Novel glycine-dependent inactivation of NMDA receptors in cultured hippocampal neurons

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Abstract: Objective Glycine acts as a co-agonist for the activation of N-methyl-D-aspartate receptors (NMDARs) by binding to glycine sites, thus potentiating glutamate-elicited responses and inhibiting NMDAR desensitization in a dose-dependent manner. The present study aimed to characterize the glycine-dependent inactivation of NMDARs and to explore its pathophysiological significance. **Methods** Primary hippocampal cell cultures from embryonic days 17–18 rats were treated with NMDA or NMDA plus glycine. Patch-clamp recording and intracellular Ca^{2+} imaging were performed to test the effects of glycine on NMDA-activated currents and increase of intracellular free Ca^{2+} respectively. Immunofluorescence staining was conducted to examine NR1 internalization. Cell damage was tested with MTT method and lactate dehydrogenase leakage. **Results** Glycine reduced the peak current and Ca^{2+} influx elicited by NMDA application at concentrations $\geq 300 \mu\text{mol/L}$. This is a novel suppressive influence of glycine on NMDAR function, since it occurs *via* the NMDAR glycine-binding site, in contrast to the classic suppression, which occurs through the binding of glycine to glycine receptors. The level of membrane NMDARs was measured to evaluate whether internalization was involved. Immunohistochemical labeling showed that incubation with high concentrations of NMDA plus glycine did not change the expression of NMDARs on the cell surface when compared to the expression without glycine; hence the possibility of NMDAR internalization primed by glycine binding was excluded. **Conclusion** In summary, the novel suppressive effect of glycine on NMDARs was mediated *via* binding to the glycine site of the NMDAR and not by activation of the strychnine-sensitive glycine-receptor-gated chloride channel or by the internalization of NMDARs. The inhibitory influence of glycine on NMDARs adds a new insight to our knowledge of the complexity of synaptic transmission.

Keywords: glycine; N-methyl-D-aspartate receptors; calcium imaging; NMDA-NR1 subunit; hippocampal neurons; inactivation

1 Introduction

The N-methyl-D-aspartate receptor (NMDAR), one of the ionotropic glutamate receptors, plays a key role in

physiological and pathophysiological processes such as the development and modification of synaptic plasticity, long-term potentiation and long-term depression^[1,2], as well as in such disorders as ischemia, epilepsy, trauma, and dementia^[1,3]. NMDARs are abundantly distributed throughout the central nervous system. Currently, they are considered to be unique among ligand-gated ion channels in their requirement for a necessary co-agonist, glycine, in

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addition to the synaptic neurotransmitter glutamate^[4,5]. Two independent binding sites have been identified for glycine and glutamate, located on the NR1 and NR2 subunits, respectively^[6,7]. It has been found that, upon binding to the glycine site, glycine potentiates the glutamate-elicited response and inhibits the desensitization of NMDARs^[4,8,9]. To study the regulatory role of glycine in detail, NMDA-mediated currents and Ca^{2+} influx were assayed in primary hippocampal neurons from juvenile rats. We were surprised to find a novel inhibitory influence of glycine on NMDARs when NMDA was administered at a high concentration. The present study was designed to characterize this glycine-dependent inactivation of NMDARs and to explore its pathophysiological significance.

2 Materials and methods

2.1 Animals and chemicals Sprague-Dawley rats were obtained from the Experimental Animal Center of Nantong University, Nantong, China. All procedures were in accordance with the Institutional Guidelines of Nantong University, which comply with international rules and policies, and approved by the Animal Care and Use Committee. Common inorganic salts (NaCl, KCl, CaCl_2 , MgCl_2 and NaOH) were from Sinopharm Chemical Reagent Co. Ltd. (Shanghai, China); dimethylformamide, paraformaldehyde, sucrose, and glucose from Xilong Chemical Co. Ltd. (Guangzhou, China); and dimethylsulfoxide (DMSO) from Shenhe Chemical Reagent Co. Ltd. (Shanghai, China). Chemicals used for neuronal culture and the acetoxymethyl-ester form of Fluo-3/AM were from Invitrogen Corp. (Carlsbad, CA). NMDA, glycine, MK-801, *DL*-2-amino-4-phosphonobutyric acid (*DL*-AP-4), dynasore, β -cytosine-*D*-arabinofuranoside, cesium methanesulfonate ($\text{CH}_3\text{O}_3\text{SCs}$), 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2H-tetrazolium bromide (MTT), EGTA, CsOH, HEPES, Mg-ATP, Na-GTP, sodium dodecylsulfate, and other chemicals (unless indicated elsewhere) were from Sigma-Aldrich Corp. (St. Louis, MO).

2.2 Cell culture and transfection Primary hippocampal cell cultures were prepared as described^[10]. Dissociated hippocampal cells from embryonic days 17–18 rats were

plated onto poly-*D*-lysine-coated 96-well plates or coverslips ($1 \times 1 \text{ cm}^2$). Cells were grown in the plating medium with 85% high-glucose Dulbecco's minimum essential medium (DMEM, 12100-046) and 15% fetal bovine serum (Hyclone, Logan, UT), and were incubated at 37°C in a humidified atmosphere of 5% CO_2 . The plating medium was replaced with fresh neurobasal medium containing 98% neurobasal medium, 2% B27 and 2 mmol/L glutamine (all from Invitrogen) every 4 days. Neurons were transfected with pcDNA3.1-GFP-NR1-1a at 7 days *in vitro* (DIV) using a Lipofectamine 2000 transfection kit (Invitrogen) to test NR1 internalization.

2.3 Patch-clamp recording of NMDA-activated currents Current responses (I_{NMDA}) were elicited in cultured hippocampal neurons (12–14 DIV) perfused with NMDA. Patch pipettes had resistances of 5–8 M Ω when filled with the following solution (in mmol/L): $\text{CH}_3\text{O}_3\text{SCs}$ 135, NaCl 8, HEPES 10, EGTA 0.5, Mg-ATP 4, and Na-GTP 0.3 (pH 7.2, adjusted with 1 mol/L CsOH, and osmolarity 300–310 mOsm adjusted with sucrose). Recordings of I_{NMDA} were performed in the following bath solution (in mmol/L): NaCl 150, KCl 3, HEPES 10, $\text{CaCl}_2 \cdot 2\text{H}_2\text{O}$ 3, and glucose 8 (pH 7.4, adjusted with 1 mol/L NaOH, and osmolarity 300–310 mOsm adjusted with sucrose), using a Multi-Clamp 700A amplifier, Digidata 1320A AD converter and pCLAMP 8.0 software (Axon Instruments, Foster City, CA). The junction potential between micropipette and bath solution was –9.9 mV, as calculated using Clampex 8.0 (in pCLAMP 8.0) and nulled just before forming a giga-seal. In most experiments, the series resistance (R_s) before compensation was 10–20 M Ω . Routinely, 70%–80% of the R_s was compensated. Only neurons with an R_s <20 M Ω were selected for further tests. All NMDA currents were recorded while the membrane potential was held at –70 mV, except as otherwise noted.

The application of NMDA onto cultured neurons was performed *via* a perfusion system (MPS-2; World Precision Instruments, Sarasota, FL). The tip (diameter 250 μm) of the drug application pipette was usually placed 100 μm away from the recorded cell. To avoid drug accumulation in the bath, each 2-s pulse of drug application was fol-

lowed by 2 min of rinsing. The ambient temperature was held between 23 and 25°C.

The degree of desensitization of NMDA currents was determined according to the following equation: desensitization score = $1 - I_{ss}/I_p$, where I_p is the peak current, and I_{ss} is the steady-state current of I_{NMDA} (Fig. 1). The degree of desensitization was positively related to the calculated desensitization score.

Fig. 1. NMDA-activated current (I_{NMDA}) recorded using whole-cell mode in a cultured hippocampal neuron. I_p , peak current; I_{ss} , steady-state current of I_{NMDA} .

2.4 Intracellular Ca^{2+} imaging Fluo-3/AM, a fluorescent Ca^{2+} indicator, was dissolved in DMSO with the final working concentration of DMSO at 0.1%. At 12–14 DIV, the cultured hippocampal neurons were loaded with 5 μ mol/L Fluo-3/AM for 45 min at 37°C in the bath solution. After washing three times in fresh bath solution, the neurons were incubated at 37°C for another 30 min to complete the de-esterification of Fluo-3/AM. The fluorescence intensity (excitation at 485 nm, emission at 525 nm) was recorded every 10 s for 5 min using a laser scanning confocal microscope (TCS SP2, Leica Microsystems, Heidelberg, Germany). All image data were collected and analyzed with Leica software. The increase of intracellular free Ca^{2+} ($[Ca^{2+}]_i$) was determined according to the following equation: $[Ca^{2+}]_i$ increase (%) = $(F_{525} - F_{base, 525}) / F_{base, 525} \times 100\%$, where F_{525} is the fluorescence intensity measured after each treatment, and $F_{base, 525}$ is the basal fluorescence intensity.

2.5 Immunofluorescence procedures and internalization assays Hippocampal neurons at 14 DIV were used for the internalization assays. Neurons were pretreated with DMSO or 80 nmol/L dynasore for 10 min, then NMDA or NMDA plus glycine were added to the medium at 37°C for 3 min. For live staining, the above neurons, which had been transfected with pcDNA3.1-GFP-NR1-1a at 7 DIV (against the extracellular region), were labeled with rabbit anti-GFP antibody (1:200; Chemicon) in cold Hank's balanced salt solution (HBSS; Gibco, Carlsbad, CA) for 30 min to visualize the surface receptors. All procedures were conducted at 4°C to prevent receptor-antibody internalization. The cells were then extensively washed with cold HBSS prior to incubation with Cy3 goat anti-rabbit IgG (H + L) (1:1 000; Invitrogen) in HBSS for 10 min. Subsequently, the cells were washed with HBSS at 4°C. For endogenous receptor staining, the neurons were fixed with 4% paraformaldehyde in phosphate-buffered saline for 10 min at room temperature (RT), then permeabilized in ice-cold methanol for 8 min at -20°C to measure the total receptors, washed in wash buffer (WB, 10 mmol/L phosphate buffer, pH 7.4) three times for 5 min at RT and incubated with rabbit anti-GFP antibody (1:200; Chemicon) in staining buffer (SB, 100 mmol/L phosphate buffer, pH 7.4, containing 0.5% Triton X-100 and 1% bovine serum albumin) overnight at 4°C. The cells were then washed three times in WB for 10 min at RT. Following incubation with secondary antibody (Alexa Fluor 488 goat anti-rabbit IgG (H + L), 1:1 000, Invitrogen) in SB for 1 h at room temperature, the cells were washed three times in WB for 10 min at RT. Immunofluorescence images were then collected and processed with a laser scanning confocal microscope (TCS SP2, Leica Microsystems). NR1 internalization assays and quantitation were performed according to the ratio of surface/total receptor expression.

2.6 Measurement of cell damage Cell damage was assessed by measuring neuronal viability with the MTT method, as well as lactate dehydrogenase (LDH) leakage. After 1 h of NMDA exposure, the culture medium was removed from the 96-well plate, and the neurons (12–14 DIV) were rinsed twice with bath solution. Then 25 μ L

MTT was added to each well to a final concentration of 0.1%. Following incubation at 37°C for 4 h, 100 μ L 20% sodium dodecylsulfate (dissolved in dimethylformamide) was mixed into each well and left for 20 h to dissolve the resultant dark blue crystals. The absorbance of each well at 570 nm (OD_{570}) was measured with a Universal Microplate Reader (ELx800, BioTek Instruments, Inc., Winooski, VT).

LDH activity was measured with a kit (Jiancheng Bioengineering Institute, Nanjing, China) according to the manufacturer's instructions. Briefly, after 1 h of NMDA exposure, 0.4 mL culture medium was mixed with 1.3 mL nicotinamide adenine dinucleotide (NADH) and 1.3 mL sodium pyruvate. Both agents were dissolved in potassium phosphate buffer (100 mmol/L K_2HPO_4 , pH adjusted to 7.5 with KH_2PO_4). The mixed solution was immediately assayed with a spectrophotometer (UV-2450, Shimadzu Corp., Kyoto, Japan) by monitoring the conversion of NADH to NAD at 340 nm at 37°C, coupled with the reduction of pyruvate to lactate. LDH activity is expressed as units/mL, with one unit representing the amount of LDH that causes a decrease of 0.001 absorbance units of NADH per minute in the presence of sodium pyruvate. The same experiments were repeated once again to measure LDH and neuronal viability.

2.7 Statistical analysis Clampfit 8.0 (Axon Instruments, Foster City, CA), ImageJ (National Institutes of Health, USA) and SigmaPlot 2001 (Jandel Scientific, Costa Madre, CA) were used for data analysis and plotting. All data are presented as mean $\pm$ SD. Data from multiple groups were analyzed with one-way ANOVA and *post hoc* Newman-Keuls test. $P < 0.05$ was considered statistically significant.

3 Results

3.1 Inhibitory influence of glycine on NMDA-elicited current Whole-cell recordings were performed on cultured hippocampal neurons at 12–14 DIV. When neurons were clamped near the resting potential in Mg^{2+} -free extracellular solution, 30 μ mol/L NMDA-evoked inward peak currents showed dose-dependent glycine activation in all neurons tested (Fig. 2A). Meanwhile, the desensitization score of I_{NMDA} decreased with increasing glycine concen-

tration (Fig. 2C). However, when the NMDA concentration was raised to 300 μ mol/L, we stumbled across the inhibition of the peak current by glycine (Fig. 2B), while the desensitization score still decreased with increasing glycine concentration (Fig. 2D). This is inconsistent with the previous view that glycine acts as a co-agonist for NMDAR activation by binding to the glycine site, potentiates the glutamate-elicited response, and decreases the NMDAR desensitization in a dose-dependent manner. Besides, we used *DL*-AP-4, a non-competitive NMDAR antagonist as a positive control to test the inhibitory effect on both 30 μ mol/L and 300 μ mol/L NMDA-evoked currents. Results showed that pre-incubation with *DL*-AP-4 for 10 min inhibited the NMDA current significantly while pretreatment for 20 min suppressed the inward current. Examples of the effect of glycine on the peak current and the desensitization of I_{NMDA} with the application of NMDA at 30 and 300 μ mol/L are shown in Fig. 2E and F, respectively.

3.2 Inhibitory influence of glycine on NMDA-elicited Ca^{2+} influx To verify the above results, we performed free- Ca^{2+} imaging by confocal microscopy in living neurons. Similar to the results of I_{NMDA} recording, glycine increased Ca^{2+} influx as a function of concentration in the presence of 30 μ mol/L NMDA (Fig. 3A, C), but decreased Ca^{2+} influx when 300 μ mol/L NMDA was applied (Fig. 3B, D). We also found that the peak $[Ca^{2+}]_i$ induced by NMDA increased in a dose-dependent manner without the addition of glycine (Fig. 3E, F). However, it changed in a different way in the presence of glycine, i.e., when the concentration of NMDA was continuously augmented in the presence of glycine at 0.3 μ mol/L, glycine elevated the amplitude of the $[Ca^{2+}]_i$ increase evoked by NMDA at 10 and 30 μ mol/L, but not at 100 μ mol/L, and it significantly reduced the amplitude of the $[Ca^{2+}]_i$ elevation elicited by NMDA at 300 μ mol/L and 1 mmol/L (Fig. 3E, F).

3.3 NMDAR internalization does not occur on the cell membrane, regardless of being primed by glycine binding

It is possible that the observed inhibitory effect was due to decreased numbers of cell-surface NMDARs; in other words, high concentrations of NMDA plus glycine may contribute to the priming of receptor internalization, which

Fig. 2. Effects of glycine (Gly) on NMDA-activated peak current and desensitization. **A:** Effects of glycine on the peak I_{NMDA} elicited by 30 $\mu\text{mol/L}$ NMDA ($n = 9$) with or without pretreatment with 1 mmol/L DL-AP-4 (AP) for 10 min and 20 min ($n = 3$). **B:** Effects of glycine on the peak I_{NMDA} elicited by 300 $\mu\text{mol/L}$ NMDA ($n = 7$) with or without pretreatment with 1 mmol/L DL-AP-4 for 10 min and 20 min ($n = 3$). **C:** Influence of glycine on the desensitization score of I_{NMDA} elicited by 30 $\mu\text{mol/L}$ NMDA ($n = 9$) and inhibition by the antagonist, 1 mmol/L DL-AP-4 ($n = 3$), as control. **D:** Influence of glycine on the desensitization score of I_{NMDA} elicited by 300 $\mu\text{mol/L}$ NMDA ($n = 7$) and inhibition by the antagonist, 1 mmol/L DL-AP-4 ($n = 3$), as control. **E:** Examples of the effect of glycine on 30 $\mu\text{mol/L}$ NMDA-evoked currents with or without DL-AP-4. **F:** Examples of the effect of glycine on 300 $\mu\text{mol/L}$ NMDA-evoked currents with or without DL-AP-4. * $P < 0.05$ vs NMDA without glycine; $\Delta P < 0.05$ vs with 0.1 $\mu\text{mol/L}$ glycine.

would lead to a decrease in NMDAR currents. Thus, we sought to determine whether or not the expression of NMDAR on the neuronal surface was reduced by 300 $\mu\text{mol/L}$ NMDA plus glycine. For this, we used NR1-GFP extracellular domain recombinant protein and live staining at 4°C, both of which block the further endocytosis of membrane receptors. However, there was no significant differ-

ence in receptor expression after the application of 300 $\mu\text{mol/L}$ of NMDA, with or without glycine. In contrast, when we pretreated the cultured hippocampal neurons with 80 $\mu\text{mol/L}$ dynasore, a cell-permeable dynamin inhibitor that prevents endocytosis, we found that the expression of membrane receptors increased significantly in comparison with non-pretreated neurons (Fig. 4A, B).

Fig. 3. Effects of glycine (Gly) on NMDA-elicited Ca^{2+} influx. **A** and **C**: Dose-dependent facilitating effect of glycine on Ca^{2+} influx elicited by 30 $\mu\text{mol/L}$ NMDA ($n = 20$). **B** and **D**: Dose-dependent inhibitory effect of glycine on Ca^{2+} influx elicited by 300 $\mu\text{mol/L}$ NMDA ($n = 20$). **E** and **F**: Effect of glycine on the dose-response of NMDA-elicited Ca^{2+} influx ($n = 20$). The effects of MK-801 and DL-AP-4 served as controls. * $P < 0.05$, ** $P < 0.01$ vs NMDA without glycine; $\Delta P < 0.05$, $\Delta\Delta P < 0.01$ vs with 0.1 $\mu\text{mol/L}$ glycine; $\phi P < 0.01$ vs 10 $\mu\text{mol/L}$ NMDA alone.

Fig. 4. Immunofluorescence images of NR1 expression in the hippocampal neuron. **A:** Immunofluorescence to NR1-GFP-Alex 488 (left column, green) and NR1-GFP-Cy3 (middle column, red) was performed in permeabilized (total NR1; left columns) and non-permeabilized (membrane NR1; middle columns) cells to analyze the patterns of NR1 expression of surface/total colocalization. Areas of overlapping (yellow, right columns) indicate the puncta of NR1 expression on the cell surface of membrane. Upper panels: 300 μmol/L NMDA; middle and lower panels: 300 μmol/L NMDA plus 10 μmol/L glycine (Gly) with (lower) and without (middle) 10-min pretreatment with 80 μmol/L dynasore at 37°C. Scale bars, 20 μm. **B:** Histogram showing the ratio of surface/total NMDARs (mean ± SEM) quantified by live staining of hippocampal neurons in primary culture. Cultures ($n = 25, 29$, and 18 in three separate experiments) were exposed to 300 μmol/L NMDA alone or with 10 μmol/L glycine with or without pretreatment with 80 μmol/L dynasore. ** $P < 0.01$ vs NMDA plus glycine without dynasore. ESC, extracellular solution.

3.4 Role of the glycine-binding site in the inhibitory influence of glycine on NMDA responses To identify the potential target of the inhibitory effect of glycine on NMDARs, we tested whether the phenomenon still occurred after pretreatment with 7-CK (an antagonist at the NMDAR glycine site) or strychnine (an antagonist of glycine receptors). We found that 7-CK blocked the glycine-mediated inhibition of the $[Ca^{2+}]_i$ increase induced by 1 mmol/L NMDA, while 7-CK itself did not have any effect

on Ca^{2+} influx. On the contrary, strychnine at 10 nmol/L did not have any influence on the inhibitory effect of glycine on the increase of $[Ca^{2+}]_i$ evoked by 1 mmol/L NMDA (Fig. 5A, B).

3.5 Influence of glycine on viability of neurons exposed to NMDA To investigate the potential pathophysiological significance of the inhibitory influence of glycine on NMDARs at higher concentrations of NMDA, we measured the viability of neurons and LDH leakage from

Fig. 5. A: Blockade by the glycine-binding site antagonist 7-CK of the inhibitory effect of glycine on Ca^{2+} influx elicited by high concentration of NMDA ($n = 16$). B: Histograms show the effects of the glycine (Gly) site inhibitor 7-CK and the glycine receptor antagonist strychnine (Stry) on Ca^{2+} influx. $**P < 0.01$ vs NMDA without glycine.

Fig. 6. Influence of glycine (Gly) on the cell-damaging effects of different concentrations of NMDA. **A:** LDH leakage from cultured hippocampal neurons with 1-h exposure to 30 μmol/L NMDA and different concentrations of glycine ($n = 8$). **B:** Cell viability after exposure to 30 μmol/L NMDA and different concentrations of glycine ($n = 10$). **C:** LDH leakage from hippocampal neurons exposed to 300 μmol/L NMDA and different concentrations of glycine ($n = 8$). **D:** Neuronal viability after exposure to 300 μmol/L NMDA and different concentrations of glycine ($n = 10$). * $P < 0.05$, ** $P < 0.01$ vs control (Con); $^{\Delta}P < 0.05$, $^{\Delta\Delta}P < 0.01$ vs NMDA alone.

hippocampal neurons exposed to combinations of different concentrations of NMDA and glycine. We found that glycine dose-dependently increased the LDH leakage (Fig. 6A) and reduced the viability (Fig. 6B) when NMDA was applied at 30 μmol/L. Therefore, we postulate that glycine can facilitate NMDA-induced neurotoxicity. In contrast, when NMDA was used at 300 μmol/L, glycine reversed the NMDA-induced LDH leakage dose-dependently, and the viability was significantly elevated by glycine at 10 μmol/L (Fig. 6D, $P < 0.05$). These results indicate that glycine can protect hippocampal neurons against the excitotoxicity induced by higher concentrations of NMDA.

4 Discussion

Glycine is well known as a co-agonist in the activation of NMDARs. However, in the present study, we discovered that glycine not only played a co-agonist role but

also exerted an inhibitory influence on NMDARs if the concentration of NMDA was raised to 300 μmol/L in cultured rat hippocampal neurons. Glycine dose-dependently inhibited the peak current and Ca^{2+} influx elicited by 300 μmol/L NMDA. To our knowledge, this inhibitory effect of glycine on NMDARs has not been reported previously, and its significance as well as the specific mechanism involved has not been identified.

Many mechanisms are known to diminish the response of NMDARs, including surface membrane receptor internalization, Ca^{2+} -dependent inactivation, glycine-dependent desensitization, and glycine-independent desensitization^[11–13]. Therefore, we designed a series of experiments to investigate the problem. Because some previous studies have raised the possibility that NMDARs undergo regulated transport to and from the cell surface^[14,15], we measured the expression of surface NMDARs while applying a

high concentration of NMDA with or without glycine. Our results ruled out the possibility of membrane protein internalization.

On the other hand, Ca^{2+} -dependent inactivation is a reversible decrease in peak current, which can be induced by a rise in extracellular Ca^{2+} concentration, without agonist activation of the receptor itself^[16,17]. Desensitization is a decrease in the current response induced by the persistent presence of a glutamate site agonist^[8]. Thus, it is intriguing to consider that the inhibitory phenomenon we revealed in the present study, which is in contrast to the glycine-dependent desensitization, may be a glycine-dependent inactivation of NMDARs.

Glycine-dependent desensitization is a well-known phenomenon. While binding to the glycine site of NMDARs, glycine potentiates glutamate-elicited responses and inhibits the desensitization of NMDARs^[4,8,9]. We also found glycine-dependent desensitization. When NMDA was applied at 30 $\mu\text{mol/L}$, glycine elevated the peak current and Ca^{2+} influx and decreased the desensitization of I_{NMDA} . These results are consistent with previous reports^[4,8,9]. Glycine, a co-agonist for activation of NMDARs, is also an inhibitory neurotransmitter in the central nervous system.

Next, we sought to exclude the possibility of the involvement of glycine receptor activation by using strychnine, an antagonist of this receptor. We found that strychnine did not have any notable influence on the inhibitory effect of glycine on the increase of $[\text{Ca}^{2+}]_i$ evoked by 1 mmol/L NMDA. To identify the potential target of this glycine-dependent inactivation of NMDARs, the cultured hippocampal neurons were treated with 7-CK, an antagonist of the glycine site of NMDARs. Result showed that 7-CK blocked the inhibition by glycine of the Ca^{2+} influx evoked by 1 mmol/L NMDA. We suggest that this inactivating effect of glycine is mediated by binding to the glycine site of NMDARs.

Consequently, our results strongly suggest that the inhibitory effect we found is a novel effect of glycine, which is determined by the concentration of NMDA and glycine binding. Moreover, the inhibitory effect appears to be mediated *via* binding to the NMDAR glycine site, and not by

the activation of glycine receptors or the internalization of NMDARs primed by glycine binding.

Modulation of the NMDAR response is an important mechanism for the physiological and pathophysiological consequences of NMDAR activation^[1-3]. The concentrations of glutamate, glycine, and Ca^{2+} at the synaptic level are important determinants of the ion current flowing through the NMDAR-gated ion channel. The combination of different forms of inactivation and desensitization determines the overall degree of current response following the activation of NMDARs. Therefore, the regulation of NMDAR function, plus the effects of many other modulating factors, including polyamines, zinc, pH, redox agents and intracellular signaling molecules^[3,12], is rather complicated. The results of the present study suggest that this complexity is further increased by a novel glycine-dependent inactivation process.

In addition, we showed that continuing exposure to NMDA induced LDH release and decreased viability in cultured hippocampal neurons, and glycine facilitated this damaging effect of 30 $\mu\text{mol/L}$ NMDA. However, increasing the concentration of glycine did not increase the damaging effect of 300 $\mu\text{mol/L}$ NMDA, but decreased LDH leakage dose-dependently, indicating that a cytoprotective effect of glycine occurs when neurons are exposed to higher concentrations of NMDA. This result is in line with a previous *in vivo* experiment, in which glycine over 1 mmol/L did not significantly increase the 100 and 200 $\mu\text{mol/L}$ NMDA-evoked depolarization in the rat striatum^[18]. However, the investigators in that study did not exclude the possibility of the involvement of strychnine-sensitive glycine receptors, as glycine at the concentrations used is known to activate these receptors. Therefore, the novel inhibitory influence of glycine on NMDARs reported here has potential physiological and pathophysiological significance for protecting neurons from excessive activation of NMDARs when glutamate is released above physiological levels due to excessive glutamatergic activity.

In conclusion, our results suggest that a novel glycine-dependent inactivation occurs while NMDARs are activated by 300 $\mu\text{mol/L}$ NMDA. This inhibitory effect occurs *via*

binding to the glycine site of NMDARs but is not due to activation of the strychnine-sensitive glycine-receptor-gated chloride channel. The inhibitory influence of glycine on NMDARs adds a new insight to our knowledge about the complexity of, and feedback loops involved in, synaptic transmission.

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