Research Article

Gyogamdan, a Traditional Medicine Prescription, Ameliorated Dermal Inflammation and Hyperactive Behavior in an Atopic Dermatitis Mouse Model Exposed to Psychological Stress

Ly Thi Huong Nguyen,¹ Uy Thai Nguyen,¹ Min-Jin Choi,¹ Tae-Woo Oh,² In-Jun Yang,¹ and Heung-Mook Shin¹

¹Department of Physiology, College of Korean Medicine Dongguk University, Gyeongju 38066, Republic of Korea
²Korean Medicine-Application Center, Korea Institute of Oriental Medicine, Daegu, Republic of Korea

Correspondence should be addressed to In-Jun Yang; injuny@dongguk.ac.kr and Heung-Mook Shin; heungmuk@dongguk.ac.kr

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Psychological stress (PS) plays a significant role as an aggravating factor in atopic dermatitis (AD). The traditional medicine prescription, Gyogamdan, has been used to treat chest discomfort and mood disorders caused by PS. This study investigated the effects of an ethanolic extract of Gyogamdan (GGDE) on stress-associated AD models and the underlying mechanisms. 2,4-Dinitrochlorobenzene- (DNCB-) treated BALB/c mice were exposed to social isolation (SI) stress. The effects of orally administered GGDE (100 or 500 mg/kg) were evaluated by ELISA, western blotting, and an open field test (OFT). SI stress exaggerated the skin inflammation and induced locomotor hyperactivity in the AD mouse model. GGDE reduced the levels of IgE, TNF-α, IL-13, eotaxin, and VEGF and mast cell/eosinophil infiltration and prevented the decreases in the levels of involucrin and loricrin in the skin. GGDE also suppressed the SI-induced increases in corticotropin-releasing hormone (CRH), adrenocorticotropic hormone (ACTH), and corticosterone (CORT) in socially isolated AD mice. Furthermore, GGDE reduced traveling distances and mean speed significantly in the OFT. The in vitro experiments were performed using HaCaT, HMC-1, PC12, and BV2 cells. In the TNF-α/IFN-γ- (TI-) stimulated HaCaT cells, GGDE decreased the thymus and activation-regulated chemokine (TARC) and macrophage-derived chemokine (MDC) production significantly by inhibiting p-STAT1 and NF-κB signaling. GGDE also reduced VEGF production in HMC-1 cells stimulated with CRH/substance P (SP) by inhibiting p-ERK signaling pathway. GGDE increased the cell viability significantly and suppressed apoptosis in CORT-stimulated PC12 cells. Moreover, GGDE suppressed the LPS-induced production of NO, TNF-α, IL-1β, and IL-6 in BV2 cells. These results suggest that GGDE might be useful in patients with AD, which is exacerbated by PS.

1. Introduction

Individuals differ in terms of their perception and adaptation to psychological stress (PS). Some individuals are PS-sensitive and prone to develop PS-related pathologies, including mental disorders, whereas others are more PS-resistant [1]. A recent animal study reported that the responses to PS differ in normal and atopic dermatitis (AD) mice. PS did not affect normal mice but elicited significant changes in weight loss and anxiety-related behaviors in AD mice, suggesting that AD patients are likely to be vulnerable to PS [2].

AD and PS form a vicious cycle, each reinforcing the other negatively; AD symptoms can elicit PS, and PS can exacerbate AD symptoms [3, 4]. AD is an inflammatory skin disease characterized by redness, itching, swelling, and oozing bleeding. Sleep disturbances due to itching, anxiety about the possibility of scarring, and a chronic relapsing course can cause PS in AD patients [3, 5]. AD patients showed higher rates of stress-related disease, anxiety, and
depression than healthy people [6]. Therefore, PS management in AD patients appears to be a promising area with considerable potential for treatment and management [7, 8].

The responses of AD patients to PS occur through coordination with the hypothalamus-pituitary-adrenal (HPA) axis. PS activates the HPA axis to stimulate systemic stress hormone production, including corticotropin-releasing hormone (CRH), adrenocorticotropic hormone (ACTH), and glucocorticoids (GC, cortisol in humans and corticosterone in rodent; hereafter referred to as CORT). These stress hormones contribute to mast cell activation, increased skin vascular permeability, and decreased epidermal structural protein production in the skin [3, 9]. PS induces CRH secretion from the hypothalamus, which activates skin mast cells to increase vasodilation and vascular permeability [4]. HPA activity is regulated by a GC negative feedback from the adrenal cortex. On the other hand, prolonged chronic elevation of CORT by PS can damage the skin barrier function [5, 10]. Moreover, an abnormal HPA axis function by PS causes structural and functional changes in the prefrontal cortex, hippocampus, and amygdala regions of the brain, resulting in increased stress vulnerability and attention-deficit/hyperactivity disorder (ADHD) of AD patients [11].

According to Donguibogam, the Korean traditional medicine book, Gyogamdan (GGD) is a traditional medicine used to treat chest discomfort and mood disorders caused by psychological stress. GGD consists of *Cyperus rotundus* L. and *Poria cocos* (Schw.) Wolf, both of which have been reported to be effective in stress-induced mental and physical diseases [12, 13]. *Cyperus rotundus* L. exhibited in vivo anti-inflammatory effects without affecting gastrointestinal tract [14]. *Cyperus rotundus* L. was indicated in the herbal formula KAHF, which significantly alleviated skin severity and serum levels of inflammatory cytokines in AD patients [15]. A previous study demonstrated that a methanolic extract of *Poria cocos* (Schw.) Wolf exerted inhibitory effects on 12-O-tetradecanoylphorbol-13-acetate- (TPA-) induced dermatitis in mice model [16]. On the other hand, it is unclear if GGD has therapeutic potential in AD exaggerated by PS. Therefore, the present study evaluated the efficacy and possible action mechanism of an ethanolic extract of GGD (GGDE) on the inflammatory response in an AD-like mouse model exposed to PS. In addition, behavioral changes were also assessed to determine if GGDE regulates the psychiatric comorbidities in a murine model of AD after exposure to PS.

2. Materials and Methods

2.1. Chemicals and Reagents. High glucose Dulbecco’s modified Eagle’s medium (DMEM) was obtained from Welgene Inc. (Gyeongsangbuk, Korea), and Iscove’s modified Dulbecco’s medium (IMDM) acquired from Merck Millipore (Darmstadt, Germany). Fetal bovine serum (FBS) and antibiotics were purchased from Invitrogen Inc. (Carlsbad, CA, USA). 2,4-Dinitrochlorobenzene (DNCB), corticotropin-releasing hormone (CRH), substance P (SP), 1-thioglycerol, and dexamethasone (DEX) were procured from Sigma-Aldrich (St. Louis, MO, USA). Human VEGF and mouse IFN-γ, IL-1β, IL-6, IL-13, TNF-α, VEGF, etoxatin, and IgE ELISA kits were supplied by Koma Biotech Inc. (Seoul, Korea). Human MDC, TARC ELISA kits were obtained from R&D Systems (Minneapolis, MN, USA). The mouse CORT ELISA kits were purchased from Ario Biolaboratories (Hsinchu, Taiwan), and Mouse CRH and ACTH ELISA kits were procured from LifeSpan BioSciences (Seattle, WA, USA). Antibodies for phospho-p38, phospho-JNK, phospho-ERK, phospho-STAT1, phospho-1-κB-α, p38, JNK, ERK, STAT1, 1-κB-α, and NF-κB were acquired from Cell Signaling Technology (Danvers, MA, USA). Antibodies against loricrin, involucrin, and lamin B2 were supplied by Abcam (Cambridge, MA, USA). HRP-conjugated anti-β-actin was obtained from Sigma-Aldrich (St. Louis, MO, USA).

2.2. Plant Preparation and Analysis. GGD was purchased from Kyung Hee University Hospital (Seoul, Republic of Korea). GGD (*Poria cocos* Schw.) Wolf to *Cyperus rotundus* L. weight ratio, 1:2 (30 g) was extracted with 300 ml of aqueous EtOH (30%) at 70°C for 3 h. The extract was then filtered through Whatman #2 filter paper (Whatman International, Maidstone, UK), evaporated on a rotary vacuum evaporator, and freeze-dried (FD8508S, Ilshin, Busan, Republic of Korea) (yield 7.83% w/w). The resulting dried GGG was dissolved in distilled dimethyl sulfoxide and sterilized by passing it through a 0.22 μm syringe filter (Sartorius, Goettingen, Germany) before use. Nootkatone, α-cyperone, ergosterol, dehydrotrametenolic acid, pachymic acid, and tumulosic acid were purchased from ChemFaces Biochemical Co. Ltd. (Wuhan, China). Sample component analysis and quantification of the six standards were performed using an HPLC 1290 system (Agilent, Santa Clara, CA, USA) at the Korea Basic Science Institute (Seoul). For the analysis of nootkatone, α-cyperone, and ergosterol, the extracted samples (10 μl) were separated on Extend C18 (2.0 × 150 mm, 5 μm, Agilent) column at a flow rate of 0.4 ml/min using water (A) and acetonitrile (B) as the mobile phases. The gradient used was as follows: 50–80% (B) for 5 min, 80–100% (B) for 5 min, and equilibration for 5 min. The column was operated at 25°C. Nootkatone, α-cyperone, and ergosterol were detected at UV wavelengths of 240 nm, 254 nm, and 282 nm, respectively. For the analysis of dehydrotrametenolic acid, pachymic acid, and tumulosic acid, extracted samples (10 μl) were separated on a Kinetex C18 (4.6 × 250 mm, 5 μm, Phenomenex) column at a flow rate of 0.8 ml/min using 0.1% phosphoric acid (A) and acetonitrile (B) as the mobile phases. The gradient used was as follows: 10–100% B for 15 min and equilibration for 10 min. The column temperature was 35°C. Dehydrotrametenolic acid was detected at a UV wavelength of 210 nm; pachymic acid and tumulosic acid were detected at a UV wavelength of 243 nm. The amounts of these compounds in GGDE were determined using standard calibration curves.
2.3. Animal Experiments. BALB/c (five-week-old male) mice were purchased from Koatech Lab Animal Inc. (Seoul). All animal experimental procedures were performed in accordance with the protocols approved by the Institutional Animal Care and Use Committee of Dongguk University (IACUC-2019-7). The mice were assigned randomly to five groups (n = 5 in each group): a treatment naive control group (the CON group), DNCB only group (the AD group), DNCB plus social isolation group (the SI-AD group), DNCB plus PS plus 100 mg/kg/day of GGDE (the GGDE100 group), and DNCB plus PS plus 500 mg/kg/day of GGDE (the GGDE500 group). The animals were acclimated for one week before the experiments. All mice had free access to water and commercial pellet diet (5L79, PMI Nutrition, St Louis, MO, USA). Figure 1(a) presents a schematic diagram of the experimental schedule. Modified SI stress conditions were prepared according to a previous report of SI stress in a mouse model of allergic dermatitis [2]. The SI-AD group mice were isolated in acrylic cages (10 × 10 × 14 cm). The backs of the all mice were then shaved, and for sensitization, 200 µl of 1% DNCB (dissolved in a 3:1 mixture of acetone and olive oil) was applied to all groups except CON group three times during the first week. Immediately after the treatment period, 200 µl of 0.3% DNCB was applied to shaved dorsal areas three times per week for eight weeks in all groups except for the control group. From the 2nd week of the experiment, GGDE100 and GGDE500 group were administered 100 or 500 mg/kg of GGDE orally five times per week for seven weeks, respectively. The clinical skin severity score was measured for four symptoms: erythema/hemorrhage, edema, scarring/dryness, and erosion/excoriation, with the scoring system of 0 (no symptoms), 1 (mild), 2 (moderate), and 3 (severe). One day after the last treatment, the mice were sacrificed. The whole blood was collected, and serum was obtained by centrifugation (3000 × g, 15 min, 4°C).

2.4. Open Field Test (OFT). The OFT was used to assess locomotor activities one week before sacrifice. Before testing, mice were acclimatized to a test room for two hours and then placed in a black box (measuring 40 × 40 × 40 cm), which was wiped down with 70% alcohol between tests to remove any odors. The total distances (cm) traveled, distances traveled in the peripheral zone and the central zone, and mean speeds (cm/s) were measured in a 10-minute test. Smart V3.0 software (Panlab Harvard Apparatus, MA, USA) was used to determine the total distance traveled, distance traveled in the central and peripheral zones, and the mean speed.

2.5. Histological and Immunohistochemical Examinations. After sacrifice, the dorsal skin samples were fixed immediately in 4% paraformaldehyde and embedded in paraffin. Sections were then cut and stained with hematoxylin and eosin (H&E), Congo red, or toluidine blue. The number of eosinophils and mast cells was counted in three randomly selected areas per sample at x200 using a Lionheart FX microscope equipped with Gen5 imaging software (BioTek Instruments, Winooski, VT, USA).

2.6. Cell Culture and Treatments. HaCaT cells (human keratinocyte cell line) were cultured in DMEM supplemented with 10% FBS, 1% penicillin-streptomycin, at 37°C in a 5% CO₂ humidified environment. The medium was changed every two days during incubation, and the cells were made quiescent by starvation in serum-free medium for 24 h. Before stimulation with TNF-α and INF-γ (10 ng/ml each) for the indicated time, the cells were pretreated with GGDE (10, 100 µg/ml) or DEX (10 µM) for 1 h. The HMC-1 cells (a human mast cell line) were obtained from Merck Millipore (Darmstadt, Germany) and cultured in IMDM supplemented with 10% FBS, 1% penicillin-streptomycin, and 1.2 mM 1-thioglycerol at 37°C in a 5% CO₂ humidified environment. HMC-1 cells were incubated with 10 µg SP for 48 h and then with 1 nM CRH for 24 h. GGDE (10 or 100 µg/ml) or DEX (10 µM) was added 1 h before CRH. The PC12 cells (rat adrenal medullary pheochromocytoma cell line) were purchased from Korean Cell Line Bank (Seoul, Korea) and cultured in RPMI supplemented with 10% FBS, 1% penicillin-streptomycin, at 37°C in a 5% CO₂ humidified environment. The PC12 cells were incubated with GGDE (10 or 100 µg/ml) for 1 h and then stimulated with CORT (100 µM) for 24 h. The BV2 cells (a mouse microglial cell line) were cultured in DMEM supplemented with 10% FBS and 1% penicillin-streptomycin at 37°C in a 5% CO₂ humidified environment. BV2 microglial cells were pretreated with GGDE (10, 100, or 500 µg/ml) or DEX (10 µM) for 1 h and then stimulated with LPS (1 µg/ml) for 24 h.

2.7. Cell Viability. The cell viability was determined using an XTT assay. After treating cells with GGDE (10, 100, 500, or 1000 µg/ml) for 24 h, 50 µl of the XTT solution was added and incubated for 4 h. The absorbances were measured at 450 nm (reference wavelength 650 nm) using a microplate reader (Molecular Devices, CA, USA).

2.8. Enzyme-Linked Immunosorbent Assay (ELISA). Skin tissues were homogenized with tissue extraction reagent (Thermo Fisher Scientific, Vienna, Austria) and centrifuged at 10,000 × g for 20 min. The supernatants were collected and made up to a protein concentration of 0.5 µg/µl. The protein levels in the skin tissues and cell culture supernatants and the IgE levels in the serum were measured using commercial ELISA kits. The absorbance was measured at 450–540 nm using an automated microplate reader (Molecular Devices CA, USA).

2.9. Western Blot Analysis. Skin tissues were homogenized using a tissue extraction reagent, centrifuged at 10,000 × g for 20 min, and the supernatants were collected. HaCaT cells were washed with 1X PBS and then lysed with RIPA lysis buffer containing the protease and phosphatase inhibitors (Atto, Tokyo). After sonication, the cell lysates were
Figure 1: Continued.
centrifuged at 8000 × g for 15 min, and the supernatants were collected. The total protein levels were determined using the Bradford protein assay reagent (BioRad, CA, USA). Subsequently, 25–50 μg of the total proteins were separated by 7.5–10% SDS-PAGE electrophoresis and transferred to PVDF membranes (Merck Millipore, Darmstadt, Germany). The proteins were then blocked with 5% skim milk in 1X PBS for 2 h at room temperature. They were then incubated with the primary antibodies, followed by the secondary antibody horseradish peroxidase-conjugated anti-IgG. The blots were detected by enhanced chemiluminescence (BioRad), and the band intensities of the proteins were quantified using GelPro V3.1 software (Media Cybernetics, MD, USA).

2.10. Cell Apoptosis Analysis. The percentage of apoptotic cells was determined using the Muse Annexin V and Dead Cell kit (Millipore, Billerica, MA, USA) according to the manufacturer’s protocol. PC12 cells were incubated with GGDE (10, 100 μg/ml) for 1 h and stimulated with CORT (100 μM) for 24 h. Subsequently, 100 μl of Annexin V and Dead Cell reagent and 100 μl of a single cell suspension were mixed in a microtube and incubated in the dark for 20 min at room temperature. The cells were analyzed using a Muse Cell Analyzer (Millipore, Billerica, MA, USA).

2.11. Nitric Oxide (NO) Assay. The production of NO was measured using a Griess reagent assay. BV2 microglial cells were pre-treated with GGDE (10, 100, 500 μg/ml) or DEX (10 μM) for 1 h and stimulated with lipopolysaccharides (LPS, 1 μg/ml) for 24 h. Subsequently, 100 μl of supernatant was mixed with 100 μl of Griess reagent at room temperature. The absorbances were measured at 540 nm using an automated microplate reader (Molecular Devices CA, USA).

2.12. Statistical Analysis. The data were analyzed using a Student’s unpaired t-test. P values < 0.05 were considered significant. The in vivo results are presented as box and whisker plots with the medians as the middle lines, the 25th/75th quartiles as the box edges, and the minimum/maximum as whiskers. The in vitro data are presented as the mean ± SDs of three independent experiments.

3. Results

3.1. Effects of GGDE on AD Mice Exposed to PS. The DNCB treatment caused AD-like symptoms, such as erythema and edema, which were exacerbated by SI stress. GGDE suppressed the SI-induced increases in skin severity score, but no significant difference was observed between the SI-AD and GGDE groups (Figures 1(b) and 1(c)). No mortality was noted in the GGDE groups. Treatment with GGDE at 100 and 500 mg/kg did not show significant changes in body weights compared to the CON group (Figure 1(d)).

3.2. Effects of GGDE on Dorsal Skin Lesions in AD Mice Exposed to PS. Marked eosinophil infiltrations were observed in the SI-AD mice, and the mast cell infiltrations and epidermal thickness were significantly higher in the SI-AD mice than the AD mice. On the other hand, epidermal thicknesses in the GGDE (100 and 500 mg/kg) groups were significantly smaller than in the SI-AD group. Eosinophil and mast cell infiltrations in lesions were less severe in the GGDE mice than in the SI-AD mice (Figure 2(a)). The TNF-α, IL-13, eotaxin, and VEGF levels in the skin lysates were all significantly higher in the SI-AD mice than in the CON mice. Interestingly, although the SI-AD mice had significantly higher levels of TNF-α and VEGF than the AD mice, they also had significantly lower levels of IFN-γ. In contrast, the levels of eotaxin and VEGF were lower in the GGDE-treated
mice (100 and 500 mg/kg) than in the SI-AD mice, and the TNF-α and IL-13 levels were lower in the GGDE100 group than in the SI-AD group. Furthermore, mice in the AD and SI-AD groups had significantly higher serum IgE levels than the mice in the CON group, but the mice in the GGDE100 group had significantly lower IgE levels than those in the SI-AD group (Figure 2(b)). The negative effects of PS on the skin barrier function are well established [5, 10]. Loricrin and involucrin expression, which are positively associated with the skin barrier function, were significantly lower in the AD and SI-AD mice than in the CON mice. GGDE (100 and 500 mg/kg) prevented these decreases (Figure 2(c)).

3.4. Effects of GGDE on Inflammatory Response in HaCaT and HMC-1 Cells. The in vitro effects of GGDE were examined to understand its action mechanism. Initially, the cytotoxic effects were determined using XTT assays, and subsequent experiments were carried out at nontoxic concentrations (Figures 4(a) and 4(d)). TI (TNF-α and IFN-γ, 10 ng/ml each) stimulation increased the Th2 chemokine (TARC, MDC) levels in the HaCaT supernatants, but these increases were suppressed by GGDE (10 and 100 μg/ml). The STAT1 and NF-κB pathways play important roles as signaling mediators of the cellular responses to extracellular signals associated with the inflammatory response in AD [18]. GGDE (100 μg/ml) significantly inhibited the TI upregulated p-STAT1 expression and NF-κB p65 nuclear translocation by downregulating IκB-α phosphorylation (Figures 4(b) and 4(c)). The HMC-1 cells were treated with CRH and SP (CRH/SP) to mimic the PS-exaggerated AD condition [4]. GGDE (100 μg/ml) decreased CRH/SP-induced VEGF production and ERK phosphorylation in the HMC-1 cells, indicating that the therapeutic effect of GGDE involves mast cell suppression (Figure 4(e) and 4(f)).

3.5. Effects of GGDE on CORT-Induced Neurotoxicity in PC12 Cells. As shown in Figure 5(a), CORT reduced the viability of PC12 cells significantly. When treated with 100 μM CORT for 24 h, the cell viability decreased to approximately 60%; therefore, this concentration was used in subsequent experiments. Pretreatment with GGDE (100 μg/ml) increased cell viability in CORT-treated PC12 cells significantly (Figure 5(b)). Moreover, CORT induced apoptosis in PC12 cells, which was suppressed by the GGDE pretreatment (Figure 5(c)).

3.6. Effects of GGDE on LPS-Induced Neuroinflammation in BV2 Cells. GGDE did not show any cytotoxic effects on BV2 cells (Figure 6(a)); therefore, GGDE at concentrations of 10, 100, and 500 μg/ml was used for subsequent experiments. NO secreted by activated microglia plays an important role in neuroinflammation and neuronal death [19]. As shown in Figure 6(b), LPS increased NO production in BV2 cells, but this increase was reduced significantly by pretreatment with GGDE (100, 500 μg/ml). The levels of inflammatory cytokines TNF-α, IL-1β, and IL-6, were upregulated by LPS stimulation (Figure 6(c)). GGDE (100 or 500 μg/ml) suppressed TNF-α and IL-1β production, but the IL-6 level was decreased only by a pretreatment with GGDE at 500 μg/ml (Figure 6(c)).

3.7. Quantitative Determination of the Constituents in GGDE. The active constituents of GGD are nootkatone, α-cyperone, ergosterol, dehydrotrametenolic acid, pachymic acid, and tumulosic acid [20, 21]. Therefore, the amounts of these compounds in GGDE were determined by HPLC. Figures 7(b) and 7(c) show representative chromatogram patterns of the standard compounds and GGDE. The retention times of nootkatone and α-cyperone were 6.514 and 7.126 min, respectively. The concentrations of these two components in GGDE were 0.008 and 0.422 μg/mg, respectively (Figure 7(a)). Ergosterol, dehydrotrametenolic acid, pachymic acid, and tumulosic acid were not detected in GGDE.

4. Discussion
SI stress is a well-characterized type of PS used to investigate the impact of PS on AD [7]. In the present study, SI stress increased DNBC-induced mast cell infiltration, TNF-α, and VEGF production in dorsal skin lesions. SI stress reduced the production of IFN-γ in SI-AD mice to levels significantly lower than those of CON mice. These findings concur with studies that found that PS induced Th1/Th2 imbalance [3, 9]. Moreover, the OFT results showed that SI could be a potential factor in accelerating ADHD symptoms, such as hyper-locomotor activity in AD.

In AD, keratinocytes activated by allergens and proinflammatory cytokine secrete chemokines that induce immune cell infiltration of the skin lesions. The immune cell infiltration increases the levels of proinflammatory cytokines and maintains a chronic inflammation [22, 23]. GGDE decreased TNF-α, IL-13, eotaxin, IgE production, and reduced eosinophils infiltration in SI-AD mice. In addition, treatment of human keratinocyte HaCaT cells with GGDE suppressed Th2-attracting chemokines TARC and MDC.
Figure 2: Continued.
production and decreased STAT-1 and NF-κB activation. In particular, GGDE500 μg/ml and DEX, which was used as a positive control in this study, had a similar inhibitory effect on MDC production. These results suggest that GGDE can suppress the proinflammatory mediator and infiltration of immune cells by blocking the chemokine production in AD exposed to PS.

The PS-elicited secretion of CRH can enhance the production of VEGF in mast cells; hence, it can trigger or exacerbate AD [24]. The extent of mast cell infiltration and VEGF levels in SI-AD mice were significantly higher than in AD mice. Oral administration of GGDE reduced the serum CRH level in a dose-dependent manner. Together with the present results, it was hypothesized that GGDE might inhibit VEGF production in mast cells increased by CRH stimulation. The effects of GGDE on CRH/SP-treated HMC-1 cells were evaluated to understand the actions of GGDE at the cellular level better. SP can increase the expression of CRH receptor-1 in mast cells, and the application of SP and CRF to HMC-1 can mimic PS-associated AD conditions [4]. Consistent with the in vivo data, GGDE inhibited VEGF production in CRH/SP-treated HMC-1 cells by inhibiting p-ERK. These results suggested that the therapeutic effects of GGDE do not depend on the direct regulations of inflammatory mediators but on the indirect regulation of inflammatory response by regulating the release of CRH.

The rising levels of CORT have negative effects on skin barrier function and stratum corneum integrity [3, 25]. Increased CORT levels and decreased expression of the epidermal differentiation-related proteins were found in SI-AD mice, which were prevented by GGDE. A previous study showed that a natural ingredient, hesperidin, prevented GC-impaired skin barrier homeostasis by increasing the epidermal differentiation-related proteins [26]. These results suggest that GGDE can prevent skin barrier disruption by inflammatory cytokines and by CORT.

Patients with AD had a higher incidence of ADHD than non-AD patients [27, 28]. The underlying mechanism of this comorbidity is that AD-related PS causes hyperactivity in the HPA axis, leading to neurotoxicity of the hippocampus or prefrontal cortex. [11, 29]. High concentrations of CORT impair the HPA negative feedback loop, and prolonged high CORT levels induced further neuronal damage [5, 30]. Previous studies showed that hyperactivity is accompanied by increases in the circulating ACTH levels [31, 32]. Neuroinflammation, particularly microglial activation, has also been suggested to mediate stress-induced psychiatric disorders [33]. These results have shown that the oral administration of GGDE decreased the CORT and ACTH levels as well as the hyper-locomotion activity in SI-AD mice. In addition, GGDE has a protective effect on CORT-induced neurotoxicity in PC12 cells, which may be related to the inhibition of apoptosis. GGDE inhibited LPS-induced microglial activation by suppressing the production of NO, TNF-α, IL-1β, and IL-6 in BV2 cells. These findings show that GGDE may be a new treatment option for treating...
ADHD, the psychiatric comorbidity of stress-susceptible AD patients.

GGD is a traditional herbal prescription comprised of *Cyperus rotundus* L. and *Poria cocos* (Schw.) Wolf., which showed antiallergic and anti-inflammatory effects in previous studies [34, 35]. Topical application of *C. rotundus* extract suppressed infiltration of inflammatory cells and ear edema in TPA-induced skin inflammation in mice [36]. A preclinical study demonstrated that oral administration of *P. cocos* extract ameliorated DNCB-induced AD-like symptoms in mice by modulating the production of Th2 cytokines and the generation of Treg cells [37]. In the present study, the combination of *C. rotundus* and *P. cocos* in GGDE not only did have inhibitory effects on skin inflammation but also improved psychiatric comorbidity in AD mice exposed to SI stress, suggesting synergistic effects of this combination for the treatment of AD. HPLC showed that GGDE contains two major chemical constituents: α-cynerone and nootkatone. A herbal formula containing α-cynerone was reported to have potent antidepressant effects by regulating prefrontal cortex activation in a mouse model of depression induced by PS [38]. In contrast, nootkatone was reported to inhibit the Th2-attracting chemokines TARC and MDC in keratinocytes [39]. Overall, α-cynerone and nootkatone might at least be partially responsible for regulating the inflammation, HPA axis alteration, and hyperactive behavior in PS-exacerbated AD.

Figure 3: Effects of GGDE on hypothalamic-pituitary-adrenal (HPA) axis and behavioral changes in AD mice exposed to PS. (a) CRH, ACTH, and CORT levels in serum. (b) Behavior in the open field test. Representative open field test track sheets. Total distances (cm) traveled, distances traveled in the peripheral zone and central zone, and mean speeds (cm/s) in the 10-minute test. The results are presented as the means ± SDs (n = 5 per experiment). +P < 0.05, vs. the CON group; #P < 0.05, vs. the AD group; ∗P < 0.05, vs. the SI-AD group.
Figure 4: Continued.
Figure 4: Effects of GGDE on inflammatory mediator release in HaCaT and HMC-1 cells. (a) The HaCaT cell viabilities were assessed using XTT assays. (b) Effects of GGDE on TLR (TNF-α and IFN-γ)-induced TARC and MDC productions. (c) Western blot of p-STAT1, STAT1, p-IκB-α, IκB-α, β-actin, NF-κB p65, and lamin B in HaCaT cells. (d) HMC-1 cell viabilities were assessed using XTT assays. (e) Effects of GGDE on CRH/SP-induced VEGF production. (f) Western blot of p-ERK, ERK, p-p38, p38, p-JNK, JNK, and β-actin in HMC-1 cells. Dexamethasone (DEX) was used as a positive control. The results are presented as means ± SDs (n = 3 per experiment). *P < 0.05, vs. CON; #P < 0.05, vs. T/I-treated cells; ∗P < 0.05, vs. CRH/SP-treated cells.
ethanolic extract of GGD in an animal model. Therefore, further experiments will be needed to find appropriate concentrations to manage clinical symptoms and signs more effectively. Despite the limitations of the current study, these results provide a clue for treating patients with AD exacerbated by PS.

Figure 5: Effects of GGDE on CORT-induced neurotoxicity in PC12 cells. (a) Effects of CORT on the viability of PC12 cells. (b) Effects of GGDE on cell viability in CORT-stimulated PC12 cells. (c) Effects of GGDE on apoptosis rate in CORT-stimulated PC12 cells. The percentage of apoptotic cells was measured using Muse Annexin V and Dead Cell kit. Results are presented as means ± SDs (n = 3 per experiment). #P < 0.05, vs. CON; *P < 0.05, vs. CORT-treated cells.
Figure 6: Effects of GGDE on LPS-induced neuroinflammation in BV2 cells. (a) Effects of GGDE on the viability of BV2 cells. (b) Effects of GGDE on NO production in LPS-stimulated BV2 cells. (c) Effects of GGDE on the production of TNF-α, IL-1β, and IL-6 in LPS-stimulated BV2 cells. Results are presented as means ± SDs (n = 3 per experiment). #P < 0.05, vs. CON; *P < 0.05, vs. LPS-treated cells.
### Composition of GGDE

| Latin name          | Scientific name         | Used part                  | Amount (g) |
|---------------------|-------------------------|----------------------------|------------|
| Cyperi rhizoma      | Cyperus rotundus L.     | Rhizome                    | 20         |
| Poria sclerotium    | Poria cocos (Schw.) Wolf| Bacterium core centre white part | 10         |
| **Total**           |                         |                            | **30**     |

### Chemical components found in GGDE

| Compound                | Formula | Amount (μg/mg) |
|-------------------------|---------|----------------|
| Nootkatone              | ![Nootkatone](https://via.placeholder.com/150) | 0.008 |
| α-cyperone              | ![α-cyperone](https://via.placeholder.com/150) | 0.422 |
| Ergosterol              | ![Ergosterol](https://via.placeholder.com/150) | 0     |
| Dehydrotrametenolic acid| ![Dehydrotrametenolic acid](https://via.placeholder.com/150) | 0     |
| Pachymic acid           | ![Pachymic acid](https://via.placeholder.com/150) | 0     |
| Tumulosic acid         | ![Tumulosic acid](https://via.placeholder.com/150) | 0     |

_Figure 7: Continued._
Figure 7: Continued.
5. Conclusion

GGDE ameliorates the cutaneous inflammation, skin barrier disruption, HPA axis activation, and hyper-locomotor activity in AD mice exposed to SI. GGDE could inhibit chemokines production in keratinocytes and VEGF release in mast cells treated with SP and CRH. Moreover, GGDE has protective effects against CORT-induced neurotoxicity in PC12 cells and LPS-induced neuroinflammation in BV2 cells. These findings suggest that GGDE is a potential candidate as a treatment for patients with AD exacerbated by PS.

Abbreviations

HPA: Hypothalamic-pituitary-adrenal  
IL: Interleukin  
MDC: Macrophage-derived chemokine  
TARC: Thymus- and activation-regulated chemokine  
VEGF: Vascular endothelial growth factor.

Data Availability

The data used to support the findings of this study are included within the article.

Conflicts of Interest

The authors have no conflicts of interest to declare.

Authors’ Contributions

Ly Thi Huong Nguyen performed the experiments and the statistical analysis with the help of Uy Thai Nguyen. Min-Jin Choi performed the open field test. Tae Woo Oh designed and performed research using the BV2 mouse microglial cell line. Hung-Mook Shin and In-Jun Yang participated in the study design and helped write the manuscript.

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