 3rd days, and the escape latency time shortened with the increase in days. It indicates that EA has improved the AD models’ ability of spatial learning and memory. ELISA testing displayed that the levels of Aβ were significantly decreased, which shows a consistency with the results of immunohistochemistry and laser confocal imaging. This may be a mechanism of EAs improvement of the memory of AD mice.

The AD animal models are APPswe/PSIdE9 double transgenic mice prepared by transferring the human mutated genes APP and PS1 into mice to raise the Aβ levels in brain so as to cause a series of pathological lesions of AD. This model assumes that Aβ is the main pathological pathogenic factor. It was reported previously that the escape latency time in spatial probe test increased for 3-month-old APPswe/PSIdE9 double transgenic mice [44], while in the preliminary study of this experiment, it was found that the Morris water maze escape latency time in EA group and model group was not significantly different from that in normal control group for 5-month-old double transgenic mice. The water maze experiments of Li et al. [26] have indicated that the learning and memory disorders begin to appear at the age of 7-8 months.

This study has shown that EA can improve the spatial learning and memory of 7-month-old APP/PS1 transgenic mice in the model group which have been shown to have both learning and memory disorders.
Evidence-Based Complementary and Alternative Medicine

Figure 4: Laser confocal imaging of $\text{A}^\beta_{1-42}$ and LRP1 in hippocampus (original magnification $\times 600$): (A) stack imaging of the three; (B) imaging of cell nucleus; (C) LRP1 labeled with green fluorescence; (D) $\text{A}^\beta_{1-42}$ labeled with red fluorescence; (a) in normal control group, plaque deposits of $\text{A}^\beta_{1-42}$ were not found, while LRP1 was expressed mostly around the vascular endothelial cells; (b) in model group, there were dense-core plaques deposited, while less LRP1 was expressed; (c) in EA group, the senile plaques were relatively reduced, and there were only some diffuse plaques, while the expression of LRP1 was more than that of model group.

Our research group has previously done a study of EA on GV 20 (bài hui) and KI 1 (yòng quán) as the intervention to APPV717I AD mice. EA could intervene in the behavior of 11-month-old APP single transgenic AD mice, neuronal changes, and $\text{A}^\beta$ protein expression in brain, especially where there is a decreasing trend of $\text{A}^\beta$ sedimentation in the hippocampus microvessels [43]. EA therapy can regulate the APP/PS1 double transgenic mice hippocampus ultrastructure [45]. APP/PS1 double transgenic mice were used in this study which could cause earlier pathological manifestations such as $\text{A}^\beta$ sedimentation, senile plaques, neuronal loss, and behavioral obstacles of cognitive performance compared
to single-transgenic mice, so that the length of the EA experiment cycle can be shortened. The appearance of senile plaques on brain cortex for 7-month-old APP/PS1 transgenic mice has indicated an early intervention of EA.

The focus of this study is on the effect of EA on LRP1, which is the key receptor in Aβ clearance of vessels in brains of the models. LRP1 mainly exists in cerebral vascular endothelial cells, vascular smooth muscle cells, and other cells including neurons, astrocytes, and smooth muscle cells [46]. In this study, the laser scanning confocal microscope was applied to observe the coexpression of LRP1 and Aβ in the hippocampus, proving that LRP1 is expressed on the microvascular endothelial cells.

This experiment showed that the Aβ levels in the model group increased, while EA can decrease Aβ levels. Furthermore, LRP1 levels in the model group decreased, while EA increased LRP1 levels. Therefore, the mechanism of EA’s involvement in improving learning and memory may be its ability to increase LRP1 levels, thus increasing the clearance of Aβ.

It has been reported that Aβ can be directly transported by LRP1, and studies have also shown that Aβ can be only transported after forming complexes with other ligands of LRP1 such as ApoE [47]. Does EA regulate another mechanism of ApoE and so forth by increasing the rate and quantity of combination of Aβ and LRP1 and further decrease Aβ? Further studies of possible mechanisms are required.

In summary, this study has indicated that EA can improve learning and memory of mice. Aβ levels were shown to be significantly lower in the EA group compared to the model group, while LRP1 levels were also significantly higher in the EA group, which may indicate that the decreasing of Aβ is related to the transportation of LRP1; therefore EA may be involved in regulating LRP1 functions. Enhancing the transportation of Aβ protein and decreasing the Aβ levels may be a functional way to treat dementia using EA, but still further evidence is needed.

7. Conclusions

EA therapy can improve both learning and memory capabilities in APP/PS1 transgenic mice. The underlying mechanism may be due to the upregulation of the Aβ transport receptor LRP1, thus acting on learning and memory by contributing to decreased levels of Aβ in the hippocampus.

Ethical Approval

All procedures performed in studies involving animals were in accordance with the “Principles of Laboratory Animal Care” formulated by the National Institute of Health and the legislation of the People’s Republic of China for the use and care of laboratory animals.

Consent

Informed consent was obtained from all individual participants included in the study.

Competing Interests

The authors declare that they have no competing interests.

Authors’ Contributions

Xin Wang and Yanhuan Miao contributed equally to this work.

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