Low Non-NMDA Receptor Current Density as Possible Protection Mechanism from Neurotoxicity of Circulating Glutamate on Subfornical Organ Neurons in Rats

Wonee Chong¹, Seong Nam Kim¹, Seong Kyu Han², So Yeong Lee¹, and Pan Dong Ryu¹

¹Department of Veterinary Pharmacology, College of Veterinary Medicine, Seoul National University, Seoul 151-742. ²Department of Oral Physiology, School of Dentistry and Institute of Oral Bioscience, Chonbuk National University, Jeonju 561-756, Korea

INTRODUCTION

The subfornical organ (SFO) is one of circumventricular organs characterized by the lack of a normal blood brain barrier. The SFO neurons are exposed to circulating glutamate (60–100 μM), which may cause excitotoxicity in the central nervous system. However, it remains unclear how SFO neurons are protected from excitotoxicity caused by circulating glutamate. In this study, we compared the glutamate-induced whole cell currents in SFO neurons to those in hippocampal CA1 neurons using the patch clamp technique in brain slice. Glutamate (100 μM) induced an inward current in both SFO and hippocampal CA1 neurons. The density of glutamate-induced current in SFO neurons was significantly smaller than that in hippocampal CA1 neurons (0.55 vs. 2.07 pA/pF, p < 0.05). To further identify the subtype of the glutamate receptors involved, the whole cell currents induced by selective agonists were then compared. The current densities induced by AMPA (0.45 pA/pF) and kainate (0.83 pA/pF), non-NMDA glutamate receptor agonists in SFO neurons were also smaller than those in hippocampal CA1 neurons (2.44 pA/pF for AMPA, p < 0.05; 2.34 pA/pF for kainate, p < 0.05). However, the current density by NMDA in SFO neurons was not significantly different from that of hippocampal CA1 neurons (1.58 vs. 1.47 pA/pF, p > 0.05). These results demonstrate that glutamate-mediated action through non-NMDA glutamate receptors in SFO neurons is smaller than that of hippocampal CA1 neurons, suggesting a possible protection mechanism from excitotoxicity by circulating glutamate in SFO neurons.

Key Words: Circumventricular organs, Excitotoxicity, Hippocampus, Non-NMDA, Slice patch clamp

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Corresponding to: Pan Dong Ryu, Department of Veterinary Pharmacology, College of Veterinary Medicine, Seoul National University, 1 Gawnak-ro, Gwanak-gu, Seoul 151-742, Korea. (Tel) 82-2-880-1254, (Fax) 82-2-879-0378, (E-mail) pdryu@snu.ac.kr

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ABBREVIATIONS: SFO, Subfornical organ; AMPA, α-Amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid; NMDA, N-Methyl-D-aspartic acid; HEPES, 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid; EGTA, ethylene glycol tetraacetic acid; ACSF, artificial cerebrospinal fluid.
toxicity of circulating glutamate. In this study, to figure out the possible protection mechanism for SFO neurons from excitotoxicity induced by circulating glutamate, we compared the glutamate-induced current density in SFO neurons and hippocampal CA1 neurons in brain slice using the patch clamp technique.

METHODS

Animals

All experiments in the present study were carried out according to the protocol (SNU-100824-4) for care and use of animals approved by the Laboratory Animal Care Advisory Committee of Seoul National University. All effort was made to minimize the number of animals used and their suffering.

Brain slice patch-clamp recording

For recording the currents from SFO and hippocampal CA1 neurons in brain slice, the slice containing the SFO and hippocampus were prepared by the previously reported methods [14,20]. Sprague-Dawley rats (4–6 weeks old) were deeply anesthetized with diethyl ether and decapitated. The brain was quickly dissected out, immersed in cold artificial cerebrospinal fluid (ACSF, 0–4°C) containing (in mM) 126 NaCl, 26 NaHCO3, 5 KCl, 1.2 NaH2PO4, 2.4 CaCl2, 1.2 MgCl2, and 10 glucose and aerated with O2/CO2 (95%/5%). In each rat, coronal slices (200 μm thick) were cut from a block of the forebrain and then incubated in aerated ACSF at 32°C for 60–90 min. For recording the currents from SFO and hippocampal CA1 neurons in brain slice using the slice patch clamp technique in SFO and hippocampal CA1 neurons. The brain was quickly dissected out, immersed in cold artificial cerebrospinal fluid (ACSF, 0–4°C) containing (in mM) 126 NaCl, 26 NaHCO3, 5 KCl, 1.2 NaH2PO4, 2.4 CaCl2, 1.2 MgCl2, and 10 glucose and aerated with O2/CO2 (95%/5%). In each rat, coronal slices (200 μm thick) containing the SFO and hippocampus were cut from a block of the forebrain and then incubated in aerated ACSF at 32°C for 60–90 min. For ionic current recording, a slice was transferred to the recording chamber (0.7 ml), which was continuously perfused with aerated ACSF (37°C) at the rate of 2.7 ml/min. Whole cell currents were recorded from neurons in all regions of the SFO and CA1 region of hippocampus in the slice [13,14,21]. Pipettes were filled with a solution containing (in mM) 140 KCl, 20 HEPES, 0.5 CaCl2, 5 EGTA, 5 QX-314, and 5 MgATP (pH=7.2) with 3M KOH (about 1 ml). Pipette tip resistance range was 2–5 MΩ. Individual neuron in the slice was identified using an upright microscope equipped with Nomarski optics (X400, BX50WI, Olympus, Tokyo, Japan) as reported previously [13,14,21]. Whole cell currents were measured by using a patch clamp amplifier (Axopatch 200B, Axon Instruments, USA). Current signals were filtered at 2 kHz and digitized directly into a PC at 1 kHz with the use of pClamp (Ver 7.0) and Digidata 1200A (Axon Instruments, USA) for subsequent analysis and display. The membrane potentials were held at −70 mV. The current density was obtained by dividing the whole cell current by membrane input capacitance, which was calculated from the membrane time constant. The drugs used in this study were purchased from Sigma (MO, USA).

Statistics

All values were expressed as the mean±S.E.M. A student t-test (independent) was used to compare the mean amplitudes and mean current densities. Statistical significance was set at as p<0.05.

RESULTS

Mean membrane capacitances of hippocampal CA1 neurons (121±13.2 pF, n=6) were larger than SFO neurons (39.1±4.09 pF, n=10, p<0.05), indicating that hippocampal CA1 neurons are larger than SFO neurons in their cell surface area. Bath application of glutamate (100 μM) induced inward currents in both SFO and hippocampal CA1 neurons (Fig. 1A). As shown in Fig. 1B, the amplitude of inward current at its peak in SFO neurons was much lower than that in hippocampal CA1 neurons (20.6±6.39 pA, n=10 vs. 289±146 pA, n=6, p<0.05). When the currents were normalized by the mean membrane capacitance, the mean current density of SFO neurons (0.55±0.15 pA/pF) was much smaller than that in hippocampal CA1 neurons (2.07±0.79 pA/pF, p<0.05, Fig. 1C). These results indicate that the maximal response of glutamate in SFO neurons is less than that in hippocampal CA1 neurons in slice preparation.

To further determine the subtype of glutamate receptors mediating the glutamate-induced inward currents, we compared the whole cell currents induced by non-NMDA receptor agonists (AMPA and kainate) in SFO and hippocampal CA1 neurons. Bath application of AMPA and kainate (10, 30, and 100 μM) induced a reversible and concentration-dependent inward current as reported previously [1]. In this study, we compared the current response at the threshold concentration (10 μM) that induced an inward current in all tested cells. As shown in Fig. 2A, AMPA and kainate induced inward currents both in SFO and hippocampal CA1 neurons. The inward currents induced by AMPA (10 μM) and kainate (10 μM) in SFO neurons were significantly smaller than those in hippocampal CA1 neurons (17.3±3.79 pA, n=13 vs. 208±24.4 pA, n=11, p<0.05 for AMPA; and 24.9±3.44 pA, n=11, vs. 202±44.1 pA, n=7, p<0.05 for kainate), respectively (Fig. 2B). When we normal...
Reduced Glutamate Current in SFO Neurons

Fig. 2. Current responses elicited by non-NMDA receptor agonists (AMPA and kainate) in SFO and hippocampal CA1 neurons in the slice preparation. (A) Representative traces showing currents induced by AMPA (10 μM) and kainate (10 μM) in SFO (left) and hippocampal CA1 neuron (right), respectively. (B, C) Mean amplitude (B) and mean density (C) of inward currents induced by AMPA and kainate in SFO (n=13 and 11) and hippocampal neurons (n=11 and 7) at the peak (*p<0.05). SFO, Subfornical organ neuron; HP, Hippocampal CA1 neuron.

Fig. 3. Current responses elicited by NMDA in SFO and hippocampal neurons. (A) Left and right traces show NMDA (10 μM) induced inward current in SFO and hippocampal CA1 neurons. Mg^2+-free ACSF was applied for recording of NMDA induced current. (B, C) Mean inward current induced by NMDA (B) and its mean current density (C) in SFO (n=4) and hippocampal CA1 neurons (n=4). SFO, Subfornical organ neuron; HP, Hippocampal CA1 neuron.

ized the AMPA- and kainate-induced currents by the capacitance of each neuron, we found that the densities of AMPA- and kainate-induced currents in SFO neurons were also significantly smaller than those in hippocampal CA1 neurons (0.45±0.10 vs. 2.44±0.28 pA/pF, p<0.05; 0.83±0.20 vs. 2.34±0.32 pA/pF, p<0.05), respectively (Fig. 2C). The current density ratios of SFO neurons to hippocampal CA1 neurons were 0.18 and 0.35 for AMPA and kainate, respectively.

For comparison of the inward current induced by glutamate through the activation of NMDA receptors, we recorded NMDA (10 μM)-induced currents in the ACSF without Mg^2+, which is known to block NMDA current [22] as illustrated in Fig. 3. The mean inward currents by NMDA from SFO and hippocampal CA1 neurons were 56.9±9.02 pA (n=4) and 154±63.2 pA (n=4), respectively (Fig. 3B). As shown in Fig. 3C, the normalized mean current densities were not significantly different (1.38±0.31 [n=4] vs. 1.47±0.48 pA/pF [n=4]).

**DISCUSSION**

This study is the first to demonstrate a protection mechanism of SFO neurons from possible excitotoxicity of circulating glutamate. The density of glutamate-induced current in SFO neurons is 4 times lower than that in hippocampal neurons, and this difference is mostly due to the smaller responses mediated by non-NMDA receptor subtypes.

It is well known that glutamate-induced neurotoxicity is caused by an intracellular accumulation of Ca^{2+} via mainly AMPA and NMDA receptors [16,17,23]. Kainate can also induce neurotoxicity in various brain sites [24,25]. AMPA and kainate receptors trigger a rapid excitatory neurotransmission by promoting the influx of Na^+ into neurons. NMDA receptors are associated to a high conductance with calcium channel which is blocked by Mg^2+ in non-depolarizing condition in a voltage dependent manner. Their activation is secondary to AMPA or kainate receptor activation, which depolarizes the neuron, allowing for the release of the Mg^2+ blockade. Therefore, AMPA and kainate receptors play a major role in the initiation of the excitotoxicity process in neurons [26]. In this study, we also found that non-NMDA receptor responses are selectively lower in the SFO neurons than in the hippocampal neurons. Since the SFO neuronal cell bodies are located out of blood brain barrier, circulating glutamate can freely access the SFO neurons and its concentration at the SFO is likely to be sufficiently high to induce excitotoxicity reported in general central neurons, such as hippocampal neurons. However, SFO neurons in adult rat normally do not suffer from excitotoxicity and are sensitive to exogenously applied glutamate [27-29]. Therefore, it is likely that circulating glutamate causes much smaller inward current in SFO neurons and the threshold of the excitotoxicity of glutamate is much higher in SFO neurons than in the other central neurons.

In the experiments comparing the densities of ionotropic glutamate receptor agonist-induced currents, the current density ratios of SFO neurons to hippocampal CA1 neurons were 0.18 for AMPA and 0.35 for kainate, respectively. However, there was no significant difference in current density by NMDA between SFO neurons and hippocampal CA1 neurons. These data indicate that the lower density of glutamate-induced currents in SFO neurons is due to a selective protection mechanism, which is due to the lower density of glutamate receptors in SFO neurons.


The protection mechanism in SFO neurons is, at least in part, due to the higher threshold for the initiation of excitotoxicity by glutamate.

In SFO, there may be other protection mechanisms such as the effective removal of glutamate transporters in the SFO [6,30]. The relative significance among different protection mechanisms remains to be studied further. Since the density of glutamate subtypes was functionally assessed in this study, it would be necessary to perform a direct comparison of the densities of glutamate receptor subtypes between hippocampal CA1 and SFO neurons. In addition, the current densities of NMDA receptors recorded in ACSF without Mg2+ were not significantly different between the two neuron groups (Fig. 3). Direct measurement of NMDA receptors expressed on two neuron groups is also needed.

Taken together, this study provides a new insight that the current density of non-NMDA glutamate receptors on SFO neurons is lower than other central neurons, such as CA1 neurons, that are selectively vulnerable to ischemic excitotoxicity, and suggest that lower current density of non-NMDA glutamate receptors could be an effective protection mechanism for the SFO neurons from excitotoxicity of circulating glutamate.

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