



# IL-23 induces CLEC5A<sup>+</sup> IL-17A<sup>+</sup> neutrophils and elicit skin inflammation associated with psoriatic arthritis

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## ABSTRACT

IL-23-activation of IL-17 producing T cells is involved in many rheumatic diseases. Herein, we investigate the role of IL-23 in the activation of myeloid cell subsets that contribute to skin inflammation in mice and man. IL-23 gene transfer in WT, IL-23R<sup>GFP</sup> reporter mice and subsequent analysis with spectral cytometry show that IL-23 regulates early innate immune events by inducing the expansion of a myeloid MDL1<sup>+</sup>CD11b<sup>+</sup>Ly6G<sup>+</sup> population that dictates epidermal hyperplasia, acanthosis, and parakeratosis; hallmark pathologic features of psoriasis. Genetic ablation of MDL-1, a major PU.1 transcriptional target during myeloid differentiation exclusively expressed in myeloid cells, completely prevents IL-23-pathology. Moreover, we show that IL-23-induced myeloid subsets are also capable of producing IL-17A and IL-23R<sup>+</sup>MDL1<sup>+</sup> cells are present in the involved skin of psoriasis patients and gene expression correlations between IL-23 and MDL-1 have been validated in multiple patient cohorts. Collectively, our data demonstrate a novel role of IL-23 in MDL-1-myelopoiesis that is responsible for skin inflammation and related pathologies. Our data open a new avenue of investigations regarding the role of IL-23 in the activation of myeloid immunoreceptors and their role in autoimmunity.

## 1. Introduction

IL-23 is a pleiotropic cytokine which is involved in many autoimmune diseases including psoriasis and psoriatic arthritis [1,2]. IL-23 is critical for a plethora of immune responses, including the differentiation, survival, and expansion of Th17,  $\gamma\delta$  T cells and neutrophils [3,4]. Although the effects of IL-23 in the lymphoid cells have been extensively studied in relation to IL-17A production, less is known regarding the effects of IL-23 on myeloid cells. Despite multiple studies demonstrating the importance of IL-23 and its effector signaling in myelopoiesis and the recruitment and activation of neutrophils, the exact cellular and

molecular pathways that govern pathogenesis remain elusive [4–8].

We have previously demonstrated that IL-23 induces myelopoiesis in the bone marrow and induces Myeloid DAP12-associated lectin 1 (MDL-1) which is associated with osteoclast activation and arthritis [9–11]. MDL-1 also known as CLEC5A is a C-type lectin, (herein referred to as MDL-1) is exclusively expressed on myeloid cells and is a major PU.1 transcriptional target during myeloid differentiation regulating a number of myeloid-dependent immune and inflammatory responses including differentiation, activation and cell recruitment during inflammation [9,12–15]. Flaviviruses and *Listeria monocytogenes* stimulate MDL-1 to induce proinflammatory cytokine production [16,17],

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canonical inflammasome activation and neutrophil extracellular trap (NET) formation [18,19], while MDL-1 deficiency impairs the development of intestinal  $\gamma\delta$  T cells after *L. monocytogenes* infection [19]. MDL-1 is a key regulator of synovial injury and bone erosion during autoimmune joint inflammation [9] and dengue virus infection [20]. Furthermore, MDL-1<sup>+</sup> immature myeloid cells are responsive for ConA-induced liver inflammation [21]. All these observations suggest that MDL-1 plays a critical role in septic and aseptic inflammatory reactions.

Herein, we show that genetic ablation of MDL-1 prevents IL-23-induced skin inflammation demonstrating the importance of myelopoiesis in IL-23-induced pathology. Our data show that MDL-1 is critical for the development and maturation of promyelocytes to IL-17A<sup>+</sup> neutrophils in the skin that regulate skin pathology. Collectively our data demonstrate MDL-1 as a transcriptional target in IL-23-induced skin pathology that can be exploited for therapeutic intervention.

## 2. Material and methods

### 2.1. Study design

C57BL/6 mice were purchased from Jackson Laboratories. IL-23R<sup>GFP+/+</sup> and *Mdl-1*<sup>-/-</sup> mice were obtained from Dr. Kuchroo (Harvard Medical School) and Dr. Hsieh (National Yang-Ming University, Taiwan), respectively as previously described [16,22]. For each experiment, at least two experimental replicates were performed. Mice were selected and assigned to their groups based on their body weights at the time each experiment started. Sex- and age-matched mice (8–12 weeks) were used for all experiments under specific pathogen-free conditions.

### 2.2. Reagents

DyLight 650-Phalloidin was purchased from Thermo Fisher Scientific, and IL-23 ELISA kit was purchased from eBioscience. Monoclonal antibodies of anti-Ly6G (1A8), MDL-1 (226402) and anti- $\gamma\delta$ TCR (GL3) were purchased from R&D Systems, CD11b (M1/70) from ebioscience, and GFP conjugated FITC (FM264G), CD3e (145-2C1), CXCR3R1 (SA011F11), B220 (RA3-6B2), Ter119 (TER-119), CSF1R (AFS98), Ly6C (HK1.4), CD61 (2C9.G2), CD45 (30-F11) Ly6G (1A8), CD11b (M1/70) and IL-17A (1A8) from BioLegend.

### 2.3. Purification of GFP and IL-23 minicircle (MC) DNA and hydrodynamic gene delivery

Minicircle-RSV.Flag.mIL23.elasti.bpA or RSV.eGFP.bpA (IL-23MC) was produced as previously described, and Episomal DNA minicircle (MC) were prepared using EndoFree Megaprep plasmid purification kit (Qiagen) [11]. Fifteen micrograms of IL-23 or GFP MC DNA was injected hydrodynamically via tail vein delivery. We analyzed serum 1-day post gene transfer with an IL-23 ELISA, and collected bone marrow (BM), dorsal skin and ears for analysis on indicated days post gene transfer.

### 2.4. Spectral flow cytometry and high-dimensional analysis

Murine dorsal skin was minced and incubated with 1 U/ml of Dispase for 2 h at 37 °C, followed by addition of collagenase D for 30 min. BM cells were flushed out with PBS. Red blood cells were lysed with BD Pharm Lyse (BD Biosciences). Cells were pre-incubated with anti-CD16/32 (BD Biosciences) and then incubated with appropriate antibodies and analyzed on a CytoFLEX LX (Beckman Coulter). The data were analyzed using FlowJo (BD Biosciences). AccuCheck counting beads (Life Technologies) were used to determine absolute cell number. BM cells were analyzed on Cytex Aurora cytometer and among B220<sup>+</sup>CD3<sup>+</sup>Ter119<sup>-</sup>CSF1R<sup>+</sup> cells, clustering algorithm was conducted by FlowSOM based on the marker expression of CD11b, Ly6G, Ly6C, CD45, CX3CR1, MDL1 and CD61. As this method is highly sensitive and

vulnerable to batch effect, FlowSOM were conducted on single experiment using to avoid confounding bias [23]. The FlowSOM clusters was also displayed on top of the t-SNE axes to visualize each cluster.

### 2.5. RNA isolation and real-time PCR

RNA was isolated from mouse tissues using the RNeasy kit (Qiagen) including a DNase I digest step. Quality of RNA was analyzed with a Nanodrop spectrophotometer (Thermo Fisher Scientific). cDNA was prepared using iScript cDNA Synthesis Kit (Bio-Rad). qRT-PCR was performed using iTaq Universal SYBR Green Supermix (Bio-Rad) as previously described [24]. Relative expression of target genes was performed using the 2<sup>- $\Delta\Delta$ CT</sup> method and normalized with internal GAPDH control. Relative expression of target genes was performed using the 2<sup>- $\Delta\Delta$ CT</sup> method and normalized with internal GAPDH control as previously described [11].

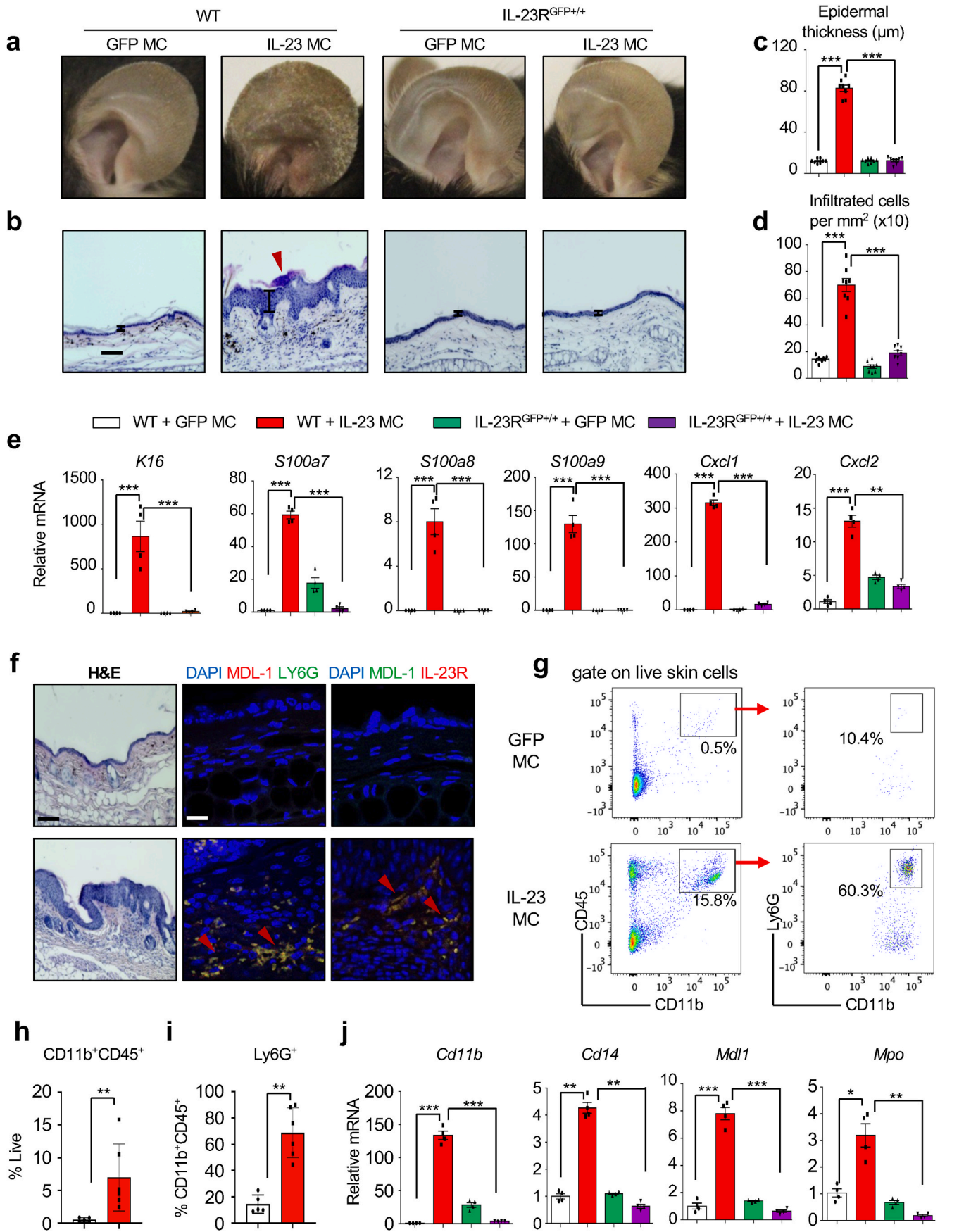
### 2.6. RNAseq and RNAseq meta-analysis

RNA was isolated from murine ears of WT or *Mdl-1*<sup>-/-</sup> mice injected with GFP MC either IL-23 MC. The 3'Tag RNA-Seq run was performed on an Illumina HiSeq 4000 and generated an average of 600,000 reads per sample. RNA-Seq reads for the 9 individual samples including three groups: control (WT + GFP MC), treatment (WT + IL-23 MC), and mutant (*Mdl-1*<sup>-/-</sup> + IL-23 MC) and three replicates each) were independently aligned to the mouse genome (ref. ID: GRCh38.p6) using the STAR v2.7.0a alignment software with the corresponding Ensembl reference genome. The read results from alignment are shown in Supplemental Table 1. The featureCounts package and edgeR package were used to count the mapped reads and for subsequent differential expression analysis. After excluding genes with low expression (less than three mapped reads), the data were normalized using the Trimmed Means of M values (TMM) method. A negative binomial distribution was used to model the variance of the read counts per gene. DEGs are defined by a fold change >2.0 or <0.5 and q-value (FDR-adjusted p-value) < 0.05. A gene-set enrichment analysis using gProfiler, in which we identified and examined the top 10 gene ontology processes for both upregulated and downregulated DEGs separately. A protein-protein interaction network was generated in which genes from interested gene groups were shown relative to detected gene clusters. The KEGG database was used in conjunction with R's GOChord package to identify relevant pathways activated by genes in our interested gene groups. The RNA-Seq data is deposited in the NCBI's Gene Expression Omnibus (GEO) database and are accessible through GEO Series under the accession number GSE133830.

For human RNA-Seq meta-analysis, RNA-Seq FASTQ files of psoriasis lesional skin were downloaded from the NCBI Sequence Read Archive (<http://www.ncbi.nlm.nih.gov/Traces/sra>). Four total datasets (Supplemental Table 5) were used. All statistical analysis was done using R software. Correlation analyses of gene expressions were performed on read counts of each identified gene normalized with DESeq2 package. Values were subsequently log transformed and winsorized. Spearman's correlation coefficients ( $r_s$ ) were calculated in R using the cor.test function, which was also used to estimate P values of the correlations by algorithm AS 89. Meta-analyses were conducted to assess findings across the multiple datasets using R package "metafor". A weighted random-effects model was used to estimate a summary effect size. Restricted maximum-likelihood estimator was selected to estimate between-study variance. Weighted estimation with inverse-variance weights was used to fit the model.

### 2.7. H&E and immunohistochemistry

Murine skin and ears were fixed in 10 % formalin in PBS and paraffin embedded for sectioning (6  $\mu$ m). Tissue sections were stained with hematoxylin and eosin Y (Sigma). Photos were visualized and analyzed by



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**Fig. 1.** IL-23-induced skin pathology depends on IL-23R signaling and correlates with increased number of epidermal MDL-1<sup>+</sup> cells. (a) Photographs of murine ears 21 days post IL-23 MC gene transfer showing the development of silvery white scales in WT mice compared to GFP MC and/or IL-23R<sup>GFP+/+</sup>. (b) H&E staining of murine ears showing epidermal hyperplasia and number of infiltrated cells. Arrow indicates neutrophil exudates (Munro's microabscess). (Images are representative of three independent experiments and 9–11 mice per each group. Scale bars, 100  $\mu$ m). (c) Quantification of epidermal thickness ( $\mu$ m) and (d) infiltrated cell number. (e) Gene expression analysis of inflammatory markers showing an elevation of *K16*, *S100a7*, *S100a8*, *S100a9*, *Cxcl-1*, and *Cxcl-2* in the ears of IL-23 MC gene transfer WT mice and/or IL-23R<sup>GFP+/+</sup> mice compared to GFP MC (control). (f) H&E and immunofluorescence images of staining with DAPI (blue), MDL-1-PE (red), LY6G-FITC (green) and DAPI (blue), MDL-1-FITC (green), IL-23R-PE (red) showing an increase of epidermal MDL-1<sup>+</sup>IL-23R<sup>+</sup> neutrophils (orange) in the ears of IL-23 MC gene transfer mice compared to GFP MC (control) mice. Scale bars, 100  $\mu$ m. (g) Flow cytometric analysis of murine dorsal skin post IL-23 MC gene transfer, showing an expansion of (h) CD11b<sup>+</sup>CD45<sup>+</sup> cells and (i) CD11b<sup>+</sup>CD45<sup>+</sup>Ly6G<sup>+</sup> cells compared to GFP MC (control) mice. (j) Gene expression analysis of murine ear post IL-23 gene transfer showing an elevation of *Cd11b*, *Cd14*, *Mdl-1*, and *Mpo* compared to GFP MC (control) and/or IL-23R<sup>GFP+/+</sup> mice. Data represent mean  $\pm$  SEM of three independent experiments. \**P* < 0.05; \*\**P* < 0.01; \*\*\**P* < 0.001 by Mann-Whitney.

Olympus BX61 microscope and BZ-II Analyzer software. Analysis and quantification were performed using ImageJ software. Histology sections were stained for immunofluorescence as previously described [25] and visualized using a confocal microscope (Nikon C1).

Skin biopsies were fixed in 10 % formalin for 48-h and paraffin embedded for sectioning (4  $\mu$ m) for either: 1) Hematoxylin and Eosin (H&E) or 2) Chloroacetate Esterase. The stained sections were digitized using a slide imaging system (Olympus VS120). For immunofluorescence, sections were blocked and stained with goat anti-IL-17 (AF-317-NA, R&D systems), mouse anti-CD66b (NB100-77808SS, Novus Biologicals), and rabbit anti-MRP8 (SA100A8), (EPR3554, Ab92331, Abcam) antibodies. Primary antibodies were detected with AF568 donkey anti-goat IgG (A11057, ThermoFisher), AF488 donkey anti-rabbit and AF647 donkey anti-mouse (715-606-150, Jackson ImmunoResearch Laboratories). Representative images were taken with a Zeiss Axio Microscope and recorded with AxioCam MRM.

## 2.8. LPS stimulation of bone marrow myeloid cells

BM myeloid cells were magnetically sorted twice using CD11b MicroBeads (130-126-725, Miltenyi Biotec). Sorted myeloid cells were stimulated with vehicle or 100 ng/ml LPS for 4 h and RNA were extracted for subsequent qRT-PCR as previously described [26].

## 2.9. Patient selection

Two healthy volunteers, six patients with PsA and nine patients with PsO who were previously untreated and sex-matched were recruited in this study. The clinical characteristic of patients with PSA are as follows: (Mean  $\pm$  SEM; Age = 54.2  $\pm$  7.17 years; DAPSA = 45.3  $\pm$  9.39; PASI = 7.92  $\pm$  1.77). The clinical characteristics of patients with PsO are as follows: (Mean  $\pm$  SEM; Age = 58.0  $\pm$  6.42 years; PASI = 2.72  $\pm$  0.73).

## 2.10. Statistical analysis

Statistical differences were analyzed by Mann–Whitney test. All results are representative of at least 3 independent experiments, unless otherwise stated. Statistically significant differences were considered as *P* < 0.05 (\**P* < 0.05, \*\**P* < 0.01, \*\*\**P* < 0.001). Data represent mean  $\pm$  SEM (standard error of the mean) of three independent experiments.

## 2.11. Study approval

All animal protocols (protocol #20010 and #20845) were approved by Institutional Animal Care and Use Committee, BIDMC/Harvard University. Human studies were reviewed and approved by BIDMC and University of Rochester, Institutional Review Boards. Written informed consent was received from each participant prior to collection of psoriatic skin biopsies.

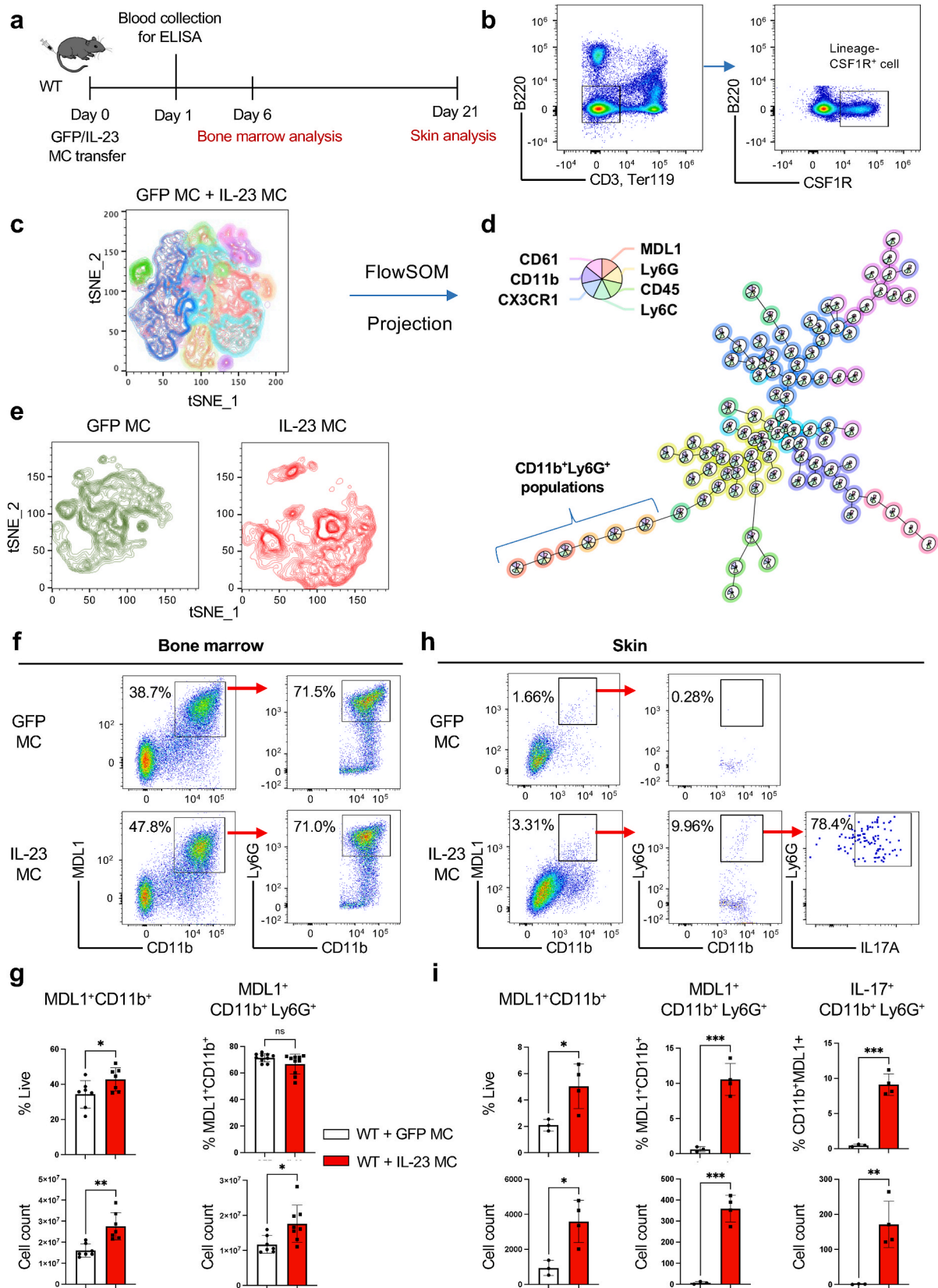
## 3. Results

### 3.1. IL-23-mediated skin pathology induces IL-23r<sup>+</sup>MDL-1<sup>+</sup> myeloid cells

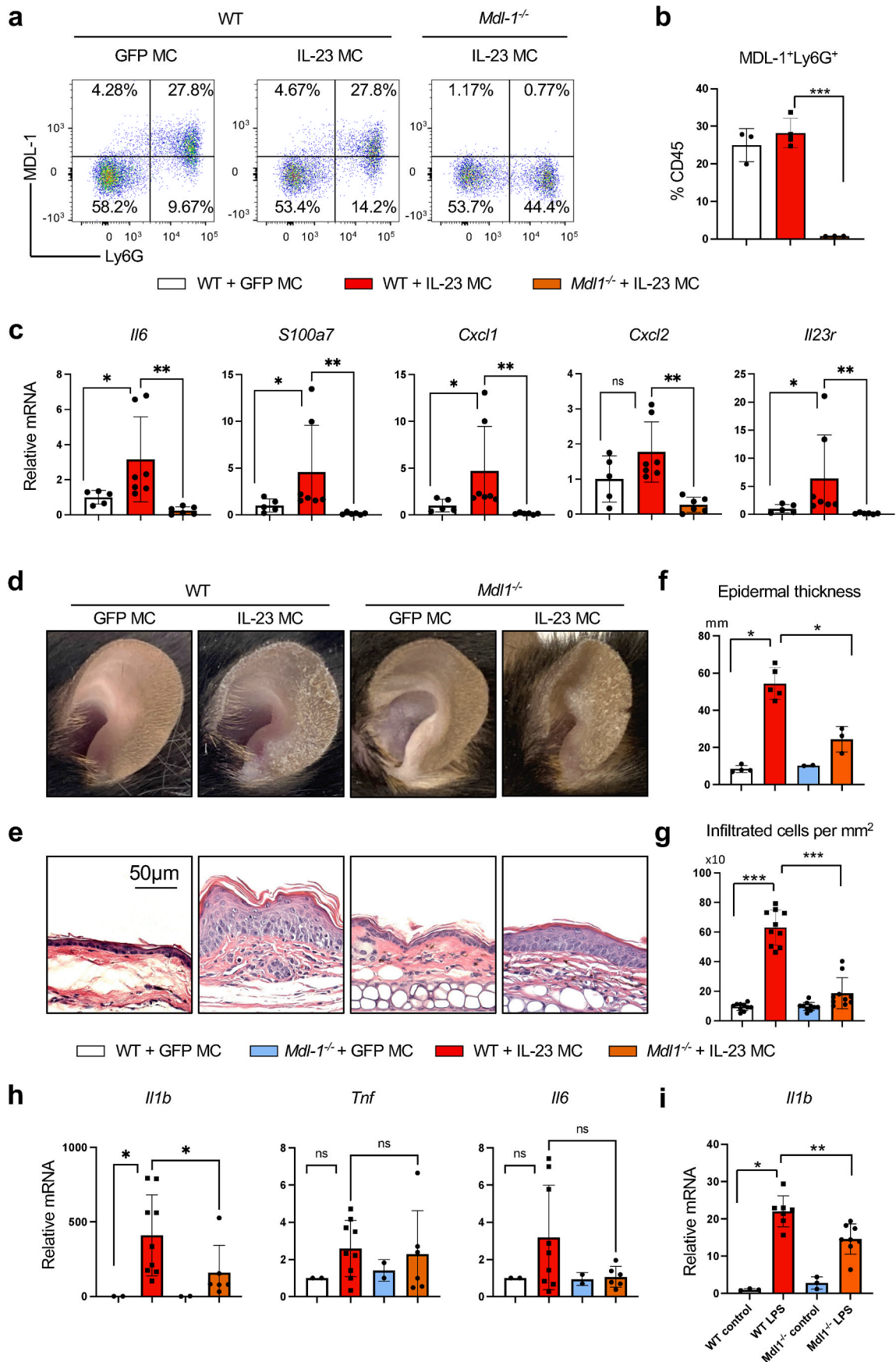
In order to investigate the effects of IL-23 in the skin we performed IL-23 gene transfer in C57BL/6 mice. (Supplemental Fig. 1a and b). IL-23 gene transfer induced erythema with silvery white scales in ears 21 days post IL-23 MC injection (Fig. 1a). Histological examination revealed a significant epidermal hyperplasia, and thickness of epidermis (Fig. 1b and c), accompanied by an increased inflammatory infiltrate with a distinct formation of neutrophil exudates (Munro's microabscesses) (Fig. 1b–d). Similar pathology to the ears was observed in the murine dorsal skin (data not shown) whereas neither control, (GFP MC), nor IL-23R<sup>GFP+/+</sup> mice (IL-23R<sup>GFP+/+</sup> is a non-functional IL-23R) showed any pathology (Fig. 1a–d). The observed pathology correlated with an elevated pattern in genes associated with keratinocyte proliferation and skin inflammation including *K16*, *S100a7*, *S100a8*, *S100a9*, *Cxcl-1*, *Cxcl-2*, *Tnf*, *Il-17*, *Il-17ra*, *Il-23r*, and *Il-22* in ears, compared with control (GFP MC) and IL-23R<sup>GFP+/+</sup> mice (Fig. 1e and Supplemental Fig. 1c). The elevated pattern in genes associated with keratinocyte proliferation and skin inflammation was also observed in the murine dorsal skin compared with control (GFP MC) and IL-23R<sup>GFP+/+</sup> mice (Supplemental Fig. 1d). Furthermore, immunofluorescence staining demonstrated the presence of epidermal MDL-1<sup>+</sup>LY6G<sup>+</sup> neutrophils and MDL1<sup>+</sup>IL-23R<sup>+</sup> cells in IL-23 MC gene transfer mice, whereas this population was undetected in GFP MC controls (Fig. 1f). Consistent with the histological phenotype, we found an expansion of the CD11b<sup>+</sup>CD45<sup>+</sup> myeloid cell population (Mean  $\pm$  SEM; WT + GFP MC = 0.54 %  $\pm$  0.14; WT + IL-23 MC = 7.01 %  $\pm$  2.09) that included CD11b<sup>+</sup>Ly6G<sup>+</sup> neutrophils (WT + GFP MC = 14.47 %  $\pm$  3.11 and WT + IL-23 MC = 68.67 %  $\pm$  7.79), by flow cytometric analysis in the skin post IL-23 gene transfer (Fig. 1g–i). In keeping with these observations, an elevation of myeloid cells markers *Cd11b*, *Cd14*, *Mdl-1*, and *Mpo* confirmed the myeloid expansion in the skin of IL-23 MC gene transfer mice compared to the control (GFP MC) and/or IL-23R<sup>GFP+/+</sup> mice (Fig. 1j).

### 3.2. IL-23 induces the expansion of MDL-1<sup>+</sup> cell subsets in the bone marrow and skin

We and others have previously demonstrated that IL-23 induces myelopoiesis in the bone marrow which leads to T-cell independent IL-17A expression and skin inflammation, however the exact mechanisms remain elusive [11,25,27,28]. To identify the effect of IL-23 in myeloid cells, we performed spectral cytometry on bone marrow cells isolated 6 days post IL-23 gene transfer using an extensive panel of myeloid specific antibodies (Fig. 2a). We focused our analysis on myeloid precursors by gating on the B220, CD3, Ter119 negative and c-fms positive population (Fig. 2b). Among this myeloid population, we integrated the data of GFP and IL-23 gene transfer and conducted an unsupervised clustering algorithm with a combination of tSNE and FlowSOM (flow cytometry data analysis using self-organizing maps) showing 10 distinct populations (Fig. 2c and d). The clustering was based on the expression level of CD11b, Ly6G, Ly6C, CD45, CX3CR1 (C-X3-C Motif Chemokine Receptor 1), MDL1 and CD61 ( $\beta$ 3-integrin) (Fig. 2d). Our data



**Fig. 2.** IL-23 induces the expansion of MDL-1<sup>+</sup> cell subsets in the bone marrow that migrate to the skin. (a) Schematic presentation of GFP/IL-23 gene transfer model in WT mice. (b) Representative Fig. of integrated flow cytometry data showing the gating strategy of lineage<sup>-</sup>CSF1R<sup>+</sup> cell used for unsupervised clustering with (c) tSNE plot and (d) FlowSOM based on seven surface molecules, showing distinct clusters. (e) Cell clusters post-GFP/IL-23 gene transfer visualized in tSNE plot. (f) Representative flow cytometry and (g) quantitative data of bone marrow derived MDL-1<sup>+</sup>CD11b<sup>+</sup> and MDL-1<sup>+</sup>CD11b<sup>+</sup>Ly6G<sup>+</sup> 6 days post-IL-23 gene transfer. (h) Representative flow cytometry (i) and quantitative data of skin derived MDL-1<sup>+</sup>CD11b<sup>+</sup>, MDL-1<sup>+</sup>CD11b<sup>+</sup>Ly6G<sup>+</sup> and MDL-1<sup>+</sup>IL-17<sup>+</sup>CD11b<sup>+</sup>Ly6G<sup>+</sup> 21 days post-IL-23 gene transfer. \**P* < 0.05; \*\**P* < 0.01; \*\*\**P* < 0.001 by Mann-Whitney.



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**Fig. 3.** IL-23 induced epidermal hyperplasia is prevented with genetic ablation of MDL-1 receptor. (a) Representative flow cytometry and (b) quantitative data of bone marrow derived MDL-1<sup>+</sup>Ly6G<sup>+</sup> cells and (c) Gene expression analysis of *Il6*, *S100a7*, *Cxcl1*, *Cxcl2* and *Il23r* of total bone marrow cells 48 h post-IL-23 gene transfer in WT and/or *Mdl-1*<sup>-/-</sup> compared to GFP (control) mice. (d) Photographs of murine ears 21 days post IL-23 gene transfer showing the development of silvery white scales in WT and/or *Mdl-1*<sup>-/-</sup> compared to GFP MC (control) mice. (e) H&E staining showing epidermal hyperplasia and inflammatory infiltrate in the murine ear. (Images are representative of three independent experiments and 9–11 mice per each group. Scale bars, 100 μm). (f) Quantification of epidermal thickness (μm) and (g) infiltrated cells per mm<sup>2</sup>. (h) Gene expression analysis of *Il1b*, *Tnf* and *Il6* in the ears of IL-23 gene transfer WT and/or *Mdl-1*<sup>-/-</sup> compared to GFP MC (control) mice. (i) Gene expression analysis of *Il1b* in sorted CD11b<sup>+</sup> bone marrow cells in the presence or absence of LPS stimulation. Data represent mean ± SEM of three independent experiments. \**P* < 0.05; \*\**P* < 0.01; \*\*\**P* < 0.001 by Mann-Whitney.

demonstrate that IL-23 induced a dynamic change in the myeloid populations by dissecting the origin of the t-SNE mapping (Fig. 2e), and particularly induced an expansion of MDL-1<sup>+</sup> cells in the bone marrow (MDL-1<sup>+</sup>CD11b<sup>+</sup> and MDL-1<sup>+</sup>CD11b<sup>+</sup>Ly6G<sup>+</sup> cells) (Fig. 2f, Supplemental Fig. 2a). Our quantitative analysis using percentages and absolute cell counts demonstrated that there was a significant increase of MDL-1<sup>+</sup>CD11b<sup>+</sup> and MDL-1<sup>+</sup>CD11b<sup>+</sup>Ly6G<sup>+</sup> cells (Fig. 2g). Next, we “followed” the inflammation for 21 days and detected the same MDL-1<sup>+</sup>CD11b<sup>+</sup>Ly6G<sup>+</sup> neutrophils in the inflamed ear skin, which were more profoundly expanded compared to the bone marrow (Fig. 2h). Again, quantitative analysis using percentages and absolute cell counts demonstrated that IL-23 induced an increase of MDL-1<sup>+</sup>CD11b<sup>+</sup>Ly6G<sup>+</sup> neutrophils in the skin compared to GFP controls (Fig. 2i). Importantly, we determined that IL-23 induced IL-17A in this MDL-1<sup>+</sup>CD11b<sup>+</sup>Ly6G<sup>+</sup> neutrophil subset compared to the GFP controls (Mean ± SEM; WT + GFP MC = 0.46 % ± 0.10; WT + IL-23 MC = 9.12 % ± 0.77) (Fig. 2h and i). Collectively, our data indicate that IL-23 induces MDL-1<sup>+</sup>CD11b<sup>+</sup> cells in the bone marrow that could migrate to the skin and are capable of producing IL-17A.

### 3.3. IL-23 induced epidermal hyperplasia is prevented with genetic ablation of MDL-1

To further investigate the requirement of MDL-1 in IL-23 induced skin pathology, we performed IL-23 MC *in vivo* gene transfer in WT and *Mdl-1*<sup>-/-</sup> mice. We first confirmed that there was significantly low number of MDL-1<sup>+</sup>Ly6G<sup>+</sup> cells in bone marrow 2 days post IL-23 MC gene transfer (Mean ± SEM; WT + GFP MC = 24.97 % ± 2.54; WT + IL-23 MC = 28.18 % ± 1.95; *Mdl-1*<sup>-/-</sup> + IL-23 MC = 0.75 % ± 0.02) (Fig. 3a and b). By 21 days post IL-23 MC gene transfer, IL-23 induced erythema with silvery white scales in ears which was not observed in GFP controls, while they were attenuated in *Mdl-1*<sup>-/-</sup> mice (Fig. 3c). Consistently, histological analysis showed attenuated epidermal hyperplasia, epidermal thickness and infiltrated cells in *Mdl-1*<sup>-/-</sup> (Fig. 3d–g). Gene expression analysis in the inflamed skin showed that there was a significant decrease in IL-1b in *Mdl-1*<sup>-/-</sup> mice (Fig. 3h). We confirmed the functional significance of this observation with *in vitro* LPS activation assays in CD11b<sup>+</sup> bone marrow cells derived from WT and *Mdl-1* and demonstrated significant differences in IL-1b, which was consistent with previous reports [19] (Fig. 3i). As previous reports showed that *Mdl-1*<sup>-/-</sup> mice had an impaired number of γδ T cells in a *Listeria* infection model [19] we measured the frequency of bone marrow and skin γδ T cells and/or IL-17A producing γδ T cells in WT and *Mdl-1*<sup>-/-</sup> mice, however, we did not detect any significant differences (Supplemental Fig. 3a and b). These data indicate that MDL-1 regulates the expansion of IL-23R<sup>+</sup> myeloid subsets critical for skin inflammation.

### 3.4. IL-23-induced skin transcriptome is consistent with inflammation and neutrophil activation

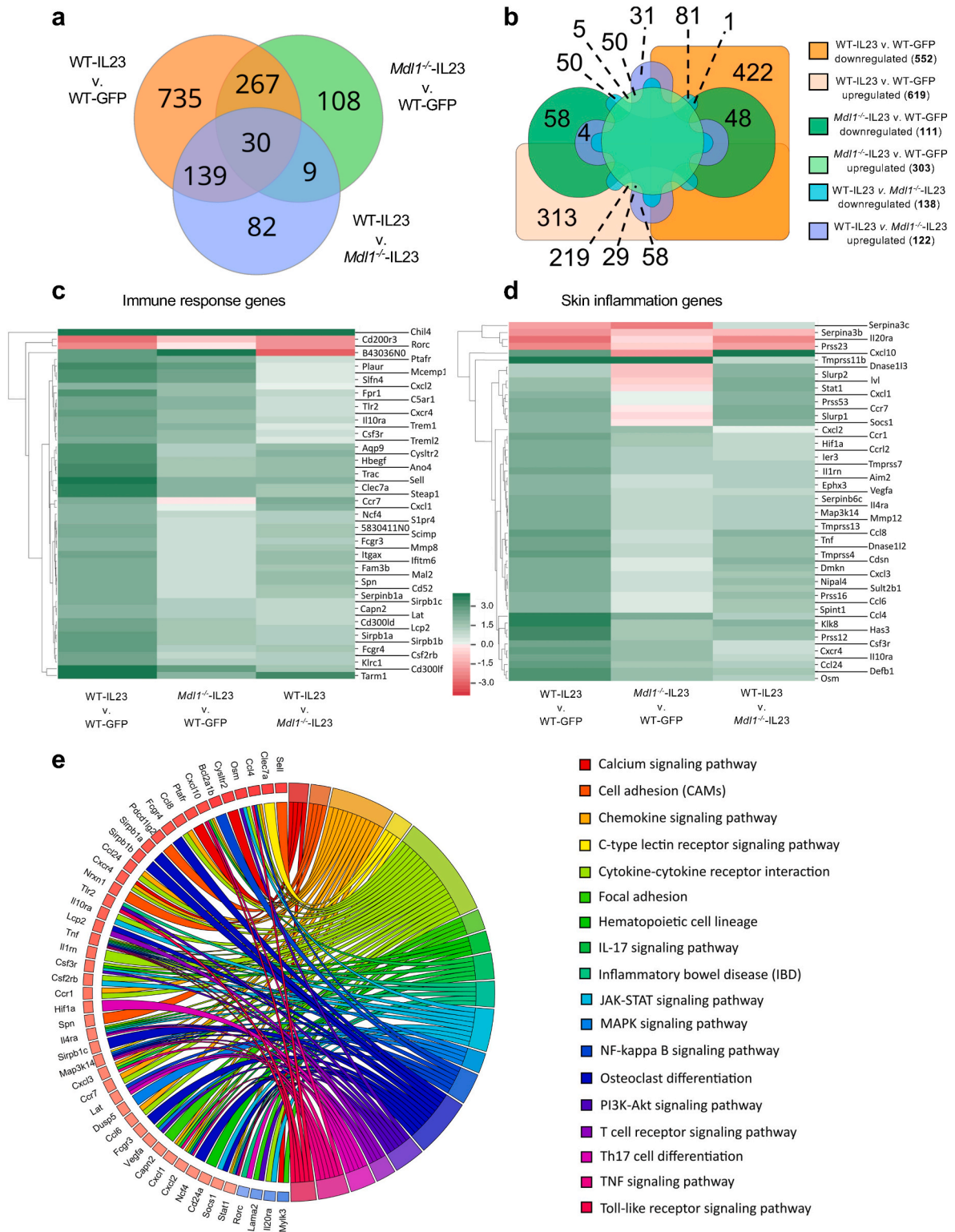
To define the transcriptomic profile of IL-23-induced skin inflammation regulated by MDL-1, we performed RNA-Seq analysis and compared three groups 1) WT + IL-23 v. WT + GFP (control), 2) *Mdl-1*<sup>-/-</sup> + IL-23 v. WT + GFP (control), and 3) WT + IL-23 v. *Mdl-1*<sup>-/-</sup> + IL-23. Sets of differentially expressed genes (DEGs) were defined on the basis of standard thresholds i.e. a fold change >2.0 or <0.5 and q-value (FDR-adjusted p-value) <0.05. We applied two complementary

dimensionality reduction techniques: principal component analysis (PCA) and t-distributed stochastic neighbor embedding (t-SNE) which illustrate potential clusters of neighboring samples and contain similar gene expression patterns (Supplemental Fig. 4a and b). After observation of the expression values for genes in each group, we observed the Pearson Correlation between all DEGs in hierarchically clustered heatmaps (Supplemental Fig. 4c and d, Dataset S1). Due to the fact that many PCC values were close to 1, we identified gene clusters in the heatmap by more closely looking at PCC values greater than 0.97 (Supplemental Fig. 4e and f, Dataset S1), and found 12 clusters in up-regulated genes and 15 clusters in down-regulated genes.

Differential expression analysis revealed 1171 (619 up-regulated and 552 down-regulated) DEGs in WT + IL-23 v. WT + GFP, 414 (303 up-regulated and 111 down-regulated) DEGs in *Mdl-1*<sup>-/-</sup> + IL-23 v. WT + GFP, and 260 (122 up-regulated and 138 down-regulated) DEGs in WT + IL-23 v. *Mdl-1*<sup>-/-</sup> + IL-23 (Fig. 4a and b). An intersection of DEGs from WT + IL-23 v. WT + GFP and *Mdl-1*<sup>-/-</sup> + IL-23 v. WT + GFP revealed a core set of 297 DEGs regulated by MDL-1 (Fig. 4a). The number of DEGs between the comparison groups is illustrated in the Venn diagram (Fig. 4a and b). To illustrate the differences and their relations to the IL-23 MC-induced skin inflammation between the groups, we performed cluster analysis of the entire sample and heatmaps of DEGs between groups were generated (Fig. 4c and d). Our data showed that IL-23 increased the expression of immune response genes were associated with myelopoiesis, neutrophils, DAP12-pathway, myeloid-T cell interactions (Fig. 4c) and skin inflammation (Fig. 4d). An increased expression of those genes was not detected in MDL-1 deficient mice, which were at similar levels to GFP control. Among the DEGs post IL-23 gene transfer, we found several genes associated with psoriasis to be highly elevated in WT but not in *Mdl-1*<sup>-/-</sup> including *Ivl*, *Dmkn*, *Defb*, *Il1er3*, *Slurp2*, *Ephx3*, *Cdsn*, *Vegfa* and *Nipal4* and *Sult2b1* previously identified as critical markers associated with ichthyosis, an abnormal scaling of the skin [29,30]. We found multiple genes associated with neutrophil recruitment and infiltration (*Plaur*, *Ptafr*, *Mmp8*), neutrophil chemotaxis and migration (*Capn2*, *Ano4*, *Ccr7*, *Fam3b*, *S1pr4*, *Cxcl1*, *Cxcl2*, *Ccr4*, *Sell*), and neutrophil activation (*C5ar1*, *Steap1*, *Fpr1*, *Serpinb1a*, *Aqp9*, *Mcamp1*), and an increased expression of genes associated with DAP12 and T cells (*Tarm1*, *Trac*, *Chil4*, *Mal2*, *Spn*, *Scart*, *Lat*, *Rorc*) that again was not detected in *Mdl-1*<sup>-/-</sup> but were at similar levels to GFP control. The overall biological processes commonly associated with those genes are illustrated in a Chord plot (Fig. 4e). The top 10 DEGs across all groups are summarized in supplementary tables (Supplemental Table 2-4).

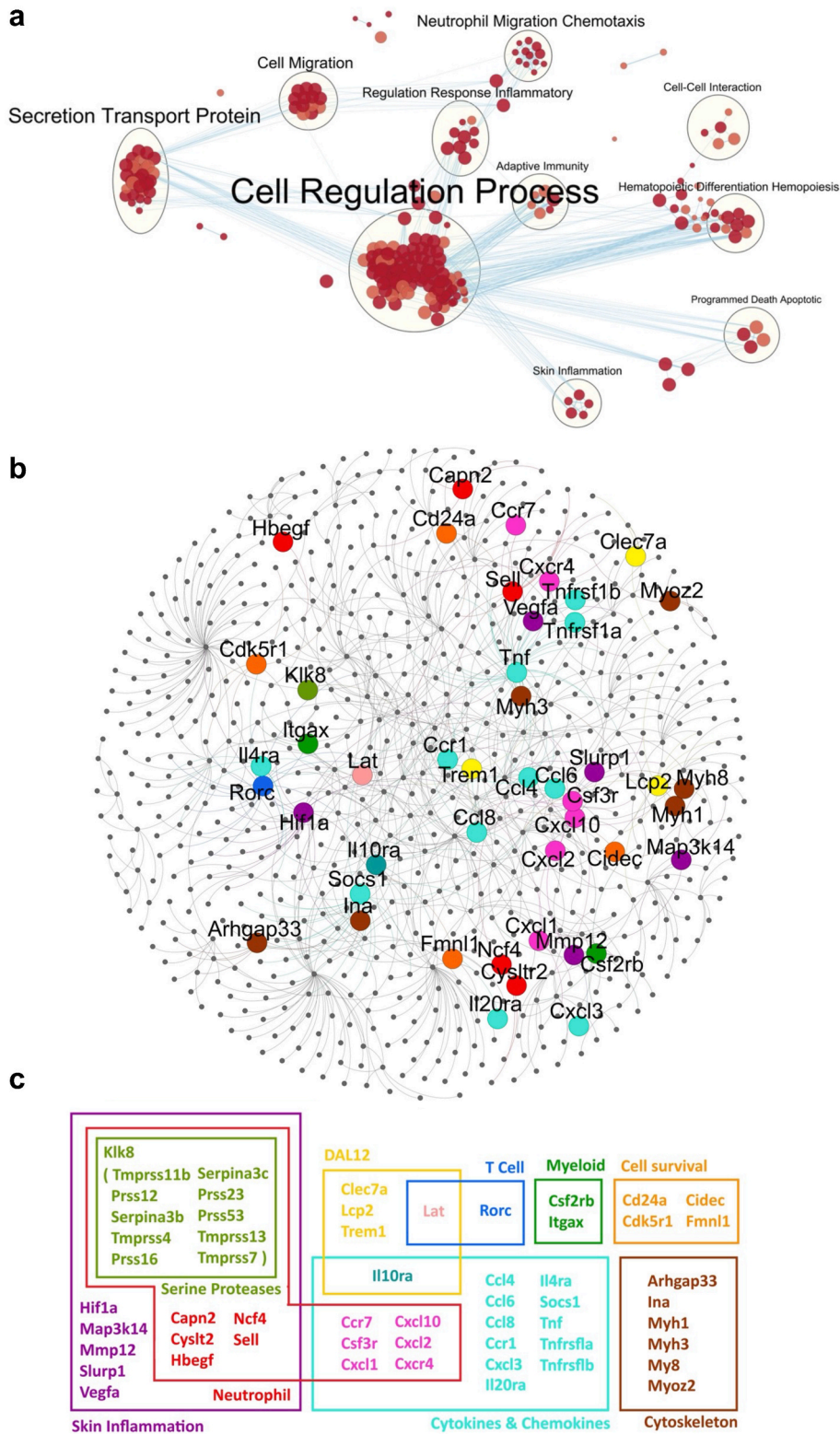
### 3.5. Genetic ablation of MDL-1 protects against IL-23-induced expression of neutrophil-associated genes, neutrophil chemotaxis, and neutrophil proteases in the skin

Next, to identify biological processes involved in IL-23 induced skin inflammation, we performed a gene-set enrichment analysis of biological processes that are statistically enriched in the shared DEGs between the WT + IL23 and *Mdl-1*<sup>-/-</sup> + IL23 groups (Fig. 5a). The immune-related biological processes were associated with hematopoietic differentiation and hemopoiesis, neutrophil migration and chemotaxis, and secretion transport protein as well as adaptive immunity. All of these cellular processes were inhibited in the MDL-1 deficient mice which showed an ameliorated phenotype. Next, we constructed a protein-

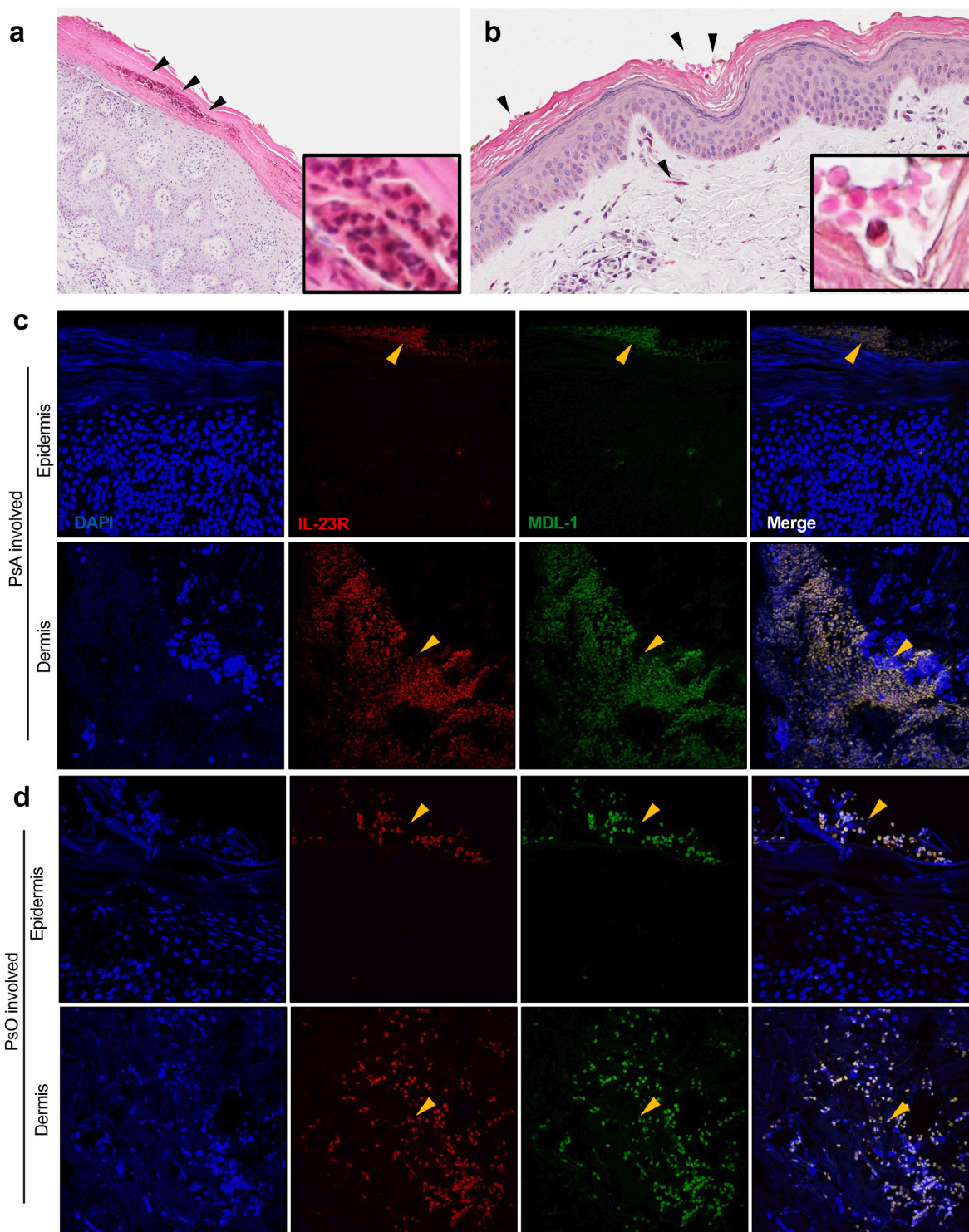


**Fig. 4.** Myeloid activation in the IL-23 induced skin transcriptome (a, b) Venn diagrams showing the overlap of DEGs between WT + IL-23 v. WT + GFP and *Mdl1*<sup>-/-</sup> + IL-23 v. WT + GFP and WT + IL-23 v. *Mdl1*<sup>-/-</sup> + IL-23. (c) The hierarchically clustered heat-map of the target DEGs associated with immune response (myeloid cells, neutrophils, T cell, DAP12) genes, (d) skin inflammation associated genes. Color-coding is based on log-transformed fold count values. DEGs are defined by fold change >2 or <0.5 and q-value (FDR-adjusted p-value) < 0.05. The samples were clustered using average linkage and 1-correlation distances. (Red to green color bar indicates low to high log fold-change values). (e) Chord diagram shows interrelationship for interested genes among KEGG pathways with 4 DEGs or more in the comparison group WT + IL-23 v. WT + GFP. Link thickness is proportional to the overlap between pathways and log-fold change is depicted from decreasing to increasing next to each gene.





**Fig. 5.** Genetic ablation of MDL-1 protects against IL-23-induced expression of neutrophil-associated genes, neutrophil chemotaxis, and neutrophil proteases in the skin. (a) Map of biological processes that are statistically enriched in the shared DEGs between the WT + IL23 and *Mdl-1*<sup>-/-</sup> + IL23 groups. Nodes are gene-sets with 10 or more genes each. Size and color intensity of each node correspond to the number of genes and its statistical enrichment, respectively. Two nodes are connected when they share a significant number of genes (Jaccard coefficient >0.25). (b) Protein-protein interaction network for the target DEGs among WT + GFP v. WT + IL-23 v. *Mdl-1*<sup>-/-</sup> + IL-23. Yellow, red, blue, dark, purple, orange, bright blue, and brown nodes indicate DAP12-, neutrophil-, T cell-, myeloid-, skin inflammation-, cell survival-, cytokines & chemokines-, and cytoskeleton-associated genes, respectively. (c) For genes that belong to different groups, the color pink was used to show an overlap between the neutrophil and cytokines & chemokines groups, light green an overlap between the DAP12 and cytokines & chemokines groups, dark blue an overlap between the DAP12 and T cell groups, and blue-green an overlap between the skin inflammation and serine proteases groups.



**Fig. 6.** *MDL-1* correlates with *IL-23R* and *IL-23* in human PsA and PsO patients. (a) Representative images of H&E staining of human psoriatic skin showing the presence of neutrophil exudates in the stratum corneum. (Arrows indicate Munro's microabscesses). (b) Chloroacetate esterase staining showing the accumulation of neutrophil polymorphs in the stratum corneum and dermis. (Arrows indicate chloroacetate esterase positive cells). Immunofluorescent staining of human skin biopsies of patients with (c) PsA and (d) psoriasis with DAPI (blue), anti-*MDL-1* (green) and anti-*IL-23R* (red) indicating *MDL-1* colocalization with *IL-23R*, (*MDL-1*<sup>+</sup>*IL-23R*<sup>+</sup> cells) (orange) in the epidermal and dermal infiltrates. Images are representative of three experiments. Scale bar, 50  $\mu$ m. (e) Gene expression correlations with *MDL-1* and *IL-23*. Linear regression lines are depicted as blue dashes. Spearman's rank-order correlations ( $r_s$ ) are listed. Within these correlations none of the data points were found to be overly influential (Cook's distance for all data points was <1). Meta-analyses were conducted to evaluate these correlations across the four independently acquired RNA-Seq datasets and the results are displayed as Forest plots. 95 % confidence intervals (CI), weighted averages and p values are shown. There was no evidence ( $p = 0.87$  and  $p = 0.75$ , respectively) of any substantial residual heterogeneity, i.e. there was no remaining variability in effect sizes that was unexplained.

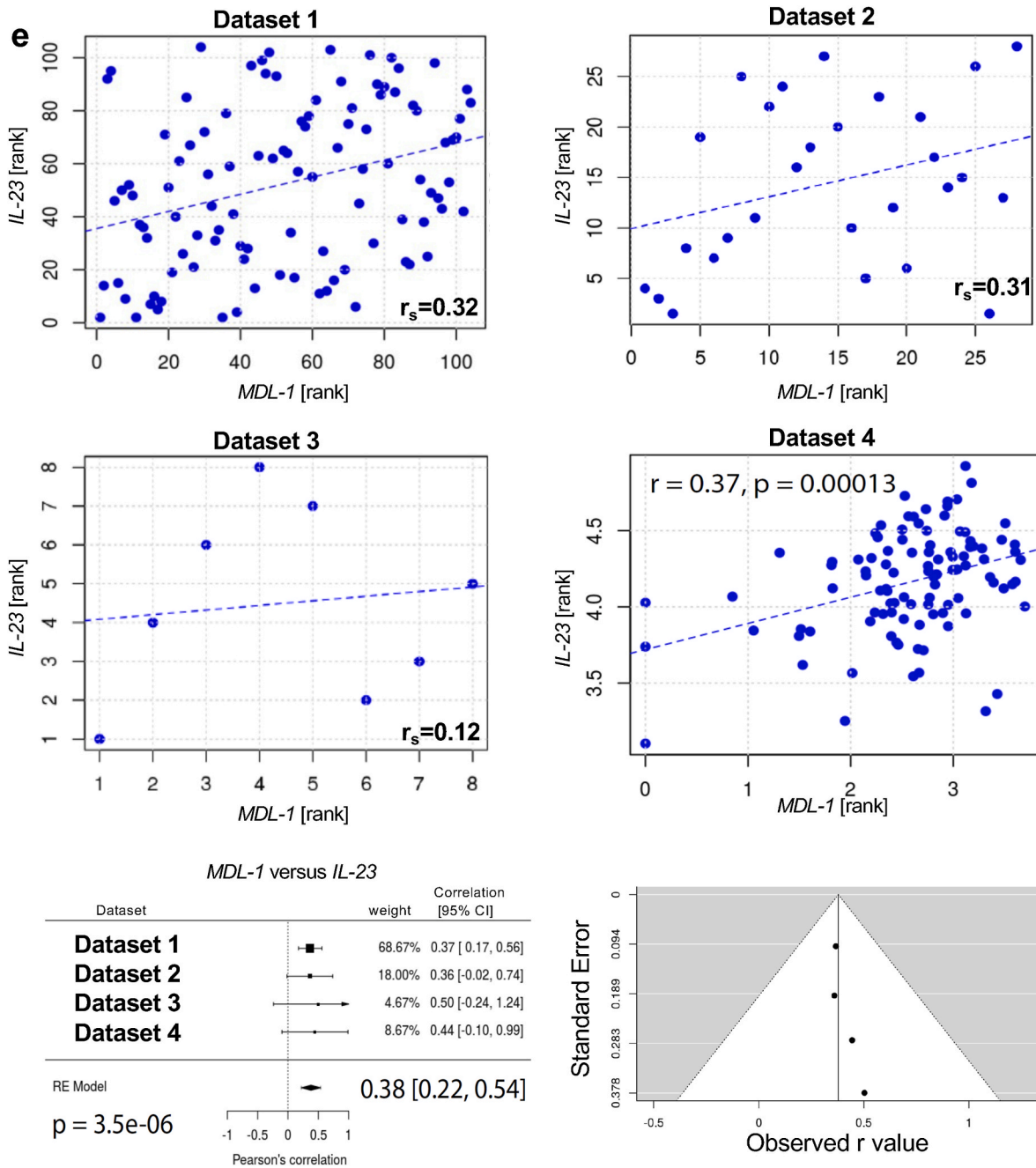


Fig. 6. (continued).

protein interaction network for the target DEGs among WT + GFP v. WT + IL-23 v. *Mdl-1*<sup>-/-</sup> + IL-23 (Fig. 5b). A tight association between neutrophil genes, neutrophil chemotaxis and especially serine proteases were linked with skin inflammation. Other DEGs such as DAP12, T cell, cell survival, cytokines & chemokines, and cytoskeleton-associated genes were also associated with skin inflammation in IL-23 gene transfer but were not detected in the MDL-1 deficient mice (Fig. 5c). Collectively these data suggest that MDL-1 is critical in the regulation of neutrophil genes associated with skin inflammation.

### 3.6. MDL-1 correlates with IL-23R and IL-23 in multiple human PsA and PsO patient cohorts

Next, we utilized human skin biopsies of PsA and PsO patients, and demonstrated by H&E the presence of epidermal hyperplasia and

prominent neutrophil exudates (Munro’s microabscesses) in the stratum corneum (Fig. 6a), a characteristic feature of psoriasis. Chloroacetate esterase staining demonstrated the accumulation of neutrophil polymorphs in the stratum corneum and dermis (Fig. 6b). We also demonstrated by immunofluorescence that a large number of the neutrophil polymorphs were also CD66b<sup>+</sup> and IL-17A<sup>+</sup> cells and accumulated in the stratum corneum and more prominently in the epidermis (Supplemental Fig. 5a), and the presence of epidermal and dermal MDL1<sup>+</sup>IL-23R<sup>+</sup> cells in the human skin of patients with PsA (Fig. 6c) and psoriasis (Fig. 6d). Importantly, this population was also present in the uninvolved skin of patients but undetected in the healthy control (Supplemental Fig. 5b and c). We also performed gene expression correlations from 4 independently acquired psoriasis RNA-Seq data sets (Supplemental Table 5) [31–34]. To understand the relationship between IL-23 and MDL-1, meta-analyses were conducted to determine the weighted averages of

their gene-gene correlations across all three RNA-Seq datasets. The meta-analyses for *IL-23* v. *MDL-1* was positive ( $r_s = 0.38$  [0.22–0.54] (Fig. 6e) and significant ( $P = 3.5E-06$ ) with no evidence ( $P = 0.87$ ) of any substantial residual heterogeneity (i.e. there was no remaining variability in effect sizes that was unexplained). As a negative control, we used *RUNX2* which in contrast, is not expressed in myeloid cells and as expected, there was no detectable correlation between *RUNX2* and *MDL-1* ( $r_s = -0.13$  [-0.60, 0.33]) (Supplemental Fig. 5d). Dash lines depict linear regressions and Spearman's rank-order correlations ( $r_s$ ) are displayed in each graph.

#### 4. Discussion

*IL-23* regulates the development of myeloid and lymphoid immune cells which in turn orchestrate a plethora of immune responses. The implication of *IL-23* signaling in psoriasis and psoriatic arthritis is well-documented and corroborated by GWAS studies, however the exact mechanisms that govern pathogenicity remain elusive [1]. The involvement of *IL-23* in the development of pathogenic *IL-17A* producing T cells, has dominated the literature and overshadowed the importance of myeloid cells which clinically characterize the disease, not only by the distinctive Munro's microabscesses (neutrophil exudates) in the inflamed skin but also by the accumulation of neutrophils in the joints, entheses and nails [35–39]. In our model these neutrophil exudates are secondary to an expansion of neutrophils in the bone marrow. We have previously shown that adoptive transfer of murine bone marrow neutrophils is sufficient to phenocopy disease [25] however the role of neutrophils in PsA are under investigation as recently reviewed [40,41]. We have previously shown that both *IL-23* and *IL-17A* gene transfer induce myelopoiesis in the bone marrow and *IL-17A* induce systemic neutrophil expansion and exhibit clinical manifestations of skin inflammation as commonly observed in psoriasis [11,25,27]. Herein we demonstrate that *IL-23*-mediated myelopoiesis is dependent on *CLEC5A/MDL-1*, a *PU.1* transcriptional target during myeloid differentiation that regulates activation and recruitment of myeloid cells during inflammation [12–15]. We also show that activation of *MDL-1* is specifically important in the maturation of *IL-17A*<sup>+</sup> neutrophils, which have previously been described to be induced by *IL-23* and *IL-6* during fungal infection [42], and observed in psoriatic skin [43,44]. Notably the importance of *IL-17A* expression by neutrophils in psoriasis is under scrutiny, as other neutrophilic functions including NETosis occur simultaneously with *IL-17A* secretion and have also been implicated in skin inflammation and thus pathogenicity should not be attributed to a single pathogenic characteristic [43]. Additionally, transcriptomic analysis of the skin following *IL-23* gene transfer showed a marked increase in neutrophil serine proteases, *KLK8*, *PRSS12*, *PRSS16*, *SERPIN3AB*, which were recently associated with PsA [45,46]. The pathogenic neutrophil-associated genes play far more critical roles than *IL-17A* alone as shown by previous studies where local *IL-17A* injections failed to induce skin inflammation [28] and adoptive transfer of *Ly6G*<sup>+</sup> cells was sufficient to induce skin pathology in the absence of exogenous *IL-17A* [25]. So collectively, regardless of the critical importance of *IL-17A* our results are consistent with multiple studies that have shown the dependence of skin inflammation on neutrophils [25,47]. In fact, we and others have shown that myelopoiesis is a critical event in autoimmune disorders [48–50]. Although we recently demonstrated *MDL-1*-induced myelopoiesis has been previously been associated with osteoclastogenesis and bone destruction [51] this is the first report to our knowledge that associates *MDL-1* with skin inflammation.

In keeping with this notion, our observations of *IL-23R*<sup>+</sup>*MDL-1*<sup>+</sup> cells were corroborated in PsO and PsA human skin biopsies and the interaction of *IL-23* with *MDL-1* in the psoriatic skin was further confirmed across multiple patient cohorts (141 healthy controls VS 149 PsO patients). Although we did observe a significant correlation between *IL-23* and *MDL-1* in these patients the correlation coefficient (R value) was not very high. This could very well be because the patient cohort included

patients with modest to severe psoriasis whereas *MDL-1* seems to play a role in the early initiation phase of the disease. The role of bone marrow myelopoiesis in the pathogenesis of PsA and the contribution of these mechanisms in maintaining the chronicity of the disease remains to be deciphered. We believe that these mechanisms may differ within the patient population of PsA and hope that our data provide a mechanistic explanation of bone marrow driven disease manifestations that can be of benefit to all or to a subtype of patients. Collectively, our data demonstrate that myelopoiesis is critical in the development of skin inflammation and identify *MDL-1* as a potential target for therapeutic interventions.

#### Author contributions

All authors were involved in drafting the article or revising it critically for important intellectual content, and all authors approved the final version to be submitted for publication. Dr. Adamopoulos had full access to all of the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis. IEA conceived and designed the research. HF, CTN, TC, G-HML, AIM, AM and IEA conducted the experiments. HF, CTN, G-HML, SLH, CTR, IT, EM, GCT and IEA analyzed and interpreted the data. HF, CTN, G-HML, EM and IEA wrote the paper, with comments provided by all other authors.

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#### Declaration of competing interest

The authors have no conflicts of interest to declare.

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#### Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.jaut.2024.103167>.

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