Maintenance of the Neuroprotective Function of the Amino Group Blocked Fluorescence-Agmatine

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Received: 10 November 2020 / Revised: 21 February 2021 / Accepted: 2 April 2021 / Published online: 29 April 2021
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Abstract
Agmatine, an endogenous derivative of arginine, has been found to be effective in treating idiopathic pain, convulsion, stress-mediated behavior, and attenuate the withdrawal symptoms of drugs like morphine. In the early stages of ischemic brain injury in animals, exogenous agmatine treatment was found to be neuroprotective. Agmatine is also considered as a putative neurotransmitter and is still an experimental drug. Chemically, agmatine is called agmatine 1-(4-aminobutyl guanidine). Crystallographic study data show that positively-charged guanidine can bind to the protein containing Gly and Asp residues, and the amino group can interact with the complimentary sites of Glu and Ser. In this study, we blocked the amino end of the agmatine by conjugating it with FITC, but the guanidine end was unchanged. We compared the neuroprotective function of the agmatine and agmatine-FITC by treating them in neurons after excitotoxic stimulation. We found that even the amino end blocked neuronal viability in the excitotoxic condition, by NMDA treatment for 1 h, was increased by agmatine-FITC, which was similar to that of agmatine. We also found that the agmatine-FITC treatment reduced the expression of nitric oxide production in NMDA-treated cells. This study suggests that even if the amino end of agmatine is blocked, it can perform its neuroprotective function.

Keywords Agmatine · Guanidine · Neuroprotection · NMDA

Introduction
Primary amine agmatine can be created naturally from α-amino l-arginine. The decarboxylation of l-arginine to produce agmatine occurs in the presence of the mammalian mitochondrial outer membrane enzyme arginine decarboxylase. Agmatine is catabolized into prototypical polyamine putrescine by agmatinase, a human ureohydrolase. Agmatine has been found in different organs of the body at lower concentrations, with enrichments in a few parts of the brain and spinal cord, packaged in the synaptic vesicles [1–3]. n the synaptic vesicles, agmatine is found to be co-localized with other classical neurotransmitters such as glutamate and vasopressin and can be released by calcium-dependent depolarization [4, 5]. With the above characteristics, agmatine is believed to be a neuromodulator that can act as a neurotransmitter, and localized in the oxytocin and vasopressin neurons [6, 7].

Agmatine was found to be neuroprotective in different excitotoxic and ischemic neurological diseases. Agmatine has been reported to reduce vascular permeability in the brain; this induces gastric protection in ischemic-injury rats.
The neuroprotective mechanism of agmatine is not yet clearly understood; however, it is believed that during the early hours of the ischemic/traumatic injury, activation of the unabated inducible nitric oxide synthase (iNOS) occurs, which increases nitric oxide (NO) production, whereas the level of arginine decarboxylase (ADC)/agmatine remains low [9–11]. In this condition, the sustained higher level of NO opens the N-methyl-D-aspartate (NMDA) channels through which the calcium influxes, causing neurotoxicity. After several hours of injury, the ADC/agmatine level increases and controls the iNOS and NMDA receptor functions [12, 13]. In addition to ischemic or traumatic injury, agmatine has been found to have a positive effect on neurological diseases such as Alzheimer’s disease, Parkinson’s disease, epilepsy, and other mental diseases [14–16]. Agmatine was also reported to be anti-apoptotic, an effect it achieves by attenuating the expression of caspase-3 and Bax, and elevating the expression of BCL2, Nrf2, and PI3K. [15, 17–20]. In a recent neurochemical profiling study, acute and sub-chronic oral treatment of agmatine was reported to be well tolerated and did not show any adverse effect in APP-swe/PS1ΔE9 transgenic (Tg) mice, and also found to cross the blood brain barrier and accumulate in the brain [21].

Agmatine was first believed to be a clonidine-displacing substance that specifically binds to α2-adrenoceptors, I1- and I2-binding sites [22]. However, later studies suggested that agmatine can function through other neurological receptors, such as N-methyl-D-aspartate receptor, 2-amino-3-(5-methyl-3-oxo-1,2-oxazol-4-yl) propanoic acid receptor, kainate receptor, acetylcholine receptor, and serotonin receptor [23]. The positively-charged guanidine and amino end of the agmatine tend to bind with the Gly and Asp residues of the protein and the Gln and Ser residues of the complementary sites of the protein, respectively [24]. These two ends in agmatine have been suggested to play a vital role in neuroprotection, but there is no specific report yet. In this study, we intended to visualize the exogenous agmatine by attaching a fluorescein isothiocyanate (FITC) to its amino end and studied the role of agmatine in vitro neuron culture. Our goal was to check, if the modified fluorescence-agmatine still conserve its neuroprotective function as the normal agmatine and how?

**Material and Methods**

**Chemistry of Agmatine-FITC**

**FITC-Agmatine**

To a stirred solution of FITC (50 mg, 0.12 mmol) and TEA (18 μL, 0.12 mmol) in DMF (1 mL) was added agmatine (0.16 mg, 0.12 mmol) at room temperature. After stirring for 3 h at room temperature, the reaction was quenched by the addition of H2O. The organic solution was washed with water and brine, dried over anhydrous Na2SO4, filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography (CH3Cl2: MeOH = 10:1) to give a product in 60% yield: 1H nuclear magnetic resonance (NMR) (400 MHz, DMSO-d6) δ.82 m, 2 H), 1.61–1.45 (m, 4 H); ESI–MS calcd for C26H39N6O6S [M]+ 519.1 found 519.1.

**Primary Neuronal Cell Culture**

Primary neuron culture was done as previously reported [25]. Before primary neuronal cell culture, the plates were coated with Poly-D-Lysine (Gibco) containing laminin overnight at room temperature, and the dishes were washed three times with autoclaved distilled water. Brains were extracted from ICR mice E14.5 (Koatech). The pregnant mice were euthanized using ether. Before incision, the abdomen was wiped with 70% ethanol to prevent possible contamination, after which the skin was cut and removed, and the abdominal wall was incised. Fetuses were decapitated with a pair of scissors, and the heads were placed in a petri dish containing hank’s balanced salt solution (HBSS). When the skull was opened, the olfactory bulbs, meninges, and hippocampus were removed systematically, and the cortex was isolated. The cells were pelleted by centrifugation at 1000×g for 3 min, and the supernatant was removed. The cells were mechanically dissociated using a Pasteur pipette (20 times). Approximately 1 × 106 cells/mL were plated on PDL-laminin-coated 6-well plates in a neurobasal medium (Gibco) containing t-glutamine (Hyclone), penicillin–streptomycin (Hyclone), and B27 supplement (Gibco), and then cultured at 37 °C in 5% CO2/95% air. After 7 days in vitro, the cells were treated with NMDA (50 μM), agmatine (100 μM), and agmatine-FITC (AgmF, 100 μM). To check the cellular uptake of the AgmF, cells were treated with AgmF and CellTracker™ Red CMTPX Dye (Thermo Fisher, cat: C34552). All animal experiments were conducted in accordance with the guidelines on the use and care of laboratory animals established by the Animal Care Committee Yonsei University.

**Immunocytochemistry**

Neurons were washed three times with iced-PBS and permeabilized with 4% PFA. The cells were then incubated with primary antibodies at 4 °C overnight. The following primary antibodies were used: DCX (1:500), NOS2 (1:200). The primary antibody was then removed, and the cells were washed...
three times for 3 min with PBST (10% with Trionx-100). The cells were incubated with FITC-conjugated anti-IgG (1:500) and rhodamine-conjugated anti-IgG (1:500) at room temperature for 2 h. After washing three times for 3 min with PBS, the cells were counterstained with DAPI (1:500) for 10 min at room temperature. They were imaged using a Zeiss LSM 700 confocal microscope (Carl Zeiss, Thornwood, NY, USA). For each immunocytochemistry experiment, 3–5 independent experiments were performed and 4–5 images from each experiment were analyzed while counting the cell number.

**Crystal Violet Staining**

To check the viability of the adherent neurons after the NMDA and AgmF treatment we performed crystal violet staining of the neurons. After treating the cells with NMDA followed by Agmatine and AgmF we washed the cells with PBS for 2 times and stained the cells with crystal violet (0.5% in water/methanol 4/1) for 5 min and then rinsed with water and left to dry overnight [26]. The images were taken by using a light microscopy. Cell number was counted from 4–5 images obtained from 4 independent experiments.

**MTT Assay**

Neurons (1 × 10^6 cells/mL) were seeded in 24-well plates to monitor all experimental conditions. The cells were then treated with NMDA (50 μM) for 1 h followed by a PBS wash, agmatine, and AgmF for 6 h. Next, cells were rinsed twice with PBS, and the culture medium was replaced with neurobasal medium (Gibco). Then, 100 μL of 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT, Sigma) solution (5 mg/mL in PBS) was added to each well. After 1 h of incubation, the medium was removed, and dimethyl sulfoxide was added to solubilize the purple formazan product of the MTT reaction. The supernatant from each well was analyzed using an enzyme-linked immunosorbent assay plate reader at a wavelength of 570 nm, with background subtraction at 650 nm. All experiments were repeated at least three times. Cell viability in the control medium without any treatment was considered 100%. Cell viability was reported as the value relative to the control group.

**Nitric Oxide Production in the Neurons**

The neurons obtained from the ICR mice were cultured in a neurobasal medium (Gibco) containing 1% l-glutamine (Hyclone), 1% penicillin–streptomycin (Hyclone), and B27 supplement (Gibco). The cells were plated at a density of 1 × 106, pre-incubated for 24 h at 37 °C, and maintained in a humidified atmosphere containing 5% CO₂. The mammalian cells were treated with NMDA (50 μM) for 1 h and then washed 3 times with PBS to remove the NMDA and treated with agmatine and AgmF for 6 h in culture media. Accumulated nitrite (NO₂⁻), as an index of nitric oxide (NO), in the media was determined using a colorimetric assay based on the Griess reaction. After 6 h cells were washed with PBS and fixed with 4% PFA and stored at 4 °C until NOS2 staining.

**Statistical Analysis**

The data are quantified as the mean ± SEM from at least three different experiments performed from separate cell preparations, and at least quadruplicate determinations were performed in each experiment. Statistical tests to determine differences between groups others were analyzed with analysis of variance followed by Tukey’s post hoc test. SigmaPlot 12.0 (Systat Software Inc., San Jose, California, USA) for Windows was used. p values less than 0.05 were considered significant.

**Results**

**Chemistry of Agmatine-FITC and Neuronal Uptake**

**General**

Analytical thin-layer chromatography was conducted on silica gel in 60 F254 glass plates. Compound spots were visualized by ultraviolet light (254 nm) and/or by staining with 10 wt% phosphomolybdic acid in ethanol. Flash column chromatography was performed using silica gel 60 (230–400 mesh). NMR spectra were recorded on a Bruker DRX-400 instrument. Mass spectra were obtained using a Waters 3100 LC/MS System. Chemical reaction showing the making of agmatine-FITC (Fig. 1a). To confirm the cellular uptake of the AgmF, we treated the cultured neurons with AgmF (green) for 6 h, and then CellTracker™ Red CMTPX Dye (red) and confocal images were taken (Fig. 1b). Green AgmF was found in both the cell body and cell neurite.

**AgmF can Protect the Neuronal Cells from NMDA-Induced Neurotoxicity**

NMDA receptors are found in the neuron cell body, and synapses are responsible for important physiological and pathological conditions of neurons. Depending on the NMDA treatment concentration, NMDA receptors can regulate the intercellular response bi-directionally. Lower and higher NMDA concentrations in NMDA treatment can activate the survival and pro-death signaling in neurons [27]. To check the role of AmgF in cortical neurons, we treated the neurons...
with a higher concentration of NMDA (50 µM) for 1 h, followed by AgmF (100 µM) for 6 h, and fixed cells were immunostained with the neuron marker DCX. We found that AgmF itself does not have any effect on neurons. Treatment with NMDA reduced the number of DCX-positive cells to 31% as expected when compared with the control (100%). However, NMDA-treated cells, followed by AgmF, were approximately 37% higher than the NMDA-only treated group (Fig. 2a, b). The above results suggest that AgmF does not have any harmful effect on neurons and is neuroprotective in high NMDA treatment.

Comparison of the Neuroprotective Effects Between Agmatine and AgmF

The physiological function of endogenous agmatine is largely unknown. However, several studies have reported that exogenous agmatine treatment can reduce neuronal injury in both in vitro and in vivo disease models. With this view, we investigated the changes in neuroprotection by agmatine when the amino end is blocked, and only the guanidine end is active. In this investigation, NMDA-treated cells were treated with agmatine and AgmF for 6 h. Crystal violet staining of the fixed cells showed that agmatine (63% of the control) had a greater number of cells (about 41% more) than the NMDA (21% of control) (Fig. 3a, b). The AgmF (about 54% of the control) treatment also has a significantly higher (about 33% more) number of cells than the NMDA-treated cells. However, the difference in the cell number between agmatine and AgmF was not significant, at approximately 8%. Again, in the MTT assay, the NMDA treatment reduced the cell viability to 54%, whereas NMDA treatment followed by agmatine and AgmF maintained the cell viability at 82% and 91%, respectively, compared to the control; this was significantly higher than that of the NMDA-only treatment (Fig. 3c). The above results suggest that there was no significant difference in the neuroprotective functions of agmatine and AgmF.

AgmF Reduced no Production by Attenuating NOS2 Expression

One of the mechanisms of neuroprotection by agmatine is the reduction of NO production. To investigate the involvement of this mechanism in AgmF-mediated neuroprotection, we performed the NO assay after treating the cells with NMDA, agmatine, and AgmF. NMDA treatment increased NO production (1.3 µM) by about five-fold compared to the control (0.3 µM). However, both agmatine and AgmF treatment significantly reduced NO production (0.7 and 0.6 µM) when compared with NMDA (Fig. 4a). Immune staining of the cells with NOS2 showed that NOS2 expression was increased by NMDA treatment; however, both agmatine and AgmF treatment followed by NMDA treatment significantly reduced NOS2 expression (Fig. 4b). The NOS+ neurons are counted and found that NMDA has increased the NOS2 expressing cells about 60% more than the control cells. However, the agmatine and the AgmF treatment have reduced the number of the NOS+ about 50% of the NMDA treated cells (Fig. 4c). The above results show that AgmF exerts neuroprotective effects by reducing NO production, which is achieved by inhibiting NOS2 expression.
Discussion

In our study, we demonstrated the procedure of making a fluorescence agmatine by binding a FITC molecule at the amino end of the agmatine and the guanidine end remains active. After producing the fluorescence-agmatine we also evaluated that it protects against NMDA-mediated injury in neuronal cultures. The NMDA treatment showed that the number of cells with neurites gradually reduced with time. The a–d are the 4× enlarged images of the corresponding white square boxed images. The treatment of AgmF with or without NMDA groups showed that the number of cells with neurites and cell bodies was significantly higher than that in the NMDA-treated cells. Scale bar: 50 μm. *P < 0.001 versus the NMDA, #P < 0.001 versus the cortical and $p < 0.01 versus NMDA + AgmF.

Agmatine-expressing cells have been found in all regions of the brain, such as the hypothalamus, frontal cortex, striatum, medulla, hippocampus, and locus coeruleus (LC), with measurable ADC activity [28]. However, the highest number of agmatine-expressing cells was found in the hypothalamus, which also had the highest ADC activity [28, 29]. There are diverse mechanisms of neuroprotection in different neurodegenerative diseases by agmatine that have been reported by different research groups [16, 30, 31] and also by us. Agmatine belongs to the class of organic compounds known as guanidines. Structurally, agmatine has two ends that consist of a guanidine moiety and an amino group [32]. In this study, we blocked the amino group by binding a FITC molecule and confirmed the binding of agmatine to FITC by NMR. The binding of FITC to the amino group did not change the neuronal uptake of agmatine. Although the actual function of endogenous agmatine has not yet been discovered, it is well known that exogenous agmatine can act as a neurotransmitter and protect neurons in different diseases such as AD, PD, ischemic stroke, traumatic brain injury, spinal cord injury, etc. [4, 14, 33–35]. In pathological conditions, increased extracellular glutamate over-activates the NMDARs, resulting in elevated Ca²⁺ and Na⁺ influx into the cell, which triggers NMDAR-mediated neuronal
excitotoxicity and causes neuronal cell death [36–38]. In our study, we found that NMDA treatment reduced the number of DCX-positive cells, but there was no change in the AgmF-treated cells. Moreover, AgmF inhibited the cytotoxic effect of NMDA. Therefore, even if the free amino end was blocked, agmatine would not be harmful to the neurons. Given that agmatine and AgmF might differ with respect to neuroprotection, we studied neuronal viability by agmatine and AgmF following NMDA treatment. The neuronal viability by AgmF was found to be similar to that of agmatine.

NMDAR over-activation by exogenous NMDA treatment produces NO by activating NOS, which is a major mediator of neuronal death [39–41]. As a result, controlling neuronal excitotoxicity and the inhibition of NO through...
In conclusion, we suggest that the guanidine end of agmatine regulates its neuroprotective function. As of today, there is no agmatine conjugated with a fluorescent compound. Our modified FITC AgmF can be used in studies in which fluorescence agmatine is needed.

Supplementary Information The online version contains supplementary material available at https://doi.org/10.1007/s11064-021-03319-9.

Acknowledgements The authors thank Dr. Hyun Ji Young for assistance with the preparation of the Agmatine-FITC. This study was supported by a National Research Foundation of Korea grant funded by the Korea government (MSIP) (NRF-2017R1A2B2005350 and NRF-2021R1A2C2008034).

Author Contributions JEL provided concept, design and overall supervision of this study. SB and AYS contributed in the experiment and writing. SB, AYS, KJY, IJS and JEL participated in the discussion and revision. All authors approved and agreed to be accountable for all aspects of the work.

Data Availability The datasets generated during and analysed during the current study are available from the corresponding author on reasonable request.

Declarations

Conflict of interest The authors declare no conflict of interest in relation to the present review paper.

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