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Single-cell transcriptomics applied to emigrating cells from psoriasis elucidate pathogenic vs. regulatory immune cell subsets

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Abstract

Background: Inflammatory cells in previous human skin single-cell data constituted only a small fraction of the overall cell population, such that functional subsets were difficult to ascertain.

Objective: Our aims were to overcome the limitation by applying single-cell transcriptomics to emigrating cells from skin and elucidate *ex vivo* gene expression profiles of pathogenic vs. regulatory immune cell subsets in psoriasis skin.

Methods: We harvested emigrating cells from human psoriasis skin after incubation in culture medium without enzyme digestion or cell sorting and analyzed cells with single-cell RNA sequencing and flow cytometry simultaneously.

Results: Unsupervised clustering of harvested cells from psoriasis and control skin identified NK cells, T-cell subsets, dendritic cell subsets, melanocytes, and keratinocytes in different layers.

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Comparison between psoriasis vs. control cells within each cluster identified: 1) Cutaneous Type 17 T-cells display highly differing transcriptome profiles depending on IL-17A vs. IL-17F expression and IFN γ vs. IL-10 expression, 2) Semimature dendritic cells are regulatory dendritic cells with high IL-10 expression but a subset of semimature dendritic cells expresses IL-23A and IL-36G in psoriasis, and 3) CCL27-CCR10 interaction is potentially impaired in psoriasis due to decreased CCL27 expression in basal keratinocytes.

Conclusion: We propose that single-cell transcriptomics applied to emigrating cells from human skin provide an innovative study platform to compare gene expression profiles of heterogeneous immune cells in various inflammatory skin diseases.

Clinical Implication: Our method can readily be extended to study modulation of leukocyte transcripts by immune-targeted treatments.

Capsule Summary:

Single-cell transcriptomics applied to emigrating cells from psoriasis skin elucidate *ex vivo* gene expression profiles of cutaneous type 17 T-cell and regulatory T-cell subsets and mature vs. semimature dendritic cells.

Keywords

Psoriasis; single-cell RNA sequencing; T-cells; dendritic cells; keratinocytes; emigrating cells

Introduction

Psoriasis vulgaris is the most common inflammatory disease in humans that is mediated by IL-17 producing Type 17 T-cells (T17 cells), which includes CD4⁺ (Th17) and CD8⁺ (Tc17) subsets¹. The heterogeneity in T17 cells is suggested by the finding that individual T-cells can synthesize only IL-17A or IL-17F (homodimers) or they can synthesize IL-17A/F (heterodimer), while each of these IL-17 isoforms has differing ability to activate IL-17 receptors on target cells². The expression of pathogenic vs. regulatory markers in IL-17A vs. IL-17F-producing T-cells is presently unknown.

The chronicity of psoriatic lesions may be caused by defective negative regulatory pathways, such as decreased function of regulatory T-cells (Tregs)³, reduced Type 1 regulatory T-cells (Tr1)⁴ and low expression of negative regulatory factors that are highly expressed in resolving delayed-hypersensitivity skin reaction lesions⁵. However, the function of Tregs has been assessed only in peripheral blood³, and the properties of Tregs and other regulatory immune cells in human skin is largely unexplored⁶.

Normal skin contains a subset of myeloid dendritic cells (DCs) that are characterized by BDCA-3 expression, IL-10 production and expression of negative immune regulators, so this population is considered to be regulatory contributing to immune homeostasis/tolerance under non-inflammatory conditions⁷. In contrast, psoriasis lesions contain an increased number of “inflammatory” CD11c⁺ DCs and mature populations marked by DC-LAMP expression⁸. The relative expression of inflammatory vs. regulatory molecules in DC populations present in psoriasis lesions has not been widely explored.

Single-cell RNA sequencing (scRNA-seq) is a powerful technique to characterize gene expression in individual leukocytes or other tissue-resident cells in human skin^{9–19}. However, leukocyte subsets were difficult to study in depth in human skin, as other skin cells such as fibroblasts and keratinocytes dominated the overall cell mixture. In this study, we have performed scRNA-seq in psoriasis and control skin using emigrating cells from skin biopsies to enrich leukocyte subsets without cell sorting or activation. We find novel and distinct patterns of gene expression within different T17 cell subsets, Tregs, mature and semimature DC subsets and keratinocytes in different layers that are disease-related and may drive key inflammatory pathways in psoriasis.

Methods

Please see the supplementary experiment section and Fig E1, E2 and E3 in the article's Online Repository for validation of our new method - scRNA-seq of emigrating cells from skin. With skin biopsies from the same sample, we compared our new method with our group's previous methods – 1) scRNA-seq of isolated cells from cryopreserved tissues by conventional enzyme digestion^{9–11}, and 2) flow cytometry analysis of emigrating cells²⁰ (Fig E1 and Fig E2). In addition, we compared our new method with existing scRNA-seq data of isolated cells from fresh skin tissues by conventional enzyme digestion (Fig E3). For extended methods, please see the supplemental materials section in the article's Online Repository. The scRNA-seq data have been deposited in NCBI's Gene Expression Omnibus and are publicly accessible through GEO Series accession number GSE151177. All the scripts used to present and analyze the data are publicly available from https://github.com/jaehwan79/KruegerLab_scRNAseq_emigrating_cells_human_skin. There is no restriction on the use of the data or code.

Patients

Human subject research was performed in accordance with the Helsinki Declaration and approved by the Institutional Review Board of the Rockefeller University, New York, NY, USA. Demographics and disease severity of the study cohort is presented in Table E1 in this article's Online Repository.

Harvesting emigrating cells from skin biopsy tissues

6 mm punch skin biopsy tissues from 13 psoriasis patients and 5 healthy volunteers were bisected, and one of the two bisected skin tissues was incubated in culture medium for harvesting emigrating cells. To split epidermis and dermis (Video E1), harvested skin tissue was immediately placed in 0.2% Dispase II (Sigma-Aldrich) and incubated in a humidified incubator at 37°C and 5% CO₂ for 3 hours. Then, epidermis and dermis were separated with forceps and sliced into small pieces with #10 blade scalpels (Video E2). Epidermis and dermis were separately incubated in RPMI-1640 medium with L-glutamine (Cytiva) supplemented with 10% human albumin serum (Sigma-Aldrich) without any enzyme in a humidified incubator at 37°C and 5% CO₂. Nonplastic adherent cells that had emigrated out of epidermis and dermis were harvested after 48 hours (Video E3). The harvested cells from epidermis and dermis were combined and filtered through a 40-µm cell strainer (Corning) (Video E4) and stored on ice. The cell numbers and viability were determined using a

Countess automated cell counter (Invitrogen) and trypan blue staining (BioRad). The cell viability of individual samples is presented in Table E1.

Single-cell RNA sequencing experiments

The 10x Genomics Chromium Single Cell 3' Reagents Kit user guide (<https://support.10xgenomics.com>) was used to prepare the single cell suspension. All the samples were sequenced by a single NextSeq 500 sequencer (Illumina) with the identical run parameters: read 1 for 28 cycles, read 2 for 55 cycles, and index for 8 cycles. Sequencing information including the sequencing saturation and depth is presented in Table E1.

Analysis of single-cell RNA sequencing data

We used Seurat R package (version 3.0) installed in R (version 3.6.2) for the downstream single-cell RNA sequencing data analysis²¹. Before data integration, single-cell data quality control was performed separately for each individual sample as previously described for human skin single-cell RNA sequencing analyses⁹⁻¹¹.

Principal component analysis and graph-based clustering analysis were performed, and differentially expressed genes of each cluster compared to all other cells were found. The average gene expression of psoriasis vs. control cells within a cluster was calculated. To compare psoriasis vs. control cells within clusters representing each type of skin immune cells, we merged adjacent clusters of common immune cell subsets (Fig E4). For immune cell cluster comparison, cells expressing target genes within each cluster were quantified by (1) number of cells expressing the target gene and (2) proportion of cells expressing the target gene.

We used iTALK R package (version 0.1.0) for the receptor-ligand interaction analysis. Differentially expressed genes between psoriasis vs. control in each cluster were found, and then the fold changes of differentially expressed genes were used to calculate receptor-ligand interaction between different cell clusters.

Multiparameter flow cytometry experiments

When emigrating cells from skin biopsy tissues were harvested and 5,000-10,000 harvested skin cells were entered into the 10X Genomics single-cell chip, the rest of harvested skin cells were simultaneously analyzed by multiparameter flow cytometry with dendritic cell panels. The harvested skin cells were washed and incubated in ice for 30 minutes with fluorochrome-conjugated monoclonal antibodies to cell-surface markers, and the cells were acquired with the BD LSR II flow cytometer (BD Biosciences) and analyzed with FlowJo software (TreeStar).

Results

Microfluidic partitioning of emigrating cells from human skin empowers single-cell transcriptomic profiling of heterogeneous immune cells under ex vivo conditions

Dimensionality reduction analysis of 23,220 single-cell data identified clusters of NK cells, CD161⁺ T-cells, CD8⁺ T-cells, CD4⁺ T-cells, Tregs, mature DCs, semimature DCs,

macrophages, melanocytes, and KCs in different layers of Stratum (S.) corneum, S. granulosum, S. spinosum, and S. basale without subclustering (Fig 1A–E). Leukocytes (NK cells, T-cells, dendritic cells and macrophages) constituted 53.0% of scRNA-seq data of psoriasis skin and 15.5% of control skin (Fig 1F and Fig E5B). The average number of T-cells per skin biopsy sample increased 7.0 times from control (67.4 ± 19.6) to psoriasis (472.9 ± 108.5), and the average number of DCs per skin biopsy sample increased 3.4 times from control (70.8 ± 33.4) to psoriasis (241.7 ± 63.7) ($p < 0.05$, Fig E5C). The top 10 most differentially expressed genes in each cluster are presented in Fig E6.

CD161⁺ T-cell and CD8⁺ T-cell clusters were segregated from other T-cell clusters (Figure 1C and 1D), but CD4⁺ T-cell and Treg clusters were not segregated and the designations were nominal (Fig 1E). Compared to CD4⁺ T-cell cluster, Treg cluster was characterized by a higher proportion of cells expressing CD25 (IL2RA) (16.1% in Treg vs. 1.7% in CD4₊ T-cell clusters), FoxP3 (16.2% in Treg vs. 0.4% in CD4₊ T-cell clusters) and CTLA4 (14.6% in Treg vs 1.7% in CD4⁺ T-cell clusters) (Fig E7). When we applied heuristic cut-off value of 1 to FoxP3 expression, FoxP3^{High} subsets in the Treg cluster expressed high levels of CD25 (IL2RA) with low levels of CD127 (IL7R) together with high levels of OX40 (TNFRSF4), TIGIT, CTLA4, and CCR10 (Fig 2 and Fig E6). When T-cell clusters were subset by cytokine mRNA expression, numerous cytokines were expressed by T-cells in psoriasis skin (Fig 2 and Fig E7) - TNFSF10 (23.9%), TNFSF12 (15.6%), CCL20 (10.1%), IL-36G (5.9%), TNF (4.2%), IL-26 (3.0%), IFN γ (2.8%), IL-17F (0.9%), IL-17A (0.6%), IL-22 (0.3%), and IL-13 (0.1%).

Cutaneous T17 cells display highly differing transcriptomes depending on IL-17A vs. IL-17F expression and IFN γ vs. IL-10 expression

T17 cells constituted 2.7% of CD4⁺ T-cell cluster, 1.3% of CD8⁺ T-cell cluster, 2.4% of CD161⁺ T-cell cluster and 0.5% of Treg cluster (Fig E7). When T17 cells were subdivided into Th17 (CD4⁺ T17) cells, Tc17 (CD8⁺ T17) cells, CD161⁺ T17 cells, and T17 cells in a Treg cluster (Fig E8), Th17 (CD4⁺ T17) cells expressed IL-17A (32.7%) or IL-17F (76.9%) (Fig E9). In contrast, Tc17 (CD8⁺ T17) cells expressed only IL-17A, and IL-17F was undetected in Tc17 cells with a cut-off value of 1. 50% of Tc17 cells expressed IFN γ , while only 13.5% of Th17 cells expressed IFN γ . 5.8% of Th17 cells expressed IL-10, while IL-10 was undetected in Tc17 cells with a cut-off value of 1. CD161⁺ T17 cells co-expressed CD8B (36.4%) or CD4 (9.1%) and expressed IL-17A (36.4%), IL-17F (72.7%) and IL-10 (9.1%).

Majority of T17 cells expressed either IL-17A or IL-17F and only 7.8% of T17 cells co-expressed IL-17A/IL-17F (Fig 3A). Excluding IL-17A/IL-17F co-producing T17 cells, IFN γ was expressed by 25.6% of IL-17A producing T17 cells, while only 3.9% of IL-17F producing T17 cells expressed IFN γ . In contrast, IL-10 was undetected by IL-17A producing T17 cells with a cut-off value of 1, but 5.9% of IL-17F producing T17 cells expressed IL-10.

IL-17A⁺ T17 cells in psoriasis skin expressed high levels of inflammatory cytokines, such as IL-26, CCL3, CCL4 and CCL5, which have been reported in pathogenic T17 cell transcripts^{22, 23} (Fig 3B). In addition, IL-17A⁺ IFN γ ⁺ T17 cells expressed high levels of

transcription factors, such as RORC and STAT4^{6, 23}, inflammatory cytokines including IL-36G and TNF¹, cytotoxic transcripts such as GZMA, GZMB, GNLY, CD8B and PRF1, and other pathogenic T17 cell transcripts^{23, 24}, such as CASP1 and LAG3.

IL-17F⁺ IL-10⁺ T17 cells expressed high levels of MAF, AHR, CD73 (NT5E) and IL1RN, which have been reported in non-pathogenic IL-10-producing T17 cell transcripts^{23, 25–30} (Fig 3B). In contrast, IL-17F⁺ IL-10⁻ T17 cells expressed high levels of inflammatory cytokines, such as IL-1 β , IL-2, IL-24, IL-34, EBI3 and LTA, and IL23R. Both IL-17F⁺ IL-10⁻ T17 cells and IL-17A⁺ IL-17F⁺ T17 cells expressed high levels of IL-22 and CCR6. In addition, IL-17A⁺ IL-17F⁺ T17 cells expressed high levels of BATF, SLC7A8 and TNFSF10.

Mature DCs in psoriasis skin express high levels of IL-23A & IL-36G and low levels of KYNU compared to mature DCs in control skin

When the average gene expression was compared between mature DC and semimature DC clusters split by psoriasis vs. control in the scRNA-seq data, mature DCs were characterized by 1) high expression of MHC class II molecules, 2) skin DC marker⁸ expression of CD86, DC-LAMP (LAMP3), CD205 (LY75), CD40, CIITA, CD80, PD1-L1 (CD274) and PD1-L2 (PDCD1LG2), and 3) expression of DC regulatory tryptophan metabolism enzyme³¹ KYNU (Fig 4A and Fig E10C).

Simultaneous flow cytometry analysis confirmed scRNA-seq findings of mature DCs (Fig 4B and Figure E10A). The Median fluorescence intensity (MFI) of HLA-DR, CD205 (LY75) and CD40 in mature DCs was higher than the MFI of HLA-DR, CD205 and CD40 in semimature DCs (Fig 4B). In addition, the proportions of CD205⁺ and CD40⁺ cells in mature DCs were higher than the proportion of CD205⁺ and CD40⁺ cells in semimature DCs (Fig E10A).

Mature DCs in psoriasis skin expressed more IL-23A and IL-36G compared to mature DCs in control skin. IL-23A or IL-36G expression in mature DCs increased 4.2 times from control skin (1.5%, 4/269) to psoriasis skin (6.2%, 148/2390) (Fig E11A and E11C). In contrast, mature DCs in psoriasis skin expressed less KYNU than mature DCs in control skin. KYNU expression in mature DCs decreased 2.1 times from control skin (21.6%, 58/269) to psoriasis skin (10.4%, 249/2390) (Fig E10C and Fig E11D).

Semimature DCs in psoriasis skin express high levels of IL-10, BDCA-3 and LILRB2, but a subset of semimature DCs also expresses IL-23A and IL-36G

Semimature DCs were characterized by 1) intermediate expression of MHC class II molecules (HLA-DRB5), 2) skin DC marker⁸ expression of CD11c (ITGAX), SIRPA, CD14, AIF1 and CD209 (DC-SIGN) and 3) High IL-10 expression (Fig 4A and Fig E10B). Among all IL-10 expressing cells in total scRNA-seq data, 53.5% (145/271) of IL-10 expressing cells were semimature DCs (Fig 5A and 5B). IL-10 expressing semimature DCs in psoriasis skin highly co-expressed BDCA-3 (THBD) and LILRB2 (ILT4). 46.5% (60/129) of IL-10 expressing semimature DCs in psoriasis skin co-expressed BDCA-3 (Fig 5C, 7D and 7F). 41.9% (54/129) of IL-10 expressing semimature DCs in psoriasis skin co-expressed LILRB2.

Psoriasis semimature DCs expressed more BDCA-3, DCR1 (TNFRSF10C), ITGAM, LILRB2, LILRB1 and LILRB4, and less BDCA-1 (CD1c)⁸ than semimature DCs in control skin (Fig 4A, 5C, 5D and Fig E10D). The BDCA-3 to BDCA-1 ratio increased 5.2 times from control (1.6) to psoriasis skin (8.4) (Fig 5E).

Simultaneous flow cytometry analysis confirmed scRNA-seq findings of semimature DCs (Fig 4B and Fig E10A). The MFI of BDCA-3, DCR1 and LILRB2 in psoriasis semimature DCs was higher than the MFI of BDCA-3, DCR1 and LILRB2 in control semimature DCs (Fig 4B). The MFI of BDCA-1 in psoriasis semimature DCs was lower than the MFI of BDCA-1 in control semimature DCs. In addition, the proportion of BDCA-3⁺ and LILRB2⁺ cells in psoriasis semimature DCs was higher than the proportion of BDCA-3⁺ and LILRB2⁺ cells in control semimature DCs (Fig E10A). The proportion of BDCA-1⁺ cells in psoriasis semimature DCs was lower than the proportion of BDCA-1⁺ cells in control semimature DCs.

When we analyzed the receptor-ligand interaction with the scRNA-seq data, inhibitory receptors LILRB2 and LILRB1 in psoriasis semimature DCs were potentially stimulated by a key component of MHC class I molecules (B2M) in CD161⁺ T-cells, CD4⁺ T-cells, melanocytes and KCs in stratum corneum³² (Fig E12A).

Like mature DCs, semimature DCs in psoriasis skin expressed more IL-23A and IL-36G than semimature DCs in control skin (Fig 4A and Fig E11). The number of cells expressing IL-23A or IL-36G in semimature DCs increased 9.8 times from control (1.2%, 1/85) to psoriasis skin (11.6%, 87/752). Semimature DCs expressing IL-23A or IL-36G and semimature DCs expressing IL-10 were independent – only 0.6% (1/157) of semimature DCs expressing IL-23A or IL-10 co-expressed IL-23A and IL-10.

scRNA-seq of emigrating cells from human skin identifies locations of keratinocyte transcriptome changes implicated in psoriasis pathogenesis

Our scRNA-seq data showed that the decrease of FLG expression in psoriasis epidermis occurs in stratum corneum³³ (Fig 6A). 40.7% (700/1720) of FLG expressing KCs were located in S. corneum (Fig 6B), and the number of KCs expressing FLG in S. corneum decreased 2.1 times from control (18.8%, 473/2513) to psoriasis (3.5%, 227/6418).

Our data showed that the increase of IL-36G expression in psoriasis epidermis occurs in suprabasal layers (S. corneum, S. granulosum and S. spinosum) and NFKBIZ, a transcriptional regulator of IL-36–driven gene expression³⁴ is co-expressed with IL-36G in stratum spinosum. 97.2% (3196/3288) of KCs expressing IL-36G in psoriasis skin was located in suprabasal layers, and 51.0% (292/573) of KCs expressing IL-36G in S. spinosum of psoriasis and control skin co-expressed NFKBIZ (Fig 6C).

Our data showed that CCL27 (chemotactic ligand for skin-associated memory T lymphocytes binding to CCR10)³⁵ and KRT15 (keratinocyte stem marker of quiescence)³⁶ were decreased in stratum basale (Fig 6D). CCL27 expression in KCs in S. basale decreased 10.5 times from control (17.3%, 189/1093) to psoriasis (1.7%, 7/424). KRT15 expression in KCs in S. basale decreased 11.3 times from control (48.1%, 526/1093) to psoriasis (4.2%,

18/424). When we analyzed the CCL27/CCL28-CCR10 interaction with the scRNA-seq data, CCL27-CCR10 interaction was potentially impaired in psoriasis due to decreased CCL27 expression in basal KCs^{37, 38} (Fig 6E).

Discussion

Recent scRNA-seq technology provided the opportunity to compare gene expression profiles of heterogeneous immune cells in the skin without the need to predetermine markers for various cell subsets. Co-authors of this paper pioneered single-cell transcriptome profiling of keratinocytes in lupus nephritis patients' skin^{10, 11} and single-cell transcriptome profiling of fibroblasts in atopic dermatitis patients' skin⁹, and many researches are now utilizing scRNA-seq for studying human skin to further understand inflammatory skin diseases¹²⁻¹⁷. However, scRNA-seq assessment of immune cells in human skin has been challenging with whole tissue enzyme dissociation where most cells are keratinocytes and fibroblasts, and immune cells constitute less than 5% of the isolated cells. To study immune cell subsets in the skin with scRNA-seq, skin immune cells isolated by enzyme digestion are often required to be sorted, activated and enriched before single-cell library construction, which may lead to changes in their gene expression or even function^{39, 40}.

In an attempt to overcome the limitation of enzyme dissociation for isolation of skin immune cells and to obtain single-cell transcriptome of human immune cells with minimal manipulation, we harvested emigrating cells from skin biopsy tissues after 48-hour incubation in culture medium without enzyme digestion. Our single-cell approach was established for flow cytometry analysis of cutaneous DCs^{7, 20, 41, 42} and T-cells^{43, 44}, but it has not been linked to single-cell transcriptomics to study heterogeneous immune cells in human skin.

With our new single-cell approach, we found cutaneous T17 cell subsets with highly differing transcriptomes depending on IL-17A vs. IL-17F expression and IFN γ vs. IL-10 expression: 1) IL-17A⁺ IFN γ ⁺ T17 cells, 2) IL-17A⁺ IFN γ ⁻ T17 cells, 3) IL-17A⁺ IL-17F⁺ T17 cells, 4) IL-17F⁺ IL-10⁻ T17 cells and 5) IL-17F⁺ IL-10⁺ T17 cells (Fig 3B). The T17 cell subset that most conforms with current pathogenic subsets^{6, 22-24} is the IL-17A⁺ IFN γ ⁺ population, synthesizing high levels of TNF, IL-26 and IL-36G and mostly within CD8⁺ T-cells that co-express cytotoxic markers (Tc17 T-cells).

In contrast, the largest subset of IL-17-producing T-cells isolated from psoriasis lesions are those that are IL-17F⁺ IL-10⁻. This population (presumptive IL-17F/F producers) constitutes 53% of T17 T-cells and is about 5-fold more frequent than cells co-producing IL-17A and IFN γ . High expression of cytokines such as IL-1B, CSF-2, LTA, IL-24 and IL-34 likely identifies a different inflammatory potential from cells exposing IL-17A and IFN γ . Interestingly, this subset has the highest expression of the IL-23 receptor, so it might be the most strongly affected by therapeutic IL-23 antagonists. Perhaps this subset also stems from inflammatory conversion of Tregs, as FoxP3 has the highest expression among all T17 cells. While a recent report found some human blood T-cells synthesized only IL-17F after culture in polarizing conditions⁴⁵, we believe this is the first report of a unique and

sizable IL-17F⁺ T-cell population in the context of an organ affected by an IL-17-driven autoimmune condition.

The population that best fits the description of non-pathogenic T17 cells^{6, 23–26, 30, 46} is the IL-17F⁺ IL-10⁺ population, but this population constitutes only 4% of overall T17 cells. IL-17F⁺ IL-10⁺ T17 cells expressed low levels of FoxP3 and high levels of AHR and MAF. Since Aryl hydrocarbon Receptor (AHR) interacts with c-Maf (MAF) to promote the differentiation of Tr1 cells⁴⁷, cutaneous IL-17F⁺ T17 cells may have plasticity to differentiate into Tr1 cells rather than Tregs.

We found distinct DC subsets in human skin, including mature DCs and semimature DCs (Fig 4). High expression of MHC and co-stimulatory molecules by mature DCs in psoriasis may be the reason for the highly T-cell stimulatory nature of these cells⁸. Semimature DCs are regulatory DCs expressing high levels of IL-10, BDCA-3 and LILRB2. Single-cell transcriptome of semimature DCs is consistent with gene expression profiles of human skin resident BDCA3⁺ regulatory DCs described by Chu, C.C. et al.⁷.

Since mature DCs in psoriasis skin express high levels of IL-23A & IL-36G and low levels of KYNU compared to mature DCs in control skin, mature DCs in psoriasis skin may lose tolerogenicity that is maintained by IDO1-KYNU-AHR loop^{31, 48, 49} in normal condition and express IL-23A and IL-36G (Fig E11). A subset of regulatory semimature DCs in psoriasis skin expresses IL-23A and IL-36G; dysfunctional regulatory DCs in psoriasis skin may contribute to the immune tolerance impairment implicated in psoriasis.

Our approach has some limitations⁵⁰: 1) 48 hour in vitro culture and cell emigration may change the status of cells and cause the change of gene expression, 2) we observed that the number of isolated skin cells and the proportion of different cell types were not consistent between samples, 3) heterogeneous skin cells with variable sizes were not reliably counted by automated cell counters. As a result, the number of cells submitted to a microfluidic platform was not consistent between samples and sequencing depth per cells became variable (Table E1), 4) some skin resident immune cells may not be migratory and might require other isolation techniques.

With novel findings of distinct immune cell transcriptome changes in psoriasis, we propose that single-cell transcriptomics applied to emigrating cells from skin provide an innovative study platform to compare gene expression profiles of heterogeneous immune cells in human skin. This method can also readily be extended to study other inflammatory skin diseases or modulation of leukocyte transcripts by immune-targeted treatments.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Disclosure of potential conflict of interest:

J.K. has received research funds from AbbVie. E.G. has received research funds from AbbVie, Celgene, Eli Lilly, Janssen, Medimmune/Astra Zeneca, Novartis, Pfizer, Regeneron, Vitae, Glenmark, Galderma, Asana, Innovaderm, Dermira, and UCB and is also a consultant for Sanofi Aventis, Regeneron, Stiefel/GlaxoSmithKline, MedImmune, Celgene, Anacor, AnaptysBio, Dermira, Galderma, Glenmark, Novartis, Pfizer, Vitae, Leo Pharma, AbbVie, Eli Lilly, Kyowa, Mitsubishi Tanabe, Asana Biosciences, and Promius. J.G.K. has received research support from Pfizer, Amgen, Janssen, Lilly, Merck, Novartis, Kadmon, Dermira, Boehringer, Innovaderm, Kyowa, BMS, Serono, BiogenIdec, Delenex, AbbVie, Sanofi, Baxter, Paraxel, Xenoport, and Kineta. The rest of the authors declare that they have no relevant conflict of interest.

Abbreviations:

AHR	Aryl hydrocarbon Receptor
CCL	CC chemokine ligand
DC	Dendritic cell
FLG	Filaggrin
KC	Keratinocyte
MFI	Median fluorescence intensity
NK	Natural Killer
S.	Stratum
scRNA-seq	Single-cell RNA sequencing
T17	Type 17 T-cell
Tc17	Cytotoxic type 17 T-cell
Th17	T helper 17 T-cell
Treg	Regulatory T-cell
Tr1	Type 1 regulatory T-cell
UMAP	Uniform Manifold Approximation and Projection

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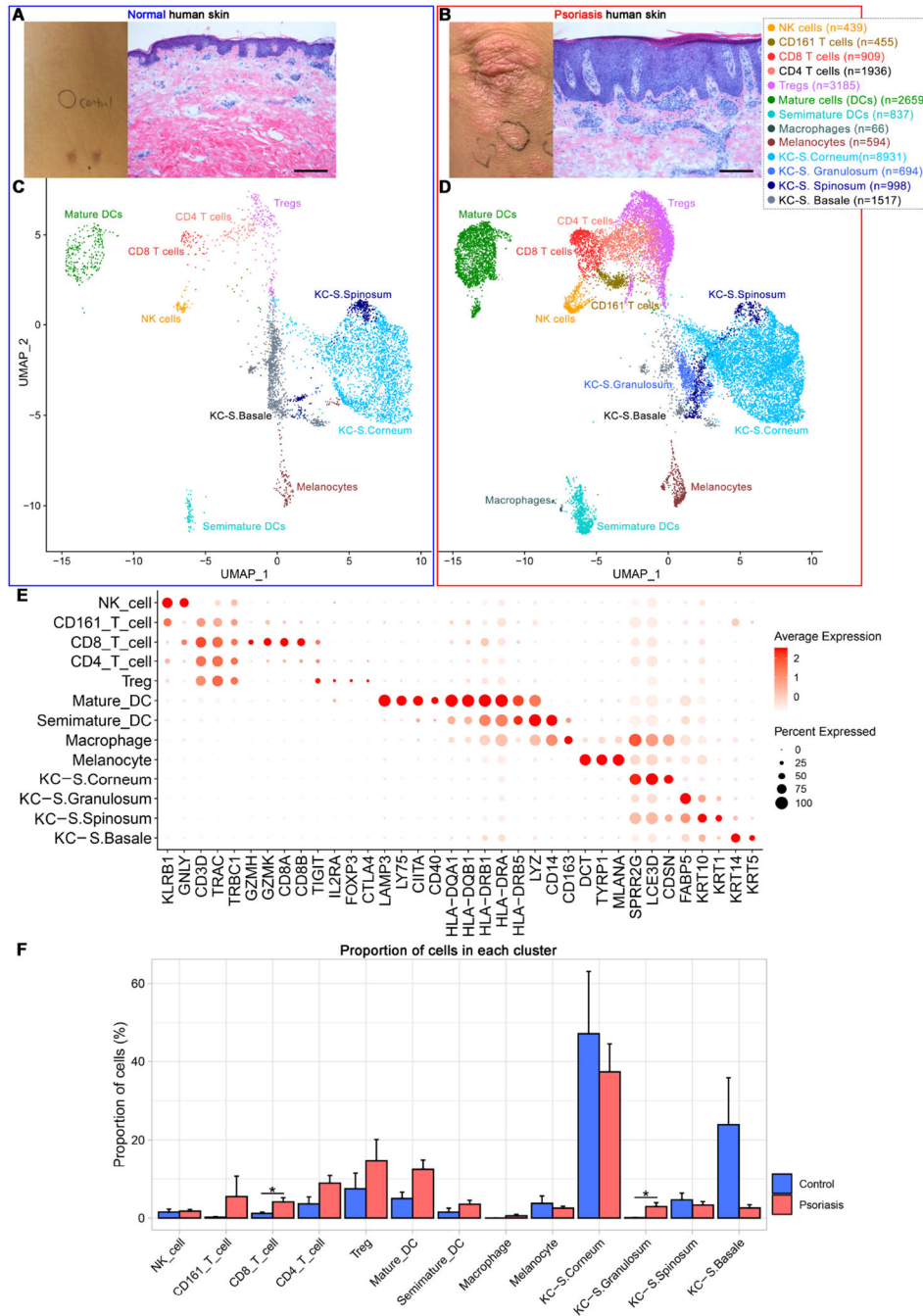


Figure 1. Single-cell transcriptomic profiling of leukocytes and keratinocytes in human psoriasis and control skin.

Clinical & microscopic figures of control (A) and psoriasis skin (B) and their leukocyte & keratinocyte scRNA-seq data presented in the Uniform Manifold Approximation and Projection plot (C and D). (E) Dot plot displaying expression levels of cluster-defining genes. (F) Cell composition of individual samples. Treg, regulatory T-cell; DCs, dendritic cells; KC, keratinocytes, S, Stratum. Scale bar in (A) and (B) = 200 μm, Proportion of cells

in **(F)** = average number of cells in cluster within individual sample / total number of cells within individual sample x 100 (%), Error bar in **(F)** = Standard Error of Mean.

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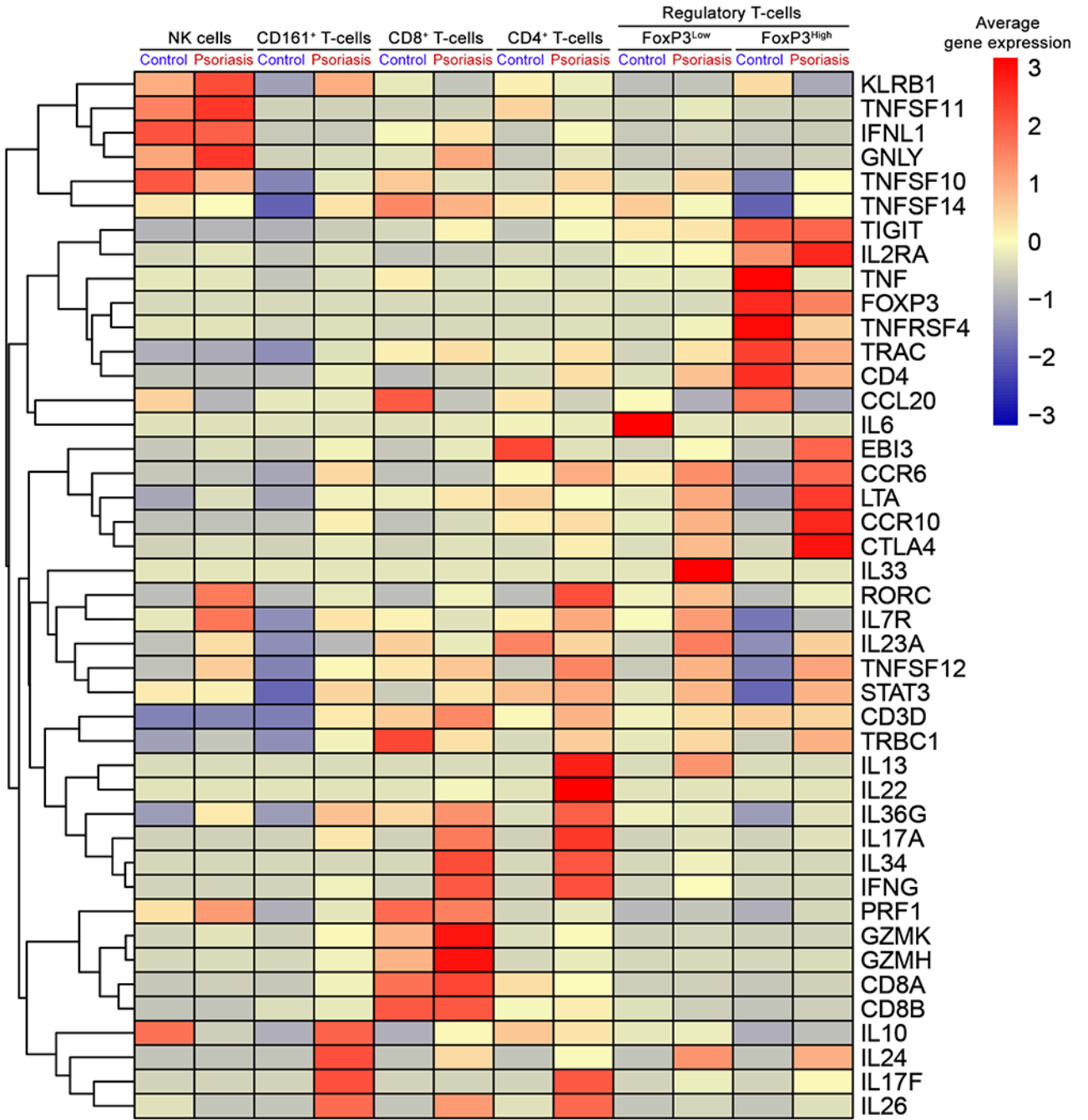


Figure 2. The average gene expression within clusters of NK cells and T-cell subsets. Heatmap of scRNA-seq analysis illustrates the average gene expression within clusters of NK cell and T-cell subsets, split by psoriasis and control. Regulatory T-cell cluster is divided into cells with FoxP3 expression high (FoxP3^{High}) and cells with FoxP3 expression low (FoxP3^{Low}) using a cut-off value of 1. Genes with similar expression patterns are linked by a complete linkage method.

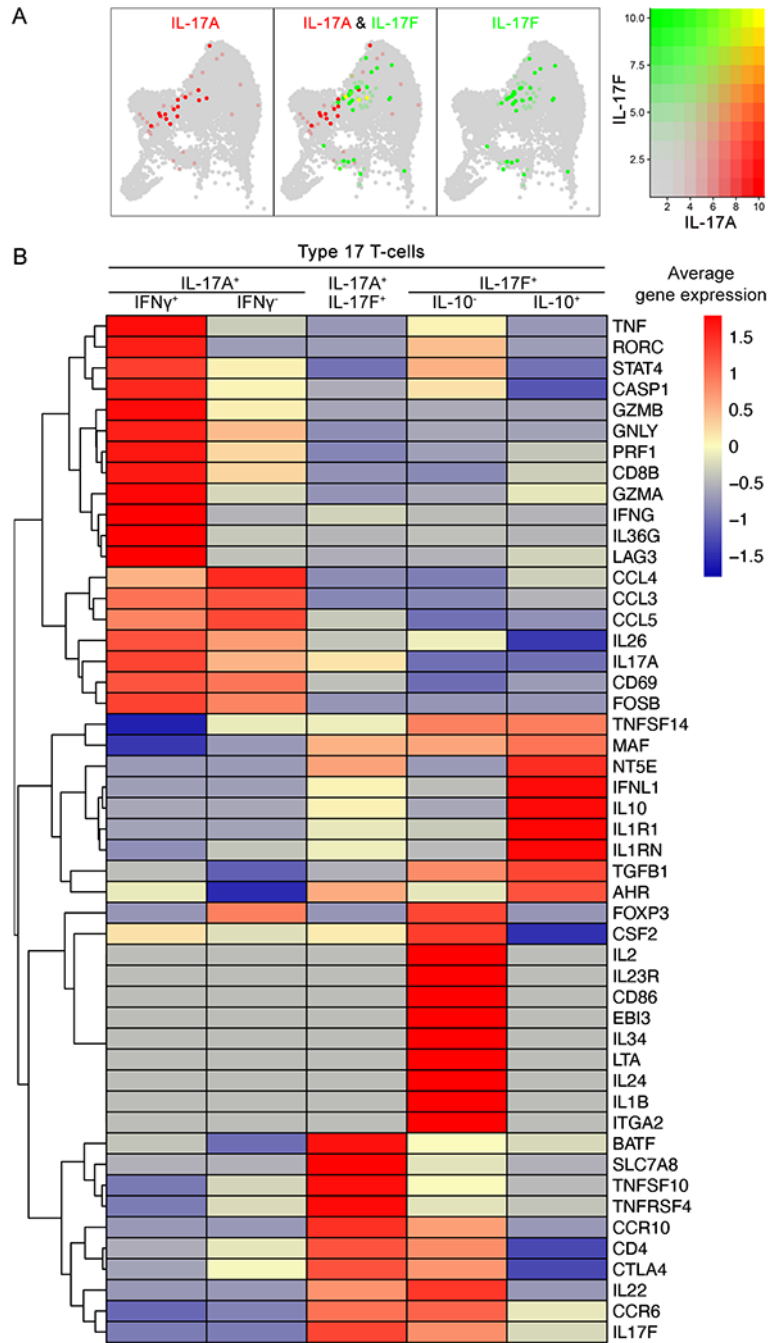


Figure 3. The average gene expression of cutaneous Type 17 T-cell (T17) cell subsets. (A) IL-17A (red), IL-17F (green) and IL-17A/IL17F co-expression (yellow) within T-cell subset clusters visualized in low-dimensional space. (B) Heatmap of scRNA-seq analysis illustrates the average gene expression of T17 cell subsets. Genes with similar expression patterns are linked by a complete linkage method.

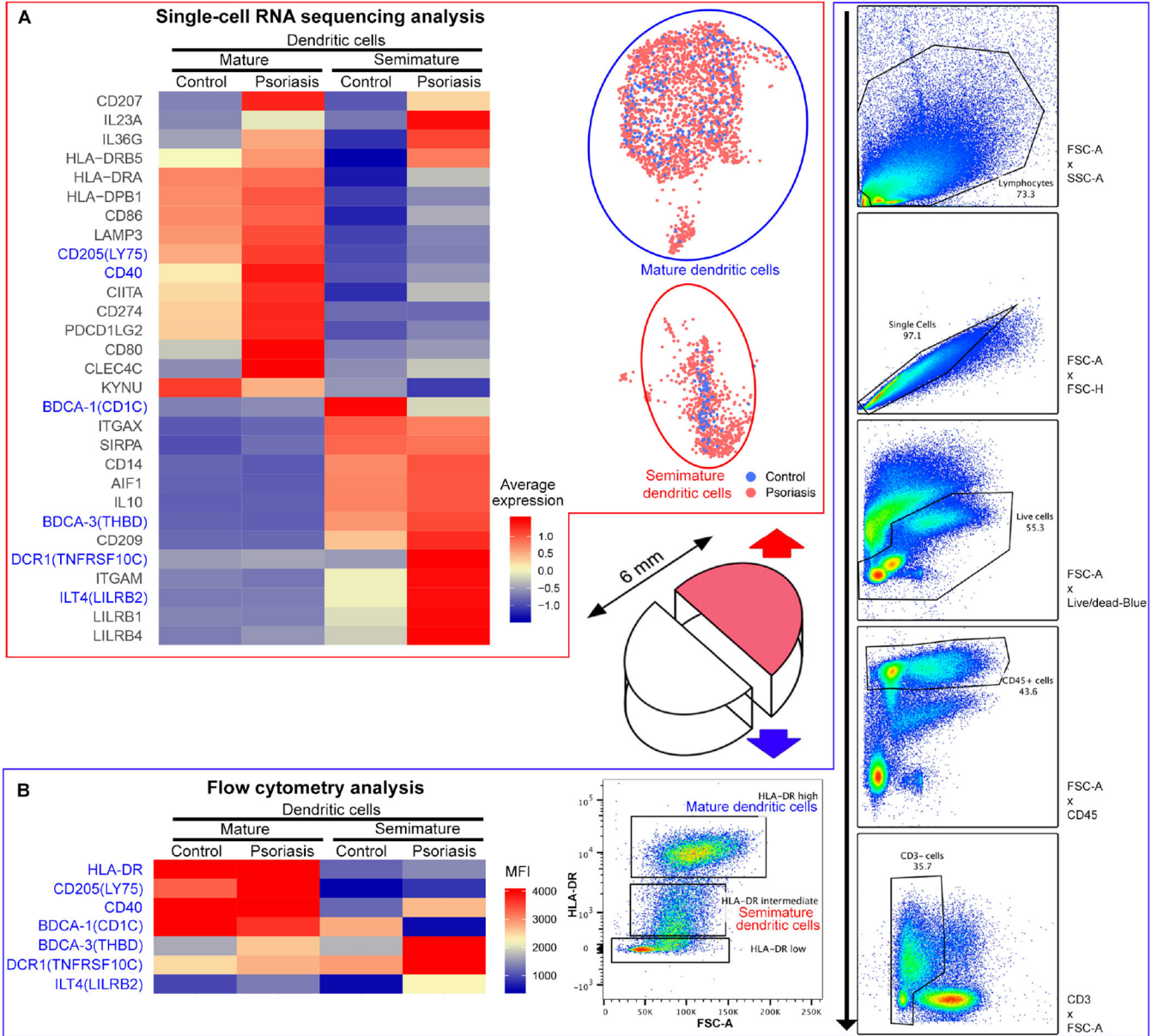


Figure 4. Simultaneous scRNA-seq and flow cytometry analyses define transcriptomic profiles of mature vs. semimature dendritic cells.

(A) Heatmap of scRNA-seq analysis illustrates the average gene expression within mature vs. semimature DC clusters, split by psoriasis and control. (B) Heatmap of flow cytometry analysis illustrates Median Fluorescence Intensity (MFI) of HLA-DR^{High} (mature) or HLA-DR^{Low} (semimature) CD45⁺ CD3⁻ HLA-DR⁺ dendritic cells in control and psoriasis skin. Consistent findings between scRNA-seq and flow cytometry analyses are marked in blue.

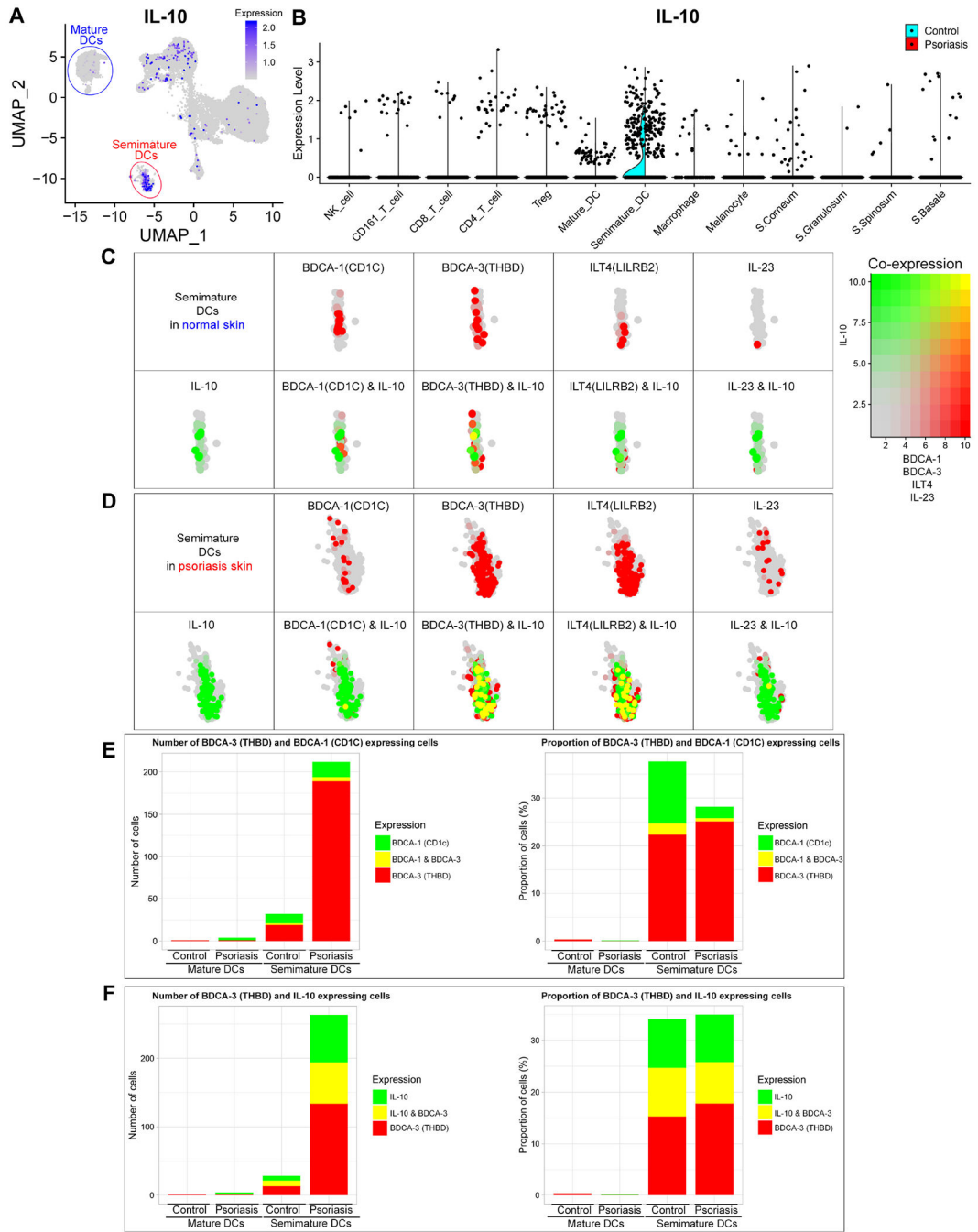


Figure 5. IL-10, BDCA-1 (CD1C), BDCA-3 (THBD), LILRB2 (ILT4) and IL-23 expression in mature dendritic cells (DCs) and semimature DCs.

IL-10 expression in total scRNA-seq data is visualized in low-dimensional space (A) and violin plot of each immune cell cluster (B). Co-expression of IL-10 & BDCA-1, BDCA-3, LILRB2 or IL-23 in normal skin semimature DCs (C) and psoriasis skin semimature DCs (D) is visualized in low-dimensional space. Cells with expression of BDCA-3 & BDCA-1 (E) and BDCA-3 & IL-10 (F) within each cluster of mature and semimature DCs in normal and psoriasis skin are quantified by number of cells and proportion of cells. Proportion of

cells = number of target gene expressing cells within cluster / total number of cells within cluster x 100 (%).

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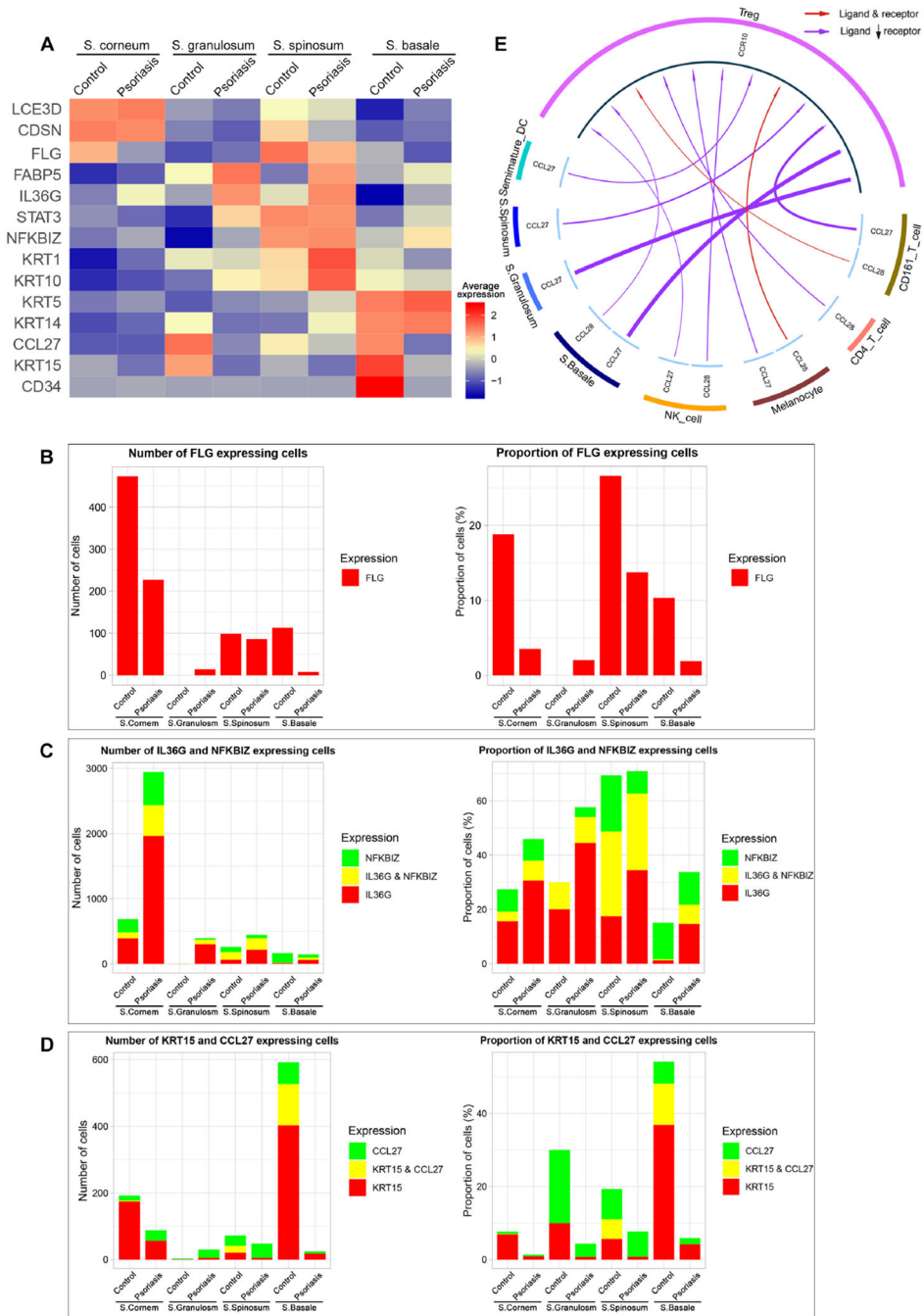


Figure 6. Gene expression within keratinocyte clusters and CCL27 (keratinocyte)–CCR10 (Treg) interactions.

(A) Heatmap of scRNA-seq analysis illustrates the average gene expression within clusters of keratinocytes (KCs) in Stratum (S.) corneum, S. granulosum, S. spinosum and S. basale, split by psoriasis and control. Cells with expression of FLG (B), IL-36G & NFKBIZ (C) and KRT15 & CCL27 (D) within each layer of KCs in control and psoriasis epidermis are quantified by number of cells and proportion of cells. Proportion of cells = number of target gene expressing cells within cluster / total number of cells within cluster x 100 (%).

(E) CCL27/CCL28-CCR10 interaction in psoriasis compared to control highlights decreased CCL27 in basal keratinocytes interacting with CCR10 in Tregs.

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