Agmatine pretreatment attenuates retinal ganglion cell death by an oxidative stress

IIZUKA YOKO

Department of Medical Science
The Graduate School, Yonsei University
Agmatine pretreatment attenuates retinal ganglion cell death by an oxidative stress

Directed by Professor Yang Woo Ick

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IIZUKA YOKO

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This certifies that the Doctoral Dissertation of IIZUKA YOKO is approved.

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Thesis Supervisor: Yang Woo Ick
------------------------------------
Thesis Committee Member#1: Seong Gong Je
------------------------------------
Thesis Committee Member#2: Lee Jong Eun
------------------------------------
Thesis Committee Member#3: Kim Chan Yun
------------------------------------
Thesis Committee Member#4: Kim Chul Hoon

The Graduate School
Yonsei University

December 2008
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<ABSTRACT>

Agmatine pretreatment attenuates retinal ganglion cell death by an oxidative stress

IIZUKA YOKO

Department of Medical Science
The Graduate School, Yonsei University

(Directed by Professor Yang Woo Ick)

Purpose; To investigate the protective effects of agmatine pretreatment on oxidative stress in immortalized and differentiated rat retinal ganglion cell line (RGC-5 cells).

Method; RGC-5 cells were differentiated by staurosporine, cultured in the presence of hydrogen peroxide (H₂O₂) with or without pretreatment of agmatine. Cell viability was determined by lactate dehydrogenase (LDH) assay. Apoptosis was examined by TUNEL assay and caspase-3 and -9 assays. Total and phosphorylated Bcl-2 family proteins (Bcl-2, Bcl-xl, Bax, and Bad) and mitogen-activated protein kinases (MAPKs; ERK p44/42, JNK, and p38) were investigated by Western blot analysis. Also mRNA expressions of Bcl-2 family were examined by quantitative real time RT-PCR.
Results; RGC-5 cells, differentiated by 1.0 μM staurosporine for 6 hours and recovered for 3 days, contained fully developed neurites and were well connected with each other. Under H₂O₂ oxidative stress, agmatine pretreatment for 2 hours enhanced cell survival dose dependently. The cytotoxicity assay showed 40.9 % cell loss, which was reduced to 27.3 % when the cells were pretreated by 100 μM agmatine for 2 hours followed by 1.0 mM H₂O₂ treatment for 16 hours. This cell loss was due to apoptotic cell death, as established by TUNEL assay, but was neither caspase-3 nor caspase-9 dependent. The effect of agmatine pretreatment was completely inhibited by yohimbine, but partially decreased by glutamate or NMDA. Total expression of Bcl-2 family proteins and MAPKs were not influenced by H₂O₂ oxidative stress. Also mRNAs of Bcl-2 family were not changed. In contrast, the phosphorylation of JNK was suppressed by agmatine pretreatment followed by H₂O₂ oxidative stress.

Conclusion; The present study shows that agmatine had neuroprotective effects on oxidative stressed differentiated RGC-5 cells. It might suggest a therapeutic strategy for many ocular diseases associated with oxidative stress.

Key words: retinal ganglion cell, agmatine, oxidative stress, neuroprotection
Agmatine pretreatment attenuates retinal ganglion cell death by an oxidative stress

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I. INTRODUCTION

Glaucoma, one of the leading causes of blindness in the world, is characterized by progressive loss of retinal ganglion cells (RGCs) including their axons, and by tissue remodeling of the optic nerve head. This is followed by visual field defects. Even though the elevated intraocular pressure is the major risk factor associated with glaucomatous visual field loss, oxidative stress has recently been proposed as another important risk factor in the pathogenesis of glaucoma.1-3

Glutamate is the principal excitatory neurotransmitter in the retina. The extracellular concentration of glutamate increases after an ischemic insult and a significant component of ischemic injury to CNS neurons results from glutamatergic excitotoxicity. Glutamate and nitric oxide (NO) promote oxidative damage by reacting with superoxide anion. RGCs express both N-methyl-D-aspartate (NMDA) and non-NMDA type glutamatergic receptors.
The oxidative stress has been shown to play an important role in ischemic retinopathy.\textsuperscript{4,5} RGCs are susceptible and vulnerable to the oxidative stress. Hydrogen peroxide (H\textsubscript{2}O\textsubscript{2}) induces apoptosis through the mitochondrial death pathway. Bcl-2 family has been shown to be involved in cytotoxicity by glutamate and NO.\textsuperscript{6,7} Also the activity of several mitogen activated protein kinases (MAPKs) have been studied in H\textsubscript{2}O\textsubscript{2} and NMDA induced apoptosis.\textsuperscript{8,9}

Primary cultures of RGCs have some limitations to study RGC pathophysiology. Despite the fact that transformed rat retinal ganglion cell line (RGC-5 cell) has a number of characteristics of normal RGCs, it is mitotically active and is not exactly same as primary RGCs. It has been shown that RGC-5 cells treated by the broad-spectrum protein kinase inhibitor staurosporine are not dividing and maintain neuronal characteristics.\textsuperscript{10-12}

Several molecules having some neuroprotective effects, to the hypoxic injury at CNS, have been reported.\textsuperscript{13-20} The neuroprotective effects by pretreatment of these molecules against hypoxia-induced neuronal death have not been well understood. Although some drugs have been reported to have a protective role in RGC, most of them have no sufficient data to support their definite role.\textsuperscript{21-24}

Agmatine, a primary amine formed by the decarboxylation of L-arginine synthesized in mammalian brain, is an agonist for the alpha 2-adrenergic and imidazoline receptors and an antagonist at NMDA receptors.\textsuperscript{25-27} Recent studies have shown that agmatine may be neuroprotective in neuronal ischemic models.\textsuperscript{28-31}

In the present study, I examined the protective effects of agmatine pretreatment on oxidative injuries of RGC-5 cells \textit{in vitro}, and characterized the mechanisms involved in these activities.
II. MATERIALS AND METHODS

1. Chemicals

Agmatine sulfate, staurosporine (from *Streptomyces staurosporeus*), glutamate, and yohimbine were purchased from Sigma-Aldrich (St. Louis, MO, USA). NMDA was purchased from Calbiochem (San Diego, CA, USA).

2. Cell culture and agmatine pretreatment

RGC-5 cells (donated by Alcon Research, Forth Worth, TX, USA) were cultured in Dulbecco’s modified Eagle’s medium (DMEM, Gibco, Carlsbad, CA, USA) with 1 g/L glucose, 100 U/mL penicillin, 100 µg/mL streptomycin, and 10% heat inactivated fetal bovine serum (FBS, Gibco). All experiments were performed at a confluence of 70 to 80%. RGC-5 cells were differentiated by exposure to 1.0 µM staurosporine. After recovery using 10% FBS-DMEM, the differentiated RGC-5 cells were incubated with various concentration of agmatine.

3. Oxidative stress

Cellular oxidative stress was induced by addition of H$_2$O$_2$ to the culture medium.

4. Cell viability assays

Total cell population and viability was quantified by lactate dehydrogenase (LDH) assay (Promega, Madison, WI, USA) according to the manufacturer's instructions. Briefly, the cells in a culture were lysed by Lysis Solution (0.9% (v/v) Triton X-100 in water). The LDH activities in 50 µL of the sample supernatant in 96 well-plates were colorized by a tetrazolium salt and the red formazan product was measured at 490 nm using
an ELISA plate reader (Vmax; Molecular Device, Sunnyvale, CA, USA). The cell population was proportional to the absorbance values.

The proportion of injured cells among total cell population was quantified by measurement of released LDH to culture media from the cells. Cytotoxicity was expressed as a percentage of the LDH activity in medium to the total (medium and cellular LDH). Data of total cell population and viability are expressed as mean ± SEM (standard error of mean).

5. Morphological analysis

Photomicrographs of cells were taken at 200×, digitalized and stored as JPEG. Axonal outgrowth of differentiated RGC-5 cells was measured and assessed by NeuronJ image software (ImageJ 1.40; http://rsb.info.nih.gov/ij). The axonal length was expressed as the mean ± SEM of 40 to 50 cells.

6. Apoptosis assays

Differentiated RGC-5 cells were plated on poly-D-lysine/Laminin-coated 12 mm φ cover slide glass (BD Biosciences, Bedford, MA, USA). The TUNEL assay was performed using Apo-BrdU in situ DNA Fragmentation Assay Kit (BioVision, Mountain View, CA, USA) with minor modification.32 Briefly, the cells were fixed in 4 % paraformaldehyde in phosphate-buffered saline (PBS), pH 7.4 for 30 minutes and permealized in 0.1 % Triton X-100 in 0.1 % sodium citrate. After washing in PBS, the cells were stored in 70 % (v/v) ethanol at -20°C overnight. After washing, the cells were carefully covered by the DNA labeling solution containing TdT enzyme and bromolated deoxyuridine triphosphate digoxigenine (Br-dUTP), and incubated in a dark and
humidified 37°C incubator for 60 minutes. After washing, the cells were covered by the anti-BrdU-FITC antibody solution and incubated in a humidified incubator at room temperature. After removing the antibody solution, the cells were covered by propidium iodide (PI) /Rnase A solution and incubated in a dark in a humidified incubator for 30 minutes. By fluorescence microscopy, apoptotic cells show green staining over an orange-red PI counter-staining. Double stained cells were visualized with a total magnification of 200×. The cells incubated in the labeling solution without TdT enzyme were used as negative control (data was not shown).

Caspase-3 and -9 activities were measured using Caspase-3 and Caspase-9 Fluorometric Assay Kits (BioVision) respectively, according to the manufacturer's instructions. Briefly, RGC-5 cells were suspended in the lysis buffer. In 96 well-plate, cell lysate 50 or 100 µg as total protein was mixed with the reaction buffer containing 10 mM DTT and added either DEVD-AFC substrate for caspase-3 or LEHD-AFC substrate for caspase-9. The reaction mixture was incubated at 37°C for 2 hours, and then read in a fluorometer equipped with a 400 nm excitation filter and 505 nm emission filter. Increase in caspase activity was determined by comparing the results with the level of untreated control. Amount of total protein was determined by Bradford protein assay (BIO-RAD, Philadelphia, CA, USA). Data are expressed as mean ± SEM of 8 to 10 independent cultures.

7. Western immunoblot

Differentiated RGC-5 cells cultured were washed twice in cold PBS, and then lysed in lysis buffer (Mammalian Cell Lysis kit, Sigma-Aldrich) containing 1.0 mM Na₃VO₃ and 1.0 mM phenylmethyonsulfonyl fluoride. The protein (20 or 30 µg) were separated on 12 % SDS-polyacrylamide gels,
transferred to a polyvinylidene difluoride membrane (Millipore, Bedford, MA, USA), and blocked by 5% nonfat dry milk in a Tween20-Tris buffered saline for one hour at room temperature. Membranes were then incubated with the following primary antibodies over night at 4°C: mouse anti-Bcl-2 (1:200; Santa Cruz Biotechnology, Santa Cruz, CA, USA), rabbit anti-phospho-Bcl-2 (1:1000; Cell Signaling Technology, Danvers, MA, USA), rabbit anti-Bcl-xL (1:1000; Cell Signaling Technology), rabbit anti-phospho-Bcl-xL (1:500; SAB, Pearland, TX, USA), rabbit anti-Bax (1:1000; Cell Signaling Technology), mouse anti-Bad (1:100; Santa Cruz Biotechnology), rabbit anti-phospho-Bad (1:1000; Cell Signaling Technology) for Bcl-2 family protein, and rabbit anti-SAPK/JNK (1:1000; Cell Signaling Technology), rabbit anti-phospho-SAPK/JNK (1:1000; Cell Signaling Technology), rabbit anti-MAPK p38 (1:1000; Cell Signaling Technology), rabbit anti-phospho MAPK p38 (1:1000; Cell Signaling Technology), rabbit anti-p44/p42 (1:1000; Cell Signaling Technology), rabbit anti-phospho-p44/p42 (1:1000; Cell Signaling Technology) for MAPK family, and mouse anti-β-actin (1:10000; Sigma-Aldrich). After treatment with the secondary goat anti-rabbit (1:2000; Cell Signaling Technology) or horse anti-mouse (1:2000; Cell Signaling Technology) IgG conjugated by horseradish peroxidase for one hour, immunoreactive bands were visualized with the ECL detection system (Thermo Scientific, Waltham, IL, USA).

8. Real time RT-PCR

Total RNA was extracted using RNeasy Mini Kit (QIAGEN, Valencia, CA, USA) and treated with DNase (QIAGEN) to remove contaminating DNA according to manufacturer directions. The cDNAs were synthesized
using Superscript III First-Strand Synthesis System for RT-PCR (Invitrogen, Carlsbad, CA, USA). A Real time PCR was performed with 50 ng cDNA per a reaction using Power SYBR Green PCR Master Mix (BIO-RAD) in a total volume of 25 µL of reaction mixture (iCycler iQ, BIO-RAD). The SYBER green data and level of target mRNA were analyzed by iCycler iQ software with a relative standard curve of β-actin. All experiments were performed in triplicate and repeated 4 times. Data were presented as mean ± SEM. Sequences of oligonucleotides used as primers are summarized in Table1.33-35

Table1. Primer sequences for real-time RT-PCR

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<tr>
<td>Bcl-xl</td>
<td>sense 5’GTA GTG AAT GAA CTC TTT CGG GAT GG 3’</td>
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</tr>
<tr>
<td></td>
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</tr>
<tr>
<td>Bax</td>
<td>sense 5’AAT ATG GAG CTG CAG AGG ATG ATT G3’</td>
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<td>Bad</td>
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<tr>
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<tr>
<td></td>
<td>antisense 5’ACC AGA GGC ATA CAG GGA CAA3’</td>
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9. Statistical analysis

The data will be analyzed by 2-tailed Student t-test or one-way ANOVA, followed by post hoc comparisons (Student-Newman-Keuls) using the Statistical Package for Social Sciences 12.0 (SPSS).
III. RESULTS

1. Differentiation of RGC-5 cells

Staurosporine changed the morphology of RGC-5 cells from flat to round shape and induced the outgrowth of neurites (Figure 1). The change in morphology was already observed after 1.0 µM staurosporine treatment for 2 and 6 hours. The neurites were growing through recovery period in 10 % FBS-DMEM for 3 days. These neurites had multiple and long branches, and sufficiently contacted with those from neighbor cells. In 3-days recovered cells, the average of axonal length was 39.5 ± 1.74 µm and 59.6 ± 2.72 µm for the 2 and 6 hours treated cells, respectively (Figure 2A). The absorbance values of the total cellular LDH at 490 nm were increasing, but there were no significant differences between the 2 and 6 hours treated cells on 3-day recovery (Figure 2B). Both of the neurites outgrowth and total cell population decreased after 3 days. Based on these results, RGC-5 cells differentiated by 1.0 µM staurosporine for 6 hours and recovered for 3 days were used for the following oxidative stress experiments.
<table>
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<td>4 day</td>
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<td>5 day</td>
<td><img src="image11.png" alt="Image" /></td>
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**Figure 1.** Staurosporine treatment and recovery of RGC-5 cells. After treatment by 1.0 µM staurosporine for 2 or 6 hours, RGC-5 cells were recovered in 10% FBS-DMEM up to 5 days.
Figure 2. Differentiation of RGC-5 cells by staurosporine. The RGC-5 cells were exposed to 1.0 μM staurosporine for 2 and 6 hours, and then recovered up to 5 days. Outgrowth length of the axons (A) and optical density of total cellular LDH (B) over time.
2. Oxidative stress induced cytotoxicity of differentiated RGC-5 cells

The concentration of H$_2$O$_2$ below 0.8 mM did not significantly induce cell death. Over 1.0 mM H$_2$O$_2$, both time and concentration dependent cell death were observed (Figure 3). The 16 hours incubation of the differentiated RGC-5 cells with 1.0 mM H$_2$O$_2$ was selected as the oxidative stress condition for further experiments.

![Graph](image)

**Figure 3.** Cytotoxicity of hydrogen peroxide (H$_2$O$_2$) on RGC-5 cells.

3. Protective effects of agmatine pretreatment

After the 2 hours pretreatment of agmatine, the cytotoxic effect of H$_2$O$_2$ on differentiated RGC-5 cells most effectively decreased (Figure 4A). Sixteen hours exposure to 1.0 mM H$_2$O$_2$ induced 40.9 % cytotoxicity, but it decreased to 35.8 %, 28.9 % (p<0.007), and 27.3 % (p<0.003) by 2 hours pretreatment of agmatine of 1, 10, and 100 µM, respectively. Although 1 µM agmatine did not make statistically significant difference, 10 and 100 µM agmatine significantly suppressed the cytotoxic effect of H$_2$O$_2$. And 2 hours incubation demonstrated the most effective protection (Figure 4A). The protective effects lasted for 24 hours (Figure 4B).
Figure 4. Protective effect of agmatine pretreatment to differentiated RGC-5 cells against oxidative stress by H$_2$O$_2$. (A) Agmatine treatment for 2, 4, or 6 hours followed by 1.0 mM H$_2$O$_2$ oxidative stress for 16 hours. (B) Agmatine treatment for 2 hours followed by 1.0 mM H$_2$O$_2$ oxidative stress for 16, 24, or 48 hours. * Negative control; the cells were incubated in DMEM without any treatments.
4. TUNEL staining

To characterize the effect of agmatine pretreatment on the cell death induced by oxidative stress, TUNEL assays were performed. TUNEL positive apoptotic cells, showing bright green refringence under the fluorescent microscope, marked increased in RGC-5 cells treated with 1.0 mM H$_2$O$_2$ only. Pretreatment with 100 µM agmatine before 1.0 mM H$_2$O$_2$ application significantly decreased apoptotic cells with green refringence (Figure 5).

![Figure 5](image)

**Figure 5.** Effect of agmatine pretreatment on apoptosis of differentiated RGC-5 cells. (A) Control; (B) 100 µM agmatine pretreatment only; (C) 1.0 mM H$_2$O$_2$ oxidative stress only; and (D) 100 µM agmatine pretreatment followed by 1.0 mM H$_2$O$_2$ oxidative stress.
5. Caspase activities on RGC-5 pretreated by agmatine

The RGC-5 cells were pretreated with 100 µM agmatine for 2 hours, and then incubated in 1.0 mM H₂O₂ for 16 or 24 hours. The caspase-3 activity slightly decreased at 24 hours (p<0.406) (Figure 6A). The caspace-9 activity decreased at the period of 16 hours after agmatine pretreatment, but it was not statistically significant (p<0.373) (Figure 6B).

(A)

(B)

Figure 6. Caspase-3 (A) and -9 (B) activities of oxidative stressed RGC-5 cells after 2 hours pretreatment with 100 µM agmatine.
6. Agmatine as an alpha 2-adrenergic receptor agonist

Effect of agmatine was examined with co-treatment of yohimbine (alpha 2 receptor antagonist) and 1.0 mM H₂O₂ (Figure 7). In RGC-5 cells without agmatine pretreatment, co-treatment of yohimbine demonstrated no difference in cytotoxicity. When the cells were pretreated by 100 µM agmatine, co-treatment of 10 nM yohimbine increased cytotoxicity by 20.4 %, thus showed that protective effect of agmatine pretreatment disappeared in the presence of yohimbine.

![Figure 7. Inhibitory effect of yohimbine to agmatine pretreatment. Negative control; the cells in DMEM without any treatments. Control; 1.0 mM H₂O₂ oxidative stress with or without 10 nM yohimbine. Pretreatment; 1.0 mM H₂O₂ oxidative stress with or without yohimbine after 100 µM agmatine pretreatments.](image)

7. Agmatine as a NMDA receptor antagonist

The protective effect of agmatine pretreatment was examined in the presence of glutamate and NMDA under 1.0 mM H₂O₂ oxidative stress for 16 hours. The cytoprotective effect of agmatine was abolished with the presence of glutamate and NMDA (Figure 8). The mean value of
cytotoxicity assays increased from 53.6 % to 66.1 % in the RGC-5 cells after the application of 1.0 µM glutamate. The agmatine pretreated RGC-5 cells also demonstrated increased cytotoxicity but less mean value after the application of 1.0 µM glutamate (Figure 8A). The application of 100 µM NMDA showed similar results (Figure 8B).

**Figure 8.** Influence of glutamate (A) and NMDA (B) on agmatine pretreatment. Negative control; the cells in DMEM without any treatments. Control; 1.0 mM H$_2$O$_2$ oxidative stress with or without 1.0 µM glutamate or 100 µM NMDA. Pretreatment; 1.0 mM H$_2$O$_2$ oxidative stress with or without glutamate or NMDA after 100 µM agmatine pretreatments.
8. **Expression of total and phosphorylated Bcl-2 family proteins**

Expression of Bcl-2 family proteins was examined by quantitative real time RT-PCR and Western blot assay. Even though the mRNA expression of Bcl-xl from 100 µM agmatine treated RGC-5 cells slightly increased at 16 hours after the application of 1.0 mM H$_2$O$_2$, other Bcl-2 family proteins (Bcl-2, Bax, and Bad) did not show significant changes (Figure 9). The protein contents of Bcl-2 family proteins from the RGC-5 cells did not demonstrate significant changes after the application of the oxidative stress (Figure 10). Bcl-xl seemed to be slightly increased by agmatine pretreatment, but the phosphorylation was not stimulated.

![Figure 9](image_url)

**Figure 9.** Results of quantitative real time RT-PCR for Bcl-2 family. (A) Bcl-2, (B) Bcl-xl, (C) Bax, and (D) Bad.
Figure 10. Expression of Bcl-2 family proteins at various incubation periods after the application of 1.0 mM H$_2$O$_2$ oxidative stress analyzed by Western immunoblot.

9. **Expression of total and phosphorylated MAPKs proteins**

Western immunoblot was performed to examine the effect of agmatine on the expression of three MAPK proteins (JNK, ERK p44/42, and p38) (Figure 11). The expression of phospho-JNK in the agmatine pretreated RGC-5 cells decreased earlier after oxidative stress than that in the RGC-5 cells without agmatine pretreatment. But agmatine pretreatment demonstrated no effect on the expression of phospho-ERK p44/42 MAPKs. Total amount of p38 was not changed thorough oxidative stress.
### IV. DISCUSSION

Researches about RGC pathophysiology have been hampered by difficulties and limitations in using primary RGCs. Cultured RGCs survived in relatively short period and their isolation is difficult. Although immortalized RGC-5 cell line expresses neuronal markers characteristic of RGC, it is mitotically active and morphologically more similar to glial cells. It has been shown that RGC-5 cells differentiated by staurosporine are non-mitotic and express sufficient neurites for a while.  

In this study,  

**Figure 11.** Expression of MAPKs proteins at various incubation periods after the application of 1.0 mM H$_2$O$_2$ oxidative stress analyzed by Western immunoblot.
RGC-5 cells differentiated by 1.0 µM staurosporine for 6 hours well survived for three days in 10 % FBS-DMEM and contained sufficient growing axons and connected with each other.

Apoptosis is programmed cell death and caspases usually mediate apoptotic neuronal death. However, there have been studies that H$_2$O$_2$ cause apoptosis by either caspase-independent or dependent pathway. In the present study, oxidative stress was induced by treatment of the differentiated RGC-5 cells with H$_2$O$_2$ and the effects of agmatine pretreatment were examined. The results indicated that the pretreatment with agmatine protected and rescued the RGC-5 cells from apoptosis dose dependently, and this effect was independent of caspase-3 and -9 activities.

Agmatine has been known as an agonist for the alpha 2-adrenergic and imidazoline receptors and an antagonist at NMDA receptors. Wheeler and Nylander and Kalapesi et al. have studied the alpha 2-adrenergic receptor on human ganglion cells and RGC-5 cell line. They demonstrated the presence of alpha 2A-adrenergic receptor on both undifferentiated and succinyl concanavaline-A differentiated RGC-5 cell lines. They also showed an increased expression of alpha 2A-adrenergic receptor on the 7-day differentiated RGC-5 cells compared to the undifferentiated and early differentiated cells. Although staurosporine was used to differentiate RGC-5 cells in the present study, it was shown that the protective effect of agmatine pretreatment was completely abolished by yohimbine (antagonist for alpha 2-adrenergic receptor) in the differentiated and 3-day recovered RGC-5 cells. This result suggests a possibility that pretreatment with agmatine protects RGC-5 cells through alpha 2-adrenergic receptor.
RGCs also express both NMDA and non-NMDA type glutamatergic receptors. NMDA-type glutamatergic excitotoxicity has been implicated as a mechanism for injuries associated with ischemia and other insults to neurons in many regions of the CNS. However, there have been contradictory results about the rule of glutamate and NMDA excitotoxicity to vulnerability of RGCs.\textsuperscript{42, 43} Although varied evidences have been shown in several species and strains of the experimental animal used, Ullian et al.\textsuperscript{42} indicated that primary rat RGCs is invulnerable to glutamate and NMDA, \textit{in vitro}. Their rat primary RGCs survived in the presence of glutamate (500 µM) or NMDA (500 µM) in relatively long period of incubation. In present experiment, the protective effect of agmatine pretreatment under H\textsubscript{2}O\textsubscript{2} oxidative stress was not completely inhibited by glutamate and NMDA. It may suggest that the agmatine pretreatment partially paralyze NMDA.

Bcl-2 family includes both anti-apoptotic (ex. Bcl-2 and Bcl-xl) and pro-apoptotic (ex. Bad and Bax) regulator proteins on mitochondrial membrane. Cytochrome c release is inhibited by Bcl-2 and Bcl-xl. Tamatani et al.\textsuperscript{6, 7} estimated that Bcl-2 or Bax were involved in cytotoxicity by glutamate and NO. Li and Osborne\textsuperscript{38} showed oxidative-induced apoptosis to an immortalized ganglion cell line is caspase independent. The results of the present study showed that Bcl-2 family proteins in the RGC-5 cells did not show dramatic changes under H\textsubscript{2}O\textsubscript{2} oxidative stress. So it is indicated that caspase independent mechanisms irrespective of Bcl-2 family proteins are involved in the H\textsubscript{2}O\textsubscript{2} induced oxidative damage of RGC-5 cells.

Exogenous H\textsubscript{2}O\textsubscript{2} causes intracellular reactive oxygen species accumulation and mitochondrial dysfunction, and activates MAPK pathway.
To elucidate the mechanism of agmatine effects, Hong et al.\textsuperscript{31} recently demonstrate that agmatine is neuroprotective against hypoxia-induced damage of RGC-5 cells through the JNK and NF-κB signaling pathways. The present study demonstrated that JNK was suppressed in agmatine pretreated RGC-5 cells after the application of oxidative stresses. Although I have not examined NF-κB signaling pathways, results of this study indicates that MAPK seems to be involved in the mechanism of the neuroprotective effects of agmatine.

In present study, I demonstrated the protective effects of agmatine against oxidative stress induced apoptosis of RGC-5 cells. Although more well-designed studies are required to elucidate the precise mechanism of neuroprotective effects of agmatine, the present results indicate several underlying mechanisms of the neuroprotective effects of agmatine to RGCs.

V. CONCLUSION

The aim of this study was to evaluate the neuroprotective effects of agmatine pretreatment on oxidative stressed RGCs \textit{in vitro} and to investigate the underlying mechanism. The RGC-5 cells were sufficiently differentiated by exposure to staurospirine and the oxidative stress was induced by incubation with H$_2$O$_2$. By agmatine pretreatment, the cell viability was increased and apoptosis was decreased under H$_2$O$_2$ oxidative stress. These protective effects seemed to be associated with MAPKs through alpha 2-adrenergic and NMDA receptors. Finally, agmatine might be a therapeutic strategy for many ocular diseases associated with oxidative stress.
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< ABSTRACT (IN KOREAN)>

아그마틴 전처치가 산화성 스트레스에 의해 유발되는
망막신경절 세포사에 미치는 영향

<지도교수 양 우익>

연세대학교 대학원 의과학과

이주카 요코

본 연구에서는 쥐의 망막신경절세포주(RGC-5 cell)를 대상으로
산화스트레스에 대한 L-아르기닌의 탈탄산화에 의해 형성되는
폴리아민인 아그마틴의 전처치 보호효과를 살펴보고자 하였다.
스타우로스포린의 첨가에 의해 분화시킨 RGC-5 세포에 아그마틴
전처치를 시행하고 세포배양액에서 산화수소를 첨가하여
산화스트레스를 유발하였다. 세포사멸은 유산염탈수소효소 분석
을 이용하여 평가하였으며, 세포고사는 TUNEL 분석과 caspase-3 및 -9
분석으로 평가하였다. 또한, Bcl-2 family (Bcl-2, Bcl-xL, Bax와 Bad)와
MAPKs (ERK p44/42, JNK, p38) 단백질의 총량과 인산화 정도는
웨스턴면역분석법을 이용하여 확인하였고, Bcl-2 family의 mRNA
발현은 real time RT-PCR을 이용하여 검토하였다. 1 μM
스타우로스포린에 6시간 동안 노출하여 분화시킨 RGC-5 세포를 2시간 동안 100 µM 아그마틴으로 전처치한 후 1.0 mM 과산화수소에 16시간 동안 노출시켜 산화스트레스를 유발하였을 때, 세포사멸은 40.9 %에서 27.3 %로 감소하였다. TUNEL분석을 이용하여 이러한 산화스트레스로 인한 RGC-5 세포의 사멸이 세포고사에 의한 것임을 알 수 있었으며 아그마틴 전처치에 의해 감소됨을 알 수 있었다. 하지만, 아그마틴 전처치에 의한 caspase-3 및 -9의 활성 변화는 관찰되지 않았다. 이러한 아그마틴 전처치의 효과는 yohimbine에 의해 완전하게 억제되었지만, 글루타민산염 또는 NMDA에 의해서는 부분적으로만 억제되었다. 또한, Bcl-2 family 및 MAPKs 단백질의 총량은 산화스트레스의 영향을 받지 않았고, Bcl-2 family의 mRNA도 변화하지 않았다. 하지만 아그마틴 전처치 후 산화스트레스를 유발하였을 때는 MAPKs 중 JNK의 인산화가 억제되었다. 본 연구를 통하여 분화된 RGC-5 세포에서 산화스트레스를 유발하였을 때 아그마틴 전처치가 신경보호 효과를 나타냄을 알 수 있었다.

핵심되는 말: 망막신경절세포, 아그마틴, 산화스트레스, 신경보호