

Fig. 1. Schematic model of epithelial–mesenchymal transition (EMT). Several factors, such as TGF- β , EGF and Wnt- β catenin signaling cascade, induce the expression of transcription factors, SNAI1, SNAI2 and TWIST1 in epithelial cells, which in turn led to the downregulation of E-cadherin (CDH1) expression, the upregulation of N-cadherin (CDH2), vimentin and fibronectin and the acquisition of morphological and functional features of mesenchymal cells.

MET) are indispensable for the differentiation and migration of specialized cell types and the acquisition of the complex three-dimensional structure of internal organs. However, EMT has recently been implicated to participate in tissue repair, cancer progression and organ fibrosis.

In this article, we review the current knowledge of EMT, focusing on its regulation in skin morphogenesis, wound repair, and cutaneous fibrosis, and discuss a future perspective of EMT-targeted therapeutics.

2. EMT in skin morphogenesis

EMT plays an important role in the gastrulation in the early development of vertebrates. Carver et al. studied embryos deficient for the Snai1 (*Snai1*), essential gene for EMT. *Snai1*^{-/-} mice were embryonic lethal. At E6.5, *Snai1*^{-/-} mutant embryos were indistinguishable from the embryos of heterozygous and wild-type littermates. At E7.5, however, the homozygous mutant embryos were smaller than the embryos of their littermates. By E8.5, the *Snai1*^{-/-} embryos were severely retarded compared to those of littermates and were resorbed. At E7.5, *Snai1*^{-/-} mutant mesoderm cells expressed mesoderm markers, however, they failed to show the characteristic mesenchymal morphology and had an appearance of a columnar epithelium. *Cdh1* expression was not downregulated in the *Snai1*^{-/-} mutant mesoderm cells, while it was decreased in those in the littermates [3].

To examine EMT in skin morphogenesis, Kong et al. performed an immunohistochemical study with antibodies against Cdh1, α -smooth muscle (α -SMA), and fibronectin using mouse skin sections. They showed that cells expressing epithelial markers, Cdh1, zonula occludens 1 protein (ZO-1) and occludin were present in the dermal area during skin development with the highest expression from P2 to P5. The co-expression of an epithelial marker, Cdh1, and a mesenchymal marker, α -SMA, was observed in the dermis around P3–P6, suggesting some dermal cells might have developed through EMT [4].

Snai1 mRNA and protein are expressed transiently at the hair bud stage of hair follicle morphogenesis in mice. In contrast, Franci et al. detected *Snai1* expression only in mesenchymal cells of the skin, particularly in the mesenchymal cells clustered below the hair buds of embryonic skin and in the dermal papillae of adult hair follicles. The discrepancy between the two studies has not been resolved [5,6].

Snai1 expression leads to the transcriptional downregulation of *Cdh1*, which results in junctional remodeling and bud formation. Transforming growth factor (TGF)- β 2 is necessary to transiently induce *Snai1* and activate the Ras-mitogen-activated protein kinase (MAPK) pathway in the bud [5].

Snai1 misexpression under the keratin 14 promoter in the epidermis of the transgenic mice led to a downregulation of *Cdh1* and α -catenin, along with epidermal hyperproliferation and a reduction in an intercellular adhesion [5].

Slug (SNAI2) is another member of the Snail zinc-finger transcription factor family that plays crucial roles in EMT. Although *Snai1* expression is essential for normal embryonic development in mice, *Snai2*-null embryos are viable. *Snai2*^{-/-} mice were small and had a diluted coat with the characteristic white forehead blaze. The histological sections of testis from *Snai2*^{-/-} mice revealed testicular atrophy and a reduced number of Leydig cells. Moreover, *Snai2*^{-/-} mice suffered from severe anemia. The activation of c-kit by stem cell factor (SCF) specifically induced *Snai2* expression, and Kit⁺ cells derived from *Snai2*^{-/-} mice exhibited migratory defects. In this context, the human piebaldisms cases provide an interesting finding. Southern blot analysis revealed that 3 patients had evident heterozygous deletions of the *SNAI2* gene encompassing the entire coding region out of 17 unrelated patients with piebaldism, who lack apparent *KIT* mutations [7–9].

At the beginning of periderm formation at E12, *Snai2* was expressed in essentially all skin epithelial cells and in the underlying mesenchyme. *Snai2* staining was observed in progressively fewer dermal cells as primitive mesenchymal cells matured. In the placode, hair germ, and peg stages of hair follicle development, *Snai2* expression was prominent in the thickened and invaginating epithelium but was absent from the underlying mesenchymal cells that ultimately form the dermal papillae [10].

In adult mice, *Snai2*-expressing keratinocytes were clustered around hair follicles. In developing follicles, *Snai2* was stably expressed in the outer root sheath, hair matrix cells and some dermal papilla cells.

3. EMT in skin tumors

3.1. EMT in squamous cell carcinoma

Recently, we reported a case of squamous cell carcinoma that mimicked atypical fibroxanthoma and expressed both vimentin and SNAI1. Atypical fibroxanthoma, first described by Helwig in 1963, typically takes place on the head and neck of the elderly patients as a solitary nodule that is often ulcerated. Histologically, atypical mitotic figures and multinucleated cells are commonly seen in this tumor seemingly originating from mesenchymal stromal cells. Squamous cell carcinoma occasionally mimics atypical fibroxanthoma and can express vimentin, a mesenchymal marker. EMT in spindle cell squamous cell carcinoma has been reported by another Japanese group [11–14].

Hoot et al. reported a role of transforming growth factor (TGF)- β -Smad2 signaling pathway in the formation and progression of squamous cell carcinoma. Mice with keratinocyte-specific Smad2 deletion exhibited an accelerated formation and malignant progression of chemically induced squamous cell carcinomas compared with wild-type mice. Moreover, they revealed that the lack of SMAD2 expression in human squamous cell carcinoma was associated with reduced expression of CDH1, higher expression of SNAI1 and poor differentiation, indicating the involvement of EMT [15].

Regulation of TP63 isoforms may be one of the targets of SNAI1 and SNAI2 in cervical squamous cell carcinoma. *TP63* is a *TP53*-related gene that contains two alternative promoters, which give rise to transcripts that encode proteins with (TP63) or without

(Δ Np63) an amino-transactivating domain. The expression of Δ N and TAp63 isoforms was down- and up-regulated *in vitro* by both SNAI1 and SNAI2, respectively. In SCC tissues, Δ N and TAp63 isoform expressions were, respectively, reduced and increased, when displaying a high immunoreactivity for SNAI1 and/or SNAI2. Δ Np63 silencing reduced cell–cell adhesion and increased the migratory properties of cancer cells. Therefore, the regulation of TP63 by SNAI1 and SNAI2 may play an important role in SCC tumor progression.

Herfs et al. revealed the reduced density of CD1a⁺ Langerhans cells and downregulation of cell-surface CDH1 in an epithelial metaplasia of the uterine cervix. Langerhans cells are intraepithelial dendritic cells that initiate immune responses against antigens, including tumor antigens, on both skin and mucosal surfaces. Langerhans cells *in vitro* adhered poorly to keratinocytes transfected with either SNAI1 or SNAI2 DNA. Therefore, downregulation of CDH1 in keratinocytes by TGF- β -SMAD-SNAI1 signaling pathway not only leads to the EMT, but also the local immunodeficiency with a reduced function of Langerhans cells [16,17].

3.2. EMT in malignant melanoma

To determine whether human melanoma cells have decreased levels of CDH1, Tang et al. examined several human malignant melanoma cell lines for CDH1 expression. Of nine melanoma cell lines studied, six showed negative staining and three (SK-MEL-1, SK-MEL2 and SK-MEL-28) showed positive staining. Of the three CDH1 expressing cell lines, only SK-MEL-1 had CDH1 at a level comparable to that of normal melanocytes, while SK-MEL2 and SK-MEL-28 had decreased CDH1 expression.

Poser et al. analyzed the expression level of SNAI1 in melanoma cell lines. Significant SNAI1 expression was found in all 8 melanoma cell lines which had a reduced CDH1 expression, whereas no SNAI1 expression was detected in primary human melanocytes. Transient transfection of a SNAI1 expression plasmid into human primary melanocytes led to the significant downregulation of CDH1, whereas transient and stable transfection of an anti-sense SNAI1 construct induced reexpression of CDH1 in melanoma cell lines [18,19].

To explore a regulatory system for SNAI1 expression, Palmer et al. analyzed chromatin structural changes associated with SNAI1 transcription in melanoma cells. A robust DNase hypersensitive site (HS) was found in the 3' region of A375 melanoma cells, in which SNAI1 was highly expressed, but not in keratinocytes or primary melanocytes. A physical interaction between the 3' enhancer and promoter was observed in SNAI1-expressing cells, demonstrating a direct role for the enhancer in SNAI1 expression. This finding suggested that the SNAI1 promoter is constitutively packaged in a poised chromatin structure that can be activated in melanoma cells by a tissue specific enhancer, which physically contacts the promoter [20].

3.3. EMT in extramammary Paget's disease

Hirakawa et al. performed an immunohistochemical study and revealed that EMT markers, N-cadherin (CDH2) and vimentin, were not expressed by Paget cells at the early stage of carcinoma *in situ*, but expressed in invasive extramammary Paget's disease. They also revealed that CDH2 and vimentin expressions in extramammary Paget's disease significantly correlated with reduced overall survival. Cytoplasmic CDH1 expression was detectable by the early stages of tumor progression in subpopulation of patients with a poor outcome. Kaplan–Meier analysis showed that cytoplasmic CDH1 was a prognostic factor for reduced overall survival in extramammary Paget's disease, indicating that

EMT-like process plays a key role in promoting tumor invasion within the primary sites [21].

4. EMT in wound healing

Successful wound healing is a complex process involving cells of the epidermis, dermis, vasculature, and the immune system. Two distinct cellular mechanisms directly contribute to the closure of skin wounds: keratinocyte-driven reepithelialization, which is achieved through a combination of migration and reduced intercellular adhesion in the epidermis proximal to the wound margin, and fibroblast-mediated contraction of the newly formed connective tissue bed beneath the wound site, pulling the edges of the wound closer together [22].

The migratory activity and the reduced intercellular adhesion of keratinocytes observed in reepithelialization recapitulate several aspects of EMT. Savagner et al. analyzed the expression of Snai2 during the wound healing by *in situ* hybridization. Snai2 expression was elevated in keratinocytes bordering cutaneous wounds in mice *in vivo*, in keratinocytes migrating from mouse skin explants *ex vivo*, and in human keratinocytes at wound margins *in vitro*. Epithelial cell outgrowth from skin explants of Snai2 deficient mice was severely compromised. Overexpression of Snai2 in keratinocytes caused increased cell spreading and desmosomal disruption. Moreover, *in vitro* wound healing was markedly accelerated in keratinocytes that ectopically expressed Snai2, suggesting a critical role for Snai2 in epithelial keratinocyte migration and wound healing [23].

The reepithelialization of excisional wound healing was significantly impaired in Snai2 deficient mice. Furthermore, 8 out of 20 Snai2 null mice but no wild type mice developed non-healing cutaneous ulcers in response to chronic UV irradiation. This knockout study also revealed an important role of Snai2 in modulating EMT in successful wound repair [24].

Epidermal growth factor (EGF) treatment of human keratinocytes induced activating phosphorylation of ERK5 and up-regulation of SNAI2 transcription. Ectopic activation of ERK5 led to an increased SNAI2 mRNA expression level and faster wound healing. In contrast, expression of shRNA specific for ERK5 in keratinocytes decreased SNAI2 expression and motility response to EGF and led to an altered more compact morphology, along with disruption of the desmosome organization of keratinocytes. The results indicated an EGFR/ERK5/SNAI2 pathway in the control of EMT in wound healing by EGF [25,26].

5. EMT in skin fibrosis

5.1. EMT in postmenopausal frontal fibrosing alopecia

It had been thought that fibrosis is formed by the pathological activation of interstitial fibroblasts and the resultant conversion to myofibroblasts producing the fibrotic collagen network. In a renal fibrosis, however, elegant cell tracing studies have shown that a significant portion of these myofibroblasts arises from the conversion of epithelial cells through an EMT process [27].

Moreover, Kim et al. revealed that EMT also takes place during the development of pulmonary fibrosis. To examine the capacity of alveolar epithelial cells for EMT, they generated mice expressing β -galactosidase (β -gal) exclusively in lung epithelial cells. In an established model of pulmonary fibrosis, overexpression of active TGF- β 1, β -gal-positive cells expressing mesenchymal markers accumulated. The increase in vimentin-positive cells within lungs was nearly all β -gal-positive, indicating epithelial cells as the main source of mesenchymal expansion in this model. *Ex vivo*, primary alveolar epithelial cells cultured on provisional matrix compo-

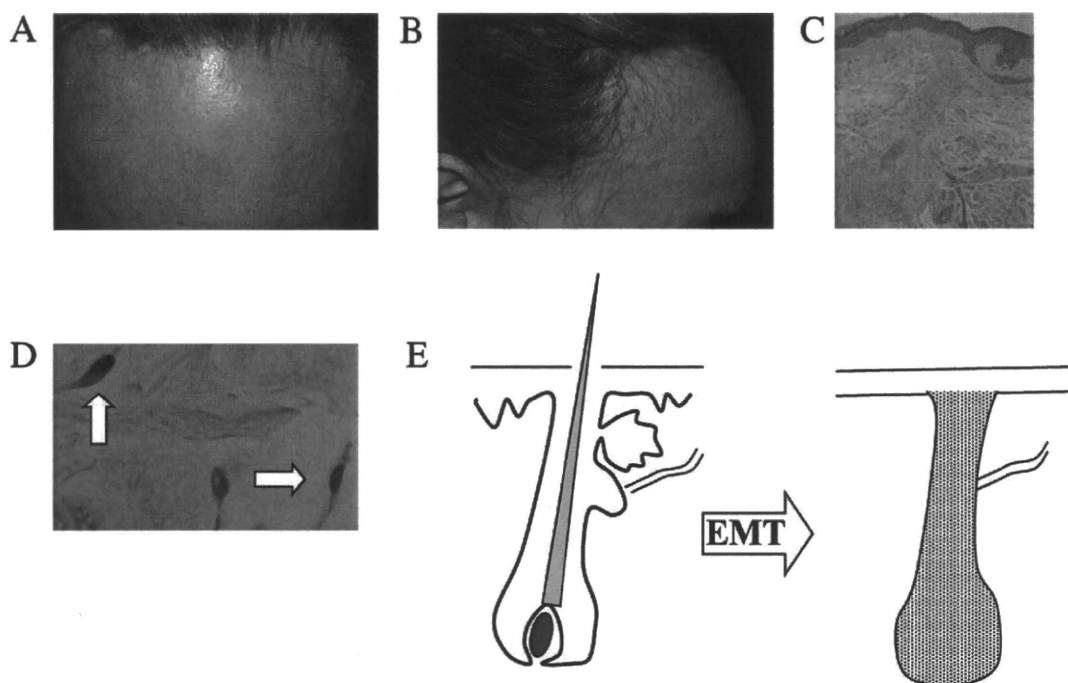


Fig. 2. (A and B) Frontotemporal recession with uniformly pale skin in a postmenopausal frontal fibrosing alopecia patient. (C) Histology of a biopsy specimen taken from the temporal region of the patient. An arrector pili muscle inserting into the fibrotic tissue surrounded by sparse lymphocytes. (D) SNAI1 positive cells found in the fibrotic dermis (arrows). (E) Proposed schematic formation of postmenopausal frontal fibrosing alopecia. EMT in hair follicles leads to the fibrosing alopecia.

nents, fibronectin or fibrin, underwent robust EMT via integrin-dependent activation of endogenous latent TGF-β1 [28].

Postmenopausal frontal fibrosing alopecia is characterized by a progressive frontotemporal recession due to a scarring hair loss. Histological examination shows a reduction or loss of terminal hair follicles and a replacement by fibrosis around the affected hair follicles. Lymphocytic infiltration is usually observed in a lichenoid pattern in the upper dermis. Altal frontal fibrosing alopecia is considered as a variant of lichen planopilaris, its exact pathogenesis still remains to be clarified [29,30].

We reported a case of postmenopausal frontal fibrosing alopecia showing the expression of SNAI1 in the fibrotic dermis, suggesting a role of EMT in the pathogenesis (Fig. 2A–D). The

existence of SNAI1 positive cells in the fibrotic dermis of our case suggested that the fibroblasts are in part derived from the hair-follicle cells via an EMT process (Fig. 2E). SNAI1 up-regulates several chemokines which may play a role in lichenoid reaction in frontal fibrosing alopecia [31].

5.2. EMT in systemic sclerosis

Systemic sclerosis (SSc) is an autoimmune disorder characterized by extensive fibrosis associated with collagen synthesis increased by the myofibroblasts and its accumulation. There are a wide variety of hypothesis on the origin of myofibroblasts in SSc patients. Postlethwaite et al. suggested that conventional, circu-

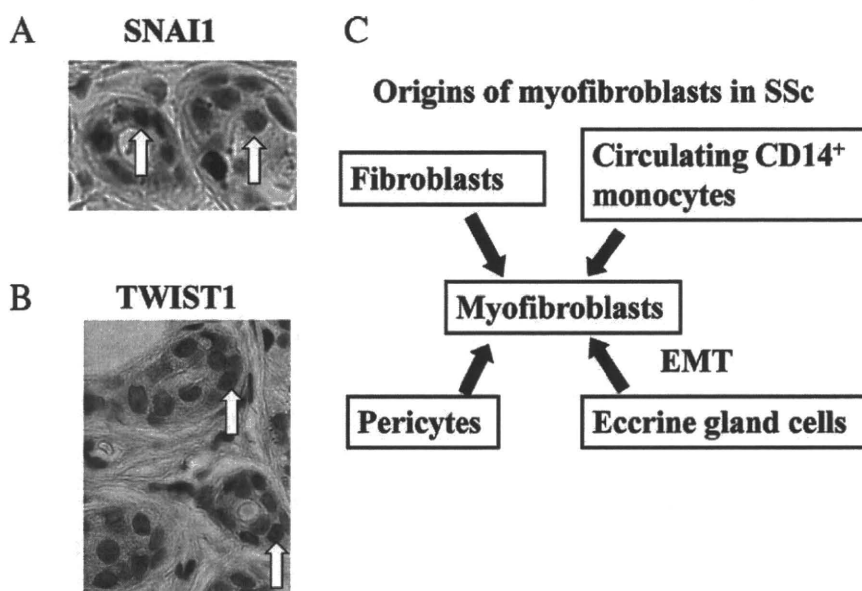


Fig. 3. (A) Expression of SNAI1 in the eccrine glands in a systemic sclerosis patient (arrows). (B) Expression of TWIST1 in the eccrine glands in a systemic sclerosis patient (arrows). (C) Several hypotheses on the origin of myofibroblasts which play central roles in the pathogenesis of systemic sclerosis (SSc). Myofibroblasts have been proposed to be derived from fibroblasts, circulating CD14⁺ monocytes, pericytes and eccrine gland cells.

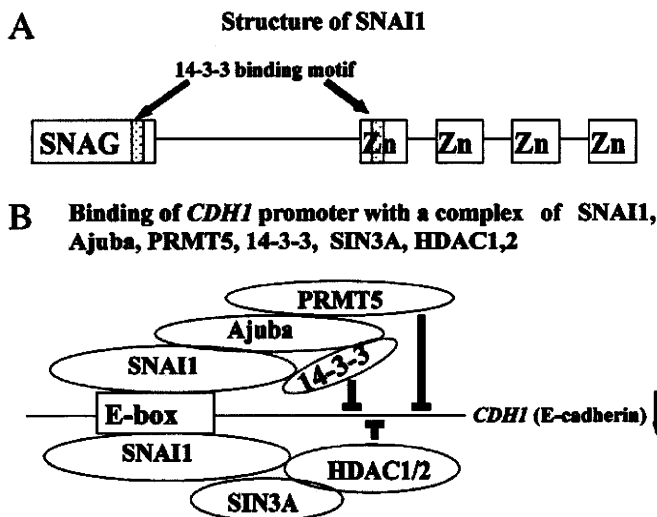


Fig. 4. (A) Structure of SNAI1. SNAI1 contains an NH₂-terminal SNAG (Snail/Slug and Gfi-1) repression domain and COOH-four COOH-terminal tandem C₂-H₂ zinc finger motifs. (B) SNAI1 binds to the E-box of *CDH1* promoter with zinc finger motifs. SNAI1 is associated with other proteins and forms a multiprotein complex.

lating CD14⁺ monocytes transdifferentiate into SSc myofibroblasts. Another group postulated a link between vascular damage and the formation of myofibroblasts from pericytes in SSc. Pericytes are primitive mesenchymal cells and can differentiate into myofibroblasts, cartilage and bones [32,33].

To examine the expression of EMT markers in the skin of three SSc female patients, we performed an immunohistochemistry against forearm skin specimens using antibodies against EMT markers, SNAI1 and TWIST1. Both SNAI1 and TWIST1 expressions were observed in the eccrine glands (Fig. 3A and B) in each dSSc patient. Neither SNAI1 nor TWIST1 expression was observed in two age-matched females' skins. These results suggested that SNAI1 and TWIST1 expression may be induced in eccrine glands at the early and/or progressive stage of dSSc. A part of the cells of eccrine glands, with a tissue-stem cell like character or not, might undergo EMT and differentiate into the myofibroblasts, contributing, at least partially, to the pathogenesis of SSc (Fig. 3C) [34,35].

6. Structure of EMT inducer, SNAI1, and its associated molecules

The SNAI1 protein contains an NH₂-terminal SNAG (Snail/Slug and Gfi-1) repression domain and COOH-four COOH-terminal tandem C₂-H₂ zinc finger motifs (Fig. 4A). SNAG motif was first identified as the growth factor independence-1 (*GFI-1*) proto-oncogene encoded protein. The SNAG domain comprises the first 21 amino acid residues in the N terminus, a potent and transferable repression motif.

Ayyanathan et al. searched SNAG domain-associated proteins by performing yeast two hybrid systems using SNAG domain as a bait. They identified Ajuba, a multiple LIM domain protein that can function as a corepressor for the SNAG domain. Ajuba interacts with the SNAG domain *in vitro* and *in vivo*, colocalizes with it, and enhances SNAG-mediated transcriptional repression (Fig. 4B) [36].

Moreover, the same group identified protein arginine methyltransferase 5 (PRMT5) as an effector recruited to SNAI1 through an interaction with AJUBA that functions to repress the SNAI1 target gene, *Cdh1*. PRMT5 binds to the non-LIM region of AJUBA and is translocated into the nucleus in a SNAI1- and AJUBA-dependent manner. The depletion of PRMT5 promotes *Cdh1* expression [37].

Hou et al. also defined functional 14-3-3 binding motifs in SNAI1 and AJUBA, which selectively bind 14-3-3 isoforms. In

SNAI1, an NH₂-terminal motif in the SNAG domain cooperates with a COOH-terminal high-affinity motif for binding to 14-3-3 proteins (Fig. 4A). Coordinate mutation of both motifs abolishes 14-3-3 binding and inhibits SNAI-mediated gene repression and EMT differentiation. SNAI1, 14-3-3 proteins, and Ajuba form a ternary complex that is detected through chromatin immunoprecipitation at the endogenous *Cdh1* promoter [38].

Snai1 is also associated with histone deacetylase 1 (HDAC1) and HDAC2, and the corepressor Sin3A through the SNAG domain of *Snai1*. *Snai1* requires HDAC activity to repress *Cdh1* promoter and overexpression of *Snai1* is correlated with deacetylation of histones H3 and H4 (Fig. 4B). Thus *Snai1* repressed *Cdh1* by the recruitment of chromatin remodeling and histone modifying activities [39].

7. Regulation of EMT

EMT is controlled by various factors through both transcriptional and post-transcriptional ways. The regulation of EMT was excellently reviewed by Peinado et al. EMT is induced by TGF- β -bone morphogenetic protein (BMP)-SMAD pathway, Wnt- β catenin signaling, and receptor tyrosine kinases (RTKs). RTKs are activated by different signals, such as fibroblast growth factor (FGF), platelet derived growth factor (PDGF) and epidermal growth factor (EGF) (Table 1). Comparative analysis of *SNAI1* and *SNAI2* promoters from different species has identified conserved and functional response elements, such as AP1 and AP4 sites, SMAD-binding elements, LEF1 binding sites and two conserved E-boxes. *SNAI1* or *Snai1* expression is also controlled by a mucin, *Helicobacter pylori* infection, a zinc-finger transcription factor, Wilms' tumor-1 (Wt1), polycomb group protein B lymphoma MoMLV insertion region 1 homolog (Bmi-1), and phosphoinositide-dependent protein kinase (PDK) 1. *SNAI1* is also regulated posttranscriptionally through ubiquitination and degrading processes by molecules, such as GSK3 β and CK1 ϵ . In the following sections, we focus on recent findings on EMT inducers and repressors [40–48].

Recently several lines of evidence have shown that sex hormones, androgens and estrogens, play crucial roles in mediating EMT processes. Androgens exert their biological effects by binding to the androgen receptor and inducing its transcriptional activity. Androgens induced the EMT pattern in prostate tumor epithelial cell with SNAI1 activation and lead to significant changes in prostate cancer cell migration and invasion potential. Expression levels of androgen receptors inversely correlated with androgen-mediated EMT in prostate tumor epithelial cells, pointing to a low androgen receptor content required for the EMT phenotype. On the contrary, estrogen receptor β plays an important role in maintaining an epithelial phenotype and repressing mesenchymal characteristics in prostate carcinoma. Poorly differentiated tumors exhibit diminished estrogen receptor

Table 1
Summary of reported inducers and repressor of EMT.

Inducer	Repressor
Androgen	Estrogen-ER β
Estrogen-ER α	GSK3
TGF- β 1, Wnt- β catenin	CK1
<i>Helicobacter pylori</i>	
MUC4 mucin, EGF	
Wilms' tumor 1 (Wt1)	
Bmi-1	
Hypoxia-HIF	
VEGF, FGF, PDGF	
RAS-MAPK	

β expression. Loss of estrogen receptor β is sufficient to promote an EMT [49,50].

The estrogen receptor α signaling plays an important role in human breast cancer formation and progression. Human breast cancers exhibit a strong direct correlation between estrogen receptor α and CDH1 expression by immunohistochemistry. When estrogen receptor α was knocked down in the estrogen receptor α positive cell lines, SNAI2 increased, CDH1 decreased, and cells became spindly and exhibited increased Matrigel invasion. When estrogen receptor α was overexpressed in the estrogen receptor α -negative cell lines, 17 β -estradiol decreased SNAI2 and increased CDH1 expression, and decreased invasiveness of tumor cells in Matrigel [51].

8. Conclusion

In this review, we focused on the important roles of EMT in wound healing, tumor formation and cutaneous fibrosis. In addition, we reviewed the recent findings on the regulation of EMT by various factors. The concept of EMT has been widely accepted just for a decade, and there may remain other skin diseases in which EMT plays a crucial role in the pathogenesis. Appropriate control of EMT in the skin by the modulation of EMT inducers and repressors can be a new therapeutic target in future for skin diseases, including skin ulcer, cutaneous malignant tumors and systemic fibrosis.

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Topical Cholecystokinin Depresses Itch-Associated Scratching Behavior in Mice

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Cholecystokinin (CCK) serves as a gastrointestinal hormone and also functions as a neuropeptide in the central nervous system (CNS). CCK may be a downregulator in the CNS, as represented by its anti-opioid properties. The existence of CCK in the peripheral nervous system has also been reported. We investigated the suppressive effects of various CCKs on peripheral pruritus in mice. The clipped backs of ICR mice were painted with CCK synthetic peptides and injected intradermally with substance P (SP). The frequency of SP-induced scratching was reduced significantly by topical application of sulfated CCK8 (CCK8S) and CCK7 (CCK7S), but not by nonsulfated CCK8, CCK7, or CCK6. Dermal injection of CCK8S also suppressed the scratching frequency, suggesting that dermal cells as well as epidermal keratinocytes (KCs) are the targets of CCKs. As determined using real-time PCR, mRNA for CCK2R, one of the two types of CCK receptors, was expressed highly in mouse fetal skin-derived mast cells (FSMCs) and moderately in ICR mouse KCs. CCK8S decreased *in vitro* compound 48/80-promoted degranulation of FSMCs with a transient elevation of the intracellular calcium concentration. These findings suggest that CCK may exert an antipruritic effect via mast cells and that topical CCK may be clinically useful for pruritic skin disorders.

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INTRODUCTION

Itch or pruritus is an unpleasant cutaneous sensation associated with the immediate desire to scratch (Ikoma *et al.*, 2006; Steinhoff *et al.*, 2006). Histamine was long considered the only mediator of pruritus, but this assumption has changed dramatically in the past two decades. Peripheral itch can be evoked in the skin either directly, by mechanical and thermal stimuli, or indirectly, through chemical mediators (Ikoma *et al.*, 2006). Itch may also be generated in the central nervous system (CNS) independent of peripheral stimulation. Different pruritic diseases involved different itch mediators, including histamine, neuropeptides, proteases, prostaglandins, serotonin, acetylcholine, cannabinoids, opioids, bradykinins, cytokines, biogenic amines, neurotransmitters, and ion channels (Steinhoff *et al.*, 2006).

Neuromediators such as neuropeptides and neurotrophins, as well as their receptors, have an important role in peripheral itch. Neuropeptides, as represented by pruritogenic neuropeptide substance P (SP), are produced by sensory nerves, but they can also be elaborated by epidermal keratinocytes (KCs), mast cells, fibroblasts, and other cutaneous immunocompetent cells (Ohkubo and Nakanishi, 1991; Scholzen *et al.*, 1998). Many other mediators are also produced by and secreted from skin-constituent cells in close contact to sensory nerves (Steinhoff *et al.*, 2006). They can activate and sensitize pruritic nerve endings and even modulate their growth, as do nerve growth factors and chemorepellents (Yamaguchi *et al.*, 2008; Yosipovitch and Papoiu, 2008). As for the receptor system, it is notable that KCs, mast cells, and fibroblasts express neurokinin-1 receptor (Ohkubo and Nakanishi, 1991; Scholzen *et al.*, 1998), which is the receptor for SP. These complicated neurophysiological interactions yield itch and render its therapeutic control difficult.

Cholecystokinin (CCK) is a peptide hormone in the gastrointestinal tract (Dufresne *et al.*, 2006), but it also serves as a neuropeptide (Ma *et al.*, 2006). Among neuropeptides, CCKs are most abundantly present in the CNS and involved in numerous physiological functions such as anxiety, depression, psychosis, memory, and feeding behavior (Dufresne *et al.*, 2006). CCKs also have anti-opioid properties in the CNS (Pommier *et al.*, 2002; Mollereau *et al.*, 2005) and exert a nociceptive effect in the spinal cord (Wiesenfeld-Hallin *et al.*, 1999). Furthermore, their existence in the peripheral nervous system has been reported (Moriarty *et al.*, 1997). In various CCKs, sulfation of the tyrosine at position 7

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Abbreviations: CCK, cholecystokinin; CCK2R, CCK2 receptor; CCK7S, sulfated CCK7; CCK8S, sulfated CCK8; CNS, central nervous system; FSMC, murine fetal skin-derived mast cell; KC, keratinocyte; SP, substance P

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from the C-terminus is a posttranslational modification that makes them biologically active peptides (Ma et al., 2006). Varying lengths of CCKs with or without sulfate have been studied for their activities as gastrointestinal hormones (Bonetto et al., 1999) and as CNS players (Rehfeld et al., 2007). Two types of CCK receptors have been identified: CCK1R (CCK-A receptor) and CCK2R (CCK-B receptor). CCK1R usually requires sulfated CCKs. CCK2R affords a binding site to both sulfated and nonsulfated ligands, but nonsulfated CCKs have affinities that are decreased 10- to 50-fold (Dufresne et al., 2006). CCKs exert their biological functions by interacting with CCK receptors located on multiple target cells in the CNS and on peripheral nerve endings (Dufresne et al., 2006; Rehfeld et al., 2007; Zheng et al., 2009). In the brain, sulfated CCK8 (CCK8S) is most abundant (Rehfeld et al., 2007) and possesses one of the strongest endogenous anti-opioid properties (Ma et al., 2006). Prolonged opioid exposure increases the expression of CCKs and CCK2R in the CNS, where CCK8S negatively modulates opioid responses and maintains homeostasis of the opioid system (Pommier et al., 2002; Agnes et al., 2008). Thus, it seems that CCKs are downregulators of opioid-mediated pruritus in CNS. However, the function of CCKs, the distribution of CCK receptors in the skin, and their effects on peripheral pruritus remain unelucidated (Ma et al., 2006).

In this study, we investigated the antipruritic effects of different lengths of CCKs with or without sulfate, which were topically applied onto the skin of mice. Results suggest that CCKs are capable of reducing SP-induced scratching at least by affecting the function of mast cells. Our findings may lead to the development of previously unreported antipruritic strategies.

RESULTS

Depressive effects of topical application of CCKs on SP-induced itch-associated response

We tested various CCKs for their ability to suppress scratching behavior induced by intradermal SP injection in ICR mice (Andoh et al., 2001). We first examined the effect of percutaneously applied CCK on the response. Because CCK8S has an inhibitory ability in the CNS (Pommier et al., 2002; Dufresne et al., 2006) and gastrointestinal tract (Miyasaka et al., 2004), CCK8S and the other CCK constructs consisting of smaller amino acids with or without sulfate (Table 1) were tested for their antipruritic activities. ICR mice were painted with various CCKs on the clipped, tape-stripped rostral part of the back skin (~1.8 cm²) and then injected with SP intradermally at the CCK-painted site.

The topical application of CCK8S significantly reduced the scratching frequency, as compared with the nonapplied control (Figure 1a). Sulfated CCK7 (CCK7S) also reduced the scratching frequency, to a level comparable to that with CCK8S, whereas nonsulfated CCK8, CCK7, and CCK6 had no ability to suppress scratching. These results showed that CCKs have an antipruritic ability, and CCK constructs bearing sulfation of position 7 are required for the ability to suppress the scratching behavior. In this itch-associated-scratching model, we also administered CCK8S intradermally together

Table 1. Amino acid sequences of CCK and its constructs

Sulfated CCK8 (CCK8S)	Asp-Tyr (SO ₃ ⁻)-Met-Gly-Trp-Met-Asp-Phe-NH ₂
Nonsulfated CCK8 (CCK8)	Asp-Tyr-Met-Gly-Trp-Met-Asp-Phe-NH ₂
Sulfated CCK7 (CCK7S)	Tyr (SO ₃ ⁻)-Met-Gly-Trp-Met-Asp-Phe-NH ₂
Nonsulfated CCK7 (CCK7)	Tyr-Met-Gly-Trp-Met-Asp-Phe-NH ₂
Nonsulfated CCK6 (CCK6)	Met-Gly-Trp-Met-Asp-Phe-NH ₂

Abbreviation: CCK, cholecystokinin.

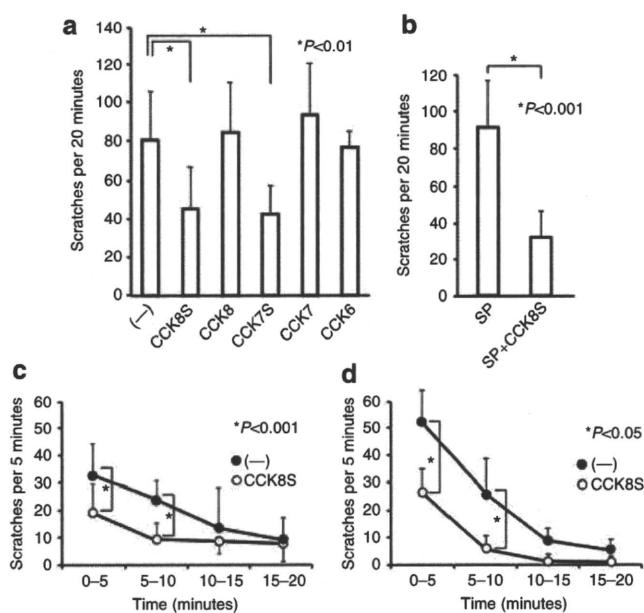


Figure 1. Suppression of substance P (SP)-induced itch by percutaneous application or intradermal injection of cholecystokinins (CCKs). (a) ICR mice were painted on the clipped skin with CCKs (0.5 nmol per site) and injected intradermally with SP (100 nmol per site). The group depicted with “(-)” represents mice injected with vehicle (acetone/olive oil) alone. The numbers of mice were as follows: CCK (-), n = 12; CCK8S, n = 10; CCK8, n = 6; CCK7S, n = 6; CCK7, n = 3; and CCK6, n = 4. (b) ICR mice were injected intradermally with SP or with SP and CCK8S. Each bar represents the mean of scratches per 20 minutes. CCK (-), n = 5 and CCK8S, n = 8. (c) Time course of scratching frequency in mice untreated (n = 12) or topically administered CCK8S (n = 10). (d) Time course of scratching frequency in mice untreated (n = 5) or intradermally injected with CCK8S (n = 8). CCK7, nonsulfated CCK7; CCK8, nonsulfated CCK8; CCK7S, sulfated CCK7; CCK8S, sulfated CCK8.

with SP to mice. The scratching frequency induced by SP was profoundly suppressed by the simultaneous injection of CCK8S (Figure 1b). When the time course of effects induced by CCK8S was examined, both topical application (Figure 1c) and intradermal injection (Figure 1d) of CCKs reduced the scratching frequency at intervals of 0–5 minutes and 5–10 minutes.

These findings raised the possibility that epidermal and dermal cells participating in the development of itch are affected by CCKs.

CCK2R expression in KCs and mast cells

Both KCs and mast cells are the possible targets of CCKs in the suppression of SP-induced pruritus. It is known that KCs express the receptors for itch-related molecules, neurokinin-1 receptor for SP, H1 receptor for histamine, and protease-activated receptor-2 for tryptase (synthetic agonist, SLIGRL-NH₂). Mast cells are well known to possess neurokinin-1 receptor (Scholzen *et al.*, 1998; Liu *et al.*, 2006). We investigated the expression of the CCK receptors CCK1R and CCK2R in KCs and mast cells. Primary cultures of KCs from ICR mice and fetal skin-derived mast cells (FSMCs) were collected, and mRNAs for CCK1R and CCK2R were quantitated by real-time PCR, RT-PCR, and western blotting analyses. The two types of cells expressed both CCK1R and CCK2R at the mRNA and protein levels (Figure 2). When the expression levels of real-time PCR analysis were normalized with the β -actin

level, the two receptors were more highly expressed in FSMCs than in KCs.

In a preliminary DNA microarray analysis of normal human epidermal KCs, we investigated the levels of gene expression from stimulation with SP (10⁻⁶ M), histamine (10 μ g ml⁻¹), or SLIGRL-NH₂ (100 nM) under high calcium (0.6 mM) concentration. The CCK2R expression by SP-stimulated KCs was most elevated; its level was 2^{8.67} times that of the control (Supplement Data online). Such an increased CCK2R expression was not observed with histamine or SLIGRL-NH₂. This observation suggested that CCK2R has an important role in the suppression of SP-stimulated KCs. To determine the effect of SP on KCs, ICR mouse KCs were incubated with SP for 30 minutes to 24 hours, and the expression of CCK2R and CCK1R was analyzed by western blotting. CCK2R was detected with antibodies as a 78 kDa band (Morisset *et al.*, 2003) and increased in KCs exposed to SP for 5 or 6 hours (Figure 3a). By contrast, CCK1R expression was not augmented by SP (Figure 3b). Thus, the expression of some receptors for CCK is enhanced by SP, depending on the receptor.

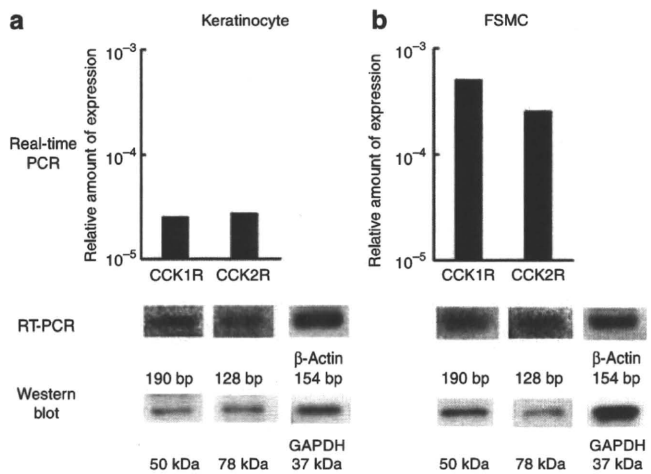


Figure 2. CCK-A receptor (CCK1R) and CCK-B receptor (CCK2R) mRNA expression in keratinocytes and mast cells. Cultured ICR mouse keratinocytes (a) and fetal skin-derived mast cells (FSMCs) (b) were examined for the expression of mRNAs for CCK1R and CCK2R using real-time PCR and RT-PCR and for the expression of their proteins using western blotting analysis. Data are expressed as the amount of expression relative to β -actin. Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) was used as control for western blotting.

Suppression of FSMC degranulation by CCK8S with transient elevation of intracellular calcium

The above findings suggested that CCKs bind to CCK2R on KCs and mast cells and downmodulate their itch-related biological functions. Given that the level of CCK2R expression was higher in FSMCs than in KCs and the application of CCKs immediately exerted its inhibitory effects on the scratching behavior, it is possible that CCKs directly affect mast cells. We therefore focused on mast cells as the primary target. The addition of compound 48/80 induced degranulation of FSMCs (Figure 4a). This promoted degranulation of FSMCs, which was significantly inhibited by CCK8S at 10⁻⁸ or 10⁻⁷ M concentration.

To test whether CCK8S evokes signal transduction in mast cells, we measured the intracellular calcium using confocal recording. The fluorescence intensity of 15 cells positively responding on addition of CCK8S was monitored with computerized color change on a display using a digital image processing system. The mast cells exhibiting green fluorescence of Fluo-4-AM at any time point between 40 and

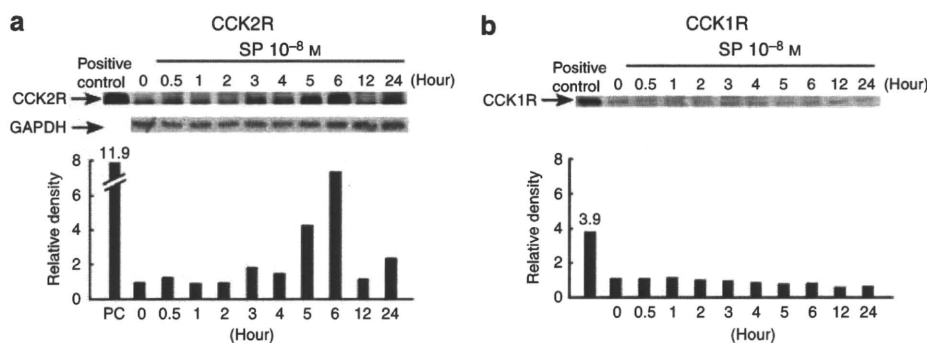


Figure 3. Western blotting of CCK-A receptor (CCK1R) and CCK-B receptor (CCK2R) in mouse keratinocytes stimulated with substance P (SP). ICR mouse keratinocytes were incubated with SP (10⁻⁸ M) for 0.5–24 hours and harvested at each time point. The expression levels of CCK2R (a) and CCK1R (b) were analyzed by western blotting. NIH/3T3 whole lysates were used as positive control. Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) was used as control for western blotting and exhibited similar intensities in the samples cultured for 0–24 hours. The intensity of each band was subjected to densitometric analysis and expressed as density relative to the nontreated control (0).

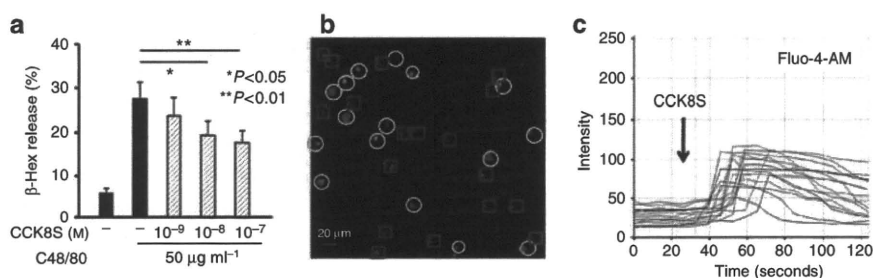


Figure 4. Suppression of fetal skin-derived mast cell (FSMC) degranulation by sulfated CCK8 (CCK8S) with transient intracellular calcium increase. (a) FSMCs were incubated with or without CCK8S in the presence of compound 48/80 (C48/80; $50 \mu\text{g ml}^{-1}$) for 15 minutes. Supernatants and cell lysates were incubated with *p*-nitrophenyl-*N*-acetyl- β -*D*-galactosaminide, and β -hexosaminidase (β -Hex) release was determined as the ratio between activity in the supernatant and cell lysate, multiplied by 100. (b) FSMCs were loaded with Fluo-4-AM (green) and stained for CD117 (c-kit, red). The percentage of CCK8S-responding FSMCs was determined from the fluorescence traces of all cells. Circles represent responding FSMCs and squares represent nonresponding FSMCs. Bar = $20 \mu\text{m}$. (c) The addition of CCK8S to FSMC culture induced an increase in intracellular calcium level.

80 seconds after CCK8S addition were defined as positively responding cells (Figure 4b and c); these are marked with a yellow circle in Figure 4b. The addition of CCK8S to the FSMC culture induced a prompt rise of calcium that persisted for more than 2 minutes and then gradually declined, indicating that CCK8S induces signaling via CCK2R in mast cells. The percentage of responding FSMCs was determined from the fluorescence traces of all cells. As shown in Figure 4b, in which circles and squares represent CCK8S-responding and nonresponding FSMCs, respectively, 45% (16 of 36 cells) of FSMCs were typically activated by CCK8S.

DISCUSSION

Despite their original name of cholecyst stimulators, CCKs have been known to serve as CNS modulators (Dufresne *et al.*, 2006) and CCK8S downmodulates opioid responses (Pommier *et al.*, 2002; Agnes *et al.*, 2008). As suggested in our present study, CCKs may also have a function of downmodulator in the peripheral itch. The topical application of CCKs suppressed the itch-related scratching behavior evoked by SP injection. The expression of CCK2R, but not CCK1R, in murine and human KCs was markedly enhanced by exposure to SP. It is likely that CCK2R, whose expression is augmented by SP, has a down-regulatory role in the excess peripheral itch.

Among the CCK constructs consisting of different numbers of amino acids with or without sulfate, CCK8S and CCK7S, but not CCK8, CCK7, or CCK6, exerted an antipruritic action. Therefore, in a comparison between CCK8S and CCK8 and between CCK7S and CCK7, sulfation of position 7 is required for the ability to suppress the scratching behavior. This is in accordance with the observation that CCK8S has an anxiogenic effect (Harro, 2006), a depressant effect, and an anti-opioid effect (Noble *et al.*, 1999), but the effects of CCK are limited by desulfation of the tyrosine (Tokunaga *et al.*, 1993).

Although the exact mode of antipruritic action of CCKs is still speculative, some informative observations were obtained from the study. Both percutaneous application and dermal injection were effective routes for CCK8S in suppressing itch, suggesting that epidermal and/or dermal constituents are the targets of CCKs. Furthermore, the administration of CCK8S rapidly inhibited the SP-induced itch. We thus investigated the mechanism of CCK suppression of itch in

in vitro studies, focusing on mast cells. CCK8S significantly reduced the degranulation of FSMCs with elevation of intracellular calcium concentration, suggesting that mast cell degranulation is inhibited by CCK8S through CCK2R. Alternatively, or additionally, KCs are possibly involved in CCR action. KCs are thought to mediate pruritus by releasing inflammatory mediators and neurotrophins. Therefore, the antipruritic effect of CCK8S on KCs, if any, seems to need a time lag to operate. As the therapeutic action of topical CCK8S is rapid, mast cells may be more likely targeted by CCK8S than KCs. The high expression of CCK2R in mast cells further supports the notion that mast cells are sensitive to CCKs. However, SP was reported to cause scratching behavior in both mast cell-deficient and wild-type mice (Hossen *et al.*, 2003), suggesting a mast cell-independent mechanism of SP-evoked itch. Given this observation, it is possible that another skin constituent, such as the sensory nerves, is affected by CCKs. The target cells and the actions of CCK8S on them warrant elucidation in the future.

The antipruritic effects of CCK8S and CCK7S on this mouse itch model may afford a therapeutic use for the CCKs for the treatment of skin diseases with peripheral itch. Given that the major target of CCKs is mast cells, urticaria, atopic dermatitis, contact dermatitis, and other types of eczematous dermatitis would be successfully treated with CCKs topically applied to the skin. The therapeutic effects of CCKs on peripheral itch may shed light on the mechanisms underlying the physiological regulation of itch, and it is notable that CCKs have potential as a topical drug for pruritic skin diseases.

MATERIALS AND METHODS

Animals

Female ICR mice were obtained from KBT Oriental (Tosu, Japan). These mice were maintained in the Laboratory Animal Research Center of the University of Occupational and Environmental Health under specific-pathogen-free conditions and used at the age of 6–8 weeks. All animal experiments were performed according to the guidelines approved by our university for the care and use of animals.

Reagents

SP, CCK8S, nonsulfated CCK8, CCK7S, and nonsulfated CCK6 were purchased from the Peptide Institute (Osaka, Japan). Nonsulfated

CCK7 was purchased from Funakoshi (Tokyo, Japan). Compound 48/80 was purchased from Sigma (St Louis, MO).

Itch-associated scratching behavior and topical application of CCKs

The hair was clipped over the rostral part of the each mouse's back 1 or 2 days before the experiment. Before behavioral recording, each animal (four per observation) was put into an acrylic cage (18 × 23 × 11 cm) for at least 1 hour for acclimation. An intradermal injection of SP (100 nmol in 50 μ l of physiological saline per site) induced scratching of the skin around the injected site with the hindpaws (Andoh *et al.*, 2001). CCKs were topically applied in two ways: percutaneous application and intradermal injection. For percutaneous application, the shaved back skin was stripped with Scotch tape (3M) six times on the day before the experiment (Nishijima *et al.*, 1997). The clipped backs of the mice were painted (1.8 cm², 1.5 cm in diameter) with 20 μ l CCK (0.5 nmol per site) in acetone/olive oil (3:1) 3–4 minutes before the SP injection (100 nmol per site). For intradermal application, mice were injected intradermally with CCK (0.5 nmol per site in 20 μ l physiological saline), followed immediately by SP injection at the same site (100 nmol per site). Immediately after treatment, the animals were put back into the same cage and their behavior was videotaped for 40 minutes using a digital video camera (Andoh *et al.*, 2001). Using the video, the frequency of scratching toward the injected site by the hindpaws was recorded for the first 20 minutes.

Murine KCs

The skin of ICR newborn mice was peeled within 24 hours after birth and incubated with 0.05% collagenase (collagenase from *Clostridium histolyticum* type 2; Sigma) for 2 days at 37 °C in a 5% CO₂ atmosphere. Epidermal sheets were collected and suspended in culture medium. Cell suspension was plated into six-well plates (Corning Glass Works, Corning, NJ) and grown to subconfluence in CnT-07 medium (Funakoshi).

Murine mast cells

The method for preparation of murine FSMCs has been described previously (Yamada *et al.*, 2003). Day 14 fetal skin was dissociated by 0.25% trypsin for 20 minutes at 37 °C in Hank's balanced salt solution (Gibco, Carlsbad, CA). Erythrocytes were lysed using RBC-lysing buffer (Sigma) and, after washing, cells were resuspended in complete RPMI, which is RPMI-1640 (Gibco) supplemented with 5% fetal bovine serum (Gibco), 1% nonessential amino acids (Gibco), 1% sodium pyruvate (Gibco), antibiotic/antimycotic (Invitrogen, Carlsbad, CA), and 0.1% 2-mercaptoethanol (Gibco). Fifty milliliters of the cell suspension in each 175 cm² 75-flask (Corning Glass Works) was cultured in the presence of 20 ng ml⁻¹ recombinant mouse IL-3 (Invitrogen) and 20 ng ml⁻¹ recombinant mouse stem cell factor (Biosource, Camarillo, CA) at 37 °C without changing the medium. Nonadherent and loosely adherent cells were harvested. The cells were centrifuged and resuspended in RPMI, layered on 40% Percoll (GE Healthcare, Buckinghamshire, England), and centrifuged at room temperature. The cell pellet at the bottom was used as FSMCs. For flowcytometric analysis, cells were stained with phycoerythrin-conjugated anti-mouse Fc ϵ R1 (eBioscience, San Diego, CA) and Alexa Fluor 700-conjugated anti-mouse CD117 (c-kit; eBioscience) for 20 minutes on ice, washed twice, and then resuspended with FACS

buffer. Flow cytometric analysis was performed with FACS Canto (BD, Franklin Lakes, NJ) and FlowJo software (TreeStar, San Carlos, CA). The purity of FSMCs was 95%, as determined by both Fc ϵ R1 and CD117 positivity by flow cytometry.

Real-time quantitative RT-PCR analysis

Total RNA was extracted from the cell pellet using the PureLink RNA Mini Kit (Invitrogen). cDNA was reverse transcribed from total RNA samples using the TaqMan Reverse Transcription reagents (Applied Biosystems, Carlsbad, CA). mRNA expression was quantified by real-time PCR using SYBR Green Dye (PE Biosystems, Foster City, CA) with an ABI PRISM 7000 Sequence Detection System (Applied Biosystems) according to the manufacturer's instructions. Primers were designed using Primer Bank for the human and murine genes, and they were constructed by Invitrogen. The sequences are as follows: murine β -actin: 5'-GGCTGTATCCCCTCCATCG-3' and 5'-CCAGTTGGTAACAATGCCATGT-3'; murine CCK2R: 5'-GATGGCTGCTACGTGCAACT-3' and 5'-CGCACCACCCGCTTCTTAG-3'; and murine CCK1R: 5'-CACGCTGGTTATCACGGTG-3' and 5'-GCCATGAAGTAGGTGGTAGTC-3'. The conditions for the real-time PCR were as follows: 2.0 minutes at 50 °C, 10 minutes at 95 °C, and then 50 cycles of amplification consisting of 15 seconds at 95 °C and 1 minute at 60 °C. All heating and cooling steps were performed with a slope of 20 °C per second. Relative expression was calculated using the delta Ct method.

Western blotting

KCs from ICR mice were stimulated with SP (10⁻⁸ M) in six-well plates (Corning Glass Works) for 0.5–24 hours, harvested with a rubber policeman, and subjected to extraction with RIPA buffer (Wako, Osaka, Japan; 50 mM Tris-HCl (pH8.0), 150 mM sodium chloride, 0.5% (wt/vol) sodium deoxycholate, 0.1% (wt/vol) SDS, and 1.0% (wt/vol) NP-40 substitute). Protein samples (20 μ g) were separated by 8% SDS-polyacrylamide gel electrophoresis and electroblotted onto polyvinylidene difluoride membranes for 2 hours at 180 mA. After blocking with 5% skim milk solution, the membranes were incubated with rabbit anti-mouse CCK1R (SC-33220; 1:1,000, Santa Cruz, Santa Cruz, CA), CCK2R (SC-33221; 1:1,000, Santa Cruz) polyclonal antibodies, or GAPDH (SC-25887; 1:1,000, Santa Cruz), and the reaction was detected with horseradish peroxidase-conjugated goat anti-rabbit IgG (1:3,000, Bio-Rad, Hercules, CA). Immunoblots were visualized using the ECL Plus Western Blotting Detection Reagents (GE Healthcare) according to the manufacturer's protocol. Bands were quantified by densitometry with the help of a CS Analyzer version 2.0 (ATTO, Tokyo, Japan). NIH/3T3 whole cell lysate (Santa Cruz) was used as a positive control.

Mast cell degranulation

Degranulation of FSMCs was assessed by β -hexosaminidase assay (Yamada *et al.*, 2003; Noguchi *et al.*, 2005). FSMCs were washed with Tyrode's buffer (Sigma) and resuspended at a concentration of 2 × 10⁴ cells per well into a 96-well plate. FSMCs were incubated with or without CCK8S in the presence of compound 48/80 (50 μ g ml⁻¹) for 15 minutes at 37 °C. The plate was centrifuged for 5 minutes, and the supernatants were placed into another 96-well plate. Cell pellets were resuspended in Tyrode's buffer containing 0.5% triton X-100 (Sigma). Supernatants and cell lysates (50 μ l) were incubated with 100 μ l of 2.5 mM *p*-nitrophenyl-*N*-acetyl- β -D-galactosaminide (Sigma),

dissolved in 0.04 mM citrate buffer (pH 4.5), at 37 °C for 90 minutes. The reaction was stopped with 100 µl of 0.4 M glycine (pH 10.7; MP Biomedicals, Solon, OH). The plate was read at 405 nm using a 595 nm reference filter in a microplate reader (Bio-Rad). β-Hexosaminidase release was determined as the ratio between activity in the supernatant and the cell lysate, multiplied by 100.

Calcium influx determined by confocal imaging

Confocal imaging of cells loaded with fluorescent dyes was performed using a Zeiss LSM5 pascal (Carl Zeiss, Jena, Germany). FSMCs were loaded with Fluo-4-AM (Invitrogen; 1 µM) at 37 °C for 20 minutes, washed twice, and set aside for at least 40 minutes. FSMCs were also stained for CD117 (c-kit) with Alexa Fluor 700-conjugated anti-mouse CD117 antibody. The LSM5 multitrack configuration was used for simultaneous measurement of intracellular calcium concentration (excitation, 488 nm; emission, long-pass 505 nm filter) and the expression of CD117 (excitation, 633 nm; emission, long-pass 650 nm filter). Images were recorded (usually recording for 300 seconds) at room temperature, in a 512 × 512-pixel format. As stimulants, SP or CCKs were added directly to the bath as a small drop (10 µl; final concentration, 10⁻⁶ M SP and 10⁻⁶ M CCK8S). Image data were analyzed offline using the Zeiss LSM510 analyzing software (LSM Image Examiner, Carl Zeiss). The mast cells exhibiting green fluorescence of Fluo-4-AM at any time point between 40 and 80 seconds after CCK8S addition were defined as positively responding cells.

Gene-array analysis

Normal human epidermal KCs were cultured with SP (10⁻⁶ M), histamine (10 mg ml⁻¹), and SLIGRL-NH₂ (100 nM) for 2 hours and harvested. For DNA microarray analysis, total RNA was extracted from Normal human epidermal KCs with the RNeasy Mini Kit (Qiagen, Valencia, CA). Images were analyzed with the DNASIS Array (DNA Chip Research, Hitachi Software Engineering, Tokyo, Japan) according to the manufacturer's instructions. The results of DNA microarray analysis of the top and bottom 20 genes with respect to expression levels induced by SP, histamine, or SLIGRL-NH₂ are shown in the Supplementary Data online.

Statistical analysis

Data were analyzed using an unpaired Student's *t*-test. *P* < 0.05 was considered to be significant.

CONFLICT OF INTEREST

The authors state no conflict of interest.

ACKNOWLEDGMENTS

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SUPPLEMENTARY MATERIAL

Supplementary material is linked to the online version of the paper at <http://www.nature.com/jid>

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SHORT REPORT

Increased circulating Th17 frequencies and serum IL-22 levels in patients with acute generalized exanthematous pustulosis

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Abstract

Background/Objective Acute generalized exanthematous pustulosis (AGEP) is a diffuse pustular disorder that usually begins in intertriginous folds with widespread erythema. The causes in the majority of the cases are drugs. T cells and interleukin (IL)-8 play roles in the development of AGEP, but the mechanism remains to be elucidated. We investigated the involvement of Th17 cells and their cytokine IL-22 in the pathogenesis.

Methods Three patients with AGEP were enrolled in this study. The percentages of IL-17⁺ Th17 cells, interferon γ ⁺ T cells and IL-4⁺ T cells were measured in the patients' peripheral blood lymphocytes by intracellular cytokine staining and flow cytometry. The concentration of IL-22 in the sera was measured by enzyme-linked immunosorbent assay.

Results The percentages of Th17 cells were markedly higher in all three patients than healthy control individuals. The frequencies of interferon γ ⁺ T cells were slightly high in the patients compared with the control, and there was no definite tendency in IL-4⁺ T-cell frequencies. The concentration of IL-22 was remarkably high in all patients when compared with normal subjects with levels under detection.

Conclusion Th17 cells and their produced cytokine IL-22 were elevated in the peripheral blood of patients with AGEP. As IL-17 and IL-22 cooperatively stimulate keratinocytes to produce IL-8, IL-8 may contribute to the accumulation of neutrophils in the lesional epidermis of AGEP.

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Keywords

acute generalized exanthematous pustulosis, drug eruption, IL-17, IL-22, Th17

Conflict of interest

None.

Funding sources

None.

Introduction

Acute generalized exanthematous pustulosis (AGEP) is a diffuse pustular disorder characterized by acute, small, non-follicular, sterile pustules that usually begin in intertriginous folds with widespread oedema and erythema.¹⁻⁵ Fever and peripheral blood leucocytosis are often seen in the patients. The majority of the cases are induced by adverse drug reactions, often triggered by pristinamycin, ampicillin/amoxicillin, quinolones, chloroquine, sulphonamides, terbinafine and diltiazem.¹ Massive subcorneal infiltration of neutrophils and dermal and epidermal infiltration of T cells are the histopathological features.

In AGEP, lymphocyte transformation test (LTT) usually shows high levels of stimulation index (S.I.) towards causative drugs, suggesting that drug-specific T cells mediate AGEP² as seen in the other types of drug eruptions.⁶ Drug-specific CD4⁺ and CD8⁺ T cells are present in the blood and play an important role by producing neutrophil chemoattractant interleukin-8 (IL-8)/CXCL8.³⁻⁵ However, the pathogenesis of AGEP cannot be explained merely with T-cell-secreting IL-8, because the mechanism of subcorneal collection of neutrophils remains to be an issue. One possibility is that IL-8-secreting T cells invade into the epidermis, leading to the infiltration of neutrophils. Alternatively, a certain population

of T cells specific for a given drug might stimulate keratinocytes to produce IL-8, and the keratinocyte-derived IL-8 might be responsible for subcorneal neutrophil aggregation. In fact, the elevated expression of IL-8 was observed in keratinocytes as well as infiltrating mononuclear cells.⁵

IL-17-producing Th17 cell is a newly established CD4⁺ T helper cell subset, and dysregulated Th17 responses mediate a variety of systemic and skin inflammatory conditions, such as psoriasis^{7,8} and atopic dermatitis.⁹ Th17 cells coexpress IL-22, and its receptor is expressed on epithelial tissues.⁷ IL-17 and IL-22 cooperatively enhance some immunological responses and exert a strong synergistic effect on the production of IL-8 by keratinocytes.⁹ Recently, an increased frequency of Th17 cells has been reported in a case of AGEF.¹⁰ To address the possible participation of Th17 cells in the pathogenesis of AGEF, we examined the frequency of Th17 cells and the level of IL-22 in three patients with AGEF.

Materials and methods

Three patients with AGEF, 32-year-old woman (case 1), 68-year-old woman (case 2) and 59-year-old man (case 3), were enrolled

in this study. The diagnosis was made on the basis of clinical manifestations, i.e. an erythematous, burning eruption with small pustules on the peripheries distributed to the inner thighs, knees, lower legs and upper limbs (Fig. 1a). In all patients, the total leucocyte counts and CRP values were elevated with neutrophilia. Histopathologically, there were subcorneal neutrophilic pustules and a dermal lymphocytic infiltrate. The causative drugs and LTT were as follows: case 1, acetaminophen (final concentration, 10 µg/L; S.I., 2.5); case 2, roxithromycin (Fig. 1b; final concentration, 5 µg/L; S.I., 2.1) and case 3, doripenem (final concentration, 1.8 µg/L; S.I., 1.7). Discontinuation of the causative drugs and oral or topical administration of corticosteroids improved the eruptions in 2 weeks. Roxithromycin as causative drug of AGEF is known,¹¹ and doripenem-induced AGEF in case 3 was reported previously.¹²

In the three patients, the percentages of circulating Th17 cells and serum levels of IL-22 were measured. The method for measurement of Th17 cells as well as Th1 and Th2 cells was described previously.⁹ Briefly, blood samples were taken from the patients 5 or 6 days after the onset of eruption. Intracellular

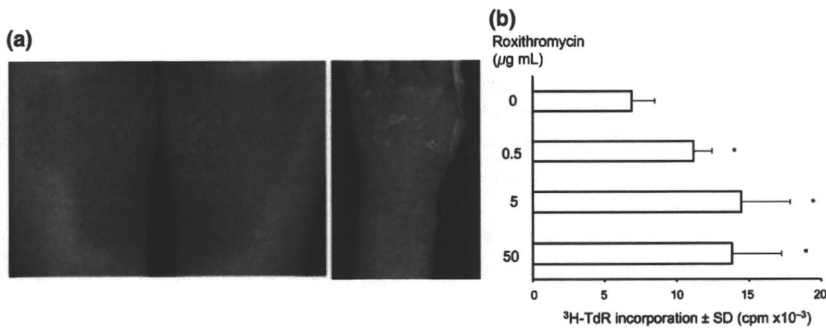


Figure 1 Representative clinical appearance and LTT to drug. (a) Clinical appearance of case 1, showing an erythematous, scaly, pustular eruption present on the thighs (left) and upper extremities (right). (b) LTT of case 2, showing an elevation of ³H-thymidine (TdR) incorporation in response to roxithromycin added to the 72-h culture of patient's PBMC. **P* < 0.05, compared with the non-addition control.

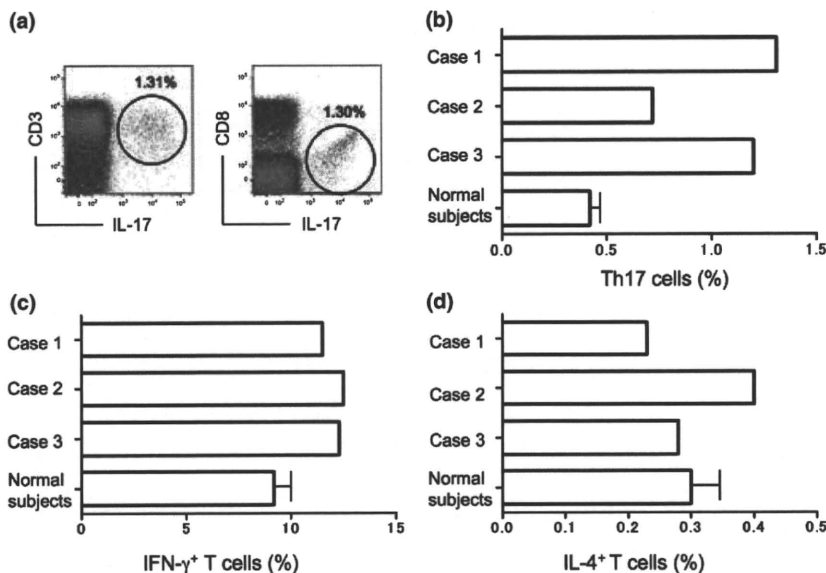


Figure 2 Th17 frequencies in PBMC of the three patients. (a) Representative flow cytometric data, showing that Th17 cells are IL-17⁺, CD3⁺ and CD8⁻. (b) Percentage of IL-17⁺CD3⁺CD8⁻ cells (Th17 cells). (c) Percentage of IFN-γ⁺CD3⁺ cells. (d) Percentage of IL-4⁺CD3⁺ cells.

cytokines of PBMC were stained according to the protocol of Cytostain (Immunotech, Marseille, France) with a few modifications. Freshly isolated peripheral blood mononuclear cells (PBMC; 2×10^6 cells/mL) were incubated in complete RPMI-1640, 10 ng/mL of phorbol 12-myristate 13-acetate (PMA; Sigma Chemical Co., St. Louis, MO, USA), 10^{-6} mol/L of ionomycin (Wako, Osaka, Japan) and 0.7 μ L of Golgistop (BD Biosciences) for 8 h. The cells were washed and directly stained with PerCP-conjugated anti-CD8 mAb (BD Biosciences, San Diego, CA, USA) and subsequently with APC-conjugated anti-CD3 mAb (BD Biosciences) for 20 min at 4 °C. After washing, 100 μ L of Cytofix/Cytoperm buffer (BD Biosciences) was added to each well and incubated for 20 min at room temperature and washed with Perm/Wash solution as manufacturer's protocol (BD Biosciences). They were stained with PE-labelled anti-IL-17 or IL-4 and FITC-labelled anti-IFN- γ monoclonal antibody. Fluorescence profiles were analysed by flow cytometry in FACSCanto.

The serum concentration of IL-22 was measured by using human IL-22 quantikine enzyme-linked immunosorbent assay (ELISA) kit (R & D Systems, Minneapolis, MN, USA).

Results

We detected circulating lymphocytes intracellularly positive for IL-17, IFN- γ or IL-4 by flow cytometry. As CD4 expression is downregulated by stimulation with PMA and ionomycin, we estimated CD3⁺CD8⁻ cells as CD4⁺ T cells.⁹ The IL-17⁺ cell population was found exclusively in CD3⁺CD8⁻ T cells (Fig. 2a), representing Th17 cells. The frequencies of Th17 cells in all three patients were elevated compared with the normal subjects (Fig. 2b). The percentage of IFN- γ ⁺ T cells was slightly higher in the three patients than the normal subjects ($n = 12$; five men and seven women; mean age, 48.8 years) (Fig. 2c), and both CD4⁺ and CD8⁺ fractions showed the similar IFN- γ ⁺ cell-elevated tendency (data not shown). IL-4⁺ T cells were exclusively CD4-positive, and there was no definite tendency of its percentage in the patients compared with the control (Fig. 2d).

In parallel, we quantified the serum concentration of IL-22. IL-22 was elevated in all patients, compared with the normal individuals ($n = 5$; two men and three women; mean age, 45.2 years) whose IL-22 levels were under detection (Fig. 3).

Discussion

It is well known that IL-8 is the most important chemokine for neutrophil infiltration. AGEP is a disease in which both neutrophils and T cells are involved,²⁻⁵ and IL-8-producing T cells have been thought to mediate AGEP.³⁻⁵ Our study confirmed the increase of Th17 cells found in PBMC of a patient with AGEP and the possible involvement of Th17 cells in the pathogenesis of AGEP.¹⁰ Furthermore, we demonstrated the elevation of their produced cytokine IL-22. As IL-17 and IL-22 cooperatively stimulate keratinocytes to produce IL-8,⁹ our finding suggests that keratino-

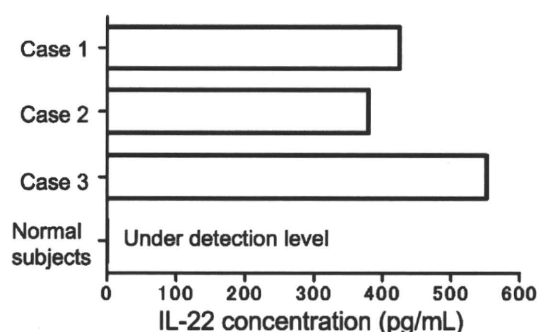


Figure 3 IL-22 levels in sera of the three patients. The concentration of IL-22 was measured with ELISA kit according to the manufacturer's directions.

cyte-derived IL-8 contributes to the accumulation of neutrophils in the lesional epidermis of AGEP. It is thus likely that not only T cells²⁻⁴ but also keratinocytes are the source of IL-8 in AGEP. The subcorneal infiltrate of neutrophils, a hallmark of AGEP, can be explained with this notion. Th17 cells are deeply involved in the pathogenesis of psoriasis,⁷ and the resultant production of IL-8 by keratinocytes may lead to Munro's microabscess and even the pustular form of psoriasis. In this concept, AGEP may share the pathomechanism with psoriasis. Th17 cells play a role for not only systemic but also for local cutaneous inflammation through the upregulation of neutrophil-chemoattracting and mobilizing chemokines in AGEP.

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CORRESPONDENCE

Increased expression of mRNAs for IL-4, IL-17, IL-22 and IL-31 in skin lesions of subacute and chronic forms of prurigo

Prurigo lesions are divided into acute, subacute, and chronic forms [1]. Subacute prurigo tends to affect middle-aged women with symmetrical distribution. Chronic prurigo consists of small, irritable papules, preferentially on the abdomen and other sites. These two forms persist for a long period and may evolve into nodular lesions. Prurigo is severely itchy and characterized histologically by infiltrates of lymphocytes and eosinophils and acanthosis of the epidermis. Th2 cells expressing interleukin (IL)-4, IL-5 and IL-10 infiltrate into the lesional skin of subacute and chronic prurigos [2]. However, all features of prurigo cannot be ascribed to Th2 cytokines, and other cytokines may participate in the pathogenesis. IL-31 is pathogenic for pruritic skin diseases, such as atopic dermatitis and prurigo nodularis [3]. On the other hand, IL-22 is a critical cytokine for proliferation of epidermal keratinocytes [4]. We therefore investigated the expression of these cytokines in the skin lesions of prurigo.

The ethical committee of our university approved this study. Papulonodular lesions of subacute and chronic prurigos (ca. 3 mm in diameter) were biopsied and subjected to real-time polymerase chain (PCR) analysis. Seven patients (M:F=4:3; mean age, 68.7 ± 10.9) were enrolled. As controls, skin specimens (center of lesional skin) of 7 patients with plaque psoriasis (M: F=4:3; $65.5 \text{ years} \pm 15.9$) and 5 normal individuals were used. Total RNA was extracted with an RNA extraction kit (Promega, Madison, WI). RNA was reverse transcribed and amplified by random hexamer with primers and probes from TaqMan (Applied Biosystems, Foster City, CA). The result for each gene was normalized to the quantity of β -actin (*ACTB*) detected in the sample. In parallel, we measured the percentage of IL-17⁺CD4⁺ T (Th17) cells

in peripheral blood mononuclear cells (PBMC) from the patients, as described previously [5]. Intracellular IL-17 of PBMC was stained according to the protocol of Cytostain (Immunotech, Marseille, France), with some modifications.

The level of *IL4* mRNA expression was higher in prurigo skin than in normal and psoriatic skin, and there was no significant difference between normal and psoriatic skin (figure 1A). *IL31* mRNA was expressed at a significantly higher level in prurigo than psoriasis and normal skin (figure 1B). mRNAs for IL-17 (figure 1C) and IL-22 (figure 1D) were highly expressed in both prurigo and psoriasis lesions, compared to normal skin. A significant correlation was found between the expression levels of *IL4* and *IL31* ($P=0.0429$), and a correlation tendency was observed between *IL17* and *IL22* ($P=0.1096$) in patients with prurigo. The percentages of peripheral blood Th17 cells (IL-17⁺CD3⁺CD4⁺CD8⁻ cells) were also examined. Figure 1E represents a typical flowcytometric analysis of Th17 cells. The frequencies of Th17 cells were significantly higher in prurigo as well as in psoriasis than in normal control (figure 1F).

In addition to the previous observation that Th2 cytokines are highly expressed in prurigo lesions [2], the present study showed that *IL31*, *IL17*, and *IL22* are also increased in prurigo. IL-31 is produced by a certain subset of Th2 cells and is involved in pruritic disorders [3], and *IL31* and *IL4* were correlated with each other in our prurigo study, as seen in atopic dermatitis [6]. Prurigo shared the high expression levels of IL-17 and IL-22 with psoriasis. Epidermal keratinocytes proliferate in response to IL-22 [4], leading to acanthosis. Accordingly, we found that *IL17* and *IL22* were elevated in prurigo lesions. Prurigo may be characterized by the increment of IL-17/IL-22 and IL-31. Since these two categories of cytokines seem to be derived from different T cell populations, Th17/Th22 and Th2, it is considered that the characteristics of prurigo, i.e. pruritus and epidermal hyperplasia, are mediated by the unique combination of pathogenic T cells. ■

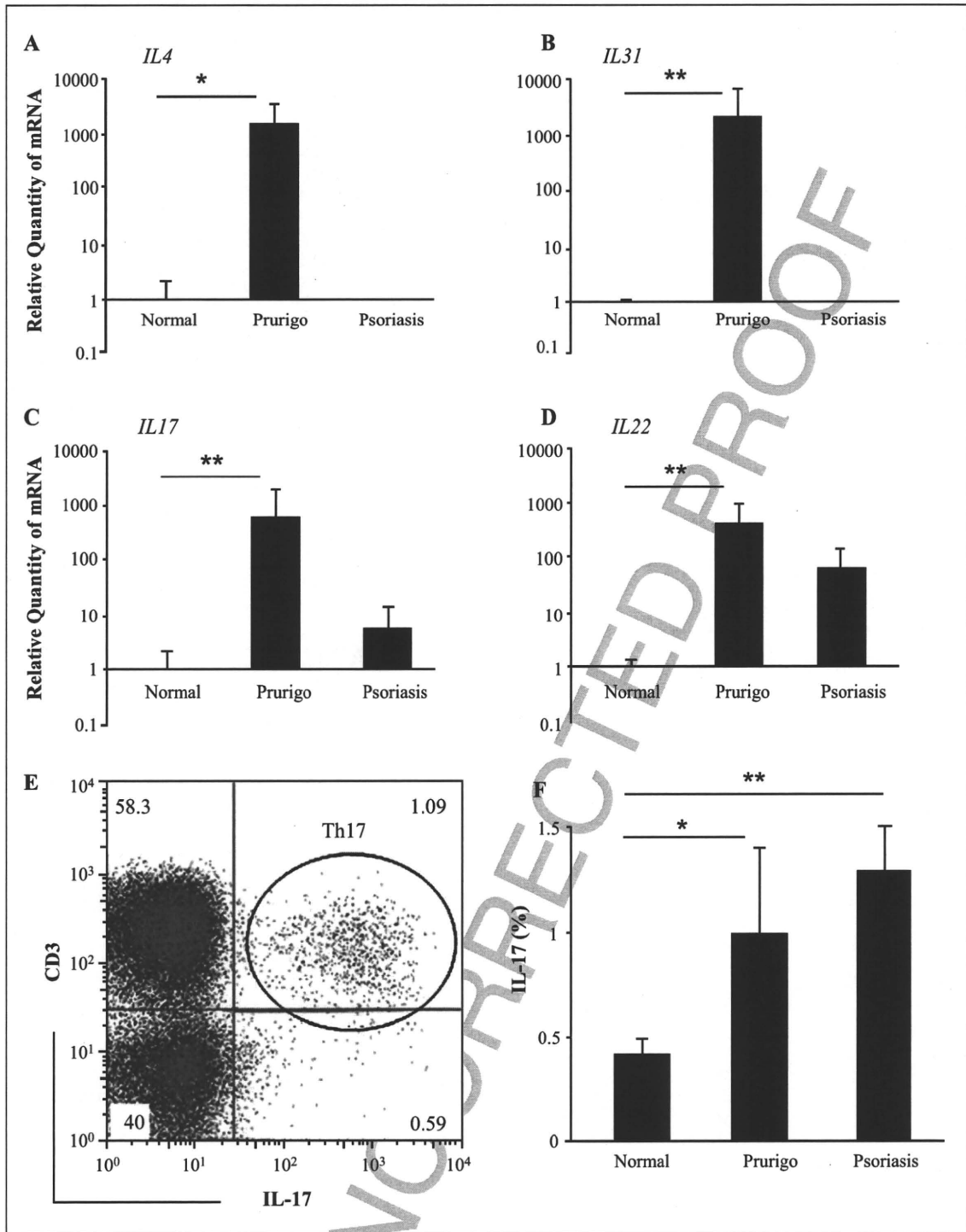


Figure 1. Expression of cytokines in the skin and percentages of Th17 cells in the blood. **A, B, C, D**) Biopsy specimens were taken from lesional skin of patients with prurigo and psoriasis, and from normal skin of healthy subjects. They were examined for the expression of the indicated four cytokines by real-time RT-PCR. The data represent the means \pm SD. UDL: under detection level. In IL-4 of psoriasis, 6 cases were UDL and one case was 5.15. In IL-31 of psoriasis, 5 cases were UDL and 2 cases were 2.77 and 3.07. * $P=0.024$, ** $P < 0.01$. **E, F**) PBMC from the patients and normal volunteers were subjected to intracellular cytokine staining and flow cytometry. A representative flow cytometric data is shown (**E**). The mean \pm SD of Th17 cell percentages in each group (**F**). * $P=0.050$, ** $P=0.0019$.

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UNCORRECTED PROOF

Regulation of Inducible Nitric-oxide Synthase by the SPRY Domain- and SOCS Box-containing Proteins*[§]

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Inducible nitric-oxide synthase (iNOS, NOS2) plays a prominent role in macrophage bactericidal and tumoricidal activities. A relatively large amount of NO produced via iNOS, however, also targets the macrophage itself for apoptotic cell death. To uncover the intrinsic mechanisms of iNOS regulation, we have characterized the SPRY domain- and SOCS box-containing protein 1 (SPSB1), SPSB2, and SPSB4 that interact with the N-terminal region of iNOS in a D-I-N-N-N sequence-dependent manner. Fluorescence microscopy revealed that these SPSB proteins can induce the subcellular redistribution of iNOS from dense regions to diffused expression in a SOCS box-dependent manner. In immunoprecipitation studies, both Elongin C and Cullin-5, components of the multi-subunit E3 ubiquitin ligase, were found to bind to iNOS via SPSB1, SPSB2, or SPSB4. Consistently, iNOS was polyubiquitinated and degraded in a proteasome-dependent manner when SPSB1, SPSB2, or SPSB4 was expressed. SPSB1 and SPSB4 had a greater effect on iNOS regulation than SPSB2. The iNOS N-terminal fragment (residues 1–124 of human iNOS) could disrupt iNOS-SPSB interactions and inhibit iNOS degradation. In lipopolysaccharide-treated macrophages, this fragment attenuated iNOS ubiquitination and substantially prolonged iNOS lifetime, resulting in a corresponding increase in NO production and enhanced NO-dependent cell death. These results not only demonstrate the mechanism of SPSB-mediated iNOS degradation and the relative contributions of different SPSB proteins to iNOS regulation, but also show that iNOS levels are sophisticatedly regulated by SPSB proteins in activated macrophages to prevent overproduction of NO that could trigger detrimental effects, such as cytotoxicity.

Ubiquitination and the subsequent proteasomal degradation of regulatory proteins represent a key mechanism for various cellular processes, including cell cycle control (1), apoptosis (2), cellular stress regulation (3), and antigen presentation (4). The ubiquitination of proteins is performed by a set of three enzymes: E1, E2, and E3 (5). E3 is a ubiquitin ligase that confers specificity to ubiquitination by recognizing target substrates and mediating the transfer of ubiquitin from E2 to the substrate. The E3 ubiquitin ligases are a large, diverse group of proteins, and many of them can act alone, but some RING-type E3 ubiquitin ligases, such as the ECS³ (Elongin B/C-Cullin-5 (Cul5)-SOCS (suppressor of cytokine signaling)-box protein)-type E3 ubiquitin ligase, are found as components of much larger multi-protein complexes (6). The SOCS box is ~40 residues in length and contains a BC box and a Cul5 box, which binds Elongin C and Cul5, respectively (7). The SOCS box has been found in more than 50 proteins encompassing nine different families that are defined by the type of domain or motif they possess upstream of the SOCS box (6, 8). The SPRY domain- and SOCS box-containing proteins, SPSB1 to SPSB4 (also known as SSB-1 to SSB-4), are characterized by a central SPRY domain essential for protein interaction and a C-terminal SOCS box, suggesting that SPSB proteins may function as part of the ECS E3 ubiquitin ligase complex with the SPRY domain determining the substrate for ubiquitination (9). It has been shown that the *Drosophila* SPSB, called "GUSTAVUS," interacts with the DEAD-box RNA helicase VASA (10), and human SPSB1, SPSB2, and SPSB4 interact with human prostate apoptosis response protein-4 (hPar-4) (11). The identification of similar sequences in VASA (D-I-N-N-N) and hPar-4 (E-I-N-N-N) suggests that the SPRY domains of SPSB1, SPSB2, and SPSB4 likely recognize a common peptide epitope. In contrast to the extensive analysis of SPRY domain structures, the biochemical and physiological functions of the SPSB family of proteins remain largely unknown.

Nitric oxide (NO) is an important multifunctional biomolecule involved in various physiological and pathological processes (12, 13). NO is endogenously synthesized from L-arginine

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³ The abbreviations used are: ECS, Elongin B/C-Cullin-5-SOCS box protein; SOCS, suppressor of cytokine signaling; SPSB, SPRY domain- and SOCS box-containing protein; NOS, nitric-oxide synthase; iNOS, inducible NOS; YFP, yellow fluorescent proteins; CFP, cyan fluorescent protein; 7-AAD, 7-aminoactinomycin D; CHX, cycloheximide; DOX, doxycycline.