

Full-length article

Upregulation of Mark3 and Rpgrip1 mRNA expression by jujuboside A in mouse hippocampus¹

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Key words

jujuboside A; hippocampus; DD-PCR; Mark3; Rpgrip1

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Abstract

Aim: To investigate the effect of jujuboside A (JuA) on modulating gene expression in the hippocampus. **Methods:** The spontaneous activity of mice was monitored, and the differential display polymerase chain reaction was adapted to screen differentially-expressed genes modulated by JuA in the mouse hippocampus. **Results:** JuA significantly decreased the total activity intensity ($P < 0.01$ vs control) at a dosage of 80 mg/kg, and the genes MAP/microtubule affinity-regulating kinase3 (Mark3) and retinitis pigmentosa GTPase regulator interacting protein1 (Rpgrip1) were upregulated by JuA in the mouse hippocampus. **Conclusion:** JuA had an inhibitory effect on the spontaneous activity of the mice, and JuA regulated the transcription of Mark3 and Rpgrip1 in the mouse hippocampus.

Introduction

Jujuboside A (JuA) is a main effective component of jujubogenin, extracted from the seed of *Ziziphus jujuba* Mill var *spinosa* (Bunge) Hu ex H F Chou (*Ziziphus*), which is widely used in treating symptoms of insomnia and anxiety^[1,2]. However, most previously published studies have only focused on the investigation of behavioral changes. Recent experimental results have suggested that a high dose of JuA could inhibit the hyperactivity of the hippocampal CA1 area induced by penicillin sodium^[2]. JuA had inhibitory effects on the Glu-mediated excitatory signal pathway in the hippocampus and hippocampal formation *in vivo* and *in vitro*^[3,4]. Those studies suggested that JuA might play the role of an inhibitor on the central nervous system, especially on the hippocampus. However, they could not explain the functions of JuA on the hippocampus, and thus, the inhibitory effect of JuA on the hippocampus is still elusive.

In the present study, we adapted the protocol of the differential display polymerase chain reaction (DD-PCR) to screen for differentially-expressed genes modulated by JuA at the transcription level in the mouse hippocampus. We found genes modulated by JuA, and explained the partial

molecular mechanism of JuA on the hippocampus at the gene transcription level. The use of DD-PCR is universal in investigating molecular pharmacological mechanisms of the effective component of a Chinese traditional herb at the gene transcription level.

Materials and methods

Chemicals and animals JuA was provided by the National Institute for the Control of Pharmaceutical and Biological Products in China (Beijing, China) with a purity above 98%. KM (Kunming strain) mice (male, Grade II, weighing 18–22 g) were obtained from the Laboratory Animals Center of Sichuan Academy of Medical Sciences (Chengdu, Sichuan, China), and maintained in an air-conditioned room with controlled temperature (23–25 °C) and humidity (50%–70%). The mice were housed in cages under a 12 h light/dark cycle with access to food and water. All the animal experiments were carried out from 19:00 PM to 23:00 PM. The animal experiments were performed with institutional ethical approval of protocol, and all efforts were made to minimize animal suffering. The mice were randomly divided between the control and the treated groups. The treated group was given an

ip injection of JuA at 80 mg/kg, and the control group was given an equal volume of saline. Half an hour after the injection, the mice in the control ($n=10$) and treated groups ($n=10$) were observed for spontaneous activity. Meanwhile, the control group ($n=6$) and the treated group ($n=6$) were used for RNA extraction.

Spontaneous activity The mice were put in a ZZ-6 monitor (Chengdu Technology and Market Co, Ltd, Chengdu, Sichuan, China) for observation of spontaneous activity and adapted to the environment within 5 min. We then recorded the number of mouse movements within a 10 min period. After recording the number of mouse movement before any treatment, the mice were taken out and injected with JuA or saline and housed in the cages. Half an hour after the injection of JuA or saline, the frequency of mouse movement was recorded again using the same method.

RNA extraction All the animals were sacrificed by decapitation and the hippocampi were removed immediately. The total RNA of the hippocampus was isolated with Trizol (Molecular Research Center Inc, Cincinnati, OH, USA) according to the manufacturer's protocol. To remove DNA contamination, the RNA samples were treated with RQ1 RNase-free DNase (Promega, Madison, WI, USA) and extracted with phenol:chloroform. Finally, the RNA samples were resuspended in 50 μ L of nuclease-free water. The concentrations of the total RNA were measured with a biophotometer, and the OD_{260}/OD_{280} ratio of all RNA samples were up to 2.0.

DD-PCR The first strand cDNA was synthesized using the RevertAid™ First Strand cDNA Synthesis Kit (MBI Fermentas, Burlington, Ontario, Canada). For each reaction, 4 μ g of deoxyribonuclease-treated total RNA was used for the reverse transcription, in a mixture of 2 μ g 3'-anchored primers ($T_{15}A$, $T_{15}C$, and $T_{15}G$, respectively) in a final volume of 12 μ L, and heated at 70 °C for 5 min. Then 4 μ L of 5 \times reverse transcription buffer, 1 μ L of 20 U/ μ L, Ribolock™ Ribonuclease inhibitor (MBI Fermentas, Burlington, Ontario, Canada), and 2 μ L of 10 mmol/L dNTP(dATP, dCTP, dGTP, dTTP) mix were added; the samples were incubated at 37 °C for 5 min. Then 1 μ L (200 units) of RevertAid™ M-MuLV reverse transcriptase was added to each sample (with a final volume of 20 μ L). The mixture was incubated at 42 °C for 60 min, and finally heated at 70 °C for 10 min.

PCR was performed in 25 μ L reaction mixtures containing 2 μ L cDNA, 2 μ mol/L 3' anchored primers and 5' arbitrary primers (Table 1), 200 μ mol/L dNTP, and 2 units of *Taq* DNA polymerase, with an initial denaturation at 94 °C for 3 min, followed by 4 cycles at 94 °C for 1 min, 34 °C for 4 min, and 72 °C for 1.5 min, and 40 cycles at 94 °C for 1 min, 38 °C for

Table 1. Anchor and arbitrary primers of DD-PCR

Anchored primers	Arbitrary primers
M1: $T_{15}A$	S1: GATCAAGTCC
M2: $T_{15}C$	S2: TCGATACAGG
M3: $T_{15}G$	S3: GTTTTCGCAG
	S4: CTGCTTGATG
	S5: GCCATGCACG
	S6: GGAAGCAGCT

2 min, and 72 °C for 1.5 min, with a final extension at 72 °C for 10 min. After PCR amplification, 10 μ L of PCR products were separated by electrophoresis on 8% denaturing polyacrylamide gels, and the gels were stained by silver^[5].

Re-amplification, ligation, and transformation of PCR products and sequencing Differentially-expressed bands were retrieved from the polyacrylamide gels^[6] and reamplified using the corresponding primer sets in 50 μ L reaction mixtures containing 5 μ L recycled DNA, and 1 μ mol/L primers, respectively, 400 μ mol/L dNTP and 5 units of *Taq* DNA polymerase, with an initial denaturation at 94 °C for 3 min, followed by 40 cycles at 94 °C for 1 min, 38 °C for 2 min, and 72 °C for 1.5 min, with a final extension at 72 °C for 10 min.

The reamplified PCR fragments were run on a 1% agarose gel and gel-purified using 3S Spin Agarose Gel DNA Purification Kit (Shenergy Biocolor, Shanghai, China) according to the kit protocol. Products were ligated into the pUCm-T cloning vector (Shenergy Biocolor, China) according to the kit protocol, then transfected into *Escherichia coli* high efficiency DH5 α competent cells^[7] with ampicillin and blue/white selection. The positive colonies were screened by PCR with a pair of M13 primers and digestion with restriction endonuclease *Pst* I. Those colonies were sequenced by a commercial sequencing service (Beijing Sunbiotech Co, Beijing, China) and the DNA sequence was analyzed with BLASTN (Bethesda, MD, USA; NCBI, National Center for Biotechnology Information).

Semi-quantitative RT-PCR The first strand cDNA was synthesized as protocol of DD-PCR except for 3'-anchored primers replaced by the oligo(dT)₁₈ primer. PCR was performed in 25 μ L reaction mixtures containing 1.5 μ L cDNA, 2 μ mol/L gene-specific forward and reverse primers (Table 2), 200 μ mol/L dNTP, and 2 units of *Taq* DNA polymerase, with initial denaturation at 94 °C for 3 min, followed by 94 °C for 1 min, 55 °C for 1 min, and 72 °C for 1 min for the number of cycles optimized for each primer pair (Table 2) to ensure that the product intensity fell within the linear phase of amplification; final extension was at 72 °C for 10 min. RT-

Table 2. Nucleotide sequence of the forward and reverse primers, lengths of PCR products, and number of cycles performed for RT-PCR.

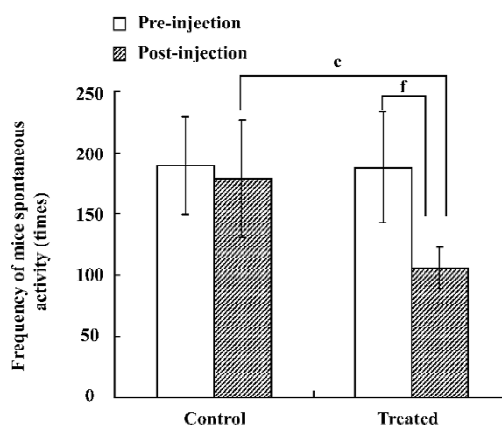
Gene name	Gene identity primers	PCR products	Cycles
β -actin	Forward: 5'ACTGTGCCCATCTACGAG3' Reverse: 5'CAGGATTCCATACCCAAG3'	332 bp	23
Mark3	Forward: 5'CACCTGGAGCATGAAGAC3' Reverse: 5'AGGGCAATGGAAACAAGAC3'	508 bp	26
Rpgrip1	Forward: 5'ACCTGGTCTCCTGTGCTAT3' Reverse: 5'TGCTGCTCTTCCCTTACT3'	409 bp	32

PCR amplification of β -actin transcript was used as the internal control to verify that equal amounts of RNA were used in each reaction. The PCR products were run on a 1% agarose gel, then photographed with a digital camera under UV illumination and analyzed by Quantity One software (Bio-Rad, Hercules, CA, USA).

Statistical analysis All data were expressed as mean \pm SD. A statistical analysis of the results was carried out by independent samples *t*-test and paired samples *t*-test. $P < 0.05$ was considered significant.

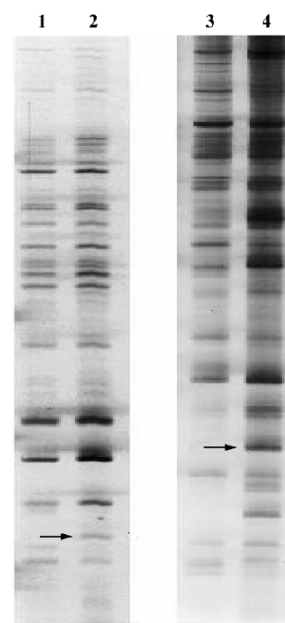
Results

Analysis of spontaneous activity of mice The number of mouse movements in the control and treated groups before any treatment was 189.4 ± 40.4 and 187.4 ± 45.0 , respectively. It was shown that there was no difference between the control and treated groups before the injection of saline and JuA ($P > 0.05$). Half an hour after the injection of saline and JuA, the number of mouse movements in the control and treated groups was 178.9 ± 47.5 and 105.7 ± 16.8 , respectively. The analysis of independent samples *t*-test between the control and treated groups after the injection of saline and JuA suggested that JuA significantly decreased the total activity intensity of the mice at a dosage of 80 mg/kg. The analysis of paired samples *t*-test between the preinjection and postinjection of JuA showed that JuA also decreased the total activity intensity of the mice significantly at the same dosage. Saline did not affect the number of mouse movements significantly (Figure 1).

**Figure 1.** Inhibitory effect of JuA on the spontaneous activity of mice. $n=10$. Mean \pm SD. ^c $P < 0.01$ vs control. ^f $P < 0.01$ vs preinjection of JuA.

Analysis of differential expression in response to JuA by DD-PCR and confirmed by semi-quantitative RT-PCR

To screen the differentially-expressed genes related to JuA in the mouse hippocampus, DD-PCR was performed (Figure 2). Following the isolation and sequencing of bands differentially-expressed between the control and treated groups and confirmed by semi-quantitative RT-PCR (Figure 3), we identified MAP/microtubule affinity-regulating kinase 3

**Figure 2.** Differentially-expressed genes identified by the mRNA DD-PCR. Lanes 1 and 3 are the control group, lanes 2 and 4 are the treated group. The differentially-expressed genes are signed (lane2 is Mark3, lane4 is Rpgrip1).

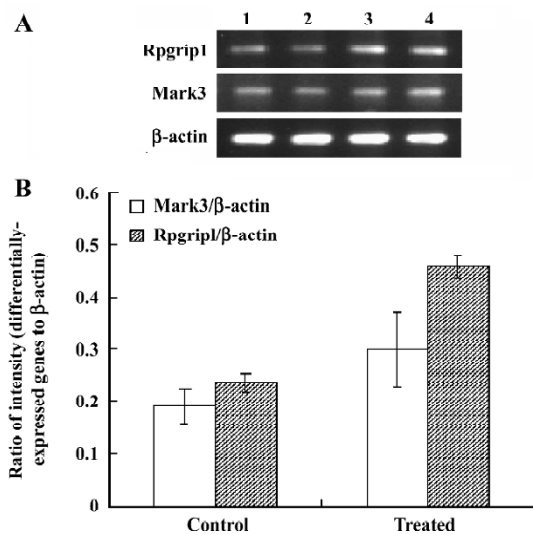


Figure 3. Analysis of semi-quantitative RT-PCR. (A) lanes 1 and 2 are the control group, lanes 3 and 4 are the treated group. (B) ratio of intensity of Mark3 and Rpgrip1 to β-actin. Mean±SD. n=2. Mark3 and Rpgrip1 gene expression was upregulated by JuA.

(Mark3) and retinitis pigmentosa GTPase regulator interacting protein 1 (Rpgrip1) as upregulated genes by JuA in the mouse hippocampus.

Discussion

We found that JuA, a component extracted from a Chinese traditional herb, significantly decreased total activity intensity in the mice. Our results were consistent with Feng et al^[8], who confirmed that JuA had an inhibitory effect on the spontaneous activity of mice and increased quiet state time.

Why does JuA inhibit the spontaneous activity of mice? Although recent studies have suggested that JuA could inhibit the hyperactivity of the hippocampal CA1 area induced by penicillin sodium, with inhibitory effects on the Glu-mediated excitatory signal pathway in hippocampus^[2,3], those studies could not explain why JuA had an inhibitory effect on the spontaneous activity of mice. In the present study, to elucidate the inhibitory effect of JuA on mice at the gene transcription level, we adapted the protocol of DD-PCR invented by Liang and Pardee^[9] for detecting differentially-expressed genes in the mouse hippocampus, and the differentially-expressed genes were confirmed by semi-quantitative RT-PCR. Finally, we identified Mark3 and Rpgrip1 as upregulated genes by JuA in the mouse hippocampus.

Mark3 is also known as C-TAK1 (Cdc25C-associated kinase 1) or PARI1^[10,11], and has been implicated in cell cycle

regulation and Ras signaling through its interactions with 2 putative substrates: the Cdc25C (cell division cycle 25C) phosphatase and the MAPK scaffold KSR1 (Kinase Suppressor of Ras 1)^[10,12]. The upregulation of Mark3 mRNA by JuA may influence cell cycle or the Ras signaling pathway indirectly. Rpgrip1, another differentially-expressed gene, is predominantly expressed and investigated in the retina^[13-15]. It is also expressed in other regions including the heart, spleen, and brain^[16], but has been poorly investigated in those regions. In the present study, we found that JuA could upregulate Rpgrip1 in the mouse hippocampus, which implies that Rpgrip1 plays an essential role in the hippocampus.

In conclusion, JuA evidently had inhibitory effects on the spontaneous activity of mice; meanwhile, Mark3 and Rpgrip1 were upregulated by JuA in the mouse hippocampus. The relevance of JuA, the reduction of mice spontaneous activity, and the upregulation of Mark3 and Rpgrip1 exist in 2 possible pathways. One pathway is that JuA can upregulate Mark3 and Rpgrip1 directly or indirectly. The upregulation of Mark3 and Rpgrip1, as well as other reasons, can inhibit the spontaneous activity of mice. The other pathway is that JuA can inhibit the spontaneous activity of mice through an unknown way, and the inhibition on the spontaneous activity of mice could induce upregulation of Mark3 and Rpgrip1. However, although there are no studies to our knowledge which suggest the relevance between spontaneous activity and Mark3 or Rpgrip1, the two possible pathways suggest that there might be an essential link between the reduction of the spontaneous activity of mice and the upregulation of Mark3 and Rpgrip1.

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