Epithelial Hyperproliferation and Transglutaminase 1 Gene Expression in Stevens-Johnson Syndrome Conjunctiva

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In Stevens-Johnson syndrome, pathological keratinization of the ordinarily nonkeratinized corneal and conjunctival mucosal epithelia results in severe visual loss. We examined conjunctiva covering cornea in five eyes in the chronic cicatricial phase of Stevens-Johnson syndrome. Normal conjunctiva from five age-matched individuals was studied also. The number of epithelial cells in Stevens-Johnson syndrome conjunctiva that were immunoreactive with a monoclonal antibody, Ki-67, to a nuclear antigen found only in proliferating cells was greater than normal (93.8/H11550 19.8 cells above 100 basal cells versus 12.8/H11550 0.5 cells above 100 basal cells; P = 0.009). In addition, although clinical inflammation was mild, massive lymphocytic infiltration was seen in the substantia propria of conjunctiva covering cornea. In situ hybridization documented transglutaminase 1 (keratinocyte transglutaminase) mRNA in suprabasal cells of the abnormally thickened conjunctival epithelium in all Stevens-Johnson syndrome patients. In contrast, no message was detected in normal conjunctival or corneal epithelia. Transglutaminase 1 is expressed during the terminal differentiation of keratinocytes where it helps synthesize cornified cell envelopes. We speculate that in Stevens-Johnson syndrome, epithelial hyperproliferation, and transglutaminase 1 gene expression lead to the pathological keratinization of ocular surface mucosal epithelia. (Am J Pathol 1999, 154:331–336)

Stevens-Johnson syndrome (SJS), erythema multiforme major, is an acute inflammatory disease that affects mucosal membranes and skin.1–5 It is a self-limited disease, and after the acute phase has passed, skin and most mucosa recover without significant scarring. However, long-term ocular consequences are devastating. During the chronic phase of the disease, for example, most SJS patients experience numerous ocular surface problems, including symblepharon, entropion, ectropion, trichiasis, dry eye, persistent conjunctival inflammation, corneal vascularization (conjunctivalization), and keratinization. Some of these problems can be managed by the use of antibiotics, corticosteroids, and/or artificial tears, however, the pathological keratinization of the ordinarily non-keratinized corneal and conjunctival mucosal epithelia is a serious and potentially debilitating problem that is difficult to manage pharmacologically.

The general term given to the pathological transition of a nonkeratinized, stratified epithelium into a keratinized epithelium is squamous metaplasia,6,7 and in the human eye, it is a process that is accompanied by the loss of conjunctival goblet cells, an increase in epithelial stratification, and an enlargement of the superficial epithelial cells.8,9 Squamous metaplasia has been described in numerous disorders of the ocular surface,6,7 including dry-eye disorders in which the aqueous layer of the tear film is deficient, eg, Sjögren syndrome,10 as well as disorders such as SJS and ocular cicatricial pemphigoid (OCP) in which the mucous layer is deficient.8 Despite the effort that has gone into understanding squamous metaplasia, the pathogenesis behind the abnormal differentiation of cells remains unknown.

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In this study, we first wanted to discover whether or not the abnormally thick conjunctiva in SJS might be a result of increased cellular proliferation, as is thought to be the case in squamous metaplasia associated with OCP.11 To investigate this, we initiated an immunohistochemical study using the murine monoclonal antibody Ki-67, an antibody that reacts with a human nuclear antigen that is present in proliferating cells but absent from quiescent cells.12 Previous work has shown that the Ki-67 nuclear antigen is expressed in the G1, S, G2, and M phases but not in the G0 phase.13 Furthermore, when compared with flow cytometry, 3H-thymidine labeling and BrdU labeling, Ki-67 immunohistochemistry allows a readier evaluation of the growth fraction of a given human cell population.14 The second feature of SJS studied was prompted by the observation that in severe cases, diseased conjunctival epithelium covering the cornea often resembles the keratinized epidermis of skin (Figure 1A). This led us to wonder whether transglutaminase 1 (keratinocyte transglutaminase; TGase1) might be involved in the pathological keratinization of ocular surface mucosal epithelia in SJS. TGase1 is an enzyme expressed during the terminal differentiation of keratinocytes to form the highly insoluble, cross-linked cell envelope at the periphery of cornified cells.15 We used in situ hybridization to investigate the expression of the TGase1 gene in normal and SJS conjunctiva.

Materials and Methods

Tissue Samples

With informed consent, we obtained conjunctiva-covering cornea from five patients with SJS (Table 1) at the time of a lamellar keratoplasty (surgical removal of the outer layer of the cornea) to improve vision. All eyes were in the chronic cicatricial phase, and the corneal surfaces were totally covered by conjunctival tissue (Figure 1A). In general, the ocular surfaces were relatively quiet, though some mild episodic inflammation was occasionally seen in four of the five patients. Impression cytology found very few goblet cells in the bulbar inferior conjunctiva of the SJS eyes (Figure 1B), a characteristic of squamous metaplasia. Again with proper informed consent, normal tissue was obtained during ocular surgery from the bulbar inferior conjunctiva from five age-matched individuals without any history of ocular surface disease. Impression cytology confirmed the presence of numerous mucin-producing goblet cells in this tissue (Figure 1C), a normal feature of conjunctiva.

Immunohistochemistry for Ki-67 Nuclear Antigen

Samples of conjunctiva were snap frozen, and 7-μm thick sections were cut. To minimize endogenous peroxidase activity, the tissue sections were incubated with 1% H2O2 for 1 hour, and to block nonspecific binding, they were incubated with 10% goat serum at room temperature for 1 hour. Subsequently, the sections were incubated with a biotinylated antiserum directed against mouse IgG (Vector Laboratories, Burlingame, CA; working dilution

Table 1. Patient Profile

<table>
<thead>
<tr>
<th>Patient number</th>
<th>Age/gender</th>
<th>Episodic inflammation</th>
<th>Visual acuity</th>
<th>Age (yrs.) at onset</th>
<th>Time (yrs.) since acute episode</th>
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</thead>
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<tr>
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<td>57/M</td>
<td>Mild</td>
<td>HM</td>
<td>55</td>
<td>2</td>
</tr>
<tr>
<td>2</td>
<td>57/M</td>
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<td>4/200</td>
<td>47</td>
<td>10</td>
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<tr>
<td>3</td>
<td>24/M</td>
<td>Mild</td>
<td>20/40</td>
<td>13</td>
<td>11</td>
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<td>Mild</td>
<td>2/200</td>
<td>44</td>
<td>30</td>
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<tr>
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<td>27/M</td>
<td>Mild</td>
<td>HM</td>
<td>7</td>
<td>20</td>
</tr>
</tbody>
</table>

HM, hand motion.
(1:200) for 2 hours at room temperature, and then with an avidin-biotin-peroxidase complex (Vector Laboratory; 1:100) for 50 minutes at room temperature. Sections were then exposed to 200 μg/ml of 3,3′-diaminobenzidine in 0.05 mol/L Tris-HCl buffer (pH 7.6) containing 50 μg/ml H2O2 (10 minutes at room temperature). Between each of these steps, sections were rinsed thoroughly with 0.1 mol/L phosphate-buffered saline. As has been done previously,11 to compare the number of proliferating cells in SJS conjunctival epithelium with the number in normal conjunctival epithelium, we counted the Ki-67-labeled cells in all epithelial layers overlying 100 basal epithelial cells. Cells in three sections of each conjunctiva were counted independently by two investigators and the data averaged.

**TGase1 in Situ Hybridization**

Paraffin-embedded tissues were sectioned into 4-μm thick sections for in situ hybridization. The RNA probes were synthesized according to our previous reports.16,17 Briefly, a 2-kb KpnI fragment of human TGase1 cDNA was inserted into plasmid pGEM4Z to construct pdM-K2. After linearizing pdM-K2 with EcoRI and BamHI, the antisense and sense cRNAs were transcribed in vitro using digoxigenin (DIG)-labeled UTP by T7 and SP6 RNA polymerase, respectively, according to the manufacturer’s manual (DIG RNA Labeling Kit SP6/T7; Boehringer Mannheim, Mannheim, Germany). The RNA probes were fragmented by limited alkaline hydrolysis. In situ hybridization was performed according to our previous protocol.17 First, the sections were deparaffinized in xylene and rehydrated through a graded ethanol series. After proteinase K digestion (18 μg/ml), the sections were postfixed with 4% (w/v) paraformaldehyde in phosphate-buffered saline for 10 minutes and treated with 0.1 mol/L triethanolamine-HCl (pH 8.0) for 1 minute. Following acetylation for 10 minutes, the sections were dehydrated, air-dried then incubated overnight at 45°C in hybridization buffer composed of 50% formamide, 10 mmol/L Tris-HCl (pH 7.5), 1 mg/ml yeast tRNA (Sigma), 1× Denhardt’s solution (Sigma), 10% PEG6000, 600 mmol/L NaCl, 0.25% sodium dodecyl sulfate, 1 mmol/L EDTA, and 0.2 μg/ml probe. After hybridization, the sections were washed at 45°C for 1 hour in 50% formamide and 2× SSC, then digested with 20 μg/ml RNase (Sigma) in 10 mmol/L Tris-HCl (pH 8.0), and 500 mmol/L NaCl at 37°C for 10 minutes. Hybridized DIG-labeled probes were visualized with a Nucleic Acid Detection Kit (Boehringer Mannheim).

**Results**

Immunohistochemistry documented numerous Ki-67-positive cells in SJS conjunctiva, all of which were located in the basal epithelial cell layer or the one or two layers of cells immediately above this (Figure 2A). Occasional localized groups of Ki-67-positive cells seen more superfi-
cially (Figure 2A) were due to in-folds of the basement membrane. Normal conjunctival epithelium contained fewer Ki-67-positive cells than SJS conjunctiva (Figure 2B). This difference was statistically significant (Mann-Whitney U-test; \( P = 0.009 \)), SJS conjunctiva containing 93.8 ± 19.8 (mean ± SE) Ki-67-positive cells above 100 basal epithelial cells compared with 12.8 ± 0.5 in normal conjunctiva (Figure 2C). The number of proliferating cells in normal conjunctiva counted in our experiment (12.8 ± 0.5 Ki-67-labeled cells above 100 basal cells) is larger than the number of proliferating cells (1.6 ± 0.2 tritiated thymidine-labeled cells per 100 basal cells) reported by Thoft and associates.\(^{11}\) Presumably, this is because the Ki-67 antibody labels proliferating cells in more stages of the cell cycle (G1, S, G2, and M phases)\(^{13}\) than does tritiated thymidine (S phase only).\(^{11}\)

Although clinically observable conjunctival inflammation in our SJS patients was either absent or mild, massive lymphocytic infiltration was seen in the substantia propria of the invading conjunctiva in all cases (Figure 3A). This was not the case in normal tissue in which only few lymphocytes were found subepithelially in conjunctiva (Figure 3B). It is worth noting that the lymphocyte infiltration in SJS was often especially prominent around vessels (Figure 3A).

Importantly, in situ hybridization clearly demonstrated the presence of TGase1 mRNA in SJS conjunctival epithelium (Figure 3C). This message was invariably located in a band of epithelial cells located either in or above the suprabasal region. TGase1 mRNA was detected in the conjunctiva of all five SJS patients studied, however, the intensity of the message varied; it was strong in patients 1, 2, and 4, but weaker in patients 3 and 5. In contrast to SJS, the TGase1 gene was not expressed in any of the normal, nonkeratinized conjunctival (Figure 3D) or corneal (not shown) epithelia examined here. It was noteworthy that in SJS conjunctiva the TGase1 signal was often particularly strong in the suprabasal epithelium near high concentrations of subepithelial inflammatory cells. This relationship was especially clear in patient 1.

**Discussion**

We have shown that, in SJS, conjunctiva that covers cornea contains significantly more Ki-67 immunopositive cells than normal (Figure 2). Our results indicate that these Ki-67 positive cells exist mainly in the basal and suprabasal regions of the conjunctival epithelium. As the Ki-67 monoclonal antibody detects a nuclear cell antigen that is present only in proliferating cells,\(^{12-14}\) we conclude that conjunctival epithelial cell hyperproliferation is a feature of SJS. Previous investigations have shown that conjunctival hyperproliferation is also a feature of OCP, as indicated by the increased mitotic rate found by tritiated thymidine autoradiography.\(^{11}\) Goblet cell frequency is also less in OCP conjunctiva.\(^{11}\) This inverse relationship between mitotic rate and epithelial goblet cell content has been demonstrated in animal experiments also,\(^{18}\) leading to the hypothesis that hypermitosis interferes with the normal differentiation of conjunctival epithelial cells in conditions like OCP, and results in ep-
thelial abnormalities such as the lack of goblet cells. Similarly in SJS conjunctiva, the goblet cell deficiency (Figure 1B) might be a consequence of epithelial hyperproliferation (Figure 2A).

Pathological keratinization of ocular surface mucosal epithelia, the end stage of squamous metaplasia, is often observed in cicatricial ocular surface diseases such as SJS and OCP. Where it occurs it represents a serious clinical problem because it is invariably associated with a worsening of vision and an instability of the tear film. At present, the disease mechanism behind this keratinization is not understood. The present results, however, suggest that, in SJS at least, a link between TGase1 gene expression and conjunctival epithelial cell keratinization might exist. TGase1, keratinocyte transglutaminase, is an enzyme coded for by a gene sublocalized to chromosome 14q11.2 that catalyzes \( \epsilon-(\gamma\text{-glutamyl}) \text{ lysine} \) cross-links of proteins to form the cell envelope at the periphery of cornified cells. It is present in the epidermis of skin, and, along with TGase3, is involved with epidermal keratinization. Specifically, TGase3 catalyzes small oligomer formation of loricrin, a component of the cornified cell envelope, by intrachain cross-linking, whereas TGase1 forms very large oligomeric loricin complexes by interchain cross-linking. Indeed, experiments with knockout mice have revealed that TGase1 is essential for the distribution of the cell envelope precursor protein at the cell periphery and that the function of TGase1 cannot be compensated for by TGase3 nor other TGase isoforms. In view of this, the present study used TGase1 as a representative marker of keratinization. We did not detect TGase1 gene expression in the mucous epithelia of normal conjunctiva (Figure 3D) and cornea (not shown). In contrast, TGase1 mRNA was clearly present in suprabasal cells in the conjunctival epithelium in SJS. The obvious implication is that the pathological keratinization of the conjunctival epithelium in SJS might be caused by the expression of TGase1, an enzyme whose mRNA is not ordinarily present in mucosal ocular surface epithelia.

As stated, this study suggests that conjunctival epithelial hyperplasia and keratinization in SJS are based on increased proliferation coupled with TGase1 gene expression. What might cause the hyperproliferation and unusual expression of the TGase1 gene? After considering possible disease mechanisms, we suspect that inflammatory processes might be involved. Previously, other investigators have speculated that conjunctival inflammation might influence goblet cell loss in diseases like SJS. We now speculate that epithelial keratinization in SJS might also be related to inflammatory activity. This is based on the notion that epithelial keratinization in SJS is a consequence of TGase1 gene expression, along with the hypothesis that the expression of the TGase1 gene might be induced by cytokines released by infiltrating cells. In support of this we point out that TGase1 mRNA expression can be induced by a number of factors, including Ca\(^{2+}\), ganglioside GQ1b, 12-O-tetradecanoyl-phorbol-13-acetate, and interferon-\( \gamma \) (IFN-\( \gamma \)). We feel that of these inducers of the TGase1 gene, IFN-\( \gamma \) is the most likely to be involved with the pathogenesis of SJS, especially in light of our previous immunohistochemical work (unpublished data) that has indicated that the subepithelial infiltrating cells in SJS conjunctiva are immunoreactive with antibodies to IFN-\( \gamma \).

Our proposed pathogenesis for SJS, ie, conjunctival epithelial hyperproliferation accompanied by subepithelial cellular infiltration and the release of IFN-\( \gamma \) leading to TGase1 gene expression and subsequent keratinization, is similar in some respects to the disease mechanism behind the keratinizing skin disease, psoriasis. For example, epidermal hyperproliferation and incomplete epidermal differentiation are hallmarks of psoriasis. We also know that, in psoriasis, TGase1 gene expression is altered, and the activity of the enzyme in the psoriatic epidermis is increased. Moreover, Th1 type CD4-lymphocytes and numerous inflammatory cytokines are present in the lesional skin in psoriasis. Our previous work has indicated that the subepithelial infiltrating cells in SJS conjunctiva are also Th1-type CD4-lymphocytes. The cytokines found in psoriatic epidermis include IFN-\( \gamma \), as stated earlier, a potent inducer of squamous differentiation and TGase1 expression that we have found previously in SJS conjunctiva. It is worth pointing out that if inflammatory events do indeed cause hyperproliferation and TGase1 gene expression in SJS conjunctiva, it is likely that, as is the case in psoriasis, several cytokines, not just IFN-\( \gamma \), would be involved.

We propose that epithelial hyperproliferation and TGase1 gene expression underlie conjunctival hyperplasia and pathological keratinization in SJS, and possibly other forms of dry eye as well. This identifies the downregulation of TGase1 gene expression as a potential treatment. Furthermore, if, as we suspect, inflammatory cytokines such as IFN-\( \gamma \) are found to induce TGase1 gene expression, anti-inflammatory medication might prove effective in controlling SJS even in its chronic cicatrical phase.

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