

The binding site for acute corticosterone effects on *N*-methyl-D-aspartate receptor-mediated Ca²⁺ signals in mouse hippocampal slices

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ABSTRACT

We have examined acute effects of corticosterone (CORT) on *N*-methyl-D-aspartate (NMDA) receptor-mediated Ca²⁺ signals in adult mouse hippocampal slices. We found so far that the 30 min preincubation of CORT induced a significant decrease of the peak amplitude of NMDA-induced Ca²⁺ elevation in the CA1 region. The membrane non-permeable bovine serum albumin-conjugated CORT also induced a similar effect in the CA1 region. Therefore the acute CORT effects should be induced via putative surface CORT receptors. A possible candidate is a classical intracellular glucocorticoid receptor (GR). To confirm this speculation, we here examined the effects of dexamethasone (DEX: an agonist of GR) and CORT with RU38486 (an antagonist of GR) on NMDA-induced Ca²⁺ signals. As a result, DEX induced a similar effect to the suppressive CORT effect in the CA1 region, and RU38486 inhibited the suppressive CORT effect. These results indicate that the surface CORT receptor should be GR bound to plasma membrane.

Keywords: Corticosterone; Ca²⁺ Signal; NMDA Receptor; Glucocorticoid Receptor; Hippocampus

1. INTRODUCTION

Corticosterone (CORT) is a principal glucocorticoid synthesized in the rodent adrenal cortex and secreted in response to stress. There are a series of studies about the

chronic and genomic effects of corticosteroids in the hippocampus [1,2]. The stress-induced increase in CORT secretion is known to produce neuronal cell damage. Exogenous application of a high dose of CORT has been shown to elicit the neuronal atrophy in the hippocampus [3]. Rats exposed to restraint stress for 3 weeks exhibited the neuronal atrophy identical to that seen in rats treated with a high dose of CORT for 3 weeks [4]. In addition to these classical genomic effects, which are actuated via intracellular steroid receptors, glucocorticoids act acutely on neuronal excitability [5,6]. The long-term potentiation (LTP) of the population spike amplitude was also acutely (within 1 h) suppressed by a high concentration of glucocorticoids [7]. It has also been demonstrated that CORT dosage for 20 min significantly suppresses the development of LTP in the CA1 region of 4-week-old rat hippocampal slices [8,9].

It is well known that Ca²⁺ influx via *N*-methyl-D-aspartate (NMDA) receptors plays a crucial role in the induction of LTP. The acute effects of CORT (appearing within 30 min) on NMDA receptor-mediated Ca²⁺ signal transduction, however, have not been well elucidated in the hippocampus. In our previous study, we examined the acute CORT effects on NMDA-induced Ca²⁺ signals in mouse hippocampal slices by using Ca²⁺ imaging technique. As a result, the 30 min preincubation of CORT induced a significant decrease of the peak amplitude of NMDA-induced Ca²⁺ elevation in the CA1 region [10]. The membrane non-permeable bovine serum albumin-conjugated CORT (BSA-CORT) also induced a similar suppressive effect in the CA1 region. Therefore the acute CORT effect should be induced via putative surface CORT receptors. A possible candidate for surface CORT

receptors is a classical intracellular glucocorticoid receptor (GR). If this speculation is confirmed, it will bring a new view into the steroid research because it leads to the conclusion that GRs may drive both classical genomic pathways and non-genomic pathways. However, there is not enough evidence on the speculation by physiological experiments although it is supported in part by some biochemical experiments [11-13]. In the present study, to confirm the speculation, we examined the effects of dexamethasone (DEX; an agonist of GR) and CORT with RU38486 (an antagonist of GR) on NMDA-induced Ca^{2+} signals in mouse hippocampal slices.

2. MATERIALS AND METHODS

2.1. Chemicals

CORT, DEX, RU38486, cremophor EL and fura-2/AM were purchased from Sigma (St. Louis, MO, USA). All other chemicals were of the highest purity commercially available.

2.2. Slice Preparation

Brain slices (coronal, 300 μm thick) were prepared from 7-week-old male ddY mice after exposure to an overdose diethylether anesthesia. In order to stabilize the plasma glucocorticoid level, mice used in all experiments were decapitated at the same moment (10 am) in the circadian cycle. Following decapitation, the brain was quickly removed and placed in ice-cold oxygenated artificial cerebrospinal fluid (ACSF) (composition in mM: NaCl 124, KCl 5, CaCl_2 2, NaHCO_3 22, MgSO_4 2, NaH_2PO_4 1.24, glucose 10, pH 7.4, bubbled with 95% O_2 /5% CO_2). Slices were then prepared using a microslicer (DTK-1000; Dosaka-EM, Kyoto, Japan). The slices were recovered in ACSF at 30°C for 60 min and held at room temperature until use. All experiments using animals were conducted in accordance with the institutional guidelines.

2.3. Ca^{2+} Measurement

Measurement of intracellular Ca^{2+} concentration $[\text{Ca}^{2+}]_i$ was performed using the Ca^{2+} -sensitive indicator fura-2. Prior to Ca^{2+} signal measurements, the slices were loaded for 30 min at room temperature with 10 μM fura-2/AM (from 1 mM stock solution in dimethyl sulfoxide (DMSO)) in the presence of 0.01% cremophor EL in 7.2 mL of ACSF. After loading with fura-2, the slices were washed in ACSF for 30 min, and then preincubated with CORT, DEX and RU38486 solutions for 30 min. These solutions were prepared at the appropriate dilution with low- Mg^{2+} ACSF (control solution) (composition in mM: NaCl 124, KCl 5, CaCl_2 2, NaHCO_3 22, MgSO_4 0.1, NaH_2PO_4 1.24,

glucose 10, pH 7.4, gassed with 5% CO_2 /95% O_2) from the stock solution in DMSO. The final concentration of DMSO was less than 0.05% in each case, and 0.05% DMSO was also contained in the control solution.

For fluorescence measurements of $[\text{Ca}^{2+}]_i$, a digital fluorescence microscope system, consisting of an inverted microscope (TE 300; Nikon, Tokyo, Japan) equipped with a xenon lamp for excitation and a CCD camera (C4742-95; Hamamatsu Photonics, Hamamatsu, Japan), was used. Preincubated brain slices were placed in a chamber on the microscope stage. The whole hippocampus fell within the microscope field by using a 4 \times fluorescence objective (Nikon, Tokyo, Japan). The slices were then perfused with the preincubation solution kept at 30°C. At 150 s after the onset of recording, the perfusion solution was replaced to the solution containing 1 mM NMDA.

For Fura-2 measurements, the excitation wavelength was varied discretely between 340 nm and 380 nm, and $[\text{Ca}^{2+}]_i$ was expressed as the ratio (F340/F380) of the 510 nm fluorescence intensity at 340 nm excitation (F340) to that at 380 nm excitation (F380). In each acquisition trial, consecutive fluorescence images were acquired at 5 s intervals for 350 s. **Figure 1** shows examples of a series of obtained images. The fluorescence images were then analyzed with AQUACOSMOS system (Ver.1.3; Hamamatsu Photonics, Hamamatsu, Japan). Eight ROIs (regions of interest), each of which consists of 5 \times 5 pixels (100 \times 100 μm), were put on the dendritic layer of each hippocampal subfield, and the fluorescence data of those ROIs were averaged.

2.4. Statistical Analysis

The data were expressed as mean \pm SEM. The significance of observed differences between groups was examined using the Tukey-Kramer posthoc multiple comparisons test when one way analysis of variance tests yielded $p < 0.05$. When comparing only one pair, two-tailed Student's t test was used for statistical analysis.

3. RESULTS AND DISCUSSION

Figure 2 shows the typical time courses of NMDA-induced Ca^{2+} signals obtained from the CA1 region. In this figure, the Ca^{2+} signal is expressed as the change ($\Delta(\text{F340}/\text{F380})$) in F340/F380 from basal level. The response to continuous NMDA exposure was characterized by a transient elevation in $[\text{Ca}^{2+}]_i$ followed by decay to a plateau within 150 s (see also **Figure 1**). The transient elevation in $[\text{Ca}^{2+}]_i$ was due to NMDA receptor-mediated Ca^{2+} influx, because no response to NMDA application was observed in the slices preincubated with 100 μM of NMDA antagonist MK-801 (data not shown). In the con-

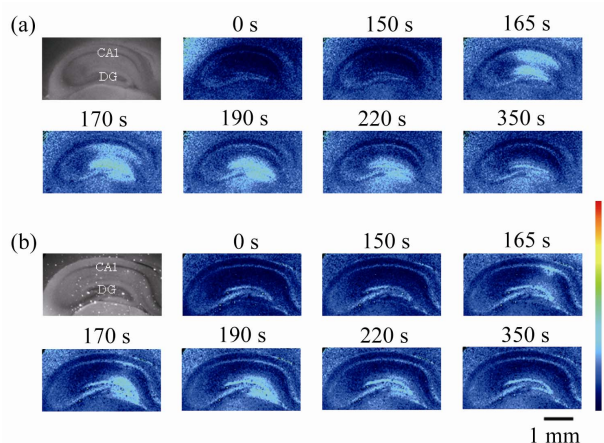


Figure 1. Fluorescence images of mouse hippocampal slices preincubated with (a) the control solution and (b) 1 μM DEX solution. Upper left one of each column is a fluorescence image of each slice. The other pseudo-colored images demonstrate $[\text{Ca}^{2+}]_i$ in the slices. The color bar shows the low (blue) to high $[\text{Ca}^{2+}]_i$ (red). The time above each pseudo-colored image shows the time from the onset of experiment. At 150 s, 1 mM NMDA was applied.

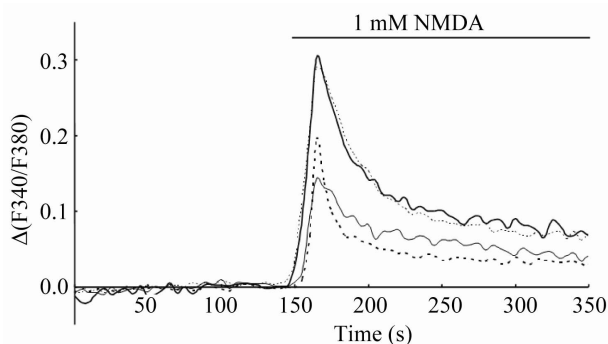


Figure 2. Typical time courses of NMDA-induced Ca^{2+} signal in the CA1 region. The vertical scale ($\Delta(\text{F340}/\text{F380})$) is the change in F340/F380 from basal level. Thick line; control, thin line; 1 μM CORT dosage, thick-dotted line; 1 μM DEX dosage, thin-dotted line; 1 μM CORT dosage with 5 μM RU38486.

control slices preincubated with the control solution for 30 min, the peak amplitudes of $[\text{Ca}^{2+}]_i$ elevation, which are the maximal values of $\Delta(\text{F340}/\text{F380})$, were 0.26 ± 0.02 ($n = 15$) in the CA1 region (**Figure 3**) and 0.27 ± 0.01 ($n = 15$) in the dentate gyrus (data not shown). In the CA1 region, the peak amplitude of $[\text{Ca}^{2+}]_i$ elevation was decreased by preincubation with 1 μM CORT (**Figure 2**). The decreased peak amplitude was $67.3\% \pm 13.1\%$ ($n = 9$, $p < 0.05$) compared with control (**Figure 3**). In contrast, the CORT dosage did not significantly affect the Ca^{2+} signal in the dentate gyrus. It was $97.2\% \pm 5.8\%$ ($n = 9$) compared with control (data not shown). Similar results were also obtained in our previous study [10]. In the previous study, we also showed that the membrane

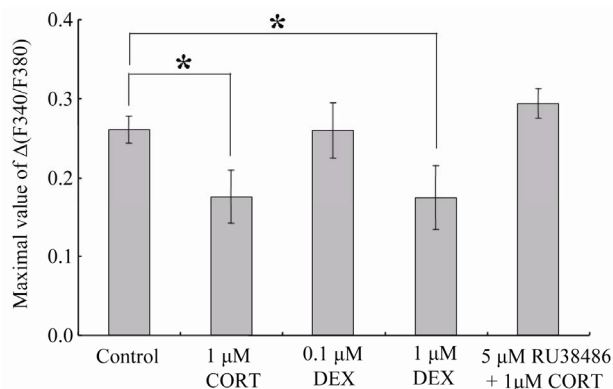


Figure 3. The peak amplitudes of NMDA-induced Ca^{2+} signal elevation in the CA1 region. The peak amplitudes are the maximal values of $\Delta(\text{F340}/\text{F380})$. The data are expressed as means \pm SEM. * $p < 0.05$ compared with control.

non-permeable BSA-CORT induced a similar suppressive effect in the CA1 region. According to the classical view of the steroid action, CORT is thought to penetrate into the cytoplasm of neurons, bind to intracellular receptors and induce genomic effects through new protein synthesis, resulting in the modulation of Ca^{2+} signals [1-4]. Our results, however, suggest that CORT may bind to putative surface CORT receptors. In addition, the possibility of genomic mechanisms was excluded as an explanation for the suppressive CORT effect, because cycloheximide, an inhibitor of protein synthesis, did not abolish the suppressive CORT effect [10].

A possible candidate for surface CORT receptors is a classical intracellular glucocorticoid receptor, GR, which may be bound to plasma membranes. To confirm this speculation, we examined the effects of DEX (an agonist of GR) on NMDA-induced Ca^{2+} signals. In the CA1 region, the peak amplitude of $[\text{Ca}^{2+}]_i$ elevation was decreased by preincubation with 1 μM DEX (**Figure 1(b)** and **Figure 2**), and as expected, the DEX dosage did not significantly affect the Ca^{2+} signal in the dentate gyrus (**Figure 1(b)**). In the CA1 region, the decreased peak amplitude was $67.0\% \pm 15.6\%$ ($n = 8$, $p < 0.05$) compared with control (**Figure 3**). However, 0.1 μM DEX induced no significant effects on the peak amplitude of $[\text{Ca}^{2+}]_i$ elevation although DEX action upon GR is almost 10 times stronger than that of CORT at the same concentration [14] (In the case of CORT, significant suppressive effects were observed from 0.4 μM [10]). Therefore we further examined the effect of 1 μM CORT with 5 μM RU38486 (an antagonist of GR) on NMDA-induced Ca^{2+} signals. As the result, the suppressive CORT effect in the CA1 region disappeared by the coexistence of RU38486 (**Figure 2**). The peak amplitude of $[\text{Ca}^{2+}]_i$ elevation was $112.7\% \pm 7.1\%$ ($n = 8$) compared with control (**Figure 3**).

4. CONCLUSIONS

We have observed so far the acute suppressive effects of CORT on NMDA receptor-mediated Ca^{2+} signals in the CA1 region of mouse hippocampal slices. From the present results, we conclude that the binding site inducing the suppressive CORT effect should be a classical intracellular glucocorticoid receptor, GR, bound to plasma membrane. Our conclusion is also supported by other reports that the immunoreactivity of antibodies against GR was associated with plasma membranes from hippocampal and hypothalamic neurons [11] and that specific CORT binding to neuronal membranes occurred in different brain areas with moderate affinity ($K_d = 120$ nM) [12,13]. Classical GRs are expressed in the cytoplasm of cultured hippocampal neurons [15]. GRs may drive both classical genomic pathways [16] and non-genomic pathways (the current study) in hippocampal neurons.

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