Protein Kinase C Mediates the Corticosterone-induced Sensitization of Dorsal Root Ganglion Neurons Innervating the Rat Stomach

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Background/Aims
Gastric hypersensitivity contributes to abdominal pain in patients with functional dyspepsia. Recent studies showed that hormones induced by stress are correlated with visceral hypersensitivity. However, the precise mechanisms underlying gastric hypersensitivity remain largely unknown. The aim of the present study was designed to investigate the roles of corticosterone (CORT) on excitability of dorsal root ganglion (DRG) neurons innervating the stomach.

Methods
DRG neurons innervating the stomach were labeled by Dil injection into the stomach wall. Patch clamp recordings were employed to examine neural excitability and voltage-gated sodium channel currents. Electromyograph technique was used to determine the responses of neck muscles to gastric distension.

Results
Incubation of acutely isolated DRG neurons with CORT significantly depolarized action potential threshold and enhanced the number of action potentials induced by current stimulation of the neuron. Under voltage-clamp mode, incubation of CORT enhanced voltage-gated sodium current density of the recorded neurons. Pre-incubation of GF109203X, an inhibitor of protein kinase C, blocked the CORT-induced hyperexcitability and potentiation of sodium currents. However, pre-incubation of H-89, an inhibitor of protein kinase A, did not alter the sodium current density. More importantly, intraperitoneal injection of CORT produced gastric hypersensitivity of healthy rats, which was blocked by pre-administration of GF109203X but not H-89.

Conclusions
Our data strongly suggest that CORT rapidly enhanced neuronal excitability and sodium channel functions, which is most likely mediated by protein kinase C but not protein kinase A signaling pathway in DRG neurons innervating the stomach, thus underlying the gastric hypersensitivity induced by CORT injection.

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Key Words
Corticosterone; Ganglia, spinal; Protein kinase C; Visceral pain; Voltage-gated sodium channel
Introduction

Functional dyspepsia (FD) affects 10-25% of the population according to various estimates. Its primary symptom is upper epigastric pain or discomfort in the absence of organic ailments. Although the pathogenesis of this heterogeneous disorder remains largely unknown, gastric hypersensitivity is thought to contribute to the abdominal pain in patients with FD. A growing body of evidence demonstrates that physical or mental distress alters gut immunity and modulates the processing of incoming sensory signals by the brain, thus leading to a potentiation of sensory signals and aberrant visceral pain perception. Corticosterone (CORT), one of the most important stress hormones, plays complex effects in neurological diseases. A recent report showed that neonatal colonic chemical irritation with acetic acid produced FD-like gastric hypersensitivity in adult rats. In addition, chronic psychological stress increased CORT levels by activation of hypothalamic-pituitary-adrenal (HPA) axis, thus increasing the perception of visceral pain. These data strongly indicate that CORT induced by physical or/and psychological stress is involved in chronic visceral hypersensitivity. However, how CORT causes visceral hypersensitivity is not fully understood.

Once released from the adrenal glands in responding to stress, CORT quickly enters the brain and peripheral nerves system, and binds to intracellular receptors. One of the primary targets of the stress hormone is the dorsal root ganglion (DRG), a region that conveys and controls somatic and visceral sensation. Exogenous application of CORT 1 μM overnight has been shown to enhance DRG neuronal excitability. The CORT-induced effect is because of its genomic effects through new protein synthesis. However, mounting evidence also shows that CORT may exert rapid effects on neurons through non-genomic pathways. CORT prolonged N-methyl-D-aspartate (NMDA)-induced Ca²⁺ elevation in cultured rat hippocampal neurons, indicating that CORT can rapidly modulate the function of NMDA receptors by non-genomic pathway. Another study shows that application of CORT for 1 hour significantly increased the density of spines of CA1 pyramidal neurons, which was abolished by co-administration of RU486, an antagonist of GRs. Blocking protein kinase C (PKC) or other single kinase, suppressed the CORT effects. These studies demonstrate the possible existence of fast-acting, non-genomic effects of CORT in neurons both functionally and structurally. Emerging evidence suggests that voltage-gated sodium channels (VGSCs) plays an important role in both stress and pain. Very recent studies indicated PKC activation leads to an increase in the membrane expression of VGSCs and facilitates sodium currents in cortical neurons, indicating that PKC may be involved in mediating VGSCs functions. However, whether and how CORT plays an acute effect on VGSCs of DRG neurons innervating the stomach is not clear.

In the light of the growing evidence, we therefore hypothesize that CORT sensitized the neuronal excitability and enhanced sodium channel activities through activation of PKC in gastric specific DRG neurons. The aim of this study was designed to explore the roles of CORT on the excitability and VGSC currents of DRG neurons innervating the stomach. In addition, the intracellular pathways mediating the CORT effects were also explored. This study might shed light on the pathogenesis of functional gastrointestinal disorders such as FD under acute or chronic conditions.

Materials and Methods

Animals

All experiments were approved by the institutional animal care and use committee at Soochow University and by the Association of Laboratory Animals in Jiangsu Province, China. Adult male Sprague-Dawley rats (6 weeks) were housed four per cage under 12 hours/12 hours dark/light cycle and in a temperature-controlled room (25 ± 1°C). Animals were allowed to access the tap water and standard food.

DiI Labeling of Dorsal Root Ganglion Neurons Innervating the Stomach

The gastric project neurons were retrogradely labeled using 1,19-dioleyl-3,3,39,3-tetramethylindocarbocyanine methane sulfonate (DiI; Invitrogen, Carlsbad, California, USA) as described previously (Fig. 1A).

Whole-cell Patch-clamp Recordings

Neuronal excitability was determined by whole-cell patch-clamp recordings as described previously. In short, DRGs (T₁-T₅) were acutely dissected out and incubated in dissecting solution with enzymes (trypsin, 1.2 mg/mL: Sigma, St. Louis, Mo, USA; collagenase D, 1.5-1.8 mg/mL: Roche, Mannheim, Baden-Württemberg, Germany) for 1.5 hours at 34.5°C. The dissection solution contained the following (mM): NaCl 130, KCl 5, KH₂PO₄ 2, CaCl₂ 1.5, MgSO₄ 6, glucose 10 and HEPES 10, pH 7.2. The osmolality of the solution was 305 mOsm. DRGs were acutely dissected out and incubated in dissecting solution with enzymes (trypsin, 1.2 mg/mL: Sigma, St. Louis, Mo, USA; collagenase D, 1.5-1.8 mg/mL: Roche, Mannheim, Baden-Württemberg, Germany) for 1.5 hours at 34.5°C. The dissection solution contained the following (mM): NaCl 130, KCl 5, KH₂PO₄ 2, CaCl₂ 1.5, MgSO₄ 6, glucose 10 and HEPES 10, pH 7.2. The osmolality of the solution was 305 mOsm. DRGs
were harvested from the enzyme solution, and washed with normal external solution, and then transferred to 2 mL of the dissecting solution containing DNase (0.5 mg/mL). Single cell suspension was then harvested by repeated trituration through flame-polished glass pipettes. Cells were put onto acid-cleaned glass coverslips. A coverslip containing the adherent DRG neurons were placed in a recording chamber with a volume of 0.5 mL, which was attached to the stage of an inverted microscope (IX71; Olympus, Tokyo, Japan) equipped for both fluorescence and phase objectives. The external solution contains (mM): NaCl 130, KCl 5, KH₂PO₄ 2, CaCl₂ 2.5, MgCl₂ 1, glucose 10 and HEPES 10, pH7.2, adjusted by NaOH, osmolarity, 295-300 mOsm. The resistance of the patch electrode was 3-5 MΩ when filled with the pipette solution containing (mM): CsF 140, MgCl₂ 1, EGTA 5, Na-GTP 3, glucose 10, and HEPES 10, pH7.2, adjusted with CsOH, osmolality 285-295 mOsm. The total VGSC currents of DiI labeled cells (Fig. 1B and 1C) were obtained in response to voltage stimulations to different testing potentials from −70 to +50 mV in 10 mV increments with a duration of 80 msec. The sodium peak current was determined as the peak of the transient component of the current at a given voltage. To reduce changes in cell size, the current density (pA/pF) was calculated by dividing the current amplitude by the cell membrane capacitance.

Implantation of Gastric Distension Balloon and Electromyographic Recordings

The electromyographic (EMG) recording of the responses of neck muscles to gastric distension (GD) was performed on a total of 14 SD rats as published previously. In brief, cells were superfused (2 mL/min) at room temperature with an external solution, which contains (mM): NaCl 60, choline chloride 80, CaCl₂ 0.1, HEPES 10, tetraethylammonium chloride 10, glucose 10, and CdCl₂ 0.1, pH 7.4 adjusted with tetraethylammonium hydroxide. The osmolality was adjusted at ~310 mOsm. The resistance of the patch electrode was 3-5 MΩ when filled with the pipette solution containing (mM): CsF 140, MgCl₂ 1, EGTA 5, Na-GTP 3, glucose 10, and HEPES 10, pH7.2, adjusted with CsOH, osmolality 285-295 mOsm. The total VGSC currents of DiI labeled cells (Fig. 1B and 1C) were obtained in response to voltage stimulations to different testing potentials from −70 to +50 mV in 10 mV increments with a duration of 80 msec. The sodium peak current was determined as the peak of the transient component of the current at a given voltage. To reduce changes in cell size, the current density (pA/pF) was calculated by dividing the current amplitude by the cell membrane capacitance.
The GD protocol was phasic ascending distension (0-20-0-40-0-60-0-80 mmHg). The distension duration for each pressure was 20 seconds and breaks for 2 minutes after each distension.

Measurements of Hind Paw Withdrawal Threshold

The 50% paw withdrawal threshold (PWT) to a static mechanical stimulus was measured using von Frey filaments as described previously. Normal rats (n = 7 in each group) were placed individually on an elevated iron mesh in a plastic cage to adapt to the testing environment for about 30 minutes. An ascending series of von Frey hairs (ranging from 0.6 g to 26.0 g) were applied to stimulate the plantar aspect of each hind paw before and after CORT administration. A trial began with the application of the 4.0 g hair and with 26.0 g as the cutoff strength of von Frey filaments. A withdrawal of hind paw upon the stimulus was defined as a positive response. If there was a positive response, a filament of a smaller force was then applied. If there was a negative response, a filament of a greater force was then applied. The tactile stimulus producing a 50% likelihood of withdrawal was determined by the “up-down” calculating method. The mean value was used as the force to produce the withdrawal response.

Figure 2. Treatment of corticosterone (CORT) enhanced excitability of dorsal root ganglion (DRG) neurons. (A) CORT application (1 μM for 1 hour) depolarized the resting membrane potential (RP) in DiI-labeled DRG cells (***P < 0.01, compared with control (CON), Mann-Whitney test). (B) CORT application resulted in a marked reduction of rheobase (***P < 0.01, compared with CON, Mann-Whitney test). (C) CORT application significantly hyperpolarized action potential (AP) threshold (***P < 0.01, compared with CON, two-sample t test). (D-F) Typical traces of APs evoked by 100, 300, and 500 pA ramp current stimulation in the absence (top) or presence of CORT (middle). Bar graphs show the numbers of AP evoked by 100, 300, and 500 pA ramp current stimulation. CORT incubation remarkably increased number of APs (***P < 0.01, compared with CON, two-sample t test).
**Reagents**

CORT was obtained from Abcam (ab143597; London, UK). Bisindolylmaleimide I (GF109203X), an antagonist of PKC, was purchased from Aladdin (gl129390; Los Angeles, CA, USA). H-89, an antagonist of protein kinase A (PKA), was purchased from Sigma (B1427; USA). Dimethyl sulfoxide (DMSO) was obtained from Sigma (D8418; Sigma). CORT was initially dissolved in DMSO as a stock solution. The final concentration of DMSO in the extracellular solution was 0.05%. No detectable effect of the vehicle was observed in our experiments.

**Data Analysis**

EMG data are presented as area under curve of the integrated EMG after baseline subtraction. All data are expressed as mean ± SEM in the present study. Statistical analysis was carried out using OriginPro 8 (OriginLab, Hampton, VA, USA) and Matlab (Mathworks, Natick, MA, USA). Normality was checked for all analyses before further analysis. Significance was determined using a two-sample t test for 2 groups. Paired sample Wilcoxon signed rank test, Dunn post hoc test following Friedman ANOVA, and the Mann-Whitney test was done where appropriate. A P-value less than 0.05 was considered to be statistically significant.

![Figure 3. Dose-responses of corticosterone (CORT) incubation. (A, B) Typical traces of action potentials (APs) evoked by 100 pA and 300 pA ramp current stimulation after CORT incubation at different concentrations (top). Bar graphs show a significant increase in numbers of AP evoked by 100 pA and 300 pA ramp current stimulation after CORT application (bottom, *P < 0.05, **P < 0.01, compared with CON, Friedman ANOVA). (C, D) Bar graphs show the time of the first AP evoked by 100 pA and 300 pA ramp current stimulation that were significantly increased after 1 μM and 10 μM CORT application (**P < 0.005, compared with CON, Friedman ANOVA).](image)
Results

Corticosterone Enhances Excitability of Gastric Dorsal Root Ganglion Neurons

We first investigated the effect of CORT on the activity of gastric projection DRG neurons. To this end, we irrigated DRG neurons with different concentrations of CORT for 1 hour. Gastric-specific DRG (including T7-T10 DRGs) neurons were identified by the fluorescent dye DiI injected into the stomach wall for patch-clamp recording (Fig. 1A-C). We measured the passive and active membrane properties of these neurons from control (CON) and CORT-treated cells. The small- and medium-sized DRG neurons were used in this study, because these neurons are responsible for pain sensation.24,29 Under current-clamp conditions, 32 cells were recorded. Resting membrane potentials (RPs) were $-48.9 \pm 0.7$ mV (n = 15) and $-42.8 \pm 0.4$ mV (n = 17) for gastric-specific DRG neurons isolated from CON and CORT-treated groups, respectively. CORT treatment significantly depolarized RPs (Fig. 2A; **$P < 0.01$, compared with CON, two-sample $t$ test). The rheobase was $77.8 \pm 7.1$ pA (n = 15) and $25.9 \pm 4.3$ pA (n = 17) for CON and CORT-treated cells, respectively. CORT reduced rheobase (Fig. 2B; **$P < 0.01$, compared with CON, two-sample $t$ test). CORT resulted in a dramatic increase in the numbers of APs evoked by 100 pA, 300 pA, and 500 pA ramp current stimulation (Fig. 2D-F, **$P < 0.01$ compared with CON, two-sample $t$ test). The numbers of APs evoked by 100 pA current ramp stimulation was $2.9 \pm 0.9$ (n = 15) and $11.5 \pm 1.5$ (n = 17) for CON and CORT-treated cells, respectively. The numbers of APs evoked by 300 pA current ramp was $12.2 \pm 1.2$ (n = 15) and $22.5 \pm 2.1$ (n = 17) for CON and CORT-treated cells, respectively. The numbers of APs evoked by 500 pA current ramp was $20.9 \pm 1.9$ (n = 16) and $32.5 \pm 2.8$ (n = 17) for CON and CORT-treated cells.
cells, respectively. CORT markedly increased the numbers of APs responding to ramp current stimulation. To examine the dose dependency, the cells were divided into 4 groups. The concentration of CORT used was as the followings: 0 μM, 0.1 μM, 1.0 μM, and 10.0 μM. Dose-response experiments showed that the enhancing effect was significant at the doses of 1.0 μM and 10.0 μM (Fig. 3; *P < 0.05, **P < 0.01, ***P < 0.005, compared with 0 μM, Friedman ANOVA; n = 11). So, 1 μM CORT was used to determine the effect of CORT on sodium currents in the following experiments.

Corticosterone Elevates Voltage-gated Sodium Current Density

We next examined the change in VGSC current density of DRG neurons before and after CORT treatment. Under voltage-clamp conditions, the current-voltage (I-V) relationship was examined (Fig. 4A and 4B). The average peak current density of sodium current obtained from CON and CORT groups were 93.9 ± 14.5 pA/pF (n = 10) and 160.24 ± 8.8 pA/pF (n = 12), respectively. The average peak current density was significantly increased after CORT treatment (Fig. 4C; **P < 0.01, compared with CON, two-sample t test). We then examined the changes in sodium current kinetics after CORT treatment. The current activation time (T a) between 10% and 90% and the half-decay inactivation time (τ in) were measured in peak sodium currents from both CON and CORT groups as described previously. T a was 2.5 ± 0.5 msec (n = 10) in control group and 1.2 ± 0.1 msec (n = 12) in CORT-treated cells. CORT significantly accelerated the activation time (Fig. 4D; **P<0.01, compared with CON, two-sample t test). However, no significant change in the τ in was observed between control and CORT groups (Fig. 4E; CON, 2.2 ± 0.2 msec, n = 10; CORT, 1.9 ± 0.1 msec, n = 11, P > 0.05).

Figure 5. Reversal of corticosterone (CORT)-induced neuronal hyperexcitability by the protein kinase C (PKC) inhibitor GF109203X (GF). (A) GF (1 nM for 15 minutes) application markedly hyperpolarized resting membrane potentials (*P < 0.01, compared with CORT, Mann-Whitney test). (B) GF application resulted in a marked increase in rheobase (**P < 0.01, compared with CORT, two-sample t test). (C) GF application significantly depolarized AP threshold (**P < 0.01, compared with CON, two-sample t test). (D-F) Typical traces of APs evoked by 100, 300, and 500 pA ramp current stimulation (top). Bar graphs show a significant decrease of numbers of AP evoked by 100, 300, and 500 pA ramp current stimulation after GF application (bottom, **P < 0.01, compared with CORT, two-sample t test).
GF109203X Reverses the Corticosterone-induced Hyperexcitability

We then investigated whether GF109203X, an inhibitor of PKC, affected the excitability of gastric-specific DRG neurons by CORT. GF109203X at concentration of 3 μM was pre-treated with acutely isolated DRG neurons for 15 minutes. Under current-clamp conditions, 28 cells were recorded. RPs were $-43.8 \pm 0.6$ mV (n = 15) and $-49.1 \pm 0.7$ mV (n = 13) for CORT and GF109203X groups, respectively. GF109203X treatment significantly hyperpolarized RPs (Fig. 5A; **P < 0.01, compared with CORT, two-sample t test). The rheobase were $23.1 \pm 3.6$ pA (n = 15) and $63.2 \pm 6.8$ pA (n = 13) for CORT and GF109203X group, respectively. GF109203X significantly increased rheobase (Fig. 5B; **P < 0.01, compared with CORT, two-sample t test). GF109203X treatment also depolarized the AP threshold (Fig. 5C; CORT, $-31.1 \pm 1.2$ mV, n = 15; GF109203X, $-25.2 \pm 1.8$ mV, n = 13, *P < 0.05, compared with CORT, two-sample t test). In addition, the numbers of APs evoked by 100 pA current ramp was $10.5 \pm 0.9$ (n = 15) and $4.5 \pm 1.2$ (n = 13) for CORT and GF109203X groups, respectively. The numbers of APs evoked by 300 pA current ramp was $21.7 \pm 2.1$ (n = 15) and $11.9 \pm 1.9$ (n = 13) for CORT and GF109203X, respectively. The numbers of APs evoked by 500 pA current ramp was $28.5 \pm 2.4$ (n = 15) and $20.7 \pm 2.0$ (n = 13) for CORT and GF109203X groups, respectively. GF109203X treatment remarkably decreased the numbers of APs evoked by 100 pA, 300 pA and 500 pA ramp current stimulation (Fig. 5D-F, **P < 0.01, compared with CORT, two-sample t test).

GF109203X Abolishes Corticosterone-induced Potentiation of Sodium Currents

Since CORT enhanced the sodium current density, we next determined whether CORT enhancement of sodium currents was PKC-dependent. Under voltage-clamp conditions, the I-V relationship was also examined (Fig. 6A and 6B). Pretreatment with GF109203X (3 μM) for 15 minutes, CORT no longer increased sodium currents. The average peak current densities obtained from

**Figure 6.** GF109203X (GF) reduced sodium current density. (A) Examples of sodium currents measured from corticosterone (CORT)-treated (top) and GF + CORT-treated neurons (bottom). Membrane potential was held at $-60$ mV and voltage steps were changed from $-70$ to $+50$ mV with 10 mV increments with a duration of 80 msec. (B) I-V curves for sodium currents of gastric DRG neurons treated with CORT or GF + CORT. (C) Bar graphs show the mean sodium peak current densities. GF application significantly reduced the peak sodium currents densities (**)P < 0.01, compared with CORT, Mann-Whitney test). (D) Bar graphs show the mean activation time (**P < 0.01, compared with CORT, Mann-Whitney test). (E) No significant alteration was induced by CORT in the half-decay inactivation time (P > 0.05, compared with CORT, Mann-Whitney test).
CORT and GF109203X were 165.3 ± 6.9 pA/pF (n = 12) and 85.1 ± 13.1 pA/pF (n = 9), respectively. The average peak current density was significantly decreased (Fig. 6C; **P < 0.01, compared with CORT, two-sample t test). The Tₐ was 1.2 ± 0.1 msec (n = 10) in CORT and 2.4 ± 0.2 msec (n = 9) in GF109203X, respectively. GF109203X significantly reversed the activation time by CORT (Fig. 6D; **P < 0.01, compared with CORT, two-sample t test). No significant change in the τᵢₘ was found between CORT and GF109203X groups (2.3 ± 0.1 msec, n = 9 in CORT vs 2.1 ± 0.2 msec, n = 10 in GF109203X + CORT; Fig. 6E).

**H-89 Did Not Alter the Corticosterone-induced Enhancement of Sodium Currents**

Some researches indicated that PKA also mediated the sodium currents.³¹ We thus determined whether sodium current is mediated by PKA in the present study. H-89, an antagonist of PKA, at a concentration of 10 μM was pretreated for 15 minutes. The I-V relationship was also examined (Fig. 7A and 7B). The average peak current density obtained from CORT and H-89 treatment was 141.7 ± 14.0 pA/pF (n = 12) and 148.9 ± 14.5 pA/pF (n = 9), respectively. The average peak current density was not altered by H-89 (Fig. 7C; P > 0.05, compared with CORT, two-sample t test). The Tₐ was 2.2 ± 0.1 msec (n = 10) in CORT and 2.5 ± 0.2 msec (n = 9) in H-89, respectively. H-89 did not significantly affect Tₐ (Fig. 7D; P > 0.05, compared with CORT, two-sample t test). No significant change in the τᵢₘ was found between CORT and H-89 (Fig. 7E; 2.3 ± 0.3 msec, n = 9 in CORT vs 2.1 ± 0.1 msec, n = 10 in H-89).

**GF109203X Suppressed Corticosterone-induced Gastric Hypersensitivity**

To record EMGs of the acromiotrapezius muscle responding to GD before and after CORT treatment, implantation of gastric balloon and electrodes were performed 1 week before injection of CORT (n = 14 rats). One-week after recovery from the surgery, GD-induced painful behaviors were examined 1 hour after CORT injection (5.0 mg/kg body weight, ip). This dose of CORT was used according to our pilot studies and a previous report.³² The re-
results demonstrated that marked effects were observed at distension pressures of 20, 40, 60, and 80 mmHg 1 hour after CORT injection (Fig. 8A and 8B; n = 7 for each group, *P < 0.05, **P < 0.01, compared with pre, paired sample Wilcoxon signed rank test). Gastric hypersensitivity induced by CORT was sustained through the next 1 hour. To determine whether PKC or PKA is involved in the development of gastric hypersensitivity in CORT-treated rats, we examined the effect of GF109203X (PKC inhibitor) and H-89 (PKA antagonist), on EMG amplitude in response to GD. H-89 had no obvious effect on EMG amplitude. However, treatment with GF109203X led to a dramatic reduction in EMG responses at 20-80 mmHg GD pressures (Fig. 8A and 8B; n = 7 for each group, *P < 0.05, **P < 0.01, compared with CORT, paired sample Wilcoxon signed rank test). In addition, intraperitoneal injection of CORT did not alter the rat hind PWT in response to von Frey filament stimulation (Fig. 8C; n = 7).

Discussion

In the present study, we demonstrated that CORT rapidly enhanced not only the neuronal excitability but also the sodium current density of gastric specific DRG neurons. In addition, systematic injection of CORT induced gastric hypersensitivity in normal rats. These findings suggest that CORT plays an important role in modulating the functions of brain and gut axis. Modulation of ion channels under acute or chronic stress has been increasingly attracting much attention. The influences of stress are elicited at least partly by CORT. The stress hormone CORT has been reported to modulate many ion channels, such as acid-sensing ion channels of cultured hippocampal neurons. In the present study, we provided direct evidence to show that CORT elevates VGSC currents of DRG neurons innervating the stomach. VGSCs have 9 subunits: Na1.1-Na1.9 with 4 β subunits. Among of them, Na1.7, Na1.8, and Na1.9 are the most abundantly expressed in DRGs and are

Figure 8. Corticosterone (CORT) produced gastric hypersensitivity. (A) Representative electromyographic (EMG) recordings from normal healthy rats in response to graded gastric distension (GD) 60 minutes after a single injection of CORT, GF109203X (GF) + CORT or H-89 + CORT. (B) Bar graph shows that CORT intraperitoneal injection produced gastric hypersensitivity (*P < 0.05, **P < 0.01 compared with pre-injection (PRE) at the same pressure, paired sample Wilcoxon signed rank test). GF PRE dramatically suppressed EMG amplitude, whereas H-89 had no significant effect EMG (*P < 0.05, **P < 0.01 compared with CORT at the pressure, paired sample Wilcoxon signed rank test). (C). CORT injection did not alter the hind paw withdraw threshold (PWT) to von Frey filament stimulation (n = 14 rats).
most closely related to pain in the periphery. In consistent with previous reports, we provided further evidence to show that VGSCs play an important role in stress and stress hormones response. Since many drugs in low concentrations have an opposite effect when compared to its high concentrations, we examined whether enhancement of CORT on excitability of DRG neurons was dose-dependent. In our experiments, CORT at 0.1 μM had no significant effect on DRG neuronal excitability, but CORT at 1 μM significantly enhanced excitability of DRG neurons innervating the stomach, which was the same as CORT at 10 μM. The active concentration of CORT in the present study is relatively lower than the real blood plasma concentration (12.15 ± 0.76 μM, n = 3). This difference may be because the DRG neurons were isolated in vitro. To the best of our knowledge, this is the first report that CORT plays a rapid effect on DRG neurons innervating the stomach of healthy rats.

The mechanism by which CORT modulates neuronal excitability and sodium channel function has yet to be investigated. In addition to genomic pathways, CORT is thought to activate ion channels in central neurons through non-genomic pathways. An earlier study showed that incubation of epinephrine (5 nM) or CORT (1 μM) overnight sensitized the DRG neurons, indicating that stress hormone-induced sensitization of DRG neurons might be through the enhanced transcription of ion channels. In the present study, however, this is least likely the case because we exposed DRG neurons with CORT for only 1 hour and the time was not long enough to affect the expression of proteins according to previous reports. Therefore, the rapid CORT effects may occur independently of the regulation of the gene expression of sodium channels in the present study. Previous studies suggest that CORT acts by signaling directly to GRs on DRG neurons or, alternatively, by enhancing levels of other pro-nociceptive mediators after immune activation in the colon.

A better understanding of the molecular mechanisms underlying stress hormone-induced alterations in neuronal excitability and sodium channel function is crucial for the development of novel beneficial therapeutic strategies. Several earlier studies have showed that protein kinase can modulate function of sodium channel by phosphorylating targeting motif. Since CORT signaling pathway can activate PKC and PKA signaling pathways, so we speculated that the CORT effects on sodium currents might be mediated by PKC and/or PKA. Although others kinase-signaling pathways cannot be excluded, we showed that CORT-induced effects are dependent of PKC but independent of PKA in gastric specific DRG neurons. There are several reasons that may explain the differences in PKA and PKC effects. The concentration of these two inhibitors may not be a major reason for the differences in PKC and PKA effect since the doses were selected according to previous reports and our pilot studies. We speculate that CORT-induced rapid effects may be tissue-, cell type-, and ion channel-specific, as others have reported that CORT-induced rapid effect on ATP currents is PKA dependent. However, this needs to be further investigated. How PKC enhances sodium channel function remains largely unknown. PKC usually regulates the transcription-translation mechanism. Since the incubation time was short (ie, 1 hour), we hypothesize that the transcriptional mechanism might not be involved in the present study. The possible mechanisms included but not excluded the phosphorylation of sodium channels and enhanced trafficking of sodium channels from the cytoplasm to the cell surface membrane. In the present study, although CORT did not alter Tm, it significantly reduced Ta of sodium currents. This result suggests that CORT-induced enhancement of sodium current density may not because of aggregation of sodium channel on cell membrane, but at least in part because of changes in channel kinetics. Since the Ta was shorter after CORT treatment, it is therefore reasonable to hypothesize that all sodium channels are open or have a trend to open at the same time. Increase in ion channel conductance was another possible mechanism, but this needs to be further investigated in the future.

It is worthy knowing that systemic administration of CORT produced gastric hypersensitivity while this did not alter the hind PWT of the normal rats. Although at the present time, we cannot provide much more evidence to distinguish possible mechanisms underlying the differentiation of visceral hypersensitivity and somatic pain sensation by CORT, this result strongly suggested that the effect induced by systematic use of CORT is visceral pain specific and it is not a non-specific or toxic effect. It is also reasonable since the visceral organs are particularly vulnerable to stress. This implies that the acute stress causes visceral responses more often and earlier than somatic responses. Consistent with previous studies that stress led to colonic hypersensitivity and functional dyspepsia-like gastric hypersensitivity in rats, which was most likely mediated by stress hormones such as CORT and epinephrine, we demonstrated for the first time that CORT injection induced gastric hypersensitivity in normal rats. However, the detailed mechanisms of systematic application CORT are not well understood. It is, however, very tempting to hypothesize that CORT in vivo application might share the same mechanism in producing gastric hypersensitivity as it effects neurons in vitro. This CORT-induced effect may explain the stress-induced epigastric pain syndrome or postprandial distress.
CORT Sensitizes DRG Neurons

**syndrome.**

In summary, we have provided a novel nociceptive signaling pathway of CORT in gastric DRG neurons. CORT incubation induces neuronal hyperexcitability and potentiates the sodium channel function, which is likely mediated by PKC but not PKA. Most importantly, systemic use of CORT leads to development of gastric hypersensitivity of healthy rats. These findings imply that stress-induced increase in CORT levels might potentiate sodium channel activities and shed light on the mechanisms of gastric hypersensitivity in patients with FD induced by adrenergic stress conditions. Our studies might also provide beneficial therapeutic strategies for the treatment of FD.

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