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In Vitro and In Situ Expression of IL-23 by Keratinocytes in Healthy Skin and Psoriasis Lesions: Enhanced Expression in Psoriatic Skin

Gamze Piskin, Regien M. R. Sylva-Steenland, Jan D. Bos, and Marcel B. M. Teunissen

Keratinocytes contribute to cutaneous immune responses through the expression of cytokines. We investigated whether human keratinocytes can express IL-23, a newly defined IFN-γ-inducing cytokine composed of a unique p19 subunit and a p40 subunit shared with IL-12. Cultured keratinocytes from normal and lesional psoriatic skin were found to express constitutively mRNA for both subunits of IL-23. Low but significant levels of the heterodimeric IL-23 protein could be detected in cell lysates and supernatants from stimulated keratinocytes by immunoblotting and ELISA. Functional analysis showed that these low levels of keratinocyte-derived IL-23 were sufficient to enhance the IFN-γ production by memory T cells. Immunostaining of skin sections confirmed expression of both subunits of IL-23 by keratinocytes in situ and also revealed expression of this cytokine in the dermal compartment. IL-23 expression was significantly higher in psoriatic lesional skin, compared with normal and psoriatic nonlesional skin. The immunostained preparations of cultured cells and IL-23 levels in culture supernatants did not show any difference between normal and psoriatic keratinocytes indicating no intrinsic aberration of IL-23 expression in keratinocytes from psoriatic skin. Double staining of cytospin preparations demonstrated that IL-23 p19 is also expressed by epidermal Langerhans cells, dermal dendritic cells, and macrophages. Psoriasis is a chronic inflammatory skin disease mediated by IFN-γ-expressing type 1 memory T cells. As IL-23 is important to activate memory T cells to produce IFN-γ, its augmented expression of IL-23 by keratinocytes and cutaneous APC may contribute to the perpetuation of the inflammation process in this disease. The Journal of Immunology, 2006, 176: 1908–1915.

The skin is the main organ connecting the body to the potentially harmful environment. To achieve homeostasis and to defend the body against microbial invaders, it serves not only as a physical barrier, but is also equipped with numerous immunological functions collectively called the skin immune system (1). In addition to the well-known immunocompetent cells, which traffic into and out of the skin (e.g., dendritic cells (DC) and T cells), keratinocytes, as the main constituent of the epidermis, also have an important contribution to the development of optimal cutaneous immune responses. Keratinocytes can, constitutively or after stimulation, produce many cytokines as reviewed elsewhere (2). Physical, chemical, and pathogenic triggers induce keratinocytes to secrete proinflammatory cytokines, such as IL-1β and TNF-α, resulting in autoactivation of these cells to produce other inflammatory cytokines and chemokines. The proinflammatory cytokines may also activate other cells in the skin, such as DC, which start to mature and migrate to lymph nodes inducing adaptive immunity (3).

Expression of type 1 cytokines is crucial for defense against intracellular microorganisms and for development of delayed-type hypersensitivity reactions. Type 1 T cells produce IFN-γ, the principle mediator of subsequent type 1-associated immune responses. Expression of IFN-γ is strictly regulated by other cytokines (4): IL-12 was initially identified as an inducer of IFN-γ expression (5); subsequently, several other cytokines were also found to contribute to this regulation, namely IL-15, IL-18, IL-21, IL-23, and IL-27, which act separately or in combination to induce type 1 cytokine responses (4, 6). Within this group of IFN-γ-inducing cytokines, IL-23 and IL-27 exhibit structural and functional similarities to IL-12. However, they differ in the timing of their contribution to the type 1 responses; IL-27 is known to be important in the early expansion of naïve T cells and IFN-γ expression by these cells, whereas IL-23 only acts on memory T cells to induce IFN-γ expression, suggesting that IL-23 might be important in the maintenance of immune responses (4). The IL-23 heterodimer is formed by the combination of the p19 and p40 subunits and possesses the most structural similarity to IL-12, compared with other IFN-γ-inducers. The p40 subunit of IL-23 is shared with IL-12, whereas the unique IL-23 p19 molecule resembles to the IL-12-specific p35 subunit. In addition to their structural similarity, IL-12 and IL-23 both act through specific receptors that share a common IL-12Rβ1 chain (7). Human keratinocytes produce both subunits of the IL-12 heterodimer, allowing these cells to promote type 1 T cell responses (8); however, it is as yet not known whether keratinocytes also express the IL-23 p19 subunit.

It has been suggested that the expression of IL-12 might play an important role in certain inflammatory skin diseases (9, 10). Psoriasis vulgaris is a chronic inflammatory type 1 cytokine-related cutaneous disease, which is characterized by infiltrates of activated memory T cells that have a high IFN-γ expression (11). An enhanced expression of IL-12 mRNA and protein was found in psoriatic lesional skin, compared with nonlesional and normal skin, and a causative relationship to the high IFN-γ expression in lesional skin was suggested (12). Considering that IL-23, not IL-12,
preferentially activates memory T cells (7) and the fact that acti-
vated proliferating memory T cells are abundantly present in psoriasis lesions (13), we proposed that IL-23 rather than IL-12 would play a major role in the sustained inflammatory reaction taking place in psoriasis plaques. This consideration prompted us to in-
vestigate the expression of IL-23 p19 in normal and lesional psoriatic skin. Special attention was paid to keratinocytes because mutual interaction and activation of keratinocytes and T cells is thought to be responsible for the perpetuation of the inflammatory process in the psoriatic lesions (14).

In this study, we show that human keratinocytes constitutively express mRNA for the two subunits of IL-23 and demonstrate that these cells are able to secrete the IL-23 heterodimer, which turned out to enhance the IFN-γ expression by memory T cells. Specific staining for IL-23 in cryostat sections confirmed the expression of this cytokine by keratinocytes in situ, and furthermore, IL-23 ex-
pression appeared to be significantly stronger in psoriatic skin compared with normal skin. Double-staining experiments indi-
cated that Langerhans cells, dermal DC, and macrophages can also express IL-23.

### Materials and Methods

#### Skin biopsy and isolation of keratinocytes

Skin biopsy specimens (5 mm) were obtained from patients with chronic plaque-type psoriasis (16 lesional and 10 nonlesional biopsy samples) and from matched skin of normal individuals (n = 13) who had an operation for abdominal or breast reduction. The study was approved by the Medical Ethical Committee of the Academic Medical Center (Amsterdam, The Netherlands). These specimens were either snap frozen for in situ stainings or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5-or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions. The frozen specimens were cut into 5- or prepared for in vitro experiments under sterile conditions.

#### Stimulation of cells

For the determination of the IL-23 heterodimer by ELISA, immature monocyte-derived DC (MoDC) generated as described elsewhere (16) were stimulated at 5 × 10^6 cells/well in a flat-bottom 96-well plate (final volume, 200 μl) with either 10 μg/ml peptide-polylysycan (Sigma-Aldrich), or 10 μg/ml synthetic dsRNA poly(E,C), or an equal number of CD40L-transfected J558 (J558CD40L) plasmacytoma cells (a gift of Dr. P. Lane, University of Birmingham, Birmingham, U.K.), or 10 ng/ml IL-1β (PBH), or 10 μg/ml IFN-γ (Roche Diagnostics), or different combinations of these stimulants for 24 h. Supernatants and cell lysates of keratinocytes (passage 3) were obtained from subconfluent cultures maintained in petri dishes and stimulated overnight with a mixture of all aforementioned stimulants to achieve a maximal response. Keratinocyte cultures of passage 3 become usually subconfluent in 3–5 days and the purity of these cell cultures was tested by RT-PCR, which showed no expression of HLA-DR (macrophages, DC, B cells), CD3 (T cells), tyrosinase (melanocytes), and para-
thyroid hormone-rat (fibroblasts) (data not shown).

#### PCR analysis

Total RNA was purified from cultured subliminated keratinocytes by using the Nucleospin RNA II kit (Machery-Nagel) following the manufacturer’s instructions. First strand cDNA was generated with help of a synthesis kit for RT-PCR (MBI Fermentas), using 9 μl of total RNA, 1 μl of oligo(dT)18, and 1 μl of D(N)6, and heating the mix at 94°C for 5 min. The primers used were: IL-23 p19 (forward) 5’-TCG GCA AGA CAA CAC CTT AGG-3’, (reverse) GCC ACC ACA TCA TTT GTA GTC-3’, IL-12 p35 (forward) 5’-AAG AGA CCA GAG TCG CCG-3’, (reverse) GGA GCA TGT TGC TGA CCG C-3’ defining a 311-bp prod-
uct; IL-12/IL-23 p40 (forward) 5’-ATT GAG GTC ATG GTG GAT GC-3’, (reverse) 5’-ATT GCT GGC ATT TTT GGC GC-3’ defining a 297-bp product; and β2-microglobulin (forward) 5’-AAG ATT CAG GGT TAC TCA CGT C-3’, (reverse) 5’-TGA TGC TGT TTA CAT GTC TGC-3’. The PCR protocol was as follows: start with a 3 min incubation at 94°C, followed by 45 cycles of sequential incubations at 94°C for 30 s, 60°C for 30 s, and 72°C for 1 min. Analysis of the PCR products was done on a 1% agarose gel containing ethidium bromide. A 100-bp DNA ladder standard (MBI Fermentas) was used as a size marker.

#### Western blot analysis

The supernatant from stimulated keratinocytes was concentrated 20-fold by means of centrifugation (Millipore), whereas adherent keratinocytes (1–2 × 10^6) were washed once with PBS before the cells were lysed in 150 μl of PBS by freeze/thaw. The 25 μl of the supernatant and cell lysate was loaded onto a 12% polyacrylamide gel (Bio-Rad) and separated by elec-trophoresis under reducing and nonreducing conditions. Next the separated proteins were transferred onto a nitrocellulose membrane (Schleicher & Schuell). Recombinant human IL-23 (R&D Systems) was used as a posi-
tive control and recombinant prestained proteins of known molecular mass (Bio-Rad) were included to enable the estimation of the molecular mass of our specific IL-23 signal. Before incubating the nitrocellulose membrane with polyclonal rabbit anti-human IL-23 p19 (a gift of Dr. J. Pihlomäinen, National Public Health Institute, Helsinki, Finland) for 18 h at 4°C. The specificity of this Ab was tested in an earlier study (17). The membrane was blocked with PBS and 5% containing 0.05% Tween 20 and 5% milk. After incubation with peroxidase-conjugated goat anti-rabbit IgG (DakoCytomation) for 1 h at room temperature, specific protein bands in the filter were visualized by the ECL Plus Western Blotting Detection System (Amer sham Biosciences) and detected by a fluorescence imager (Typhoon 9400; Amersham Biosciences).

#### Determination of the IL-23 heterodimer by ELISA

The amount of IL-23 protein in the supernatants from keratinocytes and MoDC and in keratinocyte cultures was determined by a solid-phase sand-
wich ELISA, using polyclonal goat anti-human IL-23 p19 (R&D Systems) as the coating Ab and biotin-conjugated monoclonal mouse anti-human IL-12/IL-23 p40 (BD Pharmingen) as the detecting Ab. Hence, this ELISA detects only the heterodimer form of IL-23, but not the separate subunits. Specificity of the ELISA was tested by using recombinant human IL-23 heterodimer (R&D Systems), IL-12 heterodimer (Strathmann Biotec), and IL-12/IL-23 p40 subunit (BioSource International). Only the rIL-23 het-
erodimer, but neither the IL-12 heterodimer nor the common p40 subunit, was detectable by this ELISA (data not shown) with a detection limit of 300 pg/ml.

#### Assessment of the biological activity of keratinocyte-derived IL-23

T cells from peripheral blood of healthy donors were obtained by negative selection by means of a CD4 T cell isolation kit and an AutoMACS in-
strument from Miltenyi Biotec according to the manufacturer’s protocol. Cells suspended in IMDM with 10% FCS were plated at a density of 10^5 per well in a 96-well plate (final volume, 200 μl/well) and were stimulated with plate-bound anti-CD3 plus anti-CD28. After 10 days, the memory T cells were harvested and subjected to a second round of similar stimulation, this time seeded in triplicate in the absence or presence of 10 ng/ml rIL-23 or 5% of concentrated supernatant from stimulated keratinocytes, using the monoclonal mouse anti-human IL-23 (10 μg/ml, clone 253810, R&D Sys-
tems) to block the IL-23 activity. The IFN-γ production was assessed after 65 h by a sandwich ELISA, as previously described (18).

#### Immunohistochemical staining

Immunohistochemical staining was performed to determine the expres-
sion of the p19 subunit of IL-23, the common IL-12/IL-23 p40 subunit, and the IL-12 p70 heterodimer in 5-μm skin sections, keratinocytes cultured on object glass, and cytokin prepdard from epidermal and dermal cell suspensions. Primary Abs were polyclonal rabbit anti-
human IL-23 p19, biotin-conjugated monoclonal mouse anti-human IL-
12/IL-23 p40 and monoclonal mouse anti-human IL-12 p70 (R&D Sys-
tems). After fixation with acetone at 4°C for 15 min, primary Ab at room temperature for 1 h or overnight at 4°C, biotin-conjugated goat anti-mouse (DakoCytomation) at room temperature for 30 min. After incubation of the nitrocellulose membrane with a 1:100 dilution of goat anti-mouse IgG (DakoCytomation), biotin-conjugated streptavidin (DakoCytomation) as the second step. After incubation step, except for the incubation with normal goat serum, the sections were washed three
times with Tris-buffered saline. Peroxidase activity was detected as red color using the chromogen 3-amin-9-ethylcarbazole (Sigma-Aldrich). Hematoxylin was used to perform nuclear staining. Mouse anti-human IgG1 (Dako Cytomation) was used as isotype control for both p40 and p70 Abs and polyclonal rabbit anti-human Factor XIIa Ab detecting (dermal DC; BioGenex Laboratories) was used as a control for the p19 staining.

In situ expression of the IL-23 p19 subunit was blindly assessed in three sections per biopsy, scoring separately suprabasal epidermis, basal epidermis, papillary dermis, and superficial dermis using the following arbitrary units: 0, no staining; 1, very weak staining and/or occasional single or few cells (1–5 cells/section); 2, weak staining and/or scattered cells and small groups of cells (5–30 cells/section); 3, moderate staining and/or relatively big groups of cells (30–100 cells/section); 4, strong staining and/or many cells in big groups (100–400 cells/section); or 5, very strong staining and/or very large groups of cells (>400 cells/section). Epidermal expression was defined as the sum of basal and suprabasal scores, whereas dermal expression was the sum of papillary and superficial dermal scores.

To identify the cells expressing IL-23 p19, dermal and epidermal cell cytospin preparations were double-stained with p19 and one of the following cell markers: CD1a (Langerhans cells; BD Pharmingen), HMB45 (melanocytes; DakoCytomation), CD1c (DC; BD Pharmingen), CD3 (T cells; BD Pharmingen), CD83 (mature DC; BD Pharmingen), CD36 (macrophages; Beckman Coulter), HLA-DR (APC; BD Pharmingen), and CD15 (polymorphonuclear leukocytes; DakoCytomation). After performing the IL-23 p19 staining as described, the sections were sequentially incubated with the second primary Ab, goat anti-mouse (DakoCytomation) and alkaline phosphatase-anti-alkaline phosphatase complex (DakoCytomation). The color development was achieved by using naphthol-AS-MX-phosphate (Sigma-Aldrich) for blue and 3-amin-9-ethylcarbazole for red.

Statistical analysis
The scoring of the in situ expression of IL-23 p19 was expressed as values for the mean ± SD. Data were analyzed for the statistical significance either with independent sample t test to compare the expression in normal skin with lesional and nonlesional psoriatic skin or with paired sample t test to compare the expression in nonlesional with lesional psoriatic skin. A value for p < 0.05 was considered as the level of significance.

Results
Constitutive expression of IL-23 p19 mRNA by human keratinocytes
To investigate whether human keratinocytes are able to express the IL-23 heterodimer, we extracted total RNA from cultured keratinocytes and assessed the presence of mRNA for the two subunits of this cytokine by RT-PCR. As shown in Fig. 1, nonstimulated keratinocytes from both normal and lesional lesional skin displayed mRNA for IL-23 p19 and IL-12/IL-23 p40, indicating that these cells constitutively express the two subunits of IL-23. In addition, we also demonstrated the constitutive presence of IL-12 p35 mRNA, confirming earlier studies (12, 19, 20).

Normal human keratinocytes express the IL-23 heterodimer
The RT-PCR analysis clearly revealed that keratinocytes can synthesize the mRNA for both subunits of IL-23. As a next step we wanted to determine whether these cells were able to produce these subunits at the protein level and to combine them to form the heterodimeric IL-23. The cell lysate and supernatant of stimulated normal keratinocytes were subjected to Western blot analysis, using a polyclonal rabbit Ab that is specific for IL-23 p19. In nonreduced samples, we detected a protein with a molecular mass of ~60 kDa (Fig. 2, top), which probably represents a complex composed of the p19 and p40 subunits of IL-23 and is in line with an earlier study using the same Ab (21). In the reduced samples, p19 was detectable both in the recombinant and the keratinocyte samples (Fig. 2, bottom).

Immature MoDC, a potentially rich source of IL-23, were stimulated with several different stimuli, either alone or in combination, to determine which stimulus is most powerful to provoke IL-23 production. For this purpose, we designed a specific ELISA that only detects IL-23 heterodimers. As shown in Fig. 3a, no IL-23 could be detected in supernatants of unstimulated cells. Triggering via the CD40 molecule present on the cell surface induced IL-23, whereas peptidoglycan and poly(I:C) did not. Costimulation with IL-1β resulted in an enhanced IL-23 secretion by immature MoDC. The highest expression was achieved by a combination of J558/CD40L cells, peptidoglycan, poly(I:C), IFN-γ, and IL-1β (Fig. 3a). Because this mixture of stimuli gave the strongest expression of IL-23 in MoDC, we used it to stimulate keratinocytes. In contrast to the high production of IL-23 by MoDC, we found a low but significant amount of the IL-23 heterodimer in the supernatants of stimulated keratinocytes when the supernatant was concentrated 40- to 70-fold (Fig. 3b). On the cell basis, this expression was ~1000-fold lower than in MoDC used to set up the detection system. The level of IL-23 production by keratinocytes from normal human skin or from lesional psoriatic skin appeared to be comparable.

Keratinocyte-derived IL-23 stimulates IFN-γ production in memory T cells
In the next series of experiments we questioned whether the low amount of IL-23 produced by keratinocytes was sufficient to affect

**FIGURE 1.** Keratinocytes express mRNA for the p19 and p40 subunits of the IL-23 heterodimer. Total RNA was isolated and reverse transcribed, and after 45 cycles of amplification, the products were analyzed. The mRNA integrity was controlled by amplification of β2-microglobulin. Both normal and lesional psoriatic keratinocytes were found to express mRNA for the IL-23-specific p19 subunit, in addition to the known expression of the IL-12 p35 and IL-12/IL-23 p40 subunits. This result is representative of keratinocytes from four normal individuals and four lesional psoriasis patients.

**FIGURE 2.** IL-23 heterodimer is expressed by human keratinocytes at protein level. Normal human keratinocytes were stimulated overnight with poly(I:C) and IFN-γ. The supernatants and the cell lysates were subjected to SDS-PAGE and Western blot analysis under nonreducing (top) and reducing (bottom) conditions. Recombinant human IL-23 heterodimer (100 ng) (lane 1), cell lysate (lane 2), and 20-fold concentrated supernatant of stimulated keratinocytes (lane 3) are shown, and data are representative results of independent experiments.
the IFN-\(\gamma\) production by memory T cells. To this end, we set up a T cell stimulation assay in which the IFN-\(\gamma\) production could be enhanced by addition of rIL-23, and neutralization of this effect could be achieved with anti-IL-23 (Fig. 4). Addition of concentrated supernatant from activated keratinocytes to the CD3/CD28-stimulated memory T cells provoked an enormous boost in the IFN-\(\gamma\) production (\(n = 2\)). This amplification could only partially be blocked with anti-IL-23 (Fig. 4). The isotype control mouse \(\text{IgG2b}\) did not significantly affect the IFN-\(\gamma\) production (Fig. 4), nor did the concentrated keratinocyte culture medium or the mixture used to stimulate the keratinocytes (data not shown). These results indicate that the keratinocyte-derived IL-23 is biologically active, and in addition, that other IFN-\(\gamma\) promoting cytokines are secreted by keratinocytes.

Expression of p19 and p40 subunits of IL-23 at protein level in keratinocytes

To further substantiate the evidence that human keratinocytes are able to express IL-23, we cultured keratinocytes from normal (\(n = 3\)), lesional psoriatic skin (\(n = 3\)), and nonlesional psoriatic skin (\(n = 3\)) on object glasses and subsequently stained them with Abs recognizing the unique p19 subunit or the common p40 subunit of IL-23. For reasons of comparison, we stained the cells for the presence of the p70 heterodimer of IL-12 as well. The IL-23 p19 subunit was expressed in normal, lesional, and nonlesional psoriatic keratinocytes at comparable levels (Fig. 5, a–c). The staining was stronger in small and round cells than in big cells with abundant cytoplasm, suggesting that the p19 expression was more prominent in the undifferentiated keratinocytes. Staining with the p40 Ab revealed a similar picture as the staining with the IL-23 p19 Ab (Fig. 5, d–f). However, IL-12 p70 expression, which represents the biologically active form of IL-12, was seen only in a few cells in normal and in lesional or nonlesional psoriatic keratinocytes (Fig. 5, g–i). Polyclonal rabbit Factor XIIIa Ab, which is used as a control for the polyclonal rabbit p19 Ab, did not show any staining in the keratinocytes (data not shown).

In situ expression of the IL-23 p19 subunit in skin

To confirm the expression and localization of IL-23 in situ, we stained normal (\(n = 7\)), nonlesional (\(n = 5\)), and lesional (\(n = 10\)) psoriatic skin sections with the IL-23 p19-specific rabbit polyclonal Ab. The expression of the p19 subunit in the epidermis of psoriatic lesional skin was diffuse and very strong, whereas in the dermis, perivascular cells abundantly expressed this molecule (Fig. 6a). In contrast, in normal human skin the IL-23 p19 expression in the epidermis was much weaker and in the dermis this expression was present in only a limited number of cells, mainly around the capillaries (Fig. 6c). The control stainings with Factor XIIIa, known to be specifically expressed by dermal DC, revealed a completely different staining pattern, showing positive cells in the papillary and superficial dermis and no positive cells in the epidermis.
In addition, the staining with IL-23 p19-specific polyclonal goat Ab showed the same distribution pattern and intensity of p19 as obtained with the rabbit polyclonal Ab (data not shown). The expression of IL-23 p19 in normal, nonlesional, and lesional psoriatic skin was scored in arbitrary units and was summarized in Fig. 7. Statistical analysis of these values indicates that lesional psoriatic skin had a significantly higher expression of IL-23 than normal skin and nonlesional psoriatic skin. The expression of IL-23 p19 in normal vs nonlesional psoriatic skin did not differ significantly. These results are different from the staining pattern seen in the preparations of cultured keratinocytes that showed a similar level of p19 expression in lesional, nonlesional, and normal skin. This discrepancy might be caused by the induction of IL-23 p19 expression during in vitro processing and culturing of keratinocytes.

### Identification of IL-23 p19 expressing epidermal and dermal cells

In the following experiments, we wanted to determine which cell types were responsible for the in situ expression of IL-23 p19 in the epidermis and dermis. Cytospin preparations of freshly isolated epidermal and dermal cells from normal skin were double-stained with IL-23 p19 and different cell markers. Concerning the epidermal compartment, double staining with p19 and CD1a revealed that Langerhans cells expressed IL-23 (Fig. 8a). Double staining with p19 and HMB45 indicated that melanocytes were not able to produce IL-23 (Fig. 8b). In dermal cell suspensions, IL-23 p19 expression colocalized with cells that displayed CD1c (Fig. 8c), CD36 (Fig. 8d), and HLA-DR (Fig. 8e). Few CD3\(^+\) T cells also expressed this cytokine (Fig. 8f). CD83\(^+\) cells were quite low in number, but stained for IL-23 p19 (Fig. 8g). There were no CD15\(^+\) polymorphonuclear cells present in the dermal cell suspension (data not shown).

### Discussion

In this study, we demonstrated that human keratinocytes constitutively express mRNA for the p19 and p40 subunits of IL-23 and synthesize the heterodimer protein of ~60 kDa, which is the biologically active form of this cytokine. By means of ELISA we showed that human keratinocytes can secrete low but significant levels of IL-23, and functional analysis proved that these low levels of keratinocyte-derived IL-23 are biologically active and sufficient to amplify the IFN-\(\gamma\) production by memory T cells. These results not only support the view that keratinocytes contribute to cutaneous inflammation but also indicate that they can enhance type 1 immune responses through the expression of IL-23.
We found that a powerful stimulation for IL-23 production by MoDC was provided via CD40 triggering, and this effect was enhanced by IFN-γ and in particular IL-1β. This kind of stimulation may also be biologically relevant for keratinocytes, because, like DC, they also express CD40, enabling them to interact with CD40L-expressing T cells (22, 23). IL-1β and IFN-γ are known to induce CD40 expression on DC (24, 25), whereas IFN-γ, but not IL-1β, increases the expression of CD40 by keratinocytes (26). Further, IFN-γ also stimulates expression of IL-1β by keratinocytes (27). Although it is clear that triggering by CD40L, IFN-γ, and IL-1β are appropriate stimuli for immature MoDC to produce IL-23, it cannot be excluded that other kind of costimuli are needed to boost IL-23 production by keratinocytes. In contrast, one can imagine that keratinocytes only produce low levels of IL-23 because robust secretion by the epidermal layer would be harmful, as will be discussed below.

Keratinocytes in lesional psoriatic skin expressed markedly higher levels of IL-23 compared with keratinocytes in normal skin. In a separate study we found that after successful narrow-band UV B irradiation therapy, a common treatment for moderate to severe psoriasis, the elevated levels of IL-23 were reduced to levels found in normal skin or in nonlesional skin (17). The increased expression of this cytokine in the epidermis of psoriatic lesions does not seem to be an intrinsic aberration of the keratinocytes because cultured keratinocytes from normal skin and from nonlesional and lesional psoriatic skin display similar IL-23 staining by immuno-histochemistry and secrete comparable levels of IL-23. The enhanced expression of IL-23 in psoriatic lesional keratinocytes in situ is likely to be induced by cells in their neighborhood, perhaps the activated memory T cells within the infiltrate. A key role of cutaneous T cells in the induction of psoriatic lesions is underlined in a recent study in which symptomless skin of patients with psoriasis was grafted onto immunodeficient mice lacking T cells, B cells, and NK cells (28). It appeared that due to the local proliferation of donor T cells in the transplanted skin the graft transformed into a psoriatic lesion.

The importance of IL-23 in immune responses in vivo has been clearly demonstrated in mice; IL-23-deficient mice display severely compromised T cell-dependent humoral immunity and strongly impaired delayed-type hypersensitivity responses (29). The lack of IL-23 could apparently not be compensated by endogenous IL-12, indicating that IL-23 is critical for memory T cell responses in vivo. In contrast, T cell priming is not impaired in these animals and normal levels of memory T cells are present. Interestingly, IL-23-deficient mice show a clear reduction in IL-17 production (29), which matches with the finding that IL-23, but not IL-12, is a potent inducer of the proinflammatory cytokine IL-17 in both CD4+ and CD8+ T cells (30, 31). In addition, the IL-23-deficient animals phenotypically resemble IL-17-deficient mice (29, 32). In contrast to IL-23 deficiency, engineered systemic overexpression of IL-23, as present in transgenic p19 mice, causes...
IL-23 expression by keratinocytes

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Disclosures
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