

OVOL1 Regulates Psoriasis-Like Skin Inflammation and Epidermal Hyperplasia



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Psoriasis is a common inflammatory skin disease characterized by aberrant inflammation and epidermal hyperplasia. Molecular mechanisms that regulate psoriasis-like skin inflammation remain to be fully understood. Here, we show that the expression of *Ovo1* (encoding ovo-like 1 transcription factor) is upregulated in psoriatic skin, and its deletion results in aggravated psoriasis-like skin symptoms following stimulation with imiquimod. Using bulk and single-cell RNA sequencing, we identify molecular changes in the epidermal, fibroblast, and immune cells of *Ovo1*-deficient skin that reflect an altered course of epidermal differentiation and enhanced inflammatory responses. Furthermore, we provide evidence for excessive full-length IL-1 α signaling in the microenvironment of imiquimod-treated *Ovo1*-deficient skin that functionally contributes to immune cell infiltration and epidermal hyperplasia. Collectively, our study uncovers a protective role for OVOL1 in curtailing psoriasis-like inflammation and the associated skin pathology.

Journal of Investigative Dermatology (2021) 141, 1542–1552; doi:10.1016/j.jid.2020.10.025

INTRODUCTION

Psoriasis is one of the most common inflammatory skin diseases, affecting 2–3% of the world population (Ayala-Fontanez et al., 2016). The etiology of the disease is multifactorial, encompassing both genetic and environmental components that cause dysregulations of the skin barrier and immune cell function (Benhadou et al., 2019; Bergboer et al., 2012; Lowes et al., 2014; Roberson and Bowcock, 2010). Central to skin barrier function is the epidermis, a self-renewing epithelium composed of a basal layer containing stem and progenitor cells and multiple suprabasal layers containing progressively differentiating cells that produce stratum corneum, the actual physical barrier at the skin's outmost surface (Gonzales and Fuchs, 2017). In psoriasis and other aberrant skin inflammatory processes, epidermal cells both serve as the target of immune cells and play a crucial role in shaping the immune responses (Kobayashi et al.,

2019; Pasparakis et al., 2014). The molecular mechanisms that regulate the complex cross-talk between epidermal cells and immune cells in psoriasis remain to be fully elucidated.

SNPs in *OVOL1* have been shown to associate with several inflammatory skin diseases, namely atopic dermatitis, atopic march, and acne (Hirota et al., 2012; Kang et al., 2015; Marenholz et al., 2015; Navarini et al., 2014; Paternoster et al., 2011). *OVOL1* encodes a zinc finger transcription factor, and its mouse homolog *Ovo1* is functionally required for proper embryonic epidermal development (Dai et al., 1998; Lee et al., 2014; Nair et al., 2006; Teng et al., 2007). Specifically, *Ovo1* is expressed in the transiently proliferative epidermal intermediate and spinous cells, and its germline ablation results in a thickened epidermis with expanded early differentiating cells (spinous layers) and a transient delay in barrier acquisition that is resolved by birth. Studies using cultured human keratinocytes (KCs) and clinical samples have implicated a possible functional involvement of *OVOL1* in atopic dermatitis (Furue et al., 2019; Tsuji et al., 2020, 2017). Specifically, *OVOL1* was shown to regulate the expression of epidermal terminal differentiation genes *FLG* and *LOR*. However, the in vivo function of *Ovo1/OVOL1* in skin homeostasis and inflammation has not been addressed.

Despite the long-held view of distinct mechanisms underlying each inflammatory skin disorder, overlapping molecular mechanisms that affect both psoriasis and atopic dermatitis exist (Guttman-Yassky and Krueger, 2017). In this work, we report elevated *OVOL1/Ovo1* expression in lesional psoriatic human skin and in epidermal cells of mouse skin with psoriasis-like inflammation. We show that *Ovo1* is largely dispensable for adult epidermal homeostasis, but its deletion significantly aggravates imiquimod (IMQ)-induced psoriasis-like skin defects featuring epidermal hyperplasia and neutrophil accumulation. Using bulk and single-cell (sc) RNA sequencing (RNA-seq) coupled with functional studies, we identify aberrant IL-1 α expression and signaling as an

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Abbreviations: GO, Gene Ontology; IMQ, imiquimod; RNA-seq, RNA sequencing; sc, single-cell

Received 17 June 2020; revised 22 October 2020; accepted 23 October 2020; accepted manuscript published online 14 December 2020; corrected proof published online 11 January 2021

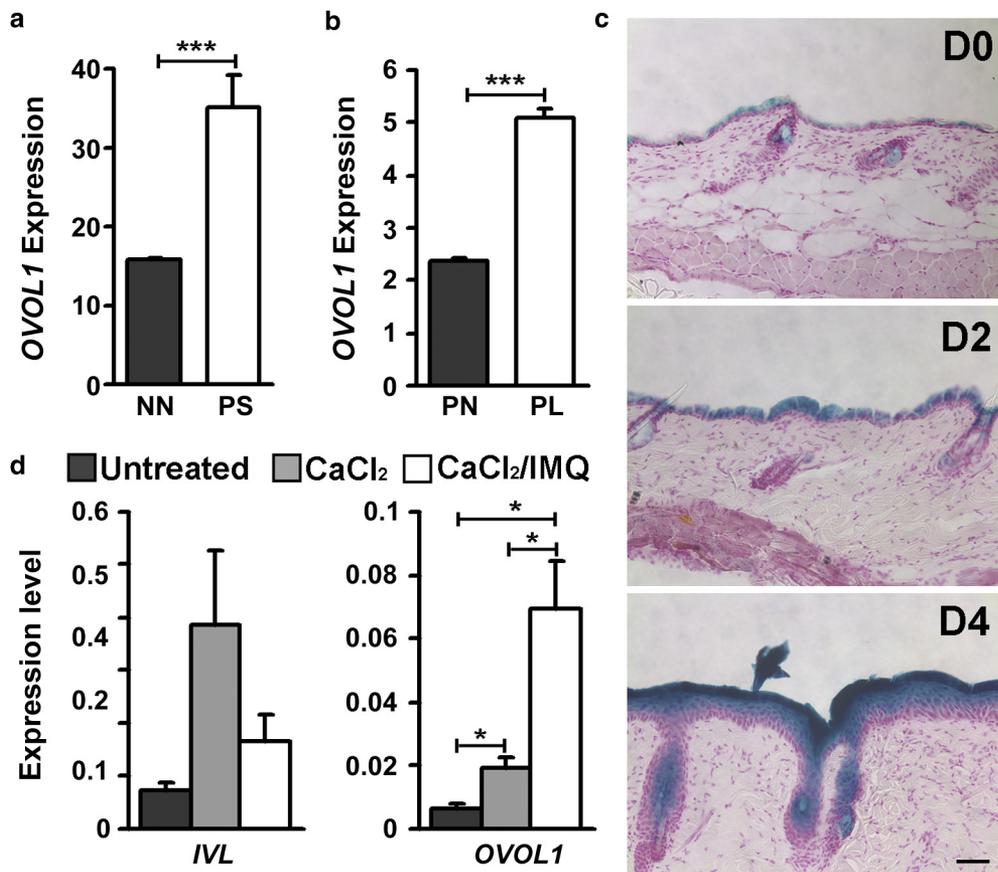


Figure 1. *OVOL1/Ovol1* expression is upregulated in psoriatic skin. (a) *OVOL1* expression in psoriatic PS taken from each patient and control NN skin samples obtained from surgical discard specimens of healthy people. $n = 5$. (b) *OVOL1* expression in PL and PN skin of human patients with psoriasis. $n = 81$. (c) X-gal staining of β -galactosidase activity in skin of *Ovol1-LacZ* mice. $n = 2-3$ (shown are images from 8-week-old mice). Bar = 100 μ m. (d) RT-qPCR analysis of the indicated genes in human foreskin keratinocytes following calcium (1.2 mM for 30 hours) and IMQ (50 μ g/ml for 6 hours) treatment. Data are from three independent experiments. *** $P < 0.005$, * $P < 0.05$. D, day after IMQ treatment; IMQ, imiquimod; IVL, involucrin; NN, normal nonpsoriatic; PL, paired lesional; PN, paired nonlesional; PS, plaque skin.

important mediator of the exacerbated inflammation and epidermal hyperplasia in *Ovol1*-deficient mice.

RESULTS

OVOL1/Ovol1 expression is upregulated in psoriatic skin

To seek clues for a functional involvement of *OVOL1/Ovol1* in psoriasis, we examined our previously published RNA-seq data on human psoriasis skin samples (Yu et al., 2020). *OVOL1* expression was found to be upregulated by >2-fold in psoriatic skin compared with normal skin (Figure 1a). We also interrogated a public dataset on paired nonlesional and lesional skin from human patients with psoriasis (Correa da Rosa et al., 2017; Suárez-Fariñas et al., 2012) and observed significantly higher *OVOL1* expression in the lesional skin (Figure 1b). To determine if this is the case in an animal model of psoriasis, we turned to IMQ-treated mouse back skin. IMQ is a toll-like receptor 7/8 agonist and its short-term (5- to 7-day-long) application induces dermatitis with phenotypic and histological resemblance to human psoriasis lesions, including hyperkeratosis, erythema, scaling, neutrophil microabscesses in epidermis, and infiltration of $\gamma\delta$ T cells and T helper type 17 cells (Gilliet et al., 2004; Schön et al., 2006; Sumida et al., 2014; Swindell et al., 2017; van der Fits et al., 2009; Wu et al., 2004). We generated *Ovol1-LacZ* mice that express β -galactosidase downstream of the *Ovol1* promoter and found that IMQ treatment results in visibly increased β -galactosidase activity in both interfollicular epidermis and hair follicles (Figure 1c). Thus,

OVOL1/Ovol1 expression is upregulated in psoriatic skin of both human and mouse.

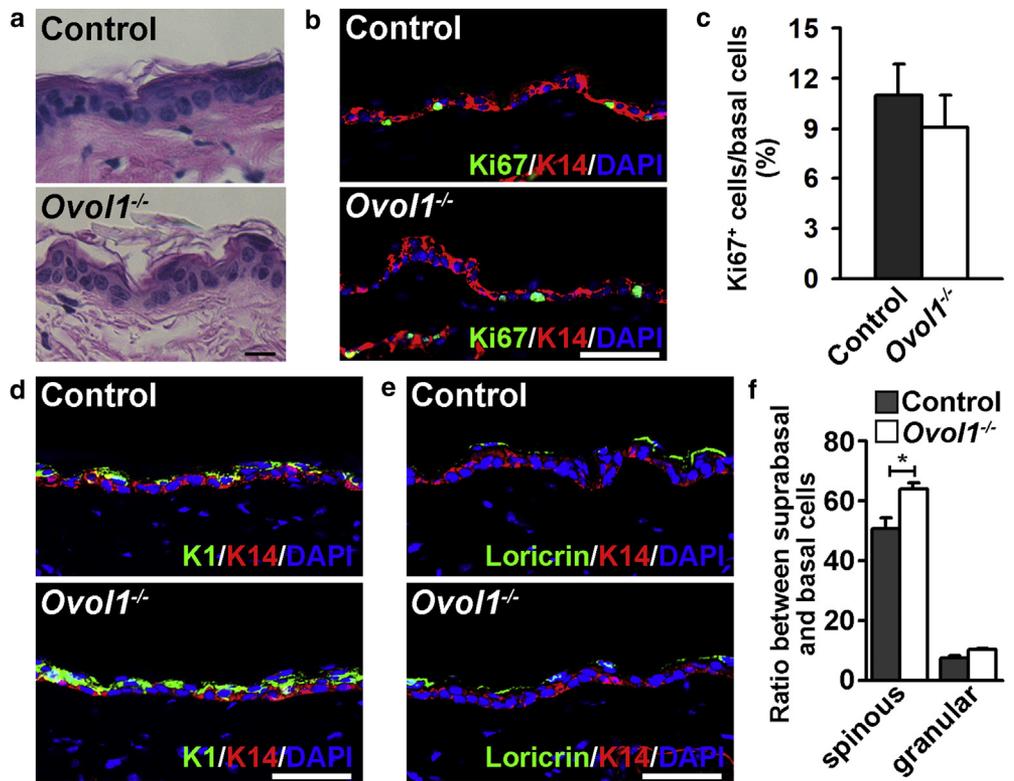
We previously showed that calcium, which is known to induce epidermal differentiation, induces *Ovol1* expression in mouse keratinocytes (Dai et al., 1998). Using human keratinocytes isolated from neonatal foreskin, we found that calcium increased the level of *OVOL1* transcript by ~3-fold, but calcium together with IMQ elicited a ~10-fold increase (Figure 1d). This finding suggests that IMQ can directly upregulate *OVOL1* expression in epidermal cells.

Ovol1 deficiency aggravates IMQ-induced epidermal hyperplasia and inflammatory response

We next investigated *Ovol1* function in adult skin by analyzing *Ovol1*^{-/-} mice on a CD1 outbred background because of perinatal lethality in a C57BL/6 strain background (Dai et al., 1998; Nair et al., 2006). Histology of, and number of Ki67⁺ proliferative cells in, *Ovol1*^{-/-} epidermis were largely similar to those in the littermate control epidermis (Figure 2a–c). However, a slight increase in the ratio between suprabasal and basal cells in *Ovol1*^{-/-} mice compared with the controls (Figure 2d–f), suggesting a very mild alteration in KC differentiation.

Next, we asked whether *Ovol1* deletion impacts IMQ-induced skin inflammation and the associated epidermal hyperplasia. Over a 6-day course of IMQ treatment, psoriasis-like features including erythema and scaling were significantly more pronounced in *Ovol1*^{-/-} mice than in control littermates (Figure 3a and b; Supplementary Figure S1a). H&E

Figure 2. Epidermal cellular defects in *Ovol1*-deficient mice. (a) H&E images of skin from 7-week-old *Ovol1*^{+/+} and *Ovol1*^{-/-} littermates. Bar = 10 μm. (b, c) Ki67 immunostaining. Shown are (b) representative images and (c) quantification of percentage of proliferating basal cells (n = 4 pairs of *Ovol1*^{-/-} and control littermates). (d) K1/K14 and (e) loricrin/K14 double immunostaining. (f) Quantitative analysis of the ratio between suprabasal (K1⁺ spinous and K14⁻ K1⁻ granular) and K14⁺ basal cells (n = 3 pairs of *Ovol1*^{-/-} and control littermates). The number of basal cells was set as 100. DAPI stains the nuclei. (b), (d), and (e) Bar = 50 μm. K, keratin.



and immunostaining analysis revealed that IMQ-treated *Ovol1*^{-/-} epidermis launched a hyperproliferative response faster than control littermates (Figure 3c–e). IMQ-induced defects in terminal differentiation were also more prominent in *Ovol1*^{-/-} skin, shown by lower levels of the differentiation marker K10 (Figure 3f). Moreover, the aberrant upregulation of LM332 protein production as an indicator of tissue repair (Iorio et al., 2015) was detected earlier in the skin of IMQ-treated *Ovol1*^{-/-} mice compared with control littermate skin (Figure 3g; Supplementary Figure S1b). Flow cytometry performed on whole skin revealed a 3.5-fold increase in CD45⁺ (immune) cells, a 16.5-fold increase in CD45⁺/Ly6G⁺/CD11b⁺ neutrophils, and a trending increase in CD45⁺/F4/80⁺/CD11b⁺/Ly6G⁻ macrophages in IMQ-treated *Ovol1*^{-/-} skin compared with control littermate skin (Figure 3h; Supplementary Figure S1c). Together, these data indicate that *Ovol1* loss leads to rapid and severe psoriasis-like symptoms in response to IMQ.

Significantly elevated inflammatory gene expression in IMQ-treated *Ovol1*^{-/-} epidermis

To elucidate the molecular nature of *Ovol1* deficiency–induced changes in skin, we performed RNA-seq analysis on dispase-isolated epidermis from *Ovol1*^{-/-} mice and control littermates that were either untreated or 6 hours after a single IMQ treatment (when epidermal hyperproliferation was not yet apparent; Figure 3d). Principle component analyses revealed that IMQ treatment and sex of the mice analyzed are major sources of variation in gene expression (Figure 4a; Supplementary Figure S2a). Dramatic differences in gene expression between IMQ-treated *Ovol1*^{-/-} and control littermate epidermis were observed (Figure 4a and

b; Supplementary Table S1). Molecular differences between untreated *Ovol1*^{-/-} and control littermate epidermis were also detected but were considerably less prominent than those with IMQ treatment (Figure 4a and b; Supplementary Table S1).

To identify the associated biological processes and pathways, we annotated genes differentially expressed between *Ovol1*^{-/-} and control littermate epidermis using Molecular Signatures Database to reveal the hallmark gene sets and using Enrichr to probe the GO_Biological_Process_2018 library (Chen et al., 2013; Kuleshov et al., 2016) (Supplementary Table S2). Gene signatures related to “inflammatory response” and “regulation of inflammatory response” were upregulated in IMQ-treated and untreated *Ovol1*^{-/-} epidermis, respectively, relative to control counterparts (Figure 4c–e). Also enriched in IMQ-treated *Ovol1*^{-/-} epidermis were Gene Ontology (GO) terms of “TNF-α signaling”, “cytokine-mediated signaling pathway”, “chemokine-mediated signaling pathway”, “regulation of cell proliferation”, “skin development”, and “keratinocyte differentiation” (Figure 4d). Specific upregulated genes in these GO terms (e.g., *Tgfa*, *Slpi*, *Aqp3*, *Ccl20*, *Krt16*, and *Lce3c/d/e*) are also upregulated in human psoriatic skin (Supplementary Table S3; Figure 1a). A number of inflammatory cytokine and chemokine genes, such as *Il1a* (3.6×, $P < 10^{-16}$) and *Tgfa* (2.3×, $P = 0.0001$), were upregulated in IMQ-treated *Ovol1*^{-/-} epidermis, whereas the expression of known psoriasis-associated cytokine genes *Il17*, *Il23*, *Il1b*, and *Il6* was not affected at this early time point. Moreover, the expression of *Flg* and *Lor*, whose human counterparts are known to be regulated by OVOL1 (Furue et al., 2019; Tsuji et al., 2017), was not significantly affected by *Ovol1* deletion in untreated and IMQ-treated epidermis. GO terms that

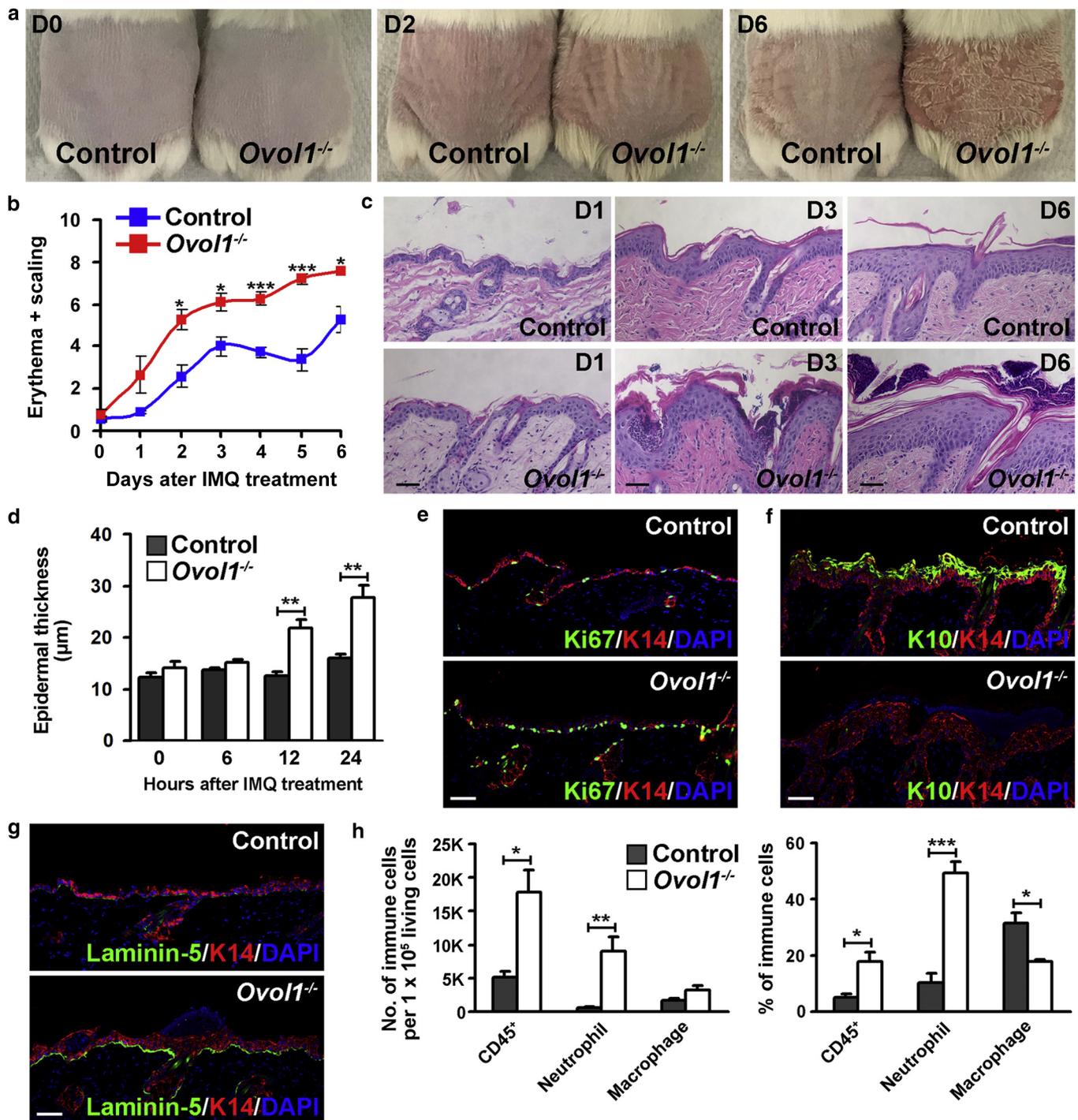


Figure 3. IMQ-induced skin phenotypes in *Ovol1*^{-/-} and control littermates. (a) External appearance, (b) cumulative scoring, and (c) skin histology before and at different days after IMQ treatment. *n* = 4–5 pairs. (d) Quantification of epidermal thickness at different time points after a single IMQ application. *n* = 3–5 per time point. (e–g) Immunostaining at 24 hours after the (e, g) first and (f) third IMQ applications. (h) Summary of flow analysis data collected at 24 hours after the second IMQ application. Left: number of the indicated cell types per 1×10^5 living cells. Right: percentages of CD45⁺ cells per living cells, neutrophils (CD45⁺/Ly6G⁺/CD11b⁺) per CD45⁺ cells, and macrophages (CD45⁺/Ly6G⁻/CD11b⁺/F4/80⁺) per CD45⁺ cells. *n* = 4 pairs. (c), (e–g) Bar = 50 µm. ****P* < 0.005; ***P* < 0.01; **P* < 0.05. D, day; IMQ, imiquimod.

were more enriched in IMQ-treated control littermate epidermis compared with *Ovol1*^{-/-} epidermis included “interferon alpha response”, “cholesterol homeostasis”, “interferon gamma response”, “skin development”, and “keratinocyte differentiation” (Supplementary Figure S2b and c). GO terms that were more enriched in untreated control

epidermis than *Ovol1*^{-/-} epidermis included “positive regulation of cAMP metabolic process” and “interleukin-18-mediated signaling pathway” (Supplementary Figure S2d). Taken together, our bulk RNA data identify enhanced inflammation and altered KC differentiation as major effects of *Ovol1* loss in epidermis.

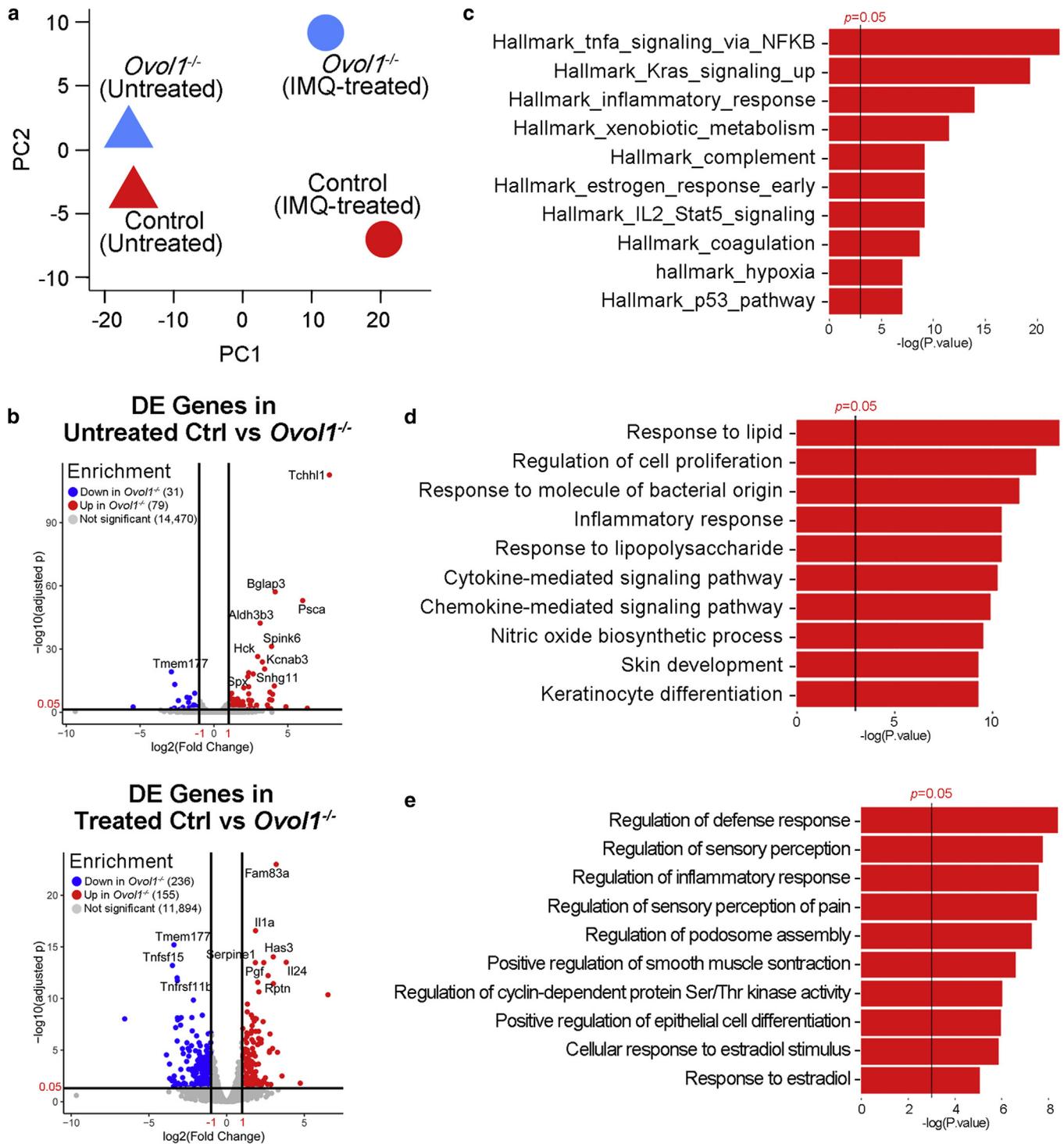


Figure 4. Gene expression in *Ovol1*^{-/-} and Ctrl littermate epidermis. (a) PCA showing epidermal gene expression differences between untreated and IMQ-treated *Ovol1*^{-/-} and Ctrl littermates. (b) Volcano plots showing DE genes between Ctrl and *Ovol1*^{-/-} epidermis from untreated and IMQ-treated mice (n = 2 per genotype: 1 male and 1 female). Genes were filtered for subsequent analysis using the following cut-off: fold change > 2; adjusted P-value < 0.05. (c, d) Functional annotation of genes that were upregulated in IMQ-treated *Ovol1*^{-/-} epidermis using Hallmark gene sets provided by (c) MSigDB and (d) GO terms in Biological Process 2018. (e) GO terms associated with genes upregulated in untreated *Ovol1*^{-/-} epidermis. Shown are the top 10 functional terms and the vertical lines correspond to p = 0.05. Ctrl, control; DE, differentially expressed; GO, Gene Ontology; IMQ, imiquimod; MSigDB, Molecular Signatures Database; PCA, principal component analysis.

scRNA-seq analysis reveals altered gene expression in epidermal basal, suprabasal, fibroblast, and immune cells of the IMQ-treated *Ovol1*^{-/-} skin

Next, we molecularly profiled the skin of *Ovol1*^{-/-} mice and control littermates at 24 hours after IMQ treatment, but this

time using the 10X Genomics scRNA-seq platform to achieve a single-cell resolution. After quality control (Supplementary Materials and Methods; Supplementary Figure S3a), a total of 23,395 cells from four distinct conditions (6,289, 5,553, 5,857, and 5,696 cells from untreated control littermate,

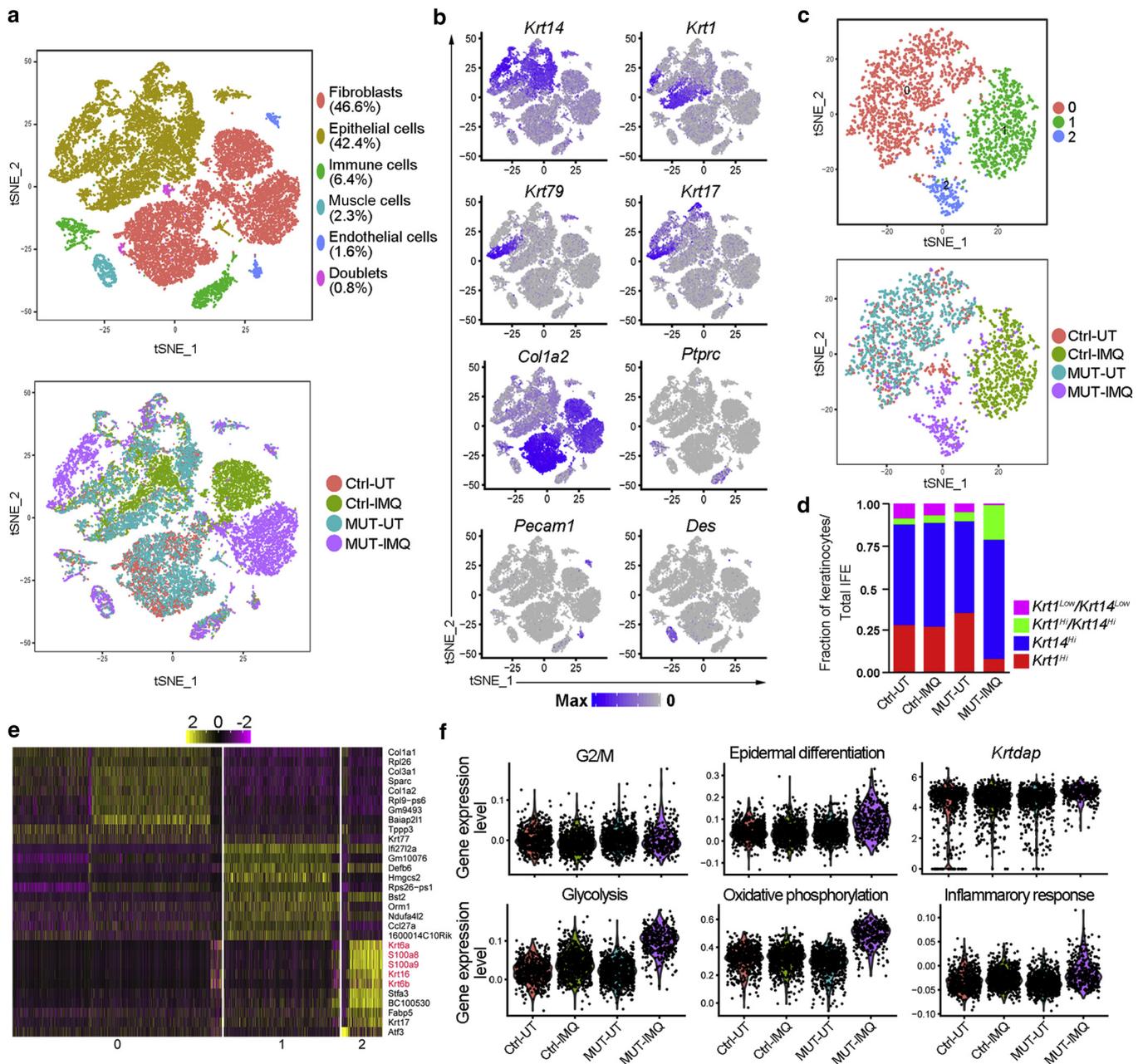


Figure 5. scRNA-seq analysis of homeostatic versus inflamed *Ovol1*^{-/-} and Ctrl littermate skin. (a) tSNE plots showing all cells under analysis that has passed quality control, labeled by cell type (top) and condition (bottom). Ctrl-UT and Ctrl-IMQ, untreated and IMQ-treated Ctrl littermate (*Ovol1*^{+/-}) skin, respectively. MUT-UT and MUT-IMQ, untreated and IMQ-treated *Ovol1*^{-/-} skin, respectively. (b) Feature plots showing expression of the indicated marker genes as in (a). (c) tSNE map of aggregated epidermal suprabasal cells, labeled by clusters (top) and conditions (bottom). (d) Quantification of *Krt14/Krt1* single- or double-positive epidermal cells under the four conditions. (e) Heatmap for the top 10 genes enriched in each cluster from (c). All marker genes are listed in [Supplementary Table S4](#). (f) Violin plots showing expression of the indicated pathway gene signatures or genes in suprabasal keratinocytes across the four conditions. Ctrl, control; IMQ, imiquimod; MUT, mutant; RNA-seq, RNA sequencing; sc, single-cell; tSNE, t-distributed stochastic neighbor embedding; UT, untreated.

untreated *Ovol1*^{-/-}, IMQ-treated control littermate, and IMQ-treated *Ovol1*^{-/-} mice, respectively) were aggregated for subsequent analysis ([Supplementary Figure S3b](#)). In the overall t-distributed stochastic neighbor embedding plot, the major cell types identified were epithelial cells (*Krt14*⁺ or *Krt1*⁺; 42.4%) and fibroblasts (*Col1a2*^{high}; 46.6%) ([Figure 5a](#), top, and [5b](#); [Supplementary Figure S3b](#) and c). Epithelial cells or fibroblasts from untreated *Ovol1*^{-/-} skin intermingled with the same cell type from untreated control littermate skin, whereas IMQ elicited differential gene expression resulting in

the formation of clusters of epithelial cells or fibroblasts that were distinct from the IMQ-treated *Ovol1*^{-/-} and control littermate skin ([Figure 5a](#), bottom).

Given that *Ovol1* is predominantly expressed in epidermal suprabasal cells, *Krt1*^{high} (suprabasal spinous cell) clusters from each sample (untreated or IMQ-treated control and *Ovol1*^{-/-}) were computationally isolated, and those from all four conditions were aggregated together in a combined dataset ([Supplementary Figure S3b](#)). Suprabasal cells from the two untreated genotypes were well mixed in cluster 0, but

those from IMQ-treated control skin were enriched in cluster 1 and those from IMQ-treated *Ovol1*^{-/-} skin were enriched in cluster 2 (Figure 5c), indicating that IMQ induced distinct gene expression changes in control and mutant suprabasal cells. We also noted that IMQ-treated *Ovol1*^{-/-} skin contained a higher number of epidermal cells that expressed both *Krt1* and *Krt14* than the other conditions, a finding validated by immunofluorescence detecting appreciable presence of keratin 1⁺/keratin 14⁺ cells in intact *Ovol1*^{-/-} skin (Figure 5d; Supplementary Figure S3d). Top marker genes of the IMQ-treated *Ovol1*^{-/-} suprabasal cells included proinflammatory genes *S100a8* and *S100a9* (Ryckman et al., 2003; Wang et al., 2018) and keratin-encoding genes *Krt6a*, *Krt6b*, and *Krt16* that have been shown to be upregulated in hyperproliferative or injured epidermis (Chen et al., 2019; Esaki et al., 2015; Lessard et al., 2013) (Figure 5e; Supplementary Table S4). Notably, the same *S100a* and keratin genes have been shown by scRNA-seq to be upregulated in the suprabasal cells of human psoriatic skin (Cheng et al., 2018). Epidermal basal cell clusters (*Krt14*^{high}/*Krt17*⁻/*CD34*⁻/*Sostdc1*⁻) were also computationally isolated and aggregated together for analysis (Supplementary Figure S3b and e). Again, IMQ elicited distinct gene expression changes in *Ovol1*^{-/-} basal cells and control littermate counterparts, featuring upregulation of inflammation- and hyperproliferation-associated genes, including *S100a8*, *S100a9*, and *Krt6a* in the former (Supplementary Figure S3e and f; Supplementary Table S5).

We next analyzed the epidermal expression of select gene signatures associated with proliferation, differentiation, metabolism, and inflammation (Supplementary Table S6). Specifically, G2/M cell cycle checkpoint genes were enriched in basal cells, whereas the epidermal differentiation process was enriched in suprabasal cells, of the IMQ-treated *Ovol1*^{-/-} skin (Figure 5f; Supplementary Figure S3g). Of note, expression of the early differentiation gene *Krt14* was slightly upregulated in basal cells of untreated, but dramatically upregulated in basal cells of IMQ-treated, *Ovol1*^{-/-} skin. Genes involved in metabolic processes (glycolysis, oxidative phosphorylation) were considerably more active in IMQ-treated *Ovol1*^{-/-} epidermal cells in both basal and suprabasal compartments. IMQ-treated *Ovol1*^{-/-} skin scored the highest for an inflammatory response-associated gene set in their suprabasal spinous cells, whereas IMQ-treated control skin scored the highest for this gene set in their basal keratinocytes, raising the possibility that *Ovol1* deletion switches the major site of KC inflammatory response from basal cells to spinous cells. Overall, the molecular changes revealed by scRNA-seq analysis suggest significantly elevated rates of cellular flux characterized by increased basal cell proliferation and early differentiation, inefficient basal-to-mature spinous transition, and enhanced metabolism- and inflammation-associated gene expression in IMQ-treated *Ovol1*^{-/-} skin epidermis.

To further investigate how *Ovol1* loss impacts IMQ-induced changes in skin, fibroblasts from the four conditions were computationally aggregated and analyzed (Supplementary Figure S3b). A total of four clusters were observed: clusters 0 and 3 contained fibroblasts from untreated control and *Ovol1*^{-/-} skin, whereas clusters 2 and 1

were enriched with fibroblasts from IMQ-treated control and *Ovol1*^{-/-} skin, respectively (Supplementary Figure S3h). Top markers in IMQ-treated *Ovol1*^{-/-} fibroblasts included inflammation-associated genes such as *Saa3* and monocyte chemoattractant *Ccl2* (Kratofil et al., 2017; Sack, 2018) (Supplementary Figure S3i; Supplementary Table S7). Moreover, IMQ induced a slight increase in the inflammatory response gene set score in control fibroblasts but a dramatic increase in *Ovol1*^{-/-} fibroblasts (Supplementary Figure S3j).

Few immune cells were recovered in our 24-hour post-IMQ scRNA-seq analysis (Figure 5a and b). However, when these cells were computationally aggregated for analysis, it was still apparent that IMQ induced distinct gene expression changes in immune cells from *Ovol1*^{-/-} and control littermate skin (Supplementary Figures S3b and S4a). *Cd3g*⁺ T cells in IMQ-treated *Ovol1*^{-/-} skin showed elevated expression of *Il17a* and *Il17f* compared with counterparts from the other three conditions (Supplementary Figure S4b–d; Supplementary Table S8), suggesting a heightened T helper type 17 response in the *Ovol1* mutant. Together, these findings show that *Ovol1* deficiency results in an overall more inflammatory skin microenvironment affecting the responses of not only epidermal cells but also fibroblasts and immune cells.

Inhibiting IL-1 α signaling suppresses IMQ-induced skin inflammation in *Ovol1*-deficient mice

IL-1 signaling represents a key proinflammatory pathway in myriad skin inflammatory processes (Cai et al., 2019; Murphy et al., 2000). The scRNA-seq data enabled us to specifically assess IL-1 pathway activity through examining the expression of ligands (*Il1a*, *Il1b*) and receptors (*Il1r1*, *Il1r2*) in different skin cell types. Expression of *Il1a* was barely detectable in the single-cell dataset, but *Il1b* expression was seen in the non-T immune cell cluster that are likely resident macrophage and dendritic cells (Supplementary Figure S5a). *Il1r1* was expressed in most of the cell types with a slight fibroblast enrichment, whereas *Il1r2* expression was highly enriched in the fibroblasts (Supplementary Figure S5a and b). *Il1r2* expression responded differently to IMQ treatment in control and *Ovol1*^{-/-} skin, characterized by increased expression in control epidermal basal and suprabasal cells and fibroblasts but decreased expression in *Ovol1*^{-/-} epidermal cells and no detectable change in *Ovol1*^{-/-} fibroblasts. In contrast, changes in *Il1r1* expression were nonremarkable (Supplementary Figure S5b). Additionally, IMQ induced overall similar changes in the expression of *Il1b* and *Il1r1/2* in immune cells of control and *Ovol1*^{-/-} mice (Supplementary Figure S5c). Because *Il1r2* encodes an IL-1 decoy receptor that inhibits IL-1 α / β signaling (Schlüter et al., 2018), our findings suggest that the absence of *Ovol1* results in a skin microenvironment that is no longer able to efficiently suppress IL-1 signaling when chemically challenged.

Despite the inability of scRNA-seq to detect *Il1a* transcripts, RNAscope analysis confirmed the bulk RNA-seq-revealed upregulation of *Il1a* mRNA in granular cells of *Ovol1*^{-/-}, but not control, skin at 6 hours after IMQ treatment (Figure 6a). Furthermore, IL-1 α protein was detected at greatly enhanced levels in loricrin-positive epidermal

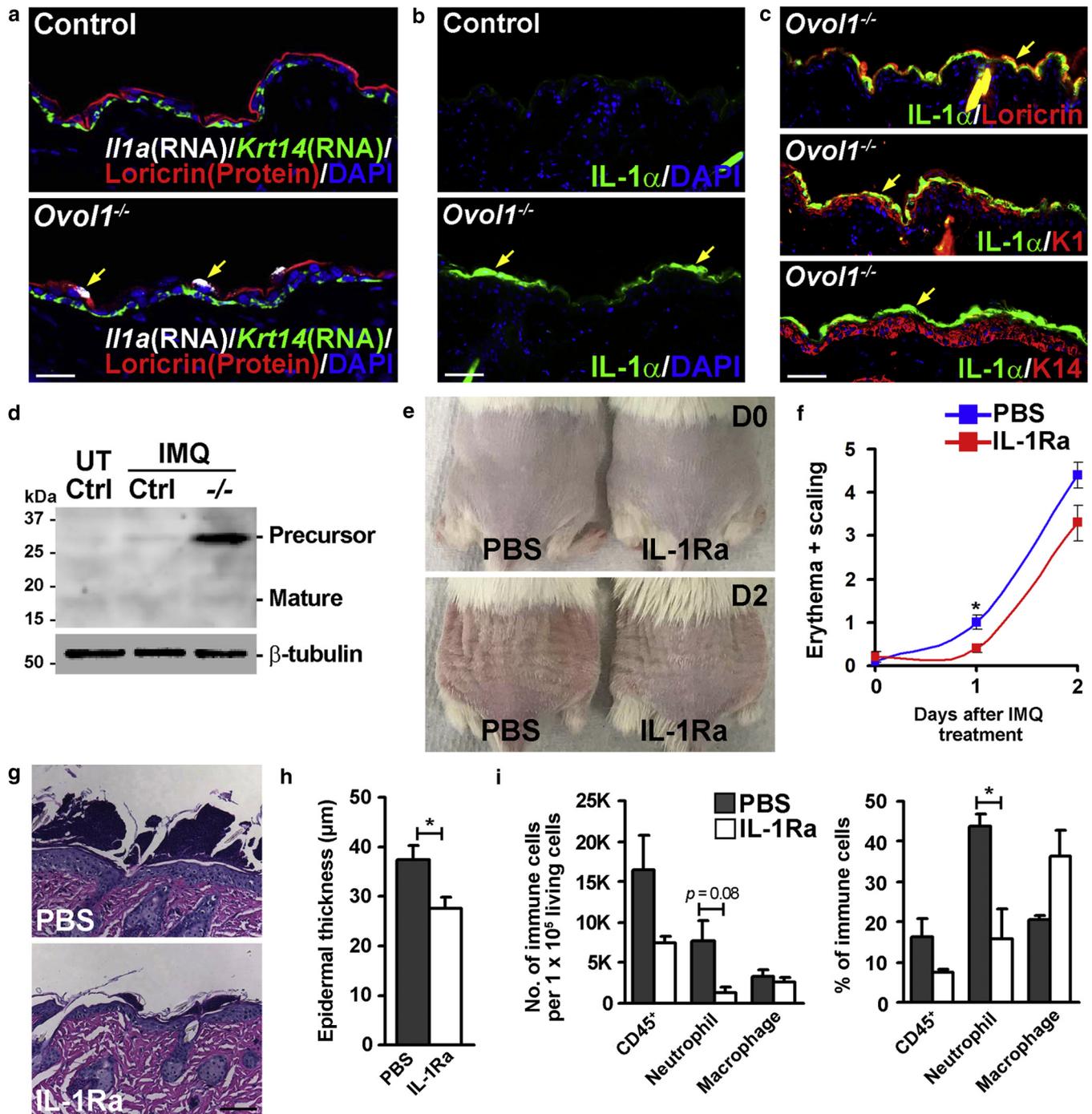


Figure 6. Expression and function of *Il1a*/IL-1 α in *Ovol1*-deficient skin. (a) RNAscope of *Ovol1*^{-/-} and control skin at 6 hours after IMQ using the indicated probes (color-matched). Loricrin antibody stains granular cells. Arrows: *Il1a* signals. Bar = 25 μ m. (b, c) Immunostaining of *Ovol1*^{-/-} and control skin at 24 hours after IMQ treatment. Bar = 50 μ m. (d) Western blotting for IL-1 α (top) and β -tubulin (bottom). (e) External appearance and (f) cumulative scoring of IMQ-treated *Ovol1*^{-/-} skin after IL-1Ra or PBS administration (n = 5 each). (g) Histology, (h) epidermal thickness, and (i) immune cell quantification of IMQ-treated *Ovol1*^{-/-} skin after IL-1Ra or PBS administration (n = 4 each). (g) Bar = 100 μ m. **P < 0.01; *P < 0.05. Ctrl, control; IMQ, imiquimod; UT, untreated.

granular cells of *Ovol1*^{-/-} skin compared with control skin at 24 hours after IMQ treatment (Figure 6b and c). Western blotting showed that the full-length IL-1 α precursor protein, but not the mature form, was predominantly elevated (Figure 6d). These results identify epidermal granular cells as the major source of IL-1 α in inflamed *Ovol1*-deficient skin.

IL-1 α is a dual function cytokine; its secreted and membrane-bound forms signal through the IL-1R1 (*Il1r1*)

receptor, and its N-terminal calpain cleavage product translocates into the nucleus to regulate chromatin structure and transcription (Rider et al., 2013; Werman et al., 2004). To ask whether the excessively produced, full-length IL-1 α in IMQ-treated *Ovol1*^{-/-} skin epidermis acts through signaling, we pharmacologically inhibited the IL-1 signaling pathway. Specifically, we administered recombinant IL-1Ra, an antagonist of IL-1 α / β through competitive receptor binding

(Petrasek et al., 2012), or PBS via daily topical skin applications at 30 minutes before IMQ application and every 24 hours until the end of experiments (24 hours after the second IMQ application). IL-1Ra treatment resulted in the normalization of psoriasis-like phenotypes of IMQ-treated *Ovol1*^{-/-} skin, manifested by reductions in erythema, scaling, epidermal thickness, and number of immune cells (neutrophils in particular) (Figure 6e–i; Supplementary Figure S6). Together, these data show that the excessive full-length IL-1 α protein produced by the epidermal granular cells in inflamed *Ovol1*^{-/-} skin functions via signaling to mediate the exaggerated inflammation, immune cell infiltration, and associated epidermal hyperproliferation.

DISCUSSION

Our work uncovers an important function of the OVOL1 transcription factor in regulating psoriasis-like skin inflammation in vivo. During normal epidermal homeostasis, *Ovol1* function is largely dispensable, possibly reflecting the existence of compensatory mechanisms. However, *Ovol1* loss predisposes skin to exacerbated psoriasis-like inflammation and epidermal hyperplasia in response to IMQ. Based on our mouse model data suggesting a protective role of *Ovol1*, we speculate that *OVOL1* may also play a protective role in human skin, such that its expression is upregulated in psoriatic skin to dampen disease progression and/or severity. Future work outside the scope of this study will be needed to address this intriguing possibility.

Ovol1 expression is upregulated in the epidermal spinous cells of IMQ-treated skin. Hence, our gene expression analysis focused on the early changes, which likely represent the more immediate effects of *Ovol1* loss in the epidermis rather than secondary consequences of disease progression. Both our bulk and scRNA-seq experiments revealed increased expression of inflammation-associated genes and epidermal differentiation genes in the absence of *Ovol1* shortly after IMQ treatment. Some of the molecular changes (e.g., *Krt16*, *Lce3c/d/e*, *S100a8/9*) are shared by human psoriatic skin, implicating the potential clinical relevance of our mouse model findings. Moreover, genes upregulated in IMQ-treated *Ovol1*^{-/-} skin at the early time points (6–24 hours after IMQ in our analysis) also show some overlap (e.g., *Lce3c/d/e*, *S100a8/9*, *Sipi*) with those upregulated in skin at 6 days after IMQ treatment (Swindell et al., 2017). This said, our data sets present unique changes that likely reflect the specific effects of *Ovol1* loss and/or the early acute events in psoriasis-like skin inflammation.

Our work also presents a snapshot of the molecular and cellular changes of IMQ-induced skin inflammation at a single-cell resolution. Not surprisingly, inflammatory responses to IMQ are reprogrammed by loss of *Ovol1* not only in epidermal cells but also in fibroblasts and immune cells. Of particular interest, although epidermal cells and fibroblasts in IMQ-treated skin are capable of suppressing IL-1 signaling, these cells in *Ovol1*-deficient skin are in a state that is highly conducive to the excessive IL-1 α signals produced by KCs. Our data provide a useful resource for dissecting the complex signaling cross-talk among the various resident cell types in inflamed mouse skin.

We have previously shown that *Ovol1* is required for efficient cell cycle exit of embryonic epidermal progenitor cells (Nair et al., 2006). *Ovol1* may function in a similar, cell-autonomous manner in IMQ-treated skin to help prevent excessive epidermal proliferation. Additionally, or alternatively, altered KC–immune cell cross-talk may contribute to the aggravated epidermal hyperplasia in IMQ-treated *Ovol1*-deficient mice through feedback mechanisms. *Ovol1*-deficient epidermal cells in IMQ-treated mice show an expansion of the *Krt14/Krt1* double-positive subpopulation and molecular features of increased G2/M cell cycle transition, differentiation, and metabolism, suggesting that loss of *Ovol1* in an inflammatory or stressed environment traps the KC in a state between active cycling and terminal differentiation.

The phenotypic rescue of *Ovol1* mutant mice by IL-1Ra administration supports the functional importance of excessive full-length IL-1 α signaling in mediating the aggravated skin inflammation and epidermal hyperplasia in these mice. Moreover, the finding that topical application of a recombinant IL-1Ra protein can be effective at symptom relief raises the possibility of such use in disease treatment. Given the association of *OVOL1* SNPs with multiple inflammatory skin diseases, our study lays the groundwork for future investigation into the functional contribution of *OVOL1* to disease pathogenesis and the underlying molecular mechanisms.

MATERIALS AND METHODS

Mice

Ovol1-LacZ (*Ovol1*^{tm1a(KOMP)^{Wtsj}) mice in a C57BL/6N background, where a cassette composed of an *Frt* site, a *LacZ* sequence, and *loxP* sites is inserted into the *Ovol1* locus, were purchased from UC Davis KOMP Repository (<https://www.komp.org>). *Ovol1*^{+/-} mice in a C57BL/6 strain background (Dai et al., 1998; Nair et al., 2006) were outcrossed onto a CD1 strain background and then intercrossed to produce homozygous mutant (*Ovol1*^{-/-}) progeny for study. CD1 *Ovol1*^{-/-} mice survive to adulthood but are sometimes smaller than control littermates, so sex- and weight-matched control and mutant littermates were used for all analyses. Information for all genotyping primers and control littermate genotypes used in each analysis is provided in Supplementary Tables S9 and S10. All animal studies have been approved and abide by regulatory guidelines of the Institutional Animal Care and Use Committee of the University of California, Irvine.}

IMQ-induced psoriasis model

Mice at 7–8 weeks of age received a daily topical dose of 62.5 mg 5% IMQ cream (Perrigo, Allegan, MI) on shaved backs for six consecutive days or as indicated. Based on a previously described objective scoring system called PASI (van der Fits et al., 2009), erythema (redness of skin) and scaling (approximated by dry, white cracks and patches on skin surface) were blindly scored independently by one or more investigators, on a score from 0 (none) to 4 (most severe). The cumulative score (erythema plus scaling) served as a measure of the severity of clinical signs (score, 0–8).

Flow cytometry

To obtain single-cell suspensions, minced samples were digested with 10 ml of a solution containing 0.25% collagenase (Sigma, St. Louis, MO; C9091), 0.01 M HEPES (Thermo Fisher Scientific, Waltham, MA; BP310), 0.001 M sodium pyruvate (Thermo Fisher Scientific; BP356), and 0.1 mg/ml DNase (Sigma; DN25) at 37 °C for 1

hour with rotation, and then filtered through a 70- μ m filter, spun down, and resuspended in 2% fetal bovine serum. A total of 5×10^5 cells were stained by incubation for 30 minutes at room temperature with the following antibodies diluted in PBS/2% fetal bovine serum: Alexa Fluor 488-conjugated anti-CD11b (BioLegend, San Diego, CA; 101217), phycoerythrin-conjugated anti-F4/80 (BioLegend; 123110), allophycocyanin-conjugated anti-CD45 (Tonbo Biosciences, San Diego, CA; clone 30-F11, 20-0451), allophycocyanin-Cy7-conjugated anti-Ly6G (Tonbo Biosciences; clone 1A8, 25-1276), 7-aminoactinomycin (BD Biosciences, San Jose, CA; 559925), and phycoerythrin-conjugated anti-CD49f (BD Biosciences; 555736).

Bulk RNA-Seq and scRNA-Seq

These experiments were performed as previously described (Haensel et al., 2020; Lee et al., 2014). For scRNA-seq, single cells were isolated from normal and inflamed back skin of adult mice, and live cells were FACS-sorted for analysis. Additional details are described in the [Supplementary Materials and Methods](#).

In vivo administration of recombinant IL-1Ra protein

Purified IL-1Ra recombinant protein was produced as previously described (Zhang et al., 2009) and was dissolved in sterilized water to a concentration of 15 mg/ml. Same-sex/same-weight *Ovol1*^{-/-} littermates received topical applications of IL-1Ra (40 mg/kg) or PBS on shaved back skin once at 30 minutes before IMQ treatment and then once every 24 hours until the end of experiments. Skin samples were fixed in 4% paraformaldehyde for H&E staining or embedded in OCT and frozen for immunostaining analysis.

Statistics and reproducibility

Most experiments were performed on at least three biological replicates or repeated at least twice. The sample size and number of independent experiments are indicated in the relevant figure legends. Disease pathology scoring was performed blindly. No data were excluded. For analysis of differences between groups, Student's unpaired *t*-test was performed with 2-tailed in Excel. *P*-values of 0.05 or less were considered statistically significant. Error bars represent mean \pm SEM. For bulk RNA-seq analysis, genes with a log₂ fold change ≥ 1 and an adjusted *P*-value < 0.05 (generated by the negative binomial generalized models and Wald test employed by DESeq2) were considered significant and used to generate GO terms.

Additional details for these methods, as well as methods for β -galactosidase assay, human foreskin KC culture and RT-PCR, histology, immunostaining, western blotting, and RNAscope are described in [Supplementary Materials and Methods](#).

Data availability statement

Datasets related to this article are deposited in the Gene Expression Omnibus database under accession code GSE158112. Code for single-cell analysis will be provided on request.

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CONFLICT OF INTEREST

The authors state no conflicts of interest.

ACKNOWLEDGMENTS

We thank Anand Ganesan and Sebastien de Feraudy for inspiration and advice, the Genomics High Throughput Facility and the Institute for Immunology FACS Core Facility at the University of California, Irvine (UCI) for expert service. This work was supported by National Institutes of Health (NIH) grants R01-AR068074 and R01-GM123731 (XD). RV, MD, and DH were partially supported by UCI National Science Foundation (NSF)-Simons Center for Multiscale Cell Fate Research through NSF grant DMS1562176 and Simons Foundation grant 594598 (QN). RV is a recipient of NSF predoctoral fellowship (DGE-1839285), and MD is partially supported by the NIH T32 Immunology Research Training Grant (AI 060573).

AUTHOR CONTRIBUTIONS

Conceptualization: XD; Data Curation: PS, RV, MD, DH, GG, QN, JW; Formal Analysis: PS, RV, MD, DH, GG, QN, JW; Funding Acquisition: XD; Investigation: PS, RV, MD, GG, ZC; Methodology: EG, EP, KK; Project Administration: XD; Resources: SA, YS, WH; Supervision: XD; Validation: PS, RV, MD, GG, ZC; Visualization: PS, RV, GG; Writing - Original Draft Preparation: XD; Writing - Review and Editing: XD, EP

SUPPLEMENTARY MATERIAL

Supplementary material is linked to the online version of the paper at www.jidonline.org, and at <https://doi.org/10.1016/j.jid.2020.10.025>.

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