Abstract. Background/Aim: Mesenchymal stem cells (MSCs) have been suggested as an alternative therapeutic option in atopic dermatitis. Palatine tonsils are lymphoepithelial tissue located around the oropharynx and have been proposed as one of the important alternative sources of MSCs. The purpose of this study was to evaluate the protective and therapeutic effects of tonsil-derived MSCs (TMSCs) in a 2,4-dinitrofluorobenzene (DNFB)-induced mouse model of atopic dermatitis (AD). Materials and Methods: The effect of TMSCs was evaluated in 20 C57BL/6J mice that were randomly divided into four groups (normal, DNFB-PBS, DNFB-TMSC7, and DNFB-TMSC16 group). TMSCs were subcutaneously injected into DNFB-sensitized mice on day 7 (DNFB-TMSC7 group) and day 16 (DNFB-TMSC16 group). Several parameters of inflammation were assessed. Results: Subcutaneously injected TMSCs significantly improved the inflammatory symptoms in a DNFB-induced AD model mice, particularly showing therapeutic effects rather than protective effects. TMSC treatment inhibited T-cell-mediated inflammatory responses by decreasing the levels of IL-6, IL-1β, TNF-α (Th1 cell marker), IL-4 (Th2 cell marker), and B-cell-mediated serum IgE. In contrast, TMSCs enhanced the anti-inflammatory cytokine TGF-β. Conclusion: In vitro and in vivo results suggest that TMSC treatment improved inflammatory skin lesions in the DNFB-induced AD mice model via immunomodulatory effects of the TMSCs. TMSCs inhibit T-cell and B-cell mediated responses, and enhance the anti-inflammatory responses.

Atopic dermatitis (AD) is a chronic inflammatory disease of the skin characterized by relapsing eczematous skin lesions with severe pruritus that impairs the quality of life (1, 2). AD affects 10-20% of children worldwide, and its incidence is increasing in industrialized countries (3). AD has multiple etiologies including genetic susceptibility, disruption of the epidermal barrier, and immune system dysregulation (2). The pathogenesis of AD is characterized by excessive Th2-derived inflammatory responses, resulting in elevated serum IgE levels and blood eosinophils (1, 4). Although AD has been described as a Th2-dominant disease, several studies have shown that AD is not exclusively explained by the Th2 immune response. AD is a biphasic inflammatory skin disorder (acute, chronic). The acute AD skin lesions show Th2 inflammation characterized by the infiltration of CD4+ T cells. Subsequently, the chronic phase demonstrates a Th1-type immune response (3, 5). Furthermore, recent studies showed that Th1, Th17, and Th22 might contribute to the pathogenesis of AD (4, 6).

Mesenchymal stem cell (MSC)-based therapy has been reported to regulate excessive inflammatory reactions in various autoimmune and inflammation-related diseases including graft-versus-host disease, inflammatory bowel disease, rheumatoid arthritis, and type I diabetes (7-9). Additionally, several studies revealed that MSCs can also improve allergic symptoms in asthma, allergic rhinitis, and dermatitis (10, 11). In general, bone marrow-derived MSCs have been widely used in research and for therapeutic purposes. However, they can also be successfully isolated from many other mesodermal tissues (12, 13). Palatine tonsils are lymphoepithelial tissue located around the
oropharynx and have been proposed as one of the important alternative sources of MSCs because of several advantages, which include the simple surgical process used to obtain the tissues (13, 14). Recently, tonsil-derived MSCs (TMSCs) showed therapeutic efficacy in allergic and autoimmune diseases such as allergic rhinitis and psoriasis by suppressing Th2 and Th17 responses (11, 15).

The purpose of this study was to investigate the protective and therapeutic effects of TMSCs in a 2,4-dinitrofluorobenzene (DNFB)-induced mouse model of AD at two different administration times, the day of the first DNFB challenge (day 7), and three days after the final DNFB challenge (day 16). Adipose tissue- and umbilical cord-derived MSCs were proposed as a therapeutic modality for AD in recent studies (1, 10, 16). However, to the best of our knowledge, this was the first study to demonstrate the efficacy of TMSCs in an AD mouse model. Several studies reported that repeated 2,4,6-trinitrochlorobenzene (TNCB) or DNFB application to mice skin induced AD-like clinical symptoms similar to those of AD in humans (3, 7, 17-19). Therefore, we evaluated the efficacy of TMSCs in AD using a well-established mouse model.

Materials and Methods

Preparation and culture of TMSCs. TMSCs were isolated and cultured according to a previously described method (13). The study protocol was approved by the Institutional Review Board of Hallym University Medical Center (2016-41). The patients or their legal guardians provided written informed consent to participate. Tonsillar tissue was obtained from a ten-year-old girl who underwent a tonsillectomy. The tonsillar tissue was minced and digested for 30 min at 37°C in high-glucose (4500 mg/l) Dulbecco Modified Eagle’s Medium (DMEM) (Welgene Inc., Gyeongsan, Republic of Korea) containing 10 mg/ml DNase (Sigma Aldrich, St. Louis, MO, USA) and 210 U/ml collagenase type I (Invitrogen, Carlsbad, CA, USA). The digested tissue was filtered through a wire mesh, and the cells were washed twice in high-glucose DMEM/20% fetal bovine serum (FBS) (Welgene Inc., Gyeongsan, Republic of Korea) and additional times in high-glucose DMEM/10% FBS. Ficoll-Paque (GE Healthcare, Little Chalfont, UK) density gradient centrifugation was used to obtain mononuclear cells. The cells (1×10^6) were plated in a 100-mm culture dish in high-glucose DMEM, 10% FBS, and 100 U/ml penicillin/streptomycin (Gibco, Grand Island, NY, USA). The non-adherent cells were removed after 48 h, and fresh culture medium was added to the adherent mononuclear cells (TMSCs). TMSCs at passage 5 were used for the experiments.

Histological analysis. All mice were euthanized on day 19 in compliance with the Animal Experiment Guidelines of the Hallym University Medical Research Institute. Ear and skin tissue samples were embedded in paraffin blocks and sectioned into 10-μm-thick slices. The sections were stained with hematoxylin and eosin (H&E) and toluidine blue according to protocols described by IHCWORLD (ihcworld.com). The thickness of the ear and the skin epidermis were measured using the ImageJ software (ImageJ 1.49v, National Institute of Health, Bethesda, MD, USA). Before euthanization and

Assessments

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under anesthesia, the spleen and inguinal lymph nodes were removed from all mice and weighed immediately (Tuning-fork vibration system, SHINKO DENSHI CO., LTD. Bunkyo-Ku, Tokyo, Japan).

In vivo tracking. A PKH26 red fluorescent cell linker (Sigma) was used to label the TMSC cell membranes. The TMSCs (2x10^6) were mixed with PKH26 solution and incubated for 5 min. Labeling was stopped with 1% bovine serum albumin (BSA, Bovogen Biologicals Pty Ltd., East Keilor, Australia) and the cells were washed with 10% DMEM. TMSCs labeled with PKH26 were injected into the subcutaneous tissue around the dorsal skin lesion using a 26-gauge sterile needle syringe in the TMSC groups. The skin tissue was harvested on day 19 and examined by fluorescence microscopy (Leica, Germany). For nuclear counterstaining, the sectioned skin tissues were incubated for 2 min with 10 μg/ml 4′-diamidino-2-phenylindole (DAPI) (Invitrogen) reagent in PBS. The cells were observed by fluorescence microscopy (Carl Zeiss Microscopy GmbH, Zeiss, Germany) under 200-fold magnification.

Immunohistochemical analysis. The sectioned tissue slides were dried in an oven for 30 min at 60°C. The deparaffinized tissues were denatured by microwaving in 10mM Tris-Hcl and 1mM EDTA (pH 9.0) and then blocked in 2% horse serum for 60 min. The slides were incubated in diluted primary antibodies for 12 h at 25°C, treated with 0.3% hydrogen peroxide in 1X PBS for 10 min, and incubated with horseradish peroxidase (HRP)-conjugated secondary antibody. HRP activity was detected with the 3,3′-diaminobenzidine (DAB) substrate. The slides were washed in PBS three times for 5 min each and counterstained with Mayer’s hematoxylin. The primary antibodies used were anti-TGF-β1 (1:100, Abcam, Cambridge, UK), TNF-α (1:50, Santa Cruz Biotechnology, Inc., Dallas, TX, USA), interleukin 6 (1:50, Santa Cruz Biotechnology, Inc.), and interleukin 4 (1:50 Santa Cruz Biotechnology). The secondary antibodies used were goat anti-mouse IgG (1:250, Thermo Fisher Scientific Inc., Waltham, MA, USA) and goat anti-rabbit IgG (1:250, Thermo Fisher Scientific). The stained skin tissues were observed by light microscopy (Carl Zeiss Microscopy GmbH). The immunopositive cells were measured using ImageJ (National Institute of Health).

Measurement of total serum IgE. Serum was prepared from whole blood by centrifugation at 1,000xg for 15 min. Capture antibody was coated on 96-well flat-bottom plates for 12 h at 4°C. The coated plates were washed and blocked with 10% PBS for 1 h at room temperature. Serial dilutions of the standard (1.6, 3.1, 6.3, 12.5, 25, 50, and 100 ng/ml) and samples (1:2 and 1:4) were incubated for 2 h at room temperature. The absorbance of the reactions was read at 450 nm within 30 min on a GloMax® Discover Microplate Reader (Madison, WI, USA). All procedures were performed in the dark.

Terminal-deoxynucleotidyl transferase-mediated nick end-labeling (TUNEL) assay. The slides were washed twice with fresh xylene (5 min each time), rehydrated in a graded ethanol series (absolute ethanol (100%) twice for 5 min, then 95%, 70%, and 50% ethanol each for 5 min), and rinsed in PBS for 5 min. The samples were permeabilized in protease K for 15 min at room temperature and rinsed in PBS three times for 5 min. The terminal deoxynucleotidyl transferase dUTP nick-end labeling (TUNEL) In situ Cell Death Detection Kit (Roche, Indianapolis, IN, USA) was used according to the manufacturer’s instructions. The slides were counterstained with 1 μg/ml DAPI (Invitrogen), incubated for another 10 min, rinsed in PBS three times for 5 min, and then sealed. A fluorescence scanning microscope (Olympus, Tokyo, Japan) was used to assess the fluorescence.

LPS-induced inflammatory response. NIH3T3 mouse embryo fibroblast cells were obtained from Hallym University (Chuncheon, Republic of Korea), and cultured in DMEM with 10% FBS in 5% CO2 at 37°C. The TMSCs were stimulated with 25 ng of lipopolysaccharide (LPS) in triplicate for 6 h. To determine the relative levels of TNF-alpha, IL-6, and IL-1β RNA transcripts, the cells were harvested using easy-BLUE (iNtRON Biotechnology, Gyeonggi-do, Republic of Korea) reagent and cDNA was synthesized according to the manufacturer’s instructions (iNtRON Biotechnology, Gyeonggi-do, Republic of Korea). cDNA was amplified by polymerase chain reaction (PCR) in a final reaction volume of 20 μl using Power SYBR Green PCR Master Mix (Thermo Fisher Scientific Inc.) and 10 pmol primer (TNF-α sense primer: GTTCTGTCCCTTCTACCTCAGT, anti-sense primer: GGTAGAGAATGGGAGACACC; IL-6 sense primer: CTAAAGCAGGAGAGTCTCAGT, anti-sense primer: CTTGCCGTAGAGATTATG; IL-1β sense primer: CTGGTCTGTAGAGTGCATCC, anti-sense primer: TGTCCTAGGGAGGAGGAC; GAPDH sense primer: ACCACAGTCTATGCCATCACC and anti-sense primer: TGCGACACTTGGTGTCTGA).

Statistical analysis. All data are presented as the mean±standard deviation. Statistical analyses were performed using the Statistical Package for Social Science software program (SPSS) (SPSS Inc., Chicago, IL, USA). p-Values were generated by the Student’s t-test, with statistical significance set at *p<0.05, **p<0.01, and ***p<0.005.

Ethical approval. Informed written consent was obtained from the legal guardians of all patients who participated in this study, and the study protocol was approved by the institutional review board of Hallym University Medical Center (2016-41). Animal studies were performed after receiving approval of the Institutional review board of Hallym University (Hallym 2019-74), Chuncheon, Republic of Korea.

Results

Gross analysis. To determine whether TMSCs could inhibit AD in mice, we induced AD-like skin lesions using DNFB and treated the mice with or without TMSCs (Figure 1). Figure 2a shows that TMSC treatment improved skin inflammation. Compared to the normal group, the DNFB-PBS group showed AD-like skin lesions, which exhibited as dried skin patches and scaling. The DNFB-TMSC7 group exhibited more severe eczematous skin lesions than the DNFB-PBS group on day 7 (before the subcutaneous administration of TMSCs). However, the lesional subcutaneous injection of TMSCs on day 7 enhanced the clinical severity of the DNFB-TMSC7 group compared to the DNFB-PBS group (Day 10). The DNFB-TMSC16 group also showed eczematous patches with scaling until day 16, before TMSC treatment. However, on day 19, these skin lesions were improved after TMSC administration. To
evaluate the protective effect of TMSCs on AD skin lesions, we compared the DNFB-PBS and DNFB-TMSC7 groups. On day 10, three days after TMSC treatment, the skin lesions of the DNFB-TMSC7 group were significantly improved compared to the DNFB-PBS group. The eczematous skin lesions were ameliorated in the DNFB-TMSC7 group, whereas the DNFB-PBS group showed scaling and dried skin patches with thickening of the skin. Although scaling and focal eczematous lesions were observed again in the DNFB-TMSC7 group on days 13, 16, and 19 because of repetitive DNFB challenges, the overall clinical severity was alleviated compared to the DNFB-PBS group.

It is well known that the weights of some immune organs such as the spleen and lymph nodes increase in response to the topical application of agents with allergenic or sensitizing potential (20). As shown in Figure 2b, an increased spleen weight was seen in the DNFB-PBS group but the spleen weights in the groups treated with TMSCs were lower.
Figure 3. Histological analyses of the ear and dorsal skin lesion on DNFB-induced AD mice. (a, b) H&E stained sections of the right ear and ear thickness (μm). (c, d) H&E stained section of the dorsal skin lesion and epidermal thickness (μm). Bar=200 μm (a, c-upper panel), Bar=50 μm (c-lower panel). DNFB: 2,4-dinitrofluorobenzene. TMSC: Tonsil-derived mesenchymal stem cell. *p<0.05, ***p<0.005.
Figure 4. Reduced mast cell infiltration in dorsal skin of the TMSCs treated AD mice. (c, d) TUNEL assay of the dorsal skin lesion on day 19. Bar=50 μm (a), Bar=200 μm (c). DNFB: 2,4-dinitrofluorobenzene; TMSC: tonsil-derived mesenchymal stem cell. *p<0.05, ***p<0.005.
Notably, DNFB-TMSC16 demonstrated significantly decreased spleen weight compared to the DNFB-PBS and DNFB-TMSC7 groups. The weight of the inguinal lymph nodes was also decreased in the TMSC treatment groups compared to the DNFB-PBS group but the difference was not significant.

**Histological analysis.** Histologic evaluation of the right ear revealed that the ear thickness was significantly increased in the DNFB-PBS group compared to the other groups (normal, DNFB-TMSC7, and DNFB-TMSC16). The DNFB-PBS group showed epidermal and dermal hyperplasia with the infiltration of inflammatory cells. However, TMSC treatment on day 7 and 16 significantly reduced the DNFB-induced ear thickness increases ($p<0.05$) (Figure 3a, b). H&E staining of the dorsal skin revealed that the epidermal hyperplasia seen in the DNFB-PBS group was attenuated by TMSC treatment (Figure 3c and d). Epidermal thickness in the DNFB-TMSC7 and DNFB-TMSC16 groups was significantly decreased compared to the DNFB-PBS group ($p<0.05$). We next performed toluidine blue staining to determine the degranulation of mast cells infiltrating the skin lesions (Figure 4a and b). Increased mast cell infiltration was observed in the dorsal skin of the DNFB-PBS group compared to the control group ($p<0.005$). However, significantly decreased mast cell infiltration was observed in the DNFB-TMSC7 ($p<0.05$) and DNFB-TMSC16 ($p<0.005$) groups compared to DNFB-PBS group. The TUNEL assay was used to investigate cell death (Figure 4c and d). In the DNFB-PBS group, TUNEL-positive cells (green) were observed in the epidermis and dermis. However, the number of TUNEL-positive cells was decreased in the DNFB-TMSC7 and DNFB-TMSC16 groups compared to the DNFB-PBS group ($p<0.005$).

**In vivo tracking of injected TMSCs.** We monitored the localization of the injected TMSC-PKH26 cells to evaluate the contribution of the TMSCs in the AD skin lesions. Fluorescence microscopy analysis on day 19 revealed that red fluorescent PKH26 dye gathered around the injected area, indicating that the TMSCs were primarily localized to the injected subcutaneous area (Figure 5). The nuclei were stained with DAPI (blue) in the merged images. Localization of TMSC-PKH26 cells was consistent with DAPI staining. Dead TMSCs were investigated using the TUNEL assay. The injected TMSCs in the DNFB-TMSC7 group survived until 12 days after injection. The DNFB-TMSC16 group showed focal apoptosis of the injected cells.

**Immunohistochemistry.** We evaluated the expression of IL-6 and TNF-α as proinflammatory cytokines (Figure 6). On day 19, the DNFB-PBS group showed significantly higher IL-6 expression compared to the other groups (normal, DNFB-TMSC7, and DNFB-TMSC16) ($p<0.005$). TMSC treatment attenuated IL-6 expression, and the DNFB-TMSC16 group demonstrated the lowest level of all groups. TNF-α expression showed a tendency similar to IL-6. The DNFB-TMSC7 and DNFB-TMSC16 groups showed decreased
TNF-α expression compared to the DNFB-PBS group. TMSCs also decreased the Th2 cytokine IL-4 (Figure 6). In contrast, TMSC treatment increased the expression level of the anti-inflammatory cytokine TGF-β (Figure 7).

Measurement of serum IgE. Elevated serum IgE is a hallmark of AD. The DNFB-PBS group showed the highest serum IgE level compared to the normal and TMSC treatment groups. On day 10, three days after TMSC injection, the serum IgE levels in the DNFB-TMSC7 group showed a significant decrease. The DNFB-TMSC16 group also showed a marked decrease in serum IgE on day 19, which was three days after the TMSC injection (day 16). However, another increase in IgE levels was observed in the DNFB-TMSC7 group on day 19. These results indicated that the serum IgE increase in AD mice was significantly attenuated by subcutaneous TMSC injection. However, repetitive DNFB challenges after TMSC injection could result in another increase in serum IgE levels (Figure 8).

LPS-induced inflammatory response: *In vitro analysis.* NIH3T3 fibroblast cells were stimulated with LPS (25 ng) to investigate whether TMSC treatment produced anti-inflammatory efficacy. As expected, LPS stimulation increased the mRNA levels of the inflammatory cytokines TNF-α, IL-1β, and IL-6. In contrast, TMSC treatment significantly decreased the mRNA levels of these three cytokines in LPS-stimulated NIH3T3 cells (p<0.005) (Figure 9).

**Discussion**

In this study, we demonstrated that subcutaneous administration of the human TMSCs could attenuate DNFB-induced AD-like skin lesions, presumably through the regulation of B cell-mediated IgE production and T-cell-derived inflammatory responses. We found that intralesional injections of TMSCs induced a significant decrease in allergic skin lesions, mast cell infiltration, IgE secretion, and the expression of inflammatory cytokines IL6 and TNF-α. In
addition, TMSC treatment increased the expression of TGF-β, which is considered an immunomodulatory factor.

Atopic dermatitis is an inflammatory skin disorder associated with asthma and allergic rhinitis that reduces the quality of life (21). The current treatment for AD include topical anti-inflammatory drugs (steroids, tacrolimus), antihistamines, and immunosuppressants. However, it is well known that the continual and long-term use of these medications usually causes side-effects (2, 20). Currently, MSC treatment has been suggested as a promising strategy for a variety of immune-mediated diseases and several studies have reported encouraging results with MSCs in the treatment of allergic diseases (1, 10, 11, 15, 16, 22). Recently, human umbilical cord blood (hUCB)- and adipose tissue (hAT)-derived MSCs were reported to alleviate atopic dermatitis (1, 4, 10, 16). In particular, the first clinical trial of hUCB for the treatment of AD was reported by Kim et al. (4). The underlying mechanisms of MSC effects on the immune system have not been clarified yet. However, it is generally believed that various mechanisms and molecules are related to the immunomodulatory potential of MSCs (11). TMSCs have also shown excellent immune modulatory effects in various disease models including muscular fibrosis (23), skin inflammation (15), B-cell-mediated immune responses (24), and autoimmune-mediated colitis (25, 26). In addition, TMSCs were shown to improve allergic rhinitis, an atopic disease (atopic dermatitis, allergic rhinitis, and asthma), by reducing Th2 cytokines and IgE secretion from B cells (11).

As expected, intralesional injection of TMSCs effectively improved AD-like skin lesions and inflammatory parameters in AD model mice. To explore the protective and therapeutic effects of TMSC, we subcutaneously injected TMSCs at different times. TMSCs were injected on the first day of the DNFB challenge (day 7) in the DNFB-TMSC7 group. After TMSC administration, repetitive DNFB challenges on days 10 and 13 were performed. In contrast, the DNFB-TMSC16 group received TMSC treatment three days after the final DNFB challenge. The current study showed that TMSC treatment rapidly improved the severity of the skin lesions. However, re-challenge with DNFB after TMSC injection seemed to make the skin lesions worse again in the DNFB-TMSC7 group. The protective effect of hUCB-derived MSCs
on AD was demonstrated in a previous study (10). To evaluate the protective effect of hUCB-derived MSCs in that study, cells were injected four times during the experimental period. Our study showed different results from the hUCB-derived MSCs, and the difference is presumed to be due to the different number of injections or injected cell numbers. Since the TUNEL assay confirmed that injected TMSCs could survive until 12 days after injection, we could expect that several TMSCs injections or different numbers of injected cells could show a protective effect on AD.

Histology also revealed that hyperplasia of the epidermis and dermal infiltration of mast cells in the DNFB-PBS group were reduced in the TMSC-injected groups, particularly in the DNFB-TMSC16 group compared to the DNFB-TMSC7 group. The TUNEL assay results were also consistent with the histologic results.

Inflammatory cytokines released from activated T cells determine the maintenance and severity of AD (27). It has been reported that hUCB-derived MSC extracts suppressed the expression of Th1- (IL-6, IL-1β, and TNF-α) and Th2- (IL-4 and IL-5)-type cytokines in LPS, TNF-α, and IFN-γ-stimulated HaCaT cells (1). The current study examined the expression of pro- and anti-inflammatory cytokines in the skin tissue of mice. We found that the intralesional injection of TMSCs decreased the expression levels of inflammatory cytokines IL-6 and TNF-α, as well as Th2 cytokine IL-4. These results indicate that the TMSCs inhibited Th1- and Th2-mediated inflammation. TMSC treatment also significantly down-regulated inflammatory cytokines (IL-6, IL-1β, and TNF-α) after the stimulation of fibroblast cells with LPS in vivo.

IgE is known to play a major role in the degranulation of mast cells and causes both acute and chronic phase skin symptoms in AD (3). Consistent with other studies, our results also demonstrated that the total serum IgE levels were significantly increased by repeated DNFB applications. However, lesional injections of TMSCs significantly decreased the total serum IgE levels in DNFB-treated mice within three days after injection. We also found that repeated DNFB treatment after TMSC injection resulted in another increase in IgE levels. Taken together, the TMSC injection protocol in this study showed effective therapeutic effects on AD. However, a protective effect on AD was not clearly identified.

The underlying anti-allergic mechanisms of TMSCs on AD have not been clarified in this study. However, Samivel et al.
demonstrated that therapeutic potency of TMSCs on allergic rhinitis could be attributed to inhibition of innate cytokine (IL-25 and IL-33) secretion and of eotaxin expression on the nasal mucosa (11). In addition, supernatant of TMSCs have shown immunomodulatory effect on the allergic rhinitis by the inhibition of T cell activation via MAP kinas, p65, and NFAT1 (28). Additional presumed mechanism of therapeutic efficacy of TMSCs on AD is anti-oxidative effect (29, 30). An important underlying mechanism that decrease epidermal health and function is keratinocyte damage caused by high levels of intracellular ROS (29). Therefore, antioxidants have been proposed as a skin protector. Recently, antioxidant effects of MSC therapy have been observed in various disease models (30). However, further investigations are needed, including in vitro studies to examine related adverse effects and the molecular mechanisms involved in the anti-allergic effects of TMSCs on AD.

In conclusion, our results indicated that the subcutaneous injection of TMSCs significantly improved inflammatory skin lesions in the DNFB-induced AD mice model, particularly showing therapeutic effects rather than protective effects. Based on the in vitro and in vivo results, the therapeutic potency of TMSCs in AD was attributed to the immunomodulatory effects of the TMSCs. These effects inhibited T-cell and B-cell-mediated inflammatory responses, decreasing the levels of IL-6, IL-1β, TNF-α (Th1 cell marker), IL-4 (Th2 cell marker), and serum IgE (B-cell mediated). In contrast, the TMSCs enhanced the anti-inflammatory cytokine TGF-β.

Conflicts of Interest

The Authors have no conflicts of interest to declare.

Authors’ Contributions

Harry Jung: Methodology, Formal analysis, Investigation, Visualization, Writing original draft; Gil Myeong Son: Methodology, Investigation, Visualization, Validation; Jae Jun Lee: Project administration, Data curation; Hae Sang Park: Conceptualization, Data curation, Supervision, Funding acquisition, Writing review & editing.

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