Kai-Xin-San, a traditional Chinese medicine formulation, exerts antidepressive and neuroprotective effects by promoting pCREB upstream pathways

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Abstract. Kai-Xin-San (KXS) is a traditional Chinese medicine that has been widely used for the treatment of emotion-related disease. However, the underlying mechanism remains largely unknown. The present study aimed to examine whether phospho-cAMP response element-binding protein (pCREB) and upstream components, such as extracellular signal-regulated kinase (ERK), phospho-ERK (pERK), phosphatidylinositol-3-kinase (PI3K), protein kinase B (Akt), glycogen synthase kinase 3β (GSK3β) and pGSK3β are associated with the antidepressive effect of KXS. In total, 24 male Wistar rats were randomly divided into three groups, including control (n=8, no treatment), induced with chronic unpredictable mild stress (CMS) (n=8), and CMS rats treated with KXS at dosage of 370 mg/kg/day orally. Primary hippocampal neuronal cultures were prepared from Wistar rats for cell survival and proliferation assays. In KXS rats, increased protein expression levels of pCREB, BDNF and tyrosine receptor kinase B (TrkB) were observed in the hippocampus and prefrontal cortex, compared with the CMS model group. Furthermore, increased expression levels of ERK, pERK, PI3K, Akt, and GSK3β were also detected in the hippocampus and prefrontal cortex of KXS-treated rats compared with CMS model rats and in primary hippocampal neuronal cells treated with KXS. These results suggest that pCREB and upstream components, including TrkB/ERK/CREB and TrkB/PI3K/CREB, may contribute to the antidepressive effect induced by KXS. Further studies are required to confirm these findings.

Introduction

It has previously been demonstrated that brain-derived neurotrophic factor (BDNF), which is a neurotrophic factor, has a key role in depression. In depressive patients with serum reduced levels of BDNF, BDNF increased after treatment with antidepressant agents (1,2). A postmortem study reported a significant decrease in BDNF levels in the hippocampi of depression-induced suicide subjects, but not in suicide subjects treated with antidepressants (3). Further studies have demonstrated that downstream components of the BDNF/tyrosine receptor kinase B (TrkB) signaling pathway, such as the phosphatidylinositol-3-kinase (PI3K)/phosphatidylinositol-3-kinase (Akt) pathway, were inhibited in depressive patients (3-6). The phospho-cAMP response element-binding protein (pCREB) may regulate the transcription of downstream genes, including BDNF, potentially by binding to response elements in the promoter regions of gene transcription, thus regulating neurotrophin, differentiation, cell viability and recognition functions (7-10).

Kai-Xin-San (KXS) is a well-known traditional Chinese medicine first recorded in Bei Ji Qian Jin Yao Fang (Thousand Formulae for Emergency Thousand Formulæ for Emergency) by Sun Si-Miao (11,12). KXS consists of ginseng (Panax ginseng C.A. Meyer), hoelen (Poria cocos F.A. Wolf), polygala (Polygala tenuifolia Willd) and acorus (Acorus tatarinowii Schott) at the ratio of 3:3:2:2 (13). It has been widely used in the treatment of depression, as well as in learning and memory deficits for thousands years (14). In our previous studies, we
have demonstrated the anti-depressive effect of KXS in rats by utilizing chronic unpredictable mild stress (CMS), the tail suspension test, the forced swim test and chronic fatigue syndrome; in addition, increased levels of BDNF and pCREB were detected, suggesting that they may have roles in the anti-depressive effect of KXS (15-18). The present study aimed to explore the mechanism of KXS via the regulation of the BDNF and pCREB pathways in rats and neuronal cells.

Materials and methods

KXS preparation. Herbs for KXS (ginseng, hoelen, polygala, and acorus) were purchased from LvYe Medicinal Material Company (Beijing, China), and the quality met the requirements outlined in the Chinese Pharmacopoeia (2010) (19). The voucher specimen was registered and deposited in the Herbarium of the Traditional Chinese Medicinal pharmacy. KXS was supplied in powder form derived from a mixture of the aqueous extract as described previously (20). Briefly, 1.5 kg ginseng, 1.5 kg hoelen, 1 kg polygala and 1 kg acorus gramineus was soaked in 50 l water for 3 h and was subsequently extracted three times using a circumfluence extraction method. The aqueous extracts were filtered and evaporated under reduced pressure. The obtained concentrates were freeze-dried to powder form and stored at 4℃. Average yield was 1 g KXS powder/4.83 g total herbs. KXS powder was standardized using a high-performance liquid chromatography-fingerprint method, as previously described (21).

Animals. In total, 24 male Wistar rats, weighing 180±10 g at 6 weeks, were obtained from the Animal Breeding Center of the People’s Liberation Army general hospital (Beijing, China). All rats were housed in a temperature- (23±2℃) and humidity-controlled (60±10%) facility with a 12-h light/dark cycle with free access to food and water. All animal experimental protocols were approved by the Animal Experimentation Ethics Committee of General Hospital of the Chinese People’s Liberation Army (Beijing, China). All procedures were performed in accordance with the Principles of Laboratory Animal Care and Chinese legislation for the use and care of laboratory animals.

Induction of CMS. CMS was induced in 16 rats following the established protocol (22). Rats received 4 weeks of stress stimulation, which included high-speed agitation (10 min), deprivation of food or water (24 h), immobilization (2 h), continuous illumination (24 h), tilted cage (12 h) and forced swimming in ice water (5 min). Rats were randomly assigned one stimulation daily at 3-5 p.m. for four weeks. The remaining eight rats were housed undisturbed without contact with the stressed animals as the control group.

Drug administration. The 16 CMS-induced rats were randomly divided into two groups treated with either water (n=8; CMS group) or KXS at 370 mg/kg orally at 9-10 am daily for three weeks (n=8; KXS group).

Sucrose-preference test. Sucrose-preference test was performed to define anhedonia prior to surgery (baseline) and after CMS induction. Rats were food- and water-deprived for 18 h, and subsequently fed with pre-weighted bottles containing 1% sucrose solution and water for 1 h. Intake was measured by weighing the bottles before and after each test. All tests were carried out in the home cage to minimize extraneous novelty and disturbance. Sucrose preference was calculated as: Sucrose preference = sucrose intake / (sucrose intake + water intake). Anhedonia was defined as a reduction in sucrose preference relative to baseline levels.

Open-field test. Locomotor activity was assessed to detect immobility or changes in motor activity in an open-field test. A metallic cubic open field arena (80x80x80 cm) was used. The floor of the box was 25 squares (5 squares long; 5 squares wide). Rats were individually placed into the center of the arena and allowed to explore freely for 3 min. The floor of the open-field apparatus was wiped cleaned with 70% ethanol between tests. The behavior, including the number of square lines crossed with all four paws, and rearing, including the number of times each rat stood on its hind limbs, were recorded.

Western blot. At the end of experiment, all rats were sacrificed with 10% chloral hydrate solution (3.5 ml/kg; i.p.). Hippocampus and prefrontal cortex tissues were collected and lysed with radioimmunoprecipitation assay buffer (Applygen Technologies, Inc., Beijing, China), schizolysised for 20 min on ice and centrifuged at 12,000 x g for 10 min at 4℃. Protein samples were heated at 95℃ for 8 min, separated by 12% SDS-PAGE and transferred to nitrocellulose membranes. Membrane were blocked for 2 h in TBST (25 mM Tris, 140 mM NaCl, 27 mM KCl and 0.02% Tween 20) containing 5% bovine serum albumin and incubated with primary antibodies specific for BDNF (ab108319, 1:200)(Abcam, Cambridge, MA, USA), TrkB (BS1431, 1:500), ERK (AP0491, 1:500), pERK (BS4621, 1:500), Akt (BS1502, 1:500), P38 (BS3678, 1:500), pGSK3β (BS4084, 1:500) and GSK3β (BS1402, 1:500; Bioworld Technology, Inc., Danvers, MA, USA) at 4℃ over night. Following three washes with TBST, membranes were incubated for 1 h at room temperature with horseradish peroxidase-labeled secondary antibodies (BS13278, 1:5,000; Bioworld Technology, Inc., Danvers, MA, USA), washed with TBST three times. Blots were developed using an electrochemiluminescence system (UVP LLC, Upland, CA, USA).

Primary hippocampal neurons culture. Primary hippocampal neuronal cultures were prepared as described previously (23), with a few modifications. A total of 36 Newborn Wistar rats (age, <24 h) were purchased from the Peking University Health Science Center (Beijing, China). Hippocampi were dissected on ice, minced and digested with 0.25% trypsin for 15 min at 37℃. Subsequently, the samples were supplemented with 5 ml fetal bovine serum (FBS) to terminate the digestion procedure. Cell suspension was passed through a 200 mesh (diameter, 74 μm) cell strainer and separated by density gradient centrifugation for 20 min at 300 x g at room temperature. The suspension was centrifuged for 5 min at 225 x g at room temperature and the pellet was resuspended in 10 ml Dulbecco’s modified Eagle’s medium (DMEM). Following the measurement of cell density using a hemacytometer, the
neuronal cells were plated into 6-well plates at 5x10^5 cell/ml for culture in DMEM supplemented with 10% FBS, 1 mmol/l glutamine, 20 mmol/l sodium pyruvate, and 10 ng/ml nerve growth factor (Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA). 200 nM K252a (Sigma Aldrich; Merck Millipore, Darmstadt, Germany) was added to cells 30 min before the exposure to KXS as an inhibitor of TrkB. Cells were used for assay analysis at day 7 when the percentage was ~85%, as assessed by microtubule-associated protein 2 (MAP-2) immunostaining.

Cell survival and proliferation assays. Cell viability was evaluated via MTT assay (24). Briefly, 20 µl MTT solution (2 mg/ml in PBS) was added at a final concentration of 0.5 mg/ml and incubated at 37°C for 4 h. Following removal of the medium, 150 µl DMSO was added. Optical densities were subsequently read using a microplate reader at 570 nm (1420 Vitor 3; Perkin-Elmer, Waltham, MA, USA). Viability was expressed as a percentage of the KXS-treated cells according to the control cells.

Statistical analysis. All data were expressed as the mean ± standard error. Differences between groups were analyzed by one-way analysis of variance followed by Dunnett’s test. Statistically analyses were performed using SPSS 17.0 software (SPSS, Inc., Chicago, IL, USA). P<0.05 was considered to indicate a statistically significant difference.

Results

Behavioral index. Table I shows the results from the open-field and sucrose-preference tests. The index of sucrose preference in the CMS group was significantly lower than that of the control group (P<0.01) and the KXS group (P<0.05). Similarly, the number of lines crossed and rearing values in the CMS group were significantly lower than that of the control group (P<0.01, P<0.05), suggested that KXS could improve the depressive symptom in rats induced by CMS.

Protein expression levels of BDNF, CREB and TrkB. Western blot analysis demonstrated that the protein expression levels of BDNF, pCREB and TrkB in the hippocampus and TrkB in the prefrontal cortex were significantly reduced in the CMS group compared with the control group (P<0.05). Furthermore, the protein expression levels of BDNF, pCREB and TrkB in the hippocampus and prefrontal cortex were increased in the KXS group compared with the CMS group (P<0.01; Fig. 1), suggested that the antidepressive effect of KXS might be involved in its regulating on BDNF and TrkB through promoting the pCREB level.
Protein expression levels of pERK/ERK, PI3K, Akt and pGSK3β/GSK3β. Fig. 2 demonstrates that the protein expression levels of pERK/ERK (P<0.01), PI3K (hippocampus, P<0.01; prefrontal cortex, Pc<0.05), and Akt (hippocampus, P<0.01; prefrontal cortex, P<0.05) in the hippocampus and prefrontal cortex, and pGSK3β/GSK3β (P<0.05) in the hippocampus of the KXS group were significantly elevated, compared with the CMS group. The results suggest that the regulation effect of KXS on BDNF and pCREB might be related to its activation on the upstream signaling pathway of CREB.

Cell viability and BDNF expression. Fig. 3 shows MAP-2 immunostaining of neuronal cells. The morphous and purity of the neuronal cells was suitable for the experiment. In cells treated with KXS, cell viability was significantly increased, as compared with the control cells (P<0.01). The level of BDNF release increased in a time-dependent manner, peaking at 6 h post-treatment (Fig. 4). KXS could increase the nerve cell viability which might be involved in the increasing level of BDNF.

Protein expression levels of TrkB/BDNF/ERK in neuronal cells. Fig. 5 indicates that the protein expression levels of BDNF, pERK and pTrkB in the KXS group were significantly higher than that of the control group (P<0.01). Such an increase was not observed when cells were treated with K252a, which is an inhibitor of TrkB. The effects of KXS on the pathway components in cells coincided with those of animal experiments.

Discussion

In CMS rats treated with KXS, increased protein expression levels of pCREB, BDNF and TrkB were detected in the hippocampus and prefrontal cortex. Consistently, in primary
hippocampal neurons, an increased level of upstream components, including ERK, pERK, PI3K, Akt and GSK3β, was observed when cells were treated with KXS. These results suggest that pCREB and upstream components may be associated with the antidepressive effect of KXS.

Depression is a complex mental disorder involving multiple factors. Impaired release of BDNF, which is a small protein utilized by the brain for development and nerve cell growth regulation and function (25), has been demonstrated to have a key role in major depression (26). The present finding of increased BDNF levels in CMS rats treated with KXS is consistent with previous reports (22). In previous animal and human studies, circulating levels of BDNF were very low and could be increased with treatment of antidepressants (27-30), which may account for its efficacy in depressive behaviors (31,32).

It has been demonstrated that ERK and PI3K pathway are the two most important pathways altered through the BDNF mediated-TrkB activation, resulting in neuroprotection effect (33). ERKs are able to activate and transfer signals to the nucleus quickly, in order to induce CREB phosphorylation and activate the nuclear transcription factor, CREB. Enhanced CREB ultimately results in increased expression of BDNF (34). These data suggest a positive feedback loop between BDNF and CREB, which may be associated with nerve cell nutrition and nerve injury repair (35). In this study, increased pERK was observed in CMS rats treated with KXS; however, such an increase was not observed when primary

Figure 3. Morphological observation and identification of primary hippocampal neurons (magnification, x200) at (A) 4 h, (B) 24 h, (C) 2 days, (D) 3 days, (E) 5 days, (F) 7 days. Primary hippocampal neurons incubated with (G) DIPA and MAP-2 at (H) day 5 and (I) day 7.

Figure 4. Effect of KXS on cell survival and BDNF release from primary hippocampal neurons. (A) For cell viability assay, cells were treated with 25 pg/ml BDNF or 50, 100 and 200 µg/ml of KXS for 48 h. *P<0.01 vs. the control group. (B) In the BDNF release test, cultured cells were treated with 100 µg/ml KXS for 0, 3, 6, 12, 24 and 48 h. **P<0.01 vs. the 0 h group. Data are presented as the mean ± standard deviation. KXS, Kai-Xin-San; BDNF, brain-derived neurotrophic factor; OD, optical density.
neurons cells were treated with K252a, which is an inhibitor of TrkB. This suggests that the KXS-induced increase of BDNF may be mediated by the TrkB-dependent ERK pathway. Activated CREB may further regulate the transcription of genes involved in neuronal cell survival.

GSK-3β is an essential pro-apoptotic factor for neuronal the apoptosis cascade and PI3K-mediated neuronal survival pathway (36). In its active form, GSK-3β is able to enhance apoptosis; however, once inactivated by phosphorylation, it cannot trigger apoptotic pathways (37). In the present study, inactivated GSK-3β by phosphorylation and an PI3K/AKT-mediated increase in BDNF was observed when rats were treated with KXS. This finding is consistent with a previous report, demonstrating that BDNF may provide neuroprotection via the activation of the PI3K/Akt pathway (38).

In conclusion, increased protein expression levels of pCREB, BDNF and TrkB were detected in the hippocampus and prefrontal cortex. Consistently, in primary hippocampal neurons, increased levels of upstream components, including ERK, pERK, PI3K, Akt, and GSK3β, were observed when cells were treated with KXS. These results suggest that pCREB and its upstream components may be associated with the antidepressive effect of KXS. Subsequent studies in other cells and animal models are required to confirm the present findings and conclusions.

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