Signalling of multiple interleukin (IL)-17 family cytokines via IL-17 receptor A drives psoriasis-related inflammatory pathways

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Data Availability Statement
The gene array dataset described in this publication has been deposited in NCBI’s Gene Expression Omnibus and is accessible through GEO Series accession number GSE158448 (https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE158448).

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Summary

Background The interleukin (IL)-23/IL-17 immune axis is of central importance in psoriasis. However, the impact of IL-17 family cytokines other than IL-17A in psoriasis has not been fully established.

Objectives To elucidate the contribution of IL-17 family cytokines in psoriasis.

Methods To address the expression and localization of IL-17 family cytokines, lesional and nonlesional skin samples from patients with psoriasis were analysed by several complementary methods, including quantitative polymerase chain reaction, immunoassays, in situ hybridization and immunohistochemistry. Mechanistic studies assessing the functional activity of IL-17 family cytokines were performed using ex vivo cultured human skin biopsies and primary human keratinocytes.

Results We demonstrated that IL-17A, IL-17F, IL-17A/F and IL-17C are expressed at increased levels in psoriasis lesional skin and induce overlapping gene expression responses in ex vivo cultured human skin that correlate with the transcriptomic signature of psoriasis skin. Furthermore, we showed that brodalumab, in contrast to ixekizumab, normalizes gene expression responses induced by the combination of IL-17A, IL-17F, IL-17A/F and IL-17C in human keratinocytes.

Conclusions Several IL-17 ligands signalling through IL-17RA are overexpressed in psoriasis skin and induce similar psoriasis-related inflammatory pathways demonstrating their relevance in relation to therapeutic intervention in psoriasis.

What is already known about this topic?
- The key role of interleukin (IL)-17A in psoriasis is well established.
- Previous studies have shown that IL-17A, IL-17F and IL-17C are overexpressed in psoriasis skin, whereas contradictory results have been published for IL-17E.
- IL-17 family cytokines induce secretion of inflammatory mediators such as antimicrobial peptides, chemokines and cytokines involved in the pathophysiology of psoriasis.

What does this study add?
- Levels of IL-17A/F are increased in lesional psoriasis skin but markedly lower than IL-17A and IL-17F.
- In ex vivo cultured human skin, a physiologically relevant model, IL-17A, IL-17F, IL-17A/F and IL-17C show functional redundancy in shaping the psoriasis transcriptome.
- IL-17RA antagonism normalizes expression of psoriasis-related genes in keratinocytes induced by the combination of IL-17 family cytokines.
The development of targeted treatments for psoriasis has demonstrated a key role of the interleukin (IL)-23/IL-17 pathway. The IL-17 family cytokines consist of six distinct homodimers (IL-17A–F), as well as an IL-17A/F heterodimer. IL-17 ligands signal through heterodimeric receptor complexes. Of the IL-17 receptors (IL-17R), IL-17RA has emerged as a common co-receptor for IL-17A, IL-17A/F, IL-17F, IL-17C and IL-17E, and interacts with another receptor subunit (IL-17RB, IL-17RC, IL-17RD or IL-17RE) to confer ligand- and cell type-dependent signalling specificity.

Previous studies have shown that IL-17A, IL-17F and IL-17C are overexpressed in psoriasis skin, whereas contradictory results have been published on whether IL-17E is expressed or increased in psoriasis skin. IL-17A is a central pathogenic cytokine in psoriasis as shown by the substantial clinical efficacy achieved in patients treated with secukinumab and ixekizumab, antibodies specifically targeting IL-17A. Recently, phase III studies with bimekizumab, targeting both IL-17A and IL-17F, have shown high efficacy, indicating that IL-17F also contributes to driving skin inflammation in psoriasis. Moreover, combined inhibition of several IL-17 family cytokines with brodalumab, targeting IL-17RA, has been shown in clinical studies to be very efficacious.

IL-17A, IL-17A/F and IL-17F, primarily produced by T helper (Th) 17 cells, act on a range of tissue cells, of which keratinocytes and fibroblasts are the main target cells in the skin. In contrast, IL-17C functions primarily in an autocrine manner acting on epithelial cells, including keratinocytes. IL-17A, IL-17F and IL-17C act in synergy with other cytokines, such as tumour necrosis factor (TNF) or IL-1, to induce expression of antimicrobial peptides, chemokines and pro-inflammatory cytokines, which promote innate immune responses, the recruitment of inflammatory cells, and keratinocyte activation and proliferation driving the psoriatic inflammation.

In this study, we analysed the mRNA and protein expression pattern of the IL-17 family cytokines in psoriasis skin with several complementary methods. Furthermore, we investigated the biological responses of the IL-17 family cytokines in ex vivo cultured human skin and their modulation by brodalumab and ixekizumab in keratinocyte cultures. Our findings demonstrate that IL-17A, IL-17F, IL-17A/F and IL-17C are upregulated in psoriasis skin and show functional redundancy in driving skin inflammation in psoriasis.

Materials and methods

Skin specimens from healthy participants and patients with psoriasis

Anonymized human skin samples were obtained from healthy participants following reconstructive surgery and from four different cohorts of patients with moderate-to-severe psoriasis (in addition, the second cohort also included a patient with mild psoriasis). Skin samples were donated after informed written consent was provided. All skin material was sampled in accordance with national legislation in the country of origin. Paired biopsies of lesional and nonlesional psoriasis skin were used for gene expression analysis by quantitative polymerase chain reaction [qPCR; cohort 1 (n = 10) and cohort 3 (n = 6)], cytokine protein analysis (cohorts 2 and 4, both n = 8) and histological analysis (cohort 2, n = 8). Each punch biopsy in cohort 2 was divided in half and used for cytokine and histological analysis.

Human skin cultures and cytokine stimulation

For ex vivo culture, skin samples from healthy participants were stored at 4–8 °C and processed within 24 h of surgery. Full-thickness (3-mm) punch biopsies (five replicates per treatment) were excised and placed in supplemented EpiLife medium (ThermoFisher Scientific, Waltham, MA, USA) in 96-well tissue-culture plates in a humidified incubator at 37 °C with 5% CO₂. Stimulation was performed for 24 h with IL-17 family cytokines as indicated.

Primary human keratinocytes were isolated from the skin of healthy adult donors (Appendix S1; see Supporting Information) and cultured in supplemented EpiLife medium (ThermoFisher Scientific) in a humidified incubator at 37 °C with 5% CO₂.

Cells were stimulated for 48 h with a combination of IL-17 family cytokines in the presence of TNF, as indicated. For all
samples treated with brodalumab, cells were preincubated with brodalumab for 30 min at 37 °C prior to cytokine addition. For all samples treated with ixekizumab, ixekizumab was preincubated with cytokine mixtures for 30 min at 37 °C before addition to the cells. Recombinant human cytokines were purchased from R&D Systems [Minneapolis, MN, USA (IL-17A: #317-ILB; IL-17C: #1234-IL; IL-17E: # 1258-IL; IL-17F: #1335-IL; IL-17A/F: #5194-IL; TNF: 210-TA)]. IL-17A, IL-17C and IL-17F and increased mRNA levels of IL17B, IL17D and IL17E (IL25) vs. nonlesional skin as shown in two independent patient cohorts (Figure 1a and Figure S1a; see Supporting Information). IL17E and its specific receptor subunit IL17RB had the lowest expression in lesional skin of the IL-17 ligands and the IL17R subunits, respectively (Figure 1a and Figure S1b).

In line with the mRNA expression pattern, protein levels of IL-17A, IL-17C and IL-17F, as well as IL-17A/F, were significantly increased in lesional skin compared with nonlesional skin (Figure 1b and Figure S1c). While the levels of IL-17A, IL-17F and IL-17C varied between the patients, IL-17C levels were typically higher than those of IL-17A and IL-17F. In addition, protein levels of IL-17A/F were markedly lower than those of IL-17A or IL-17F in all patients. We were unable to detect IL-17E protein in any of these samples, likely owing to low IL17E mRNA levels.

Distinct localization patterns of interleukin-17 family cytokines in psoriatic skin

Next, we investigated the localization and identity of the cells producing IL-17RA and all associated IL-17 ligands in psoriasis skin by ISH. IL17RA mRNA was expressed by keratinocytes in all viable layers of the epidermis, and also present in the papillary and reticular dermis (Figure 2a). IL17A and IL17F mRNA were predominantly present in cells in the epidermis and papillary dermis, and only occasionally in inflammatory infiltrates in the reticular dermis of lesional skin. The IL17A- and IL17F-expressing cells were few in number, showing high levels of mRNA for the cytokine and morphologically resembling T cells (Figure 2b, c). T-cell identity was confirmed by triple-fluorescence ISH for IL17A, IL17F and CD3 (four patients), with the vast majority of the IL17A- and IL17F-expressing cells showing co-expression of CD3 (Figure 2d). Furthermore, IL17F was almost always co-expressed with IL17A, whereas IL17A, to a lesser extent, was co-expressed with IL17F in the psoriasis skin samples (data not shown). IL17C mRNA was highly expressed in lesional skin by clusters of differentiated keratinocytes in the upper layers of epidermis (Figure 2e). Interestingly, we found that IL17E mRNA was only expressed by few keratinocytes often associated with hair follicles in lesional (Figure 2f) and nonlesional skin. Quantitative analysis of the ISH staining indicated very low expression of IL17A,

Histology on psoriasis biopsies

Single- and triple-fluorescence in situ hybridization (ISH) were performed with RNAscope® (Bio-Technne, Minneapolis, MN, USA). For multiplex fluorescence, tyramide signal amplification plus conjugated fluorophores was applied: Cy5 (CD3), Cy3 (IL17F) and fluorescein isothiocyanate (IL17A). Immunohistochemistry (IHC) was performed with goat antihuman IL-17A (R&D Systems), goat antihuman IL-17C (R&D Systems) and mouse antihuman neutrophil elastase (Agilent, Santa Clara, CA, USA). The primary antibodies for IL-17A and IL-17C were detected with BOND Polymer Refine Detection with rabbit antigoat (Agilent) as the secondary antibody and DAB as the chromogen. Neutrophil elastase was detected with BOND Polymer Refine RED Detection, with fast red as the chromogen. All stains were performed on a Leica BOND RX (Leica Biosystems, Wetzlar, Germany). Whole-slide image analysis was performed with Visiopharm Integrator Software (Visiopharm, Hoersholm, Denmark). Further details can be found in Appendix S1.

Cytokine protein analysis on psoriasis biopsies

IL-17F protein levels were measured by enzyme-linked immunosorbent assay (ELISA), using the Human IL-17F Duo-Set ELISA Kit (R&D Systems). The MSD platform (Meso Scale Discovery, Rockville, MD, USA) was used for measurement of IL-17A/F and IL-17E (Human U-PLEX, individual assay for IL-17A/F and IL-17E) and IL-17A and IL-17C (Human U-PLEX, multiplex assay). Further details can be found in Appendix S1.

Quantitative real-time polymerase chain reaction analysis

RNA extraction from lesional and nonlesional psoriasis skin and from keratinocyte cultures was followed by cDNA synthesis, and amplification of cDNA by quantitative real-time PCR using Taqman® Gene Expression Assays (see Table S1).

Gene array and bioinformatics analyses

Gene expression profiling of ex vivo cultured human skin biopsies was carried out on whole-transcript arrays (GeneChip™ Human Gene 2-1 ST Array; Thermo Fisher Scientific). Differential expression analysis was performed with the respective contrasts stimulated vs. unstimulated, using the moderated t-test of the R/limma package. Correlation analysis was carried out using the Pearson correlation coefficients (r) of the R/psych package. Further details are available in Appendix S1.
IL17C, IL17E and IL17F in nonlesional skin and a significant increase in IL17A, IL17F and IL17C, but not IL17E, mRNA in lesional skin (Figure 2g), as was also observed by qPCR analysis (Figure 1a and Figure S1a). Furthermore, we found a strong correlation between mRNA levels by ISH and protein levels in lesional skin samples from the same patients, indicating that the elevated mRNA expression of IL17A, IL17F and IL17C directly led to increased cytokine production in the skin (Figure 3a). Interestingly, when we compared IL17A mRNA and protein expression on consecutive sections of lesional skin (n = 8), we observed that while IL17A mRNA and protein are found in few cells in the epidermis and papillary dermis, the majority of cells that stained positive for IL-17A protein localized to the reticular dermis (Figure 3b, c). In addition, neutrophils (i.e. in Munro abscesses) staining positive for neutrophil elastase also stained positive for IL-17A protein in the absence of IL17A mRNA (Figure 3d–f). However, IL17C mRNA and protein co-localized in keratinocytes in the upper epidermis (Figure 3g, h).

**Global gene expression induced by interleukin (IL)-17A, IL-17F, IL-17A/F and IL-17C in human skin overlaps with the psoriasis gene signature**

To investigate the contribution of the different IL-17 family cytokines in psoriasis, we stimulated ex vivo cultured human skin biopsies for 24 h with the cytokines as indicated (Figure 4a). Global gene expression profiling was performed using Affymetrix arrays to allow direct comparison of the cytokine-induced differentially expressed genes (DEGs; vs. untreated controls) with psoriasis DEGs (comparing lesional to nonlesional skin) using the meta-analysis derived psoriasis (MAD3-PSO) transcriptome.27 IL-17A, IL-17A/F and IL-17F induced dose-dependent and qualitatively comparable gene expression responses, and increased expression of most of the top-50 upregulated MAD3-PSO genes (Figure 4a and Table S2; see Supporting Information). IL-17C induced a more modest response, but, overall, affected the same upregulated MAD3-PSO genes as for IL-17A, IL-17A/F, and IL-17F. When we
Figure 2 Cells expressing IL17A, IL17F and IL17C mRNA are predominantly localized in the epidermis of psoriatic skin. In situ hybridization (ISH) on lesional skin sections detecting (a) IL17RA mRNA, (b) IL17A mRNA and (c) IL17F mRNA. (d) Co-expression of IL17A and IL17F by CD3+ T cells, as illustrated by triple-fluorescence ISH. (e, f) As in (a–c), but for IL17C and IL17E mRNA, respectively. (g) Quantification of IL17A, IL17C, IL17F and IL17E mRNA expression by ISH in lesional and nonlesional psoriasis skin, as percentage signal area in the epidermis. Mean values (SEM) were calculated, and statistical analysis was performed by two-sided paired t-tests. Scale bars for (a–c) and (e, f) are 100 μm and 50 μm, respectively. Arrows in (b–d and f) point to locations with positive cells. *P < 0.05 and **P < 0.01. L, lesional; NL, nonlesional.
correlated all upregulated MAD3-PSO genes (log₂ fold change > 1) to the cytokine-induced gene expression profiles (using the highest concentration at 100 ng mL⁻¹), a significant correlation was found for IL-17A (r = 0.35), IL-17F (r = 0.39), IL-17A/F (r = 0.40) and IL-17C (r = 0.26; all P < 0.001) (Figure 4b). As exemplified by a panel of selected

![Figure 3](image_url)  
**Figure 3** Expression of interleukin (IL)-17 family cytokines in psoriasis skin by in situ hybridization (ISH) and immunohistochemistry (IHC). (a) Correlation analysis between cytokine levels measured by quantification of ISH and protein concentrations in skin lysates from matched samples (n = 8). (b) IL17A mRNA expression by ISH. (c) Consecutive section of (b) showing IL-17A protein expression by IHC. (d) IL17A mRNA expression by ISH. (e) Consecutive section of (d) showing IL-17A protein expression by IHC. (f) Consecutive section of (d and e) showing neutrophil elastase expression by IHC. (g) IL17C mRNA expression by ISH. (h) Consecutive section of (g) showing IL-17C protein expression by IHC. (i) Negative control staining using an isotype control antibody. Scale bars for (b) and (c) are 250 µm, and scale bars for (d–i) are 100 µm. Arrows (IL17 mRNA) and arrowheads (IL-17 protein) in (b–i) point to locations with positive cells.
Figure 4 Interleukin (IL)-17A, IL-17F, IL-17A/F and IL-17C are redundant in shaping the psoriasis transcriptome. Ex vivo cultured human skin biopsies (five replicates per treatment condition) were incubated for 24 h with individual IL-17 family cytokines at the indicated concentrations and subjected to global gene expression analysis using Affymetrix Human Gene ST 2.1 arrays. (a) Ranked heatmap of the top-50 MAD3-PSO genes and their expression (log2 fold change vs. untreated ex vivo skin cultures). (b) A correlation matrix of gene expression signatures induced by individual IL-17 family cytokines (at 100 ng mL$^{-1}$) in ex vivo cultured human skin vs. upregulated transcripts in the meta-analysis derived psoriasis transcriptome (MAD3-PSO) dataset (log2 fold change $>$ 1). The exact correlation score ($r$) and significance can be seen at the top right of the table. Crosses indicate that no significant correlation was found. (c–h) Examples of genes induced by individual IL-17 family cytokines in ex vivo cultured human skin: DEFB4A, DEFB4B, SPRR2C, S100A7A, SPRR2F and LCN2, respectively. ***$P < 0.001$. n.s., not significant.
Interleukin (IL)-17 receptor A antagonism normalizes psoriasis-relevant IL-17 family cytokine signalling

Based on the overexpression and functional redundancy of IL-17A, IL-17F, IL-17A/F and IL-17C in psoriasis, we next investigated the impact of IL-17RA antagonism vs. neutralization of IL-17A on gene expression induced by the combination of IL-17 family cytokines (using approximately equipotent concentrations) and TNF in primary human keratinocytes (Figure 5). The combined stimulation with IL-17 family cytokines in the presence of TNF led to a strong induction of DEFB4, S100A7A, CXC18, LCN2 and SPRR2F (Figure 5). Treatment with ixekizumab (anti-IL-17A antibody) was able to inhibit gene expression to a comparable level induced by the combination of IL-17F, IL-17C and TNF. However, treatment with brodalumab (anti-IL-17RA antibody) blocked the signalling of all the IL-17 family cytokines and was able to inhibit their induced gene expression approximately to the levels induced by TNF alone (Figure 5). These findings indicate that IL-17A, IL-17F, IL-17A/F and IL-17C make use of IL-17RA as functional receptor and illustrate the differences in mode of action of IL-17A neutralizing antibodies (e.g. ixekizumab) and the IL-17RA blocking antibody brodalumab.

Discussion

Our study provides a comprehensive analysis by using several complementary methods (qPCR, immunobassays, ISH and IHC) to investigate thoroughly the expression and localization of IL-17 family cytokines in psoriasis skin. Notably, for the first time, the gene and protein levels of all IL-17RA-associated IL-17 ligands were assessed on the same psoriasis skin biopsies. By the different methods employed, we found that levels of IL-17A, IL-17F and IL-17C, but not IL-17E, were significantly increased in lesional psoriasis skin, and correlated at the mRNA and protein levels. In our analysis, protein levels of IL-17C were typically higher than those of IL-17A and IL-17F, whereas levels of IL-17A/F were markedly lower in lesional skin biopsies. While protein levels of IL-17A/F in psoriasis skin have not been published previously, other studies have described increased levels of IL-17A, IL-17F and IL-17C at the mRNA and/or protein level, although protein levels of these cytokines in psoriasis skin vary widely between studies. This may be explained by differences related to local disease severity of the biopsy site, processing of the biopsies (e.g. protein recovery), detection systems and data normalization.

Although we were able to detect modest IL17F mRNA expression by ISH in keratinocytes, mostly in locations next to hair follicles, we did not find increased levels of IL-17F in lesional psoriasis skin by any of our methods, which is in contrast to findings from two independent studies. The reason for this discrepancy is not entirely clear but is possibly related to different psoriasis subtypes, sample processing and/or specificity of detection methods used in the studies. Levels of IL-17B and IL-17D were only assessed by qPCR and, in line with previous reports, were found to be downregulated in lesional psoriasis skin biopsies.

By using multiplex ISH staining, we confirmed CD3+ T cells as the main IL17A and IL17F expressing cells. Interestingly, by comparing ISH and IHC analysis of consecutive skin sections, we found that neutrophils and cells in the dermis stained positive for IL-17A protein in the absence of IL17A mRNA expression. While still being a matter of controversy, our findings, although based on a limited number of psoriasis skin sections, suggest that neutrophils accumulate and possibly internalize IL-17A but seem unable to produce IL-17A, as also supported by other studies.

Transcriptional profiling of IL-17 family cytokines have been addressed to some extent in previous studies using keratinocyte monolayer cultures or reconstructed epidermis models, but, to our knowledge, no global gene expression analysis has been generated that directly compares the different IL-17 ligands in a skin-relevant model system. We performed a global gene expression analysis comparing the IL-17 ligands in ex vivo cultured human skin biopsies. The use of whole-skin tissue allowed us to capture gene responses from different cells present in the skin and permits potential integrative or interactive effects between the different cells and cytokines that are induced. We found that the global gene expression profiles induced by IL-17A, IL-17A/F, IL-17F and, to a lower extent, IL-17C, showed a qualitatively comparable induction of genes, encoding inflammatory mediators such as antimicrobial peptides, chemokines and cytokines involved in the pathophysiology of psoriasis. Moreover, these gene expression signatures were very similar to most common upregulated genes in psoriasis and correlated significantly with the MAD3-PSO dataset. The relatively weak responses induced by IL-17C in vitro were also reported previously, not the least by comparison with the effects of IL-17C shown in mouse models. While whole-skin tissue is a more physiologically relevant model system for determining the effects of IL-17 family cytokines in cutaneous biology, a limitation is the lack of activated immune cells. Thus, in this model system, keratinocyte-induced mediators [e.g. CCL20 and C-X-C motif chemokine ligand 8 (CXC18)] do not permit amplification of the
**Interleukin (IL)-17RA antagonism normalizes psoriasis-relevant IL-17 family cytokine signalling in vitro.**

Gene expression of DEFB4, SPRR2F, LCN2, CXCL8 and S100A7A by quantitative polymerase chain reaction of primary human keratinocytes treated for 48 h with a combination of IL-17 family cytokines (IL-17A, 1 ng mL\(^{-1}\); IL-17F, 10 ng mL\(^{-1}\); IL-17A/F, 10 ng mL\(^{-1}\); IL-17C, 100 ng mL\(^{-1}\)) and tumour necrosis factor (10 ng mL\(^{-1}\)) in the absence or presence of brodalumab (anti-IL-17RA, 500 µg mL\(^{-1}\)) or ixekizumab (anti-IL-17A, 500 µg mL\(^{-1}\)) as indicated. RPLP0, PFA and ACTB were used as reference genes for normalization. Bars show mean (SD) fold change in gene expression vs. unstimulated samples (2\(^{-\Delta\Delta C\text{t}}\)). Statistical analysis was performed by unpaired t-tests. *\(p<0.05\), **\(p<0.01\) and ***\(p<0.001\). n.s., not significant.

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Supporting Information

Additional Supporting Information may be found in the online version of this article at the publisher’s website:

Appendix S1 Supplementary materials and methods.

Figure S1 Expression of interleukin-17 family cytokines and receptors in psoriasis skin biopsies.

Figure S2 Bioactivity of interleukin-17E in peripheral blood mononuclear cells and IL17R expression levels in ex vivo cultured human skin.

Table S1 Probes and primers used in quantitative polymerase chain reaction-based gene expression analysis.

Table S2 Regulation of the top-50 upregulated MAD3-PSO probes and primers used in quantitative polymerase chain reaction-based gene expression analysis.

Table S3 Differential gene expression in ex vivo cultured skin stimulated with interleukin-17 family cytokines.