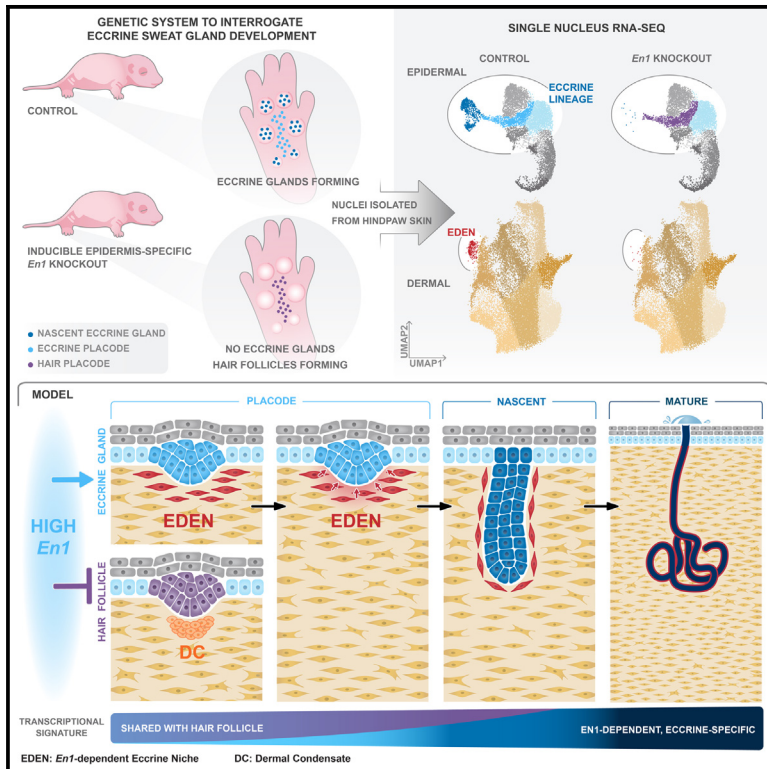


Developmental Cell

Sweat gland development requires an eccrine dermal niche and couples two epidermal programs

Graphical abstract



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In brief

Dingwall et al. demonstrate that the development of eccrine sweat glands, the major effector of the human thermoregulatory system, requires the induction of EDEN, an evolutionarily conserved dermal niche, and couples two concurrent but separable transcriptomes. These findings resolve the molecular signatures and cellular drivers of eccrine identity.

Highlights

- Eccrine gland development requires EDEN, an evolutionarily conserved dermal niche
- Eccrine and hair placode transcriptomes are predominantly congruent
- An *En1*-dependent, low-abundance signature differentiates early eccrine identity



Article

Sweat gland development requires an eccrine dermal niche and couples two epidermal programs

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SUMMARY

Eccrine sweat glands are indispensable for human thermoregulation and, similar to other mammalian skin appendages, form from multipotent epidermal progenitors. Limited understanding of how epidermal progenitors specialize to form these vital organs has precluded therapeutic efforts toward their regeneration. Herein, we applied single-nucleus transcriptomics to compare the expression content of wild-type, eccrine-forming mouse skin to that of mice harboring a skin-specific disruption of *Engrailed 1* (*En1*), a transcription factor that promotes eccrine gland formation in humans and mice. We identify two concurrent but disproportionate epidermal transcriptomes in the early eccrine anlagen: one that is shared with hair follicles and one that is *En1* dependent and eccrine specific. We demonstrate that eccrine development requires the induction of a dermal niche proximal to each developing gland in humans and mice. Our study defines the signatures of eccrine identity and uncovers the eccrine dermal niche, setting the stage for targeted regeneration and comprehensive skin repair.

INTRODUCTION

Eccrine sweat glands are the most numerous appendage found in human skin and are critically important for human thermoregulation, removing excess heat and protecting our bodies from lethal overheating.^{1–3} Despite their importance, there are no therapeutic paradigms to regenerate eccrine glands for reconstructive skin repair, leaving patients with epithelial injuries such as extensive burns with extreme, even life-threatening deficits in thermoregulation.^{1,2,4–7} Thwarting these efforts is a lack of information on how the skin forms these critical appendages in the first place.

Eccrine glands belong to the ectodermal appendage organ class, which also includes hair follicles and mammary glands.^{2,8} These organs form from thickenings of the deepest (basal) layer of the epidermis called placodes, and mature by growing down and differentiating into the underlying connective tissue dermis.^{8–11} To date, studies investigating eccrine gland development have largely implicated molecular regulators and developmental pathways shown to also be required for the formation of other ectodermal appendages, such as hair follicles and mammary glands.^{5,8,10,12–18} However, the molecular factors that distinguish epidermal progenitors as they transition through eccrine gland development are poorly understood.^{5,8,10,12,19,20}

For other ectodermal appendages such as hair follicles, teeth, and mammary glands, a major function of the placode is to induce local condensation of the underlying dermis to form a specialized niche that engages in coordinated and reciprocal interaction with the appendage progenitors to direct further development.^{3,21–25} Strikingly, eccrine gland placodes, and indeed developing eccrine glands at any stage, are not associated with the morphologically evident mesenchymal condensations that are the hallmarks of known dermal niches and an analogous niche for the eccrine gland has not been identified.^{23,31} Accordingly, it is unknown whether eccrine developmental progression requires extrinsic input from the dermis or diverges from the classical ectodermal appendage paradigm.^{8,10,12,26,28–31}

To enable precise targeting of the eccrine gland developmental program, we took advantage of an *in vivo* system that allows direct comparison within the same spatiotemporal context between the developmental program for eccrine glands and that of another major ectodermal appendage, the hair follicle. Specifically, we capitalized on the fact that in the eccrine-forming skin of mice, the palmar/plantar (volar) paw, expression of the *Engrailed 1* (*En1*) gene promotes the formation of eccrine glands and concomitantly inhibits the formation of hair follicles.^{5,12,19,32–34} Using genetic modulation of *En1* levels in this



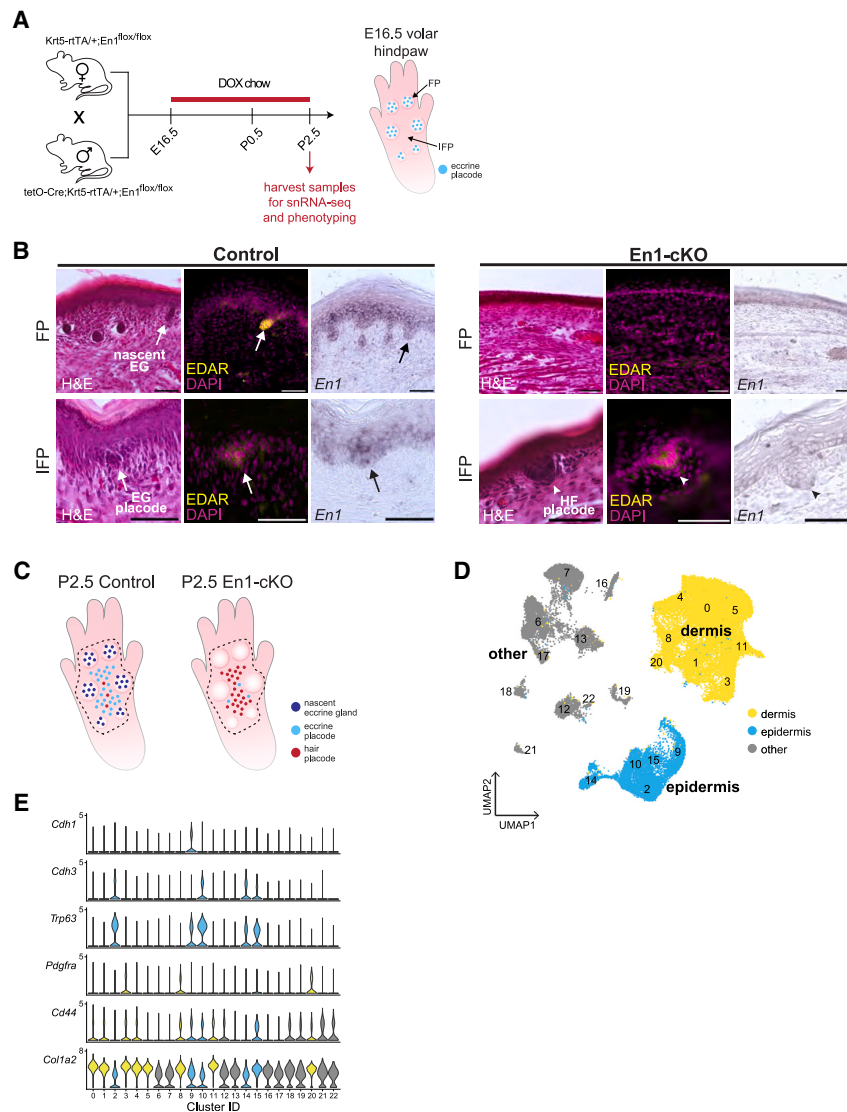


Figure 1. Single nucleus transcriptomic profiling of mouse skin after targeted disruption of eccrine placode identity

(A) Mating scheme to generate animals, dosing schedule, and time points for analyses performed in this study. All En1-cKO and Control mice were administered doxycycline (DOX) starting on embryonic day (E)16.5 until day of harvest. Volar hindpaw skin was harvested on post-natal day (P) 2.5 and analyzed as indicated. Schematic of E16.5 volar hindpaw showing appendage composition at start of Dox administration.

(B) Representative images of stained adjacent sagittal sections through the footpad and inter-footpad regions of P2.5 Control (left) and En1-cKO (right) volar skin. First section in series is stained with hematoxylin and eosin (H&E) for overall morphology; second section in series is stained with EDAR antibody (yellow) to visualize developing eccrine and hair appendages, as indicated. Section is counterstained with 4',6-diamidino-2-phenylindole (DAPI; magenta). Hair placode associated dermal condensate (arrowheads). *In situ* hybridization for *En1* (purple) on the third adjacent section in each series. Scale bars represent 50 μ m. EG, eccrine gland; HF, hair follicle; FP, footpad; and IFP, inter-footpad.

(C) Schematic of Control (left) and En1-cKO (right) volar hindpaw skin at P2.5 summarizing where nascent glands (dark blue), eccrine placodes (light blue), and hair placodes (red) are found at this stage. Dashed line indicates the region of volar hindpaw skin that was dissected and snap frozen for snRNA-seq.

(D) UMAP projection of 45,370 nuclei from merged Control and En1-cKO datasets, which were analyzed in this study. Cluster identification numbers are annotated. Epidermal and dermal nuclei are highlighted in blue and yellow, respectively. All other nuclei are in gray.

(E) Violin plots of gene expression by cluster for markers of the epidermis (*Cdh1*, *Cdh3*, and *Trp63*) and dermis (*Pdgfra*, *Cd44*, and *Col1a2*).

tissue, we shifted the composition of mouse volar skin from eccrine to hair forming and applied single nucleus transcriptomics to parse out the transcriptional and cellular identifiers that make the developing eccrine gland distinct.

RESULTS

Targeted disruption of eccrine development by inhibition of eccrine placode identity

En1 is expressed throughout the deepest (basal) layer of the epidermis in the palmar/plantar (volar) skin of the mouse paw and is focally upregulated in the earliest eccrine anlagen, or placodes, that form therein.^{12,19,35} This expression pattern is recapitulated in human fetal skin in regions where eccrine glands are developing.^{12,34} Importantly, the level of epidermal *En1* during the developmental period when eccrine placodes are specified is directly correlated with the number of eccrine glands that form in mouse and human skin, and reducing *En1* levels results in a dosage-dependent decrease in the number of eccrine

glands.^{5,12,19,33,34,36} We therefore reasoned that disrupting epidermal *En1* expression at the eccrine placode stage would lead to a specific depletion of the eccrine developmental signature from the skin.

To this end, we derived doxycycline inducible, epidermis-specific, *En1*-null, or knockout, mice (En1-cKO) by breeding mice harboring alleles for a tetracycline-responsive Cre recombinase (tetO-Cre), a basal keratinocyte-specific reverse tetracycline transactivator (Krt5-rtTA), and a conditional *En1*-null allele (*En1*^{fllox}).^{21,27,37} We induced *En1* disruption in the Keratin 5 (Krt5)-positive, basal epidermis by administering doxycycline to pregnant dams on embryonic day (E) 16.5 (Figure 1A). At this developmental stage, the first eccrine placodes are forming in the six thickened elevations, or footpads, located at the periphery of the volar hindpaw.^{10–12,19} On post-natal day (P) 2.5, we harvested the volar skin from En1-cKO (tetO-Cre/+;Krt5-rtTA/+;En1^{fllox/fllox}) and Control animals (En1^{fllox/fllox}, Krt5-rtTA/+;En1^{fllox/fllox}, and tetO-Cre/+;En1^{fllox/fllox}) and compared their volar appendage compositions. Consistent with our

previous findings in wild-type mice, developing eccrine glands at multiple stages are simultaneously present in P2.5 Control volar skin.¹⁹ In the footpads, where eccrine gland development began days earlier, nascent-stage eccrine glands that are co-positive for the pan-appendage marker EDAR and for *En1* expression, are differentiating and proliferating into the underlying dermis (Figure 1B).^{11,19} Importantly, the centrally located interfootpad volar skin of Control mice is populated by placode-stage eccrine glands that show characteristic upregulation of *En1* and are also positive for EDAR (Figure 1B, bottom left).^{10,11,19} In contrast, the footpads and the interfootpad regions of P2.5 En1-cKO mice are significantly depleted of nascent eccrine glands and eccrine placodes, respectively (Figure 1B, right panels). Moreover, we find that the interfootpad regions of En1-cKO mice are populated by hair follicle placodes (Figure 1B, bottom right). Each hair follicle placode is readily differentiated from its eccrine counterparts by an underlying dermal condensate (DC), a morphologically evident aggregate of dermal cells that is the instructive niche for each developing hair follicle and the precursor of the dermal papilla (Figure 1B, bottom right, arrowheads indicate DC).^{8,26,28–31}

Analyses of adult Control and En1-cKO mice confirm that the regimen we used to disrupt epidermal *En1* during development permanently inhibits eccrine formation (Figure S1A). We find that compared with Controls, the volar skin of adult En1-cKO mice exhibits a dramatic reduction in footpad eccrine glands coupled with a near-complete loss of these organs in the interfootpad space, as well as an increase in the number of fully formed, interfootpad hair follicles (Figures S1B–S1E; Tables S1–S3).

Our findings demonstrate that genetic manipulation of the *En1* locus in the basal epidermis can be used as a tool to specifically perturb the eccrine developmental program at its onset. Collectively, the reciprocal effect of *En1* on P2.5 eccrine and hair placode identities in the interfootpad space, and the loss of differentiating nascent eccrine glands in the footpads of En1-cKO mice at this stage, makes this an ideal experimental system to capture not only the eccrine developmental program but also the transitions that characterize its progression (Figure 1C).

Single-nucleus RNA sequencing recovers transcriptional signatures of the primary skin layers

The volar hindpaw skin of P2.5 mice is composed of many different cell types, which have differential sensitivity to *En1* and varying degrees of involvement in eccrine gland development. Accordingly, we adapted Drop-seq-based, single-nucleus RNA sequencing (snRNA-seq) to recover the transcriptional profiles of P2.5 Control and En1-cKO skin samples.^{22–24} Using the regimen described above, we dosed pregnant dams with doxycycline at E16.5 and collected the volar hindpaw skin of P2.5 Control and En1-cKO pups for snRNA-seq (Figures 1A and 1C). For each of the three profiled genotypes (two Control genotypes: En1^{flox/flox}, Krt5-rtTA/+; En1^{flox/flox}, one En1-cKO genotype: tetO-Cre/+;Krt5-rtTA/+;En1^{flox/flox}), we profiled two biological replicates, where each replicate was comprised pooled, volar hindpaw skins from five mice of the same genotype. In total, the transcriptomes of 45,370 sequenced nuclei passed quality control filtering and were interrogated in subsequent analyses.

After sample integration with Harmony, initial clustering on the merged data from all nuclei across Control and En1-cKO groups yielded 23 clusters (Figure 1D; Figure S2A).²⁵ Sample replicates showed substantial overlap both via principal components analysis and upon further dimensionality reduction via UMAP (Figure S2B). Analysis for marker gene enrichment reveals that our experiment captured the transcriptomes of nuclei from the two major compartments of the skin, namely nuclei of epidermal (blue) origin and of dermal (yellow) origin (Figures 1D, 1E, S2C, and S2D; Data S1A). Characteristic of the epidermal skin layer, we find clusters that show enrichment for the epidermal makers *Cdh1* (e-cadherin), *Cdh3* (p-cadherin), and *Trp63* (p63) (9,070 nuclei; Figures 1E and S2C; Data S1A).^{38–40} The two Control genotypes contribute 6,013 nuclei to the epidermal clusters, while the En1-cKO genotype contributes 3,057 nuclei. Nuclei from clusters of a nominally dermal origin, which are the most abundant populations captured in our analyses, express known fibroblast markers including *Pdgfr alpha* (*Pdgfra*), *Cd44*, and *Collagen type I alpha 2* (*Col1a2*) (25,859 nuclei; Figures 1E and S2D; Data S1A).⁴¹ Of these, the two Control genotypes contribute 17,398 nuclei to the dermal clusters, and the En1-cKO genotype contributes 8461 nuclei. Based on marker enrichment, we additionally identify clusters of endothelial, smooth muscle, and immune origins (Figures 1D, gray, S2E, and S2F; Data S1A). These findings demonstrate that the nuclear transcriptome reflects the cellular heterogeneity of the volar skin and that our dataset captures expression profiles from the primary skin populations from both Control and En1-cKO experimental samples.

A specialized transcriptome dominates the expression landscape of nascent-stage eccrine glands

Similar to all ectodermal appendages, eccrine glands form from *Krt5*- and *Krt14*-expressing basal keratinocytes.^{8–12} All cells of the developing eccrine gland retain expression of these basal-specific epidermal markers from the placode stage through the nascent period.^{9,11} We observe enrichment for *Krt5* and *Krt14* in clusters 2, 10, 14, and 15, suggesting that the expression signatures of eccrine epidermal progenitors are likely to be contained within this subset of clusters (Figure 2A; Data S1A). Consistent with this, we find that cluster 14 is almost entirely depleted of En1-cKO nuclei, accounting for just 1.64% of En1-cKO epidermal nuclei in the dataset but accounting for 17.71% of epidermal nuclei recovered from Controls (Figures 2B and 2C). *In situ* hybridization to detect *Trpv6* and other top marker genes of cluster 14 reveals a consistent pattern of enrichment in nascent eccrine glands in the hindpaw footpads of wild-type mice (Figures 2D, 2E, S3A, and S3B). That the nascent gland transcriptome is sufficiently diverged as to cluster separately from the other epidermal populations shows that at this stage of development, the overriding transcriptional character of eccrine epidermal progenitors is highly specialized and differentiated with respect to the rest of the epidermis.

Identification of an eccrine developmental lineage in the epidermis

We specifically interrogated the heterogeneity of the epidermal clusters with a goal of resolving finer scale expression differences and identifying transcriptional signatures of pre-nascent-stage eccrine gland progenitors including those of the

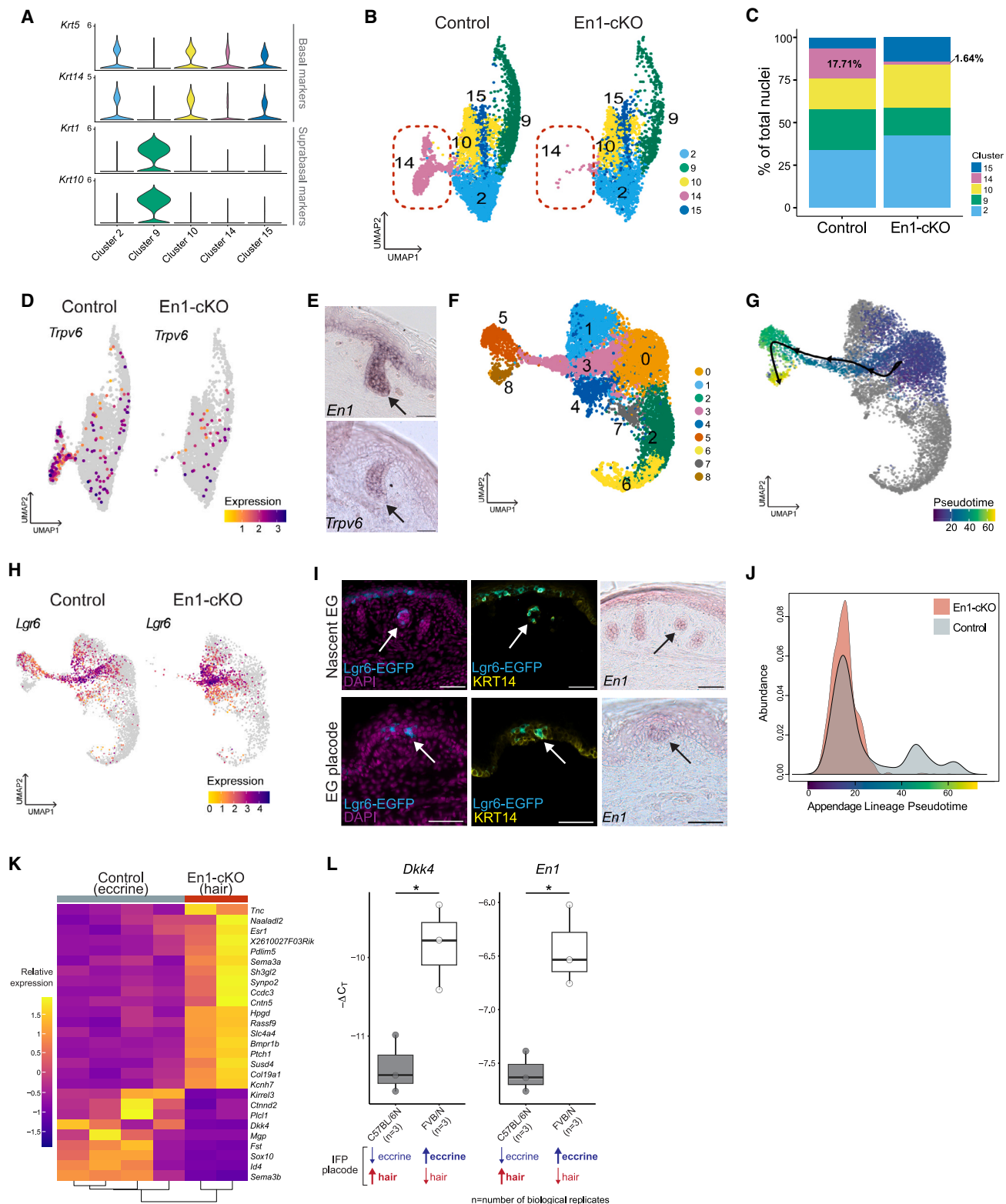


Figure 2. Transitions in the eccrine transcriptome during development

(A) Violin plots of gene expression in only epidermal clusters for markers of basal keratinocytes (*Krt5* and *Krt14*) and suprabasal keratinocytes (*Krt1* and *Krt10*). (B) UMAP projections of epidermal nuclei from original clustering (Figure 1D) colored by cluster identity and split by condition (Control and En1-cKO). Red dashed oval highlights cluster 14, which is depleted in En1-cKO.

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ecrine placodes. After clustering only the epidermal nuclei isolated from merged Control and En1-cKO genotypes, we find that these further resolve into nine epidermal subclusters (Epi0-8) (Figure 2F; Data S1B). Of these, Epi5 and Epi8 correspond to the validated nascent gland cluster 14 in the initial analysis, while nuclei in Epi2, Epi6, and Epi7 are characterized by enrichment for the suprabasal markers *Krt1* and *Krt10* (Figures S3C and S3D; Data S1B).^{9,11} Since early eccrine epidermal progenitors, including those of the placode, are basal in character, the nuclei within Epi0, Epi1, Epi3, or Epi4, which are enriched for the basal epidermal markers *Krt5* and *Krt14*, may represent these earlier stages of eccrine gland development (Figures S3C–S3E; Data S1B).

Trajectory inference performed on all epidermal nuclei identifies a lineage that originates with cluster Epi0, which bears the transcriptional hallmarks of multi-potent basal epidermal keratinocytes and ends with the validated nascent eccrine gland clusters Epi5 and Epi8 (Figures 2G and S3D; Data S1B). Epi3 represents an intermediate state along this inferred lineage and is enriched for transcripts involved in ectodermal appendage development including *Ecdar* (Figures S3D–S3F; Data S1B).

To classify the Epi3 nuclei, we investigated the expression of *Lgr6*, which is the top upregulated transcript of this subcluster compared with all other epidermal populations (Figures 2H and S3D; Data S1B). *Lgr6* is known to be expressed in hair follicle placodes, suggesting that Epi3 may capture the transcriptional signatures of the earliest appendage primordia.⁴² Consistent with this, we find that in the volar hindpaw skin of P2.5 *Lgr6-EGFP* knockin reporter mice, EGFP is expressed in the eccrine placodes of the interfootpad space and in the nascent glands of the footpads (Figures 2H and 2I; Data S1B). These collective data reveal that our snRNA-seq dataset captures a continuous eccrine developmental lineage. Notably, both Control and En1-cKO nuclei contribute to the early inferred eccrine lineage in the epidermis, and Control and En1-cKO samples cluster together in Epi0 and the early portion of Epi3, the origin and intermediate phases of the lineage, respectively (Figures 2J and S3F). These observations suggest that during the initial stages of ec-

crine development, including during the placode stage, the transcriptional identity of eccrine progenitors is highly concordant with that of early hair follicle progenitors.

An *Engrailed-1*-dependent transcriptome distinguishes the sweat gland placode from that of the hair follicle

Our findings reveal that nuclei derived from placode-stage volar appendages, both eccrine gland and hair follicle, are represented in Epi3. We reasoned that the transcriptional signature of the eccrine placode could be resolved by specifically comparing the transcriptional content of Control (ecrine-forming) versus En1-cKO (hair-forming) nuclei within this intermediate cluster. We compared pseudo-bulk expression profiles of nuclei from Epi3 for Control versus En1-cKO samples (Figure 2K). This analysis identifies only 29 transcripts that are significantly differentially expressed between the Epi3 nuclei derived from the two conditions (likelihood ratio test: adjusted $p < 0.01$; \log_2 fold change > 0.58). Of these, nine are relatively upregulated in the Control (ecrine) samples compared with the En1-cKO (hair) samples (Figure 2K).

Our targeted comparative analysis identified the secreted Wnt inhibitor *Dkk4* as a nuclear transcript that is relatively enriched in the eccrine-containing Controls versus En1-cKO samples (Figure 2K). *Dkk4* expression has previously been reported in placode-stage hair follicles and eccrine glands, which we confirmed by *in situ* hybridization in P2.5 volar skin of wild-type mice (Figure S3G).^{10,43–46} To validate the quantitative enrichment of *Dkk4* in eccrine placodes relative to hair follicle placodes, we took advantage of strain specific differences in the number of volar hair follicles and eccrine glands between C57BL/6N and FVB/N mice.¹⁹ We have previously reported that, on average, the hindpaw interfootpad region of FVB/N mice contains 39 eccrine glands and 5 hair follicles, while the same region in C57BL/6N mice contains 8 eccrine glands and 65 hair follicles.¹⁹ Using genetic mapping, we have previously implicated the higher expression of epidermal *En1* in FVB/N P2.5 volar hindpaw skin as the major causal driver for the specification of more eccrine gland placodes and fewer hair follicles placodes in this strain

(C) Percent contribution of nuclei from each epidermal cluster to total epidermal nuclei from Control and En1-cKO samples, respectively. Clusters are coded by color as depicted in (B).

(D) Feature plots showing normalized expression of cluster 14 marker *Trpv6* in Control and En1-cKO epidermal nuclei. Negative nuclei are shown in gray.

(E) *In situ* hybridization for *En1* (top) and *Trpv6* (bottom) in a wild-type footpad at P2.5. Arrow indicates nascent eccrine gland.

(F) UMAP projection of subclustered epidermal nuclei colored by subcluster identity (merged Control and En1-cKO conditions).

(G) Results of slingshot trajectory inference mapped onto the subclustered epidermal UMAP embedding that depicts an inferred lineage through epidermal subclusters 0, 3, 5, and 8. Points representing nuclei are colored according pseudotime; nuclei not involved in this lineage are colored gray.

(H) Feature plot showing normalized expression of *Lgr6*, a top marker of epidermal subcluster 3, for Control (left) and En1-cKO (right) subclustered epidermal nuclei.

(I) Immunofluorescence staining of sagittal sections through the volar hindlimb skin of *Lgr6-EGFP* mice at P2.5. Tissue sections are stained with antibodies against EGFP (cyan), which reads out endogenous *Lgr6*, and Keratin 14 (KRT14; yellow) and 4',6-diamidino-2-phenylindole (DAPI) nuclear stain (magenta). Sections through footpad skin containing nascent eccrine glands (top), and also of the interfootpad region that contains eccrine gland placodes (bottom, arrow) are shown. *In situ* hybridization for *En1* on adjacent serial sections is used to confirm eccrine gland (top and bottom, right). Scale bars represent 50 μ m.

(J) Pseudotime density distribution of Control (gray) and En1-cKO (red) nuclei for slingshot lineage shown in (E). Kolmogorov-Smirnov test of the distributions indicates differential progression along this trajectory between the conditions ($p < 2.2 \times 10^{-16}$; $D = 0.26487$).

(K) Heatmap showing relative expression of transcripts that are differentially expressed between epidermal subcluster 3 nuclei from Control and En1-cKO pseudobulk samples, which are populated by eccrine glands and hair follicles, respectively (p adjusted < 0.01 ; absolute \log_2 fold change > 0.58). Columns are pseudobulk sample replicates and rows are genes.

(L) qRT-PCR results for *Dkk4* (left) and *En1* (right) of volar hindpaw skin from C57BL/6N ($n = 3$ mice) and FVB/N ($n = 3$ mice) mice at P2.5. Each biological sample represents the right and left volar skin from one mouse. Biological samples were assayed in technical quadruplicate. The interfootpad space of C57BL/6N contains mostly hair placodes at P2.5, while that of FVB/N contains mostly eccrine placodes at this stage. Target gene expression is normalized to the reference gene *Rpl13a* and shown as $-\Delta C_T$. Significant differences between groups assessed by Mann-Whitney-Wilcoxon test (* $p < 0.05$).

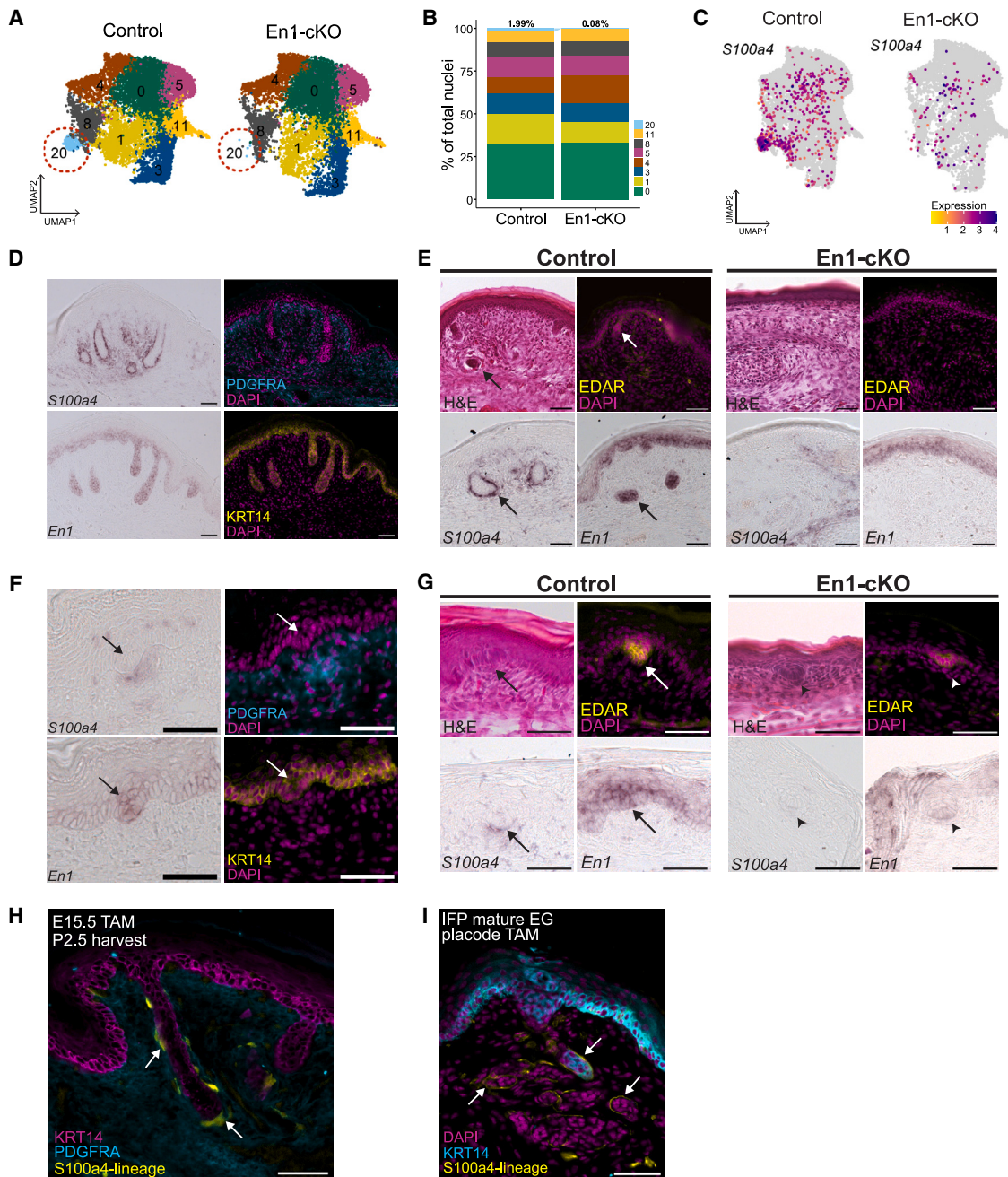


Figure 3. Identification of an *En1*-dependent dermal lineage associated with developing eccrine glands

(A) UMAP projections of dermal nuclei colored by cluster identity (original clustering from Figure 1D) and split by condition (Control and En1-cKO). Red, dotted oval highlights cluster 20, which is depleted in En1-cKO.

(B) Percent contribution of nuclei from each dermal cluster to the total dermal nuclei from Control and En1-cKO samples, respectively. Subclusters are coded by color as depicted in (A).

(C) Feature plots showing relative expression of cluster 20 marker *S100a4*, split by condition.

(D) *In situ* hybridization for *S100a4* (purple) in sagittal sections of wild-type FVB/N P2.5 footpads containing nascent eccrine glands. *En1* expression (purple) by *in situ* hybridization is shown on adjacent sections. Immunofluorescence staining for PDGFRA (cyan), which marks dermal fibroblasts, and KRT14 (yellow), which marks basal keratinocytes. All images shown are from adjacent serial sections.

(E) Representative images of *in situ* hybridization for *S100a4* (purple) and *En1* (purple) along with EDAR immunofluorescence (yellow) and hematoxylin and eosin (H&E) staining of adjacent serial sections of control (*En1*^{flox/flox}) and En1-cKO (*tetO-Cre; Krt5rtTA/+; En1*^{fl/fl}) footpad skin at P2.5. Arrows indicate nascent eccrine glands in the footpad.

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versus the P2.5 volar skin of C57BL/6N mice.¹⁹ Analysis of *Dkk4* expression by qRT-PCR in P2.5 FVB/N and C57BL/6N volar hindpaws reveals a significant upregulation of *Dkk4* in the predominantly eccrine placode-containing skin of FVB/Ns relative to that of C57BL/6N (Figure 2L; Mann-Whitney-Wilcoxon test, *Dkk4*: $p = 0.0495$, $Z = -1.964$, $r = -0.8$). Notably, the effect size estimate of *Dkk4* upregulation is equivalent to that of *En1* upregulation in FVB/N as compared with C57BL/6N (Figure 2L; Mann-Whitney-Wilcoxon test, *En1*: $p = 0.0495$, $Z = -1.964$, $r = -0.8$).

The relative increase of *Dkk4* expression in eccrine gland versus hair follicle placodes confirms the existence of a specific signature of eccrine identity that is dependent on the epidermal expression of *En1*. Taking into account the extensive transcriptional similarity we observe for Control and *En1*-cKO nuclei early in the inferred eccrine epidermal lineage, our data are consistent with the concurrent but disproportionate representation of two distinct transcriptional programs in the eccrine placode: a major transcriptome that is shared with the hair follicle placode, and a relatively underrepresented transcriptome that distinguishes the eccrine placode and is associated with its identity.

Identification of an eccrine-associated dermal lineage

Skin-specific disruption of *En1* not only depletes eccrine-associated epidermal populations but also results in dramatic depletion of *En1*-cKO nuclei from primary cluster 20, which is of dermal rather than epidermal origin (Figures 1D and 3A). Nuclei from cluster 20 account for 1.99% and 0.08% of Control and *En1*-cKO dermal populations, respectively (Figure 3B). *S100a4* is the top marker gene that differentiates cluster 20 from all other dermal populations (Figures 3C and S2D; Data S1C). Using *in situ* hybridization, we find that *S100a4* is specifically expressed in PDGFRA-positive dermal fibroblasts immediately surrounding the nascent-stage eccrine glands in the footpads of P2.5 wild-type mice (Figure 3D). This pattern is recapitulated by *Tnc*, another top marker of cluster 20 (Figures S4A, S4B, and S2D; Data S1C). The presence of a dermal sheath around nascent-stage eccrine glands of mice was previously observed histologically by Cui and colleagues.¹⁰ Consistent with the specific association of the cluster 20 transcriptional signature with dermal cells that surround the nascent glands, we find that *S100a4* expression is lost in the footpads of *En1*-cKO mice at P2.5 (Figure 3E).

Intriguingly, we also observe *S100a4* expression specifically in the PDGFRA-positive dermal cells directly beneath eccrine plac-

odes, which exhibit characteristic *En1* upregulation (Figures 3F and 3G). These *S100a4*-positive dermal cells are not morphologically distinct from the surrounding mesenchyme, consistent with previous reports that developing eccrine glands are never found in association with the dermal aggregates characteristic of the other major ectodermal appendages such as the hair follicle (Figure 3G).^{8,10,28} We find that the upregulation of *S100a4* in the eccrine placode-associated dermis is also dependent on epidermal *En1* since *En1*-cKO volar skin lacks dermal foci of *S100a4* including under the hair follicle placodes that populate the interfootpad space of this genotype (Figure 3G). By contrast, *S100a4* expression is retained in the ventral foot tendons of *En1*-cKO mice, indicating that its absence in the *En1*-cKO volar skin is specific to this context (Figure S4C). Lineage tracing of the *S100a4*-expressing dermal cells under the eccrine placode reveals that the descendants of these cells not only give rise to the nascent gland-associated dermal cells captured in cluster 20 but also to descendants that surround the mature eccrine gland (Figures 3H and 3I). We therefore conclude that epidermal *En1* expression is required for the induction of a single, eccrine-associated, dermal lineage.

Eccrine-associated dermal lineage is conserved to humans

En1 expression in the limb ectoderm initiates well before the onset of eccrine gland formation in this tissue and persists in the adult, after development is completed^{32,35,47} (Figure 4A, bottom). In contrast, we find that *S100a4* expression in the dermis proximal to the eccrine glands is restricted to the developmental period during which the epidermis is actively forming these appendages (Figure 4A, top). Accordingly, we find no evidence of *S100a4* expression either prior to eccrine placode formation or around the mature eccrine glands of adult mice (Figure 4A and S4D). This pattern is recapitulated in human skin, in which we observe *S100a4* expression in the dermal fibroblasts associated with eccrine placodes and nascent glands (120 gestational days, plantar foot skin; Figure 4B) but not in the dermal cells proximal to the mature gland (64 years old, cheek skin; Figure 4C). Thus, the eccrine-associated dermal lineage appears in conjunction with the activation of the *En1*-dependent epidermal eccrine developmental program in both mice and humans and undergoes progressive changes in gene expression over the course of eccrine gland formation. Collectively, these data implicate an evolutionarily conserved and transcriptionally distinct dermal lineage that is specifically associated with developing eccrine glands.

(F) *In situ* hybridization for *S100a4* (purple) in sagittal sections of wild-type FVB/N interfootpad skin containing eccrine placodes at P2.5. *En1* expression (purple) shown by *in situ* hybridization on adjacent sections. Immunofluorescence staining for PDGFRA (cyan) marks dermal fibroblasts and KRT14 (yellow) marks basal keratinocytes. All images shown are from adjacent serial sections. Arrows indicate eccrine placodes.

(G) Representative images of *in situ* hybridization for *S100a4* (purple) and *En1* (purple) along with EDAR immunofluorescence (yellow) and hematoxylin and eosin (H&E) staining of adjacent serial sections of control (*En1*^{fl^{ox}/fl^{ox}}) and *En1*-cKO (tetO-Cre; Krt5rtTA/+; *En1*^{fl^{ox}}) interfootpad skin at P2.5. Arrows indicate eccrine placodes in the interfootpad region. Arrowheads indicate the hair placode and underlying dermal condensate in the interfootpad region of *En1*-cKO volar skin.

(H) Fluorescence image of a representative sagittal section through the footpad of a *S100a4*-CreERT2/+; ROSA-tdTomato/+ mouse at P2.5 that was given tamoxifen at E15.5, around the time of placode formation in the footpads. *S100a4*-lineage cells are shown in yellow. Immunofluorescence staining for PDGFRA (cyan) marks dermal fibroblasts and KRT14 (magenta) marks basal keratinocytes.

(I) Fluorescence image of a representative sagittal section through the interfootpad region and mature eccrine gland of an adult *S100a4*-CreERT2/+; ROSA-mTmG/+ mouse at P22 that was given tamoxifen at P1.5 and P2.5, corresponding to the stage of placode formation in the interfootpad space. *S100a4*-lineage cells are shown in yellow. Immunofluorescence staining for KRT14 (cyan) marks basal keratinocytes and nuclei are counterstained with DAPI (magenta). Scale bars represent 50 μ m (A–I).

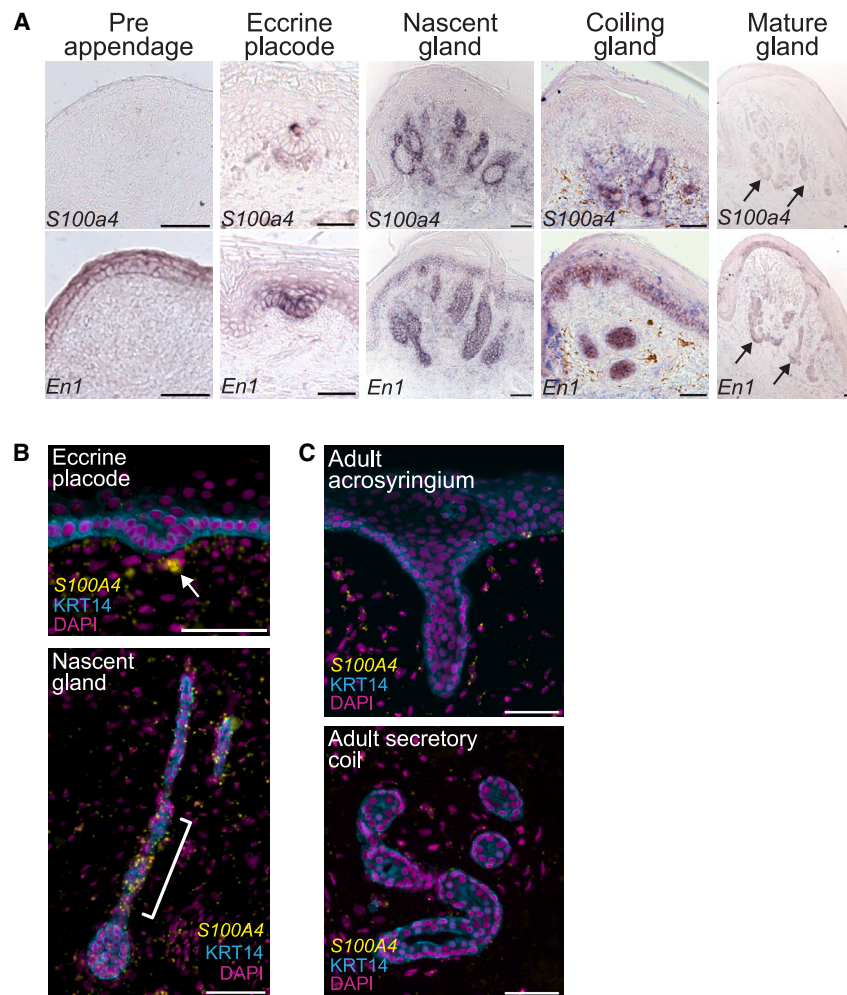


Figure 4. Evolutionary conservation of eccrine-associated dermal population

(A) *In situ* hybridization for *S100a4* (top, purple) and *En1* (bottom, purple) in volar skin at distinct stages of eccrine gland development: pre appendage (E15.5; footpad); eccrine placode (P2.5; inter-footpad); nascent eccrine gland (P2.5; footpad); coiling eccrine gland (P7; footpad); mature eccrine gland (P28; footpad).

(B) RNAscope for *S100a4* (yellow) and immunofluorescence for KRT14 (cyan) in developing human plantar skin sections (120 gestational days) containing eccrine placodes (top) and nascent eccrine glands (bottom).

(C) RNAscope for *S100a4* (yellow) and immunofluorescence for KRT14 (cyan) in adult human cheek skin sections (64 years). Top image shows the apical portion of a mature eccrine gland including the acrosyringium and part of the duct, and bottom image shows the secretory coil from the same gland. Nuclei are counterstained with DAPI (magenta) in fluorescence images (B and C). All scale bars represent 50 μ m (A–C).

An *En1*-dependent eccrine niche (EDEN) is required for eccrine gland development

In light of finding that the *En1*-dependent dermal populations derive from a single lineage, and the specific association of these populations with developing eccrine glands in the epidermis, we interrogated the snRNA-seq data for evidence of signaling between the epidermal and dermal nuclei from Control samples. We first performed subclustering (Derm0–Derm11) and lineage pseudotime inference on all dermal nuclei to resolve putative developmental relationships within the broader dermal dataset (Figures 5A and 5B). We identify an inferred dermal lineage that terminates in subcluster Derm10, which corresponds to the validated, nascent-gland-associated, dermal population represented in cluster 20 from the original analysis (Figures 3A and 5B). This lineage originates in Derm3 and successively progresses through Derm6, Derm9, and Derm2, before terminating in Derm10 (Figure 5B).

Having identified the subset of dermal nuclei that constitute the putative eccrine-associated dermal lineage, we performed cell-cell interaction modeling between this subset of dermal clusters and those which make up the validated eccrine epidermal lineage (Epi0, Epi3, Epi5, and Epi8; Figure 2G) using

CellChat.⁴⁸ We detect 185 significant receptor-ligand pairs from 32 pathways among the eccrine-associated dermal and epidermal subclusters ($p < 0.05$; Figure 5C, S5A, and S5B; Data S2). Of these, we identify 52 significant receptor-ligand interactions from 14 pathways that employ secreted signaling factors, including the WNT, BMP, TGF β , and non-canonical WNT (ncWNT) pathways (Figures S5A and S5B; Data S2). We identify receptor-ligand pairings consistent with bidirectional crosstalk between the lineages, with the

epidermal lineage signaling to the dermal lineage and vice versa (Figures 5C, S5A, and S5B). Moreover, we find that the signals mediating these interactions change over developmental time (Figures 5C, S5A, and S5B).

The signaling interactions predicted from the snRNA-seq expression data suggest that the dermal and epidermal eccrine lineages engage in reciprocal crosstalk over the course of eccrine development. This finding is intriguing given the well-established importance of dermal-ectodermal interactions for the developmental progression of other major ectodermal appendages, such as the hair follicle and mammary gland.⁸ Accordingly, we tested whether the *En1*-dependent dermal population is required for eccrine gland formation. For this purpose, we made use of the specific expression of *S100a4* in the mouse volar dermis to mark this population (Figures 3C–3G and 4A; Data S1C).

Using an inducible genetic system, we ablated *S100a4*-expressing cells at the onset of eccrine gland development using targeted expression of diphtheria toxin subunit alpha (DTA) and assessed eccrine gland density in the volar skin of Control (*S100a4*-CreERT2/+;DTA/+ or DTA/DTA) as compared with Ablated (*S100a4*-CreERT2/+;DTA/DTA or *S100a4*-CreERT2/*S100a4*-CreERT2;DTA/DTA) mice.⁴⁹ Timed pregnant females

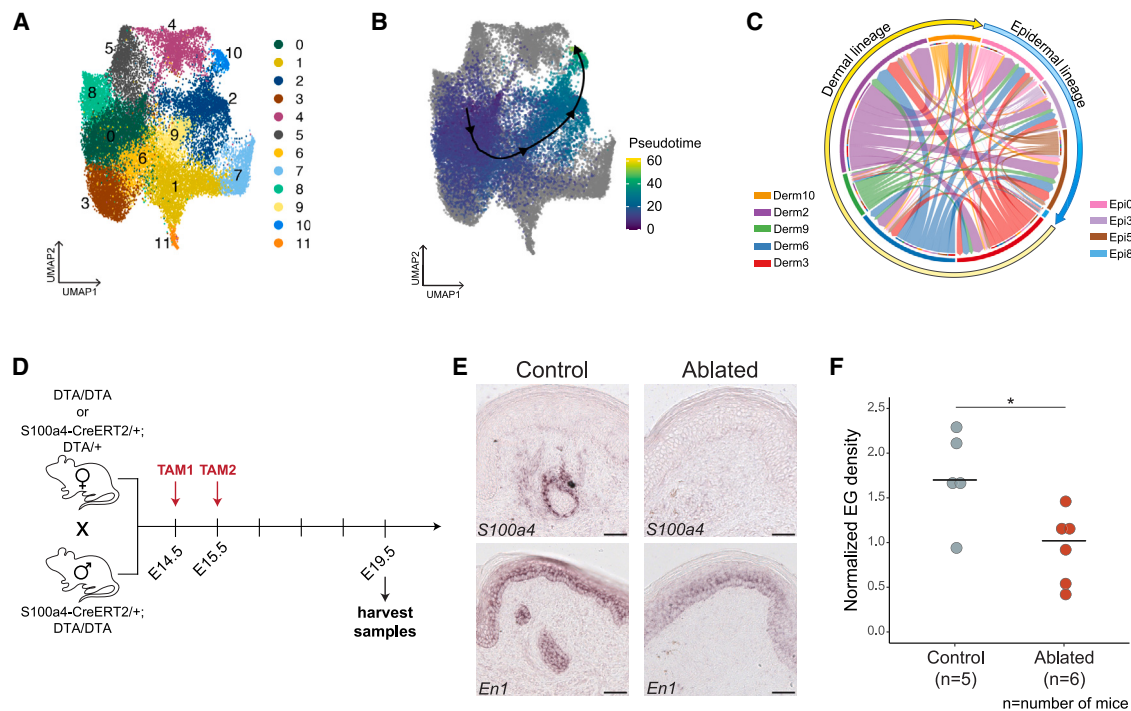


Figure 5. An *En1*-dependent eccrine niche (EDEN) is required for eccrine gland development

(A) UMAP projection of subclustered dermal nuclei colored by subcluster identity (merged Control and *En1*-cKO conditions). (B) Results of slingshot trajectory inference mapped onto the subclustered dermal UMAP embedding that depicts an inferred lineage through dermal subclusters 0, 3, 6, 9, 2, and 10. Points representing nuclei are colored according to pseudotime; nuclei not involved in this lineage are colored gray. (C) Summary chord diagram representing all significant signaling interactions between the dermal and epidermal lineage clusters inferred by CellChat ($p < 0.05$). Clusters are ordered around the circle based on their position in the inferred lineage. (D) Experimental scheme for genetic ablation of *En1*-dependent dermal population during eccrine gland development. (E) Representative images of *in situ* hybridization for *S100a4* and *En1* from adjacent serial sections of an Ablated sample (*S100a4*-CreERT2/+; DTA/DTA) compared with a littermate Control (DTA/DTA). (F) Quantification of nascent eccrine gland density in the hindlimb footpads of Ablated ($n = 6$ mice) and Control ($n = 5$ mice) mice. Each dot represents the density of eccrine glands in an individual mouse, and one foot was analyzed per mouse. Eccrine gland number for each mouse was normalized to the number of sections scored for the analyzed foot. Cross bars represent sample medians. Significance was assessed by a Mann-Whitney U test ($p = 0.0284$; $Z = 2.1909$; $r = 0.66$). All scale bars represent 50 μm .

were dosed with tamoxifen to induce Cre activation at E14.5 and E15.5 to specifically ablate *S100a4*-expressing cells around the time of footpad eccrine placode formation.^{9,11,12} At E19.5, when footpad eccrine glands are at the nascent stage, we quantified eccrine gland density in the footpads of Control and Ablated mice (Figure 5D).¹¹ This endpoint was selected due to dystocia in the dams, which necessitated endpoint cesarean section to deliver pups for analysis.

We observe a deficit of *S100a4* positive cells in the footpads of Ablated genotypes as compared with Controls (Figure 5E). Notably, *En1* expression persists in both Control and Ablated groups (Figure 5E). The impact of depleting the *S100a4* expressing dermal cells on eccrine gland development is striking. We find that eccrine gland density is on average 45% lower in Ablated compared with Control animals (Wilcoxon-Mann-Whitney test: $p = 0.0284$, $Z = 2.1909$, $r = 0.66$; Control: $n = 5$, Ablated: $n = 6$) (Figures 5E and 5F). Importantly, eccrine glands still present in Ablated animals retained normal *S100a4* expression around the gland, indicating that the population was not effectively depleted in these regions (Figure S5C). Taken together, our findings identify an *En1*-dependent ec-

crine niche (EDEN) in the dermis, direct proximity to which is required for eccrine gland development in the epidermis to proceed (Figure 6A).

DISCUSSION

In “The Variation of Animals and Plants Under Domestication,” Charles Darwin describes a family in Sindh (in present-day Pakistan) “in which ten men...were furnished, in the course of four generations...with only four small and weak incisor teeth and with eight posterior molars...have very little hair on the body, and ... suffer much during hot weather from excessive dryness of the skin⁵⁰”. One of the earliest formal accounts to suggest an underlying link between the formation of the group of organs we call ectodermal appendages, Darwin’s description also aptly captures the importance of eccrine glands in human physiology. More than 150 years later, our study uncovers a blueprint for eccrine gland development and identifies transcriptional and cellular transitions at which the ectodermal appendage developmental paradigm diverges to build these essential organs (Figure 6A).

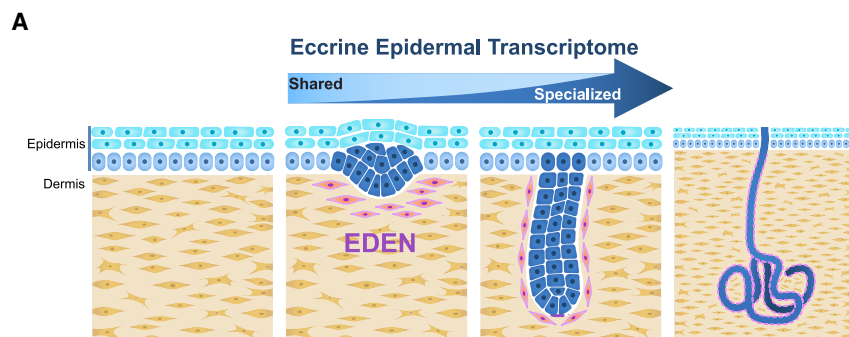


Figure 6. Transcriptional and cellular identifiers of eccrine gland development

(A) Model of eccrine gland development incorporating findings reported in this study. Eccrine gland development coordinates a progressive shift in the epidermal transcriptome from one that is predominantly shared with hair follicles, to one that is specialized and eccrine-specific, and requires the *Engrailed 1*-dependent eccrine niche (EDEN; primary niche and its derivative lineage are in pink) within the adjacent dermis. Epidermis is in blue, and dermis exclusive of EDEN is shown in yellow. Pre- and inter-appendage basal keratinocytes are light blue, suprabasal keratinocytes are cyan, and eccrine cells are dark blue. Dermal cells are dark yellow.

Duality and temporal transitions in the eccrine gland epidermal program during development

We find that the upregulation of epidermal *En1* is necessary for a transcriptional signature that distinguishes the eccrine placode from that of the hair follicle. That this signature includes upregulation of the Wnt inhibitor *Dkk4* is intriguing since previous studies have postulated that differences in the levels of canonical Wnt signaling determine the fate of ectodermal appendage placodes, with hair placodes proposed to require the highest level of Wnt signaling among the different appendage types.^{12,44,51,52}

We find that the restricted upregulation of *Dkk4* to the eccrine placode is associated with the formation of more eccrine glands in mice (Figures 2K and 2L). This suggests that spatial as well as quantitative parameters are integrated into the transcriptomic signature of the eccrine placode. Consistent with this, placode-specific *Dkk4* expression is critical for the reaction diffusion mechanism regulating the formation of skin appendages and disruption of this pattern through ectopic expression of this gene throughout the basal epidermis was reported to reduce eccrine gland density.^{10,44} Resolving the functional importance of patterned *Dkk4* upregulation and of the other *En1*-dependent transcriptional changes, we identified will set the stage for uncovering the intrinsic program that confers eccrine identity in the epidermis.

The relative paucity of the specialized eccrine transcriptome as compared with the bulk of placodal transcripts that are shared between eccrine gland and hair follicle progenitors during early development contrasts with the expression profile of later stages. Our data indicate that as development proceeds, differentiation of the eccrine cell types is coupled with a concomitant shift in the balance of the eccrine transcriptome toward a more specialized and distinctive expression profile (Figure 6A).

The extensive overlap we find in the transcriptome of hair and eccrine placodes may help to explain why previous efforts have also identified a largely shared set of signals in the early development of ectodermal appendages, irrespective of type.^{8,10,12,14,53,54} In particular, the relative dearth of eccrine placode-enriched transcripts as compared with the far greater pool of those that are generalized across hair and eccrine types has intriguing implications for the character of the elusive, first dermal signal that triggers appendage fate and formation.^{12,42,55,56} Our observations raise the possibility that the first inductive signal from the dermis, which initiates eccrine development, is also dualistic in nature. Given the relative proportions of

the two classes of placodal transcripts captured in our study, the eccrine dermal signals may show a similar skew. This would in turn reduce the power to detect the specialized dermal inducers within the context of the greater proportion of those that are generalized across the different appendage types. The identification of the subset of transcripts that are differentially enriched in eccrine versus hair follicle placodes provides a directed means to parse out the precise composition of these upstream signals from the dermis.

EDEN—A dermal niche for eccrine development

We find that the progression of the epidermal program in the eccrine placode requires the presence of EDEN, a dermal niche for eccrine glands (Figure 6A). Importantly, our data suggest that EDEN is not the source of the first dermal inductive signal that initiates placode formation. That the transcriptional identity of EDEN depends on the upregulation of ectodermal *En1*, and that EDEN is found in association with eccrine gland epidermal anlagen, but not other *En1*-expressing cells, suggests that EDEN is analogous to the dermal condensations that form in response to the placodes of other major appendages such as teeth and hair follicles.⁸ Future experiments to distinguish between the roles of *En1* in the placode as opposed to the inter-appendage epidermis in the induction of EDEN will help to clarify this. A starting point for these investigations and for functional dissection of the signals mediating interactions between the developing eccrine glands and the niche are the candidate factors identified in this study. It is also important to understand whether the EDEN lineage is required for eccrine development and homeostasis beyond the placode stage in order to establish its closest functional homology to the niches of other appendages. This is because while the hair follicle dermal condensate (and its derivative the dermal papilla) is persistently required, the dense mesenchyme that directs early mammary gland development is transient and its role supplanted by the fat pad during the later stages of mammogenesis.^{8,53}

Irrespective of the precise source in the epidermis, our data demonstrate that a cell-autonomous transcriptional repressor, *EN1*, which is expressed in the basal epidermis, is required to induce a distinct cellular identity in the underlying dermal compartment. There is precedence for this, as evidenced by the early and the separable role of ectodermal *En1* in maintaining the identity of the ventral limb bud by inhibiting mesenchymal *Lmx1b* expression.^{19,32,57} In this context, *En1* acts by inhibiting

Wnt7a in the ventral limb ectoderm.⁵⁷ Having identified a pool of *En1*-dependent transcripts in the eccrine placode and in the skin basal epidermis, our findings make it possible to determine whether *En1* effects are mediated by similar or entirely independent effectors in ectodermal appendage development. Future studies to interrogate the function of these EN1 targets are also important to understand and distinguish between EN1's roles in specifying the dermal niche, and in directly acting on the epidermal populations that make up the gland itself.

Intriguingly, we find that despite the continued expression of *En1* in adult basal keratinocytes in the eccrine-forming regions and in the mature eccrine glands themselves, the expression of *S100a4*, which specifically marks the EDEN lineage throughout the eccrine developmental period, does not persist in its descendants around the mature eccrine glands of either mice or humans.^{35,58} This suggests that the dermal niche, at least as it exists during development, may not be retained once eccrine gland formation is completed and may help to explain why eccrine glands exhibit limited ability to regenerate.⁹ Previous studies have demonstrated the existence of several, unipotent stem cell populations within the mature eccrine gland, in the duct and the secretory coils, respectively.⁹ In response to injury, these stem cells can only be mobilized to repair their respective lineages, but not the whole gland.⁹ Comparing the functional properties of EDEN lineage cells and the significance of gene expression changes therein during eccrine gland development and homeostasis may help to resolve the contribution of extrinsic factors to this regenerative limitation.

Notably, the hair follicle dermal niche is not only necessary for the regeneration of the hair follicle but is also sufficient to induce epithelial cells to form *de novo* hair follicles, even in adulthood.^{59,60} It is intriguing to speculate that EDEN may have comparable properties with respect to inducing *de novo* eccrine gland formation. Understanding whether the molecular effectors of EDEN demonstrate such eccrine-inducing capabilities has the potential to seed efforts to repair wounds and burns with skin regenerates that contain eccrine glands. Coupled with the capture of an eccrine epidermal transcriptome during development, the identification of EDEN not only addresses the long-standing developmental question of how the dermal and epidermal skin compartments of mammalian skin are differentially mobilized to build eccrine glands but also sets the stage for meeting a critical need in human regenerative medicine.

Limitations of the study

Our study analyzes the nuclear transcriptome, which represents a fraction of the total transcriptome of the cell. Therefore, there may be additional differential transcripts that are not captured by our study. Despite this, the validations we performed indicate our findings from the nuclear transcriptome are biologically relevant and are representative of the heterogeneity in developing eccrine glands.

STAR★METHODS

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SUPPLEMENTAL INFORMATION

Supplemental information can be found online at <https://doi.org/10.1016/j.devcel.2023.11.015>.

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AUTHOR CONTRIBUTIONS

Y.G.K. conceived of and supervised the study. Y.G.K. and H.L.D. designed the experiments and wrote the manuscript. H.L.D., R.R.T., A.A., B.K., P.H., Q.Q., S.M.M., J.C.D., D.A., and M.M. performed the experiments. H.L.D. and Y.G.K. analyzed and interpreted the data. H.W., I.A.G., the BDRL, and Y.G.K. contributed resources and acquired funding to support the study.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Rabbit anti-CK14 (1:10,000)	BioLegends	Cat # 905303; RRID: AB_2734678
Chick polyclonal anti-GFP (1:1000)	Abcam	Cat# ab13970; RRID: AB_300798
Goat polyclonal anti-EDAR (1:100)	R&D Systems	Cat# AF745; RRID: AB_355565
Goat polyclonal anti-PDGFRα (1:100)	R&D Systems	Cat# AF1062; RRID: AB_2236897
Goat polyclonal anti-chicken AF-488 (1:500)	Invitrogen	Cat#A-11039; RRID: AB_2434096
Donkey anti-goat AF-488 (1:200)	Abcam	Cat# ab150129; RRID: AB_2687506
Donkey anti-rabbit AF-594 (1:250)	Jackson ImmunoResearch	Cat# 711-585-152; RRID: AB_2340621
Donkey anti-rabbit AF-647 (1:250)	Jackson ImmunoResearch	Cat# 711-605-152; RRID: AB_2492288
Biological samples		
120 gestational days human plantar foot skin	Birth Defects Research Laboratory, University of Washington (UBMTA: 48286A)	N/A
64 year old human cheek skin	Skin Biology and Disease Resource Center, University of Pennsylvania	N/A
Chemicals, peptides, and recombinant proteins		
Nile Blue A	Sigma-Aldrich	Cat# N5632
Oil Red O	Sigma-Aldrich	Cat# O0625
Dispase II	Sigma-Aldrich	Cat# D4693
4',6-Diamidino-2-phenylindole	Sigma-Aldrich	Cat# D9542
Critical commercial assays		
NextSeq 500/550 75-cycle High Output v2.5 kit	Illumina	Cat# 20024906
Advantage® UltraPure PCR Deoxynucleotide Mix (10 mM each dNTP)	Clontech	Cat# 639125
Exonuclease I	NEB	Cat# M0293L
NxGen® RNase Inhibitor	Lucigen	Cat# 30281-2
Maxima H Minus Reverse Transcriptase (200 U/μL)	Life Technologies	Cat# EP0753
KAPA HiFi HotStart ReadyMix	KAPA BioSystems	Cat# KK2602
Barcoded Bead SeqB	ChemGenes	Cat# MACOSKO-2011
Nextera® XT DNA Sample Preparation Kit (96 Samples)	Illumina	Cat# FC-131-1096
Nextera® XT Index Kit (24 indexes, 96 samples)	Illumina	Cat# FC-131-1001
Agilent High Sensitivity DNA Kit	Agilent Technologies	Cat# 5067-4626
Exonuclease I	NEB	Cat# M0293L
QX200™ Droplet Generation Oil for EvaGreen	Bio-Rad	Cat# 1864006
N-Lauroylsarcosine sodium salt solution	Sigma-Aldrich	Cat# L7414-50ML
1H,1H,2H,2H-Perfluoro-1-octanol	Sigma-Aldrich	Cat# 370533-25G
SPRIselect reagent	Beckman Coulter	Cat# B23318
Fluorescent microspheres	Bangs Labs	Cat# FC06F
RNAscope 2.5 HD-RED assay kit	ACD Bio	Cat# 322350
human S100A4 RNAscope probe	ACD Bio	Cat# 422071
human PPIB RNAscope probe	ACB Bio	Cat# 313901
DapB negative control RNAscope probe	ACD Bio	Cat# 310043

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Deposited data		
Raw and processed snRNA-seq data	This paper	GEO: GSE220977
Experimental models: Organisms/strains		
Mouse: tetO-Cre: <i>Tg(tetO-cre)1Jaw/J</i>	The Jackson Laboratory; Perl et al. ³⁷	RRID: ISMR_JAX:006224
Mouse: Krt5-rtTA: <i>Tg(KRT5-rtTA)T2D6Sgkd/J</i>	Laboratory of Dr. Sarah Millar (Icahn School of Medicine at Mt. Sinai); Diamond et al. ²¹	RRID: ISMR_JAX:017519
Mouse: En1 ^{fllox} : <i>En1tm8.1Alj/J</i>	Laboratory of Dr. Alexandra Joyner (Memorial Sloan Kettering Cancer Center); Sgaier et al. ²⁷	RRID: ISMR_JAX:007918
Mouse: Lgr6-eGFP: <i>Lgr6tm2.1^{(cre/ERT2)Cle}/J</i>	Laboratory of Dr. Pantelis Rumpolas (University of Pennsylvania); Snippert et al. ⁶²	RRID: ISMR_JAX:016934
Mouse: S100a4-CreERT2: S100a4-T2A ^(CreERT2)	Dr. Mayumi Ito (New York University)	N/A
Mouse: ROSA-DTA: Gt(ROSA)26Sor ^{tm1(DTA)Lky/J}	The Jackson Laboratory; Voehringer et al. ⁴⁹	RRID: ISMR_JAX:009669
Mouse: mTmG: Gt(ROSA)26Sor ^{tm4(ACTB-tdTomato,-EGFP)Luo/J}	Laboratory of Dr. George Cotsarelis (University of Pennsylvania); Muzumdar et al. ⁶⁴	RRID: ISMR_JAX:007576
Mouse: ROSA-tdTomato: Gt(ROSA)26Sor ^{tm14(CAG-tdTomato)Hze/J}	Laboratory of Dr. Pantelis Rumpolas (University of Pennsylvania); Madisen et al. ⁶³	RRID: ISMR_JAX:007914
Oligonucleotides		
Primers for amplifying cDNA to make <i>in situ</i> hybridization probes, see Table S4	This paper	N/A
Primers for mice, see Table S4	This paper	N/A
Primers for qRT-PCR, see Table S4	This paper	N/A
Software and algorithms		
FIJI	Schindelin et al. ⁶⁵ (https://doi.org/10.1038/nmeth.2019)	N/A
Drop-seq Tools v1.12	Macosko et al. ⁶⁶ (https://doi.org/10.1016/j.cell.2015.05.002)	N/A
Seurat v4.0.2	Hao et al. ⁶⁷ (https://doi.org/10.1016/j.cell.2021.04.048)	N/A
Harmony	Korsunsky et al. ²⁵ (https://doi.org/10.1038/s41592-019-0619-0)	N/A
Slingshot	Street et al. ⁷⁰ (https://doi.org/10.1186/s12864-018-4772-0)	N/A
DESeq2	Love et al. ⁶⁹ (https://doi.org/10.1186/s13059-014-0550-8)	N/A
ClusterProfiler	Wu et al. ⁶⁸ (https://doi.org/10.1016/j.xinn.2021.100141)	N/A
CellChat	Jin et al. ⁴⁸ (https://doi.org/10.1038/s41467-021-21246-9)	N/A
Other		
Leica M680 microscope	Leica	N/A
Leica IC90E camera	Leica	N/A
Leica DM5500B microscope	Leica	N/A
Leica DFC 500 camera	Leica	N/A
Leica DFC 360X camera	Leica	N/A
Qubit 3.0	Invitrogen	N/A
Applied Biosystems QuantStudio 7 Flex real-time PCR machine	ThermoFisher Scientific	REF# 4485701
Bioanalyzer	Agilent	N/A
Illumina NextSeq 500	Illumina	N/A

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Dox Diet, Grain-Based, Doxycycline (6 gm/kg), Green, 1/2" Pellets Gamma irradiated	Bio-Serv	Cat# S4096
PDMS co-flow microfluidic droplet generation device	uFluidix	N/A

RESOURCE AVAILABILITY

Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the corresponding author, Yana G. Kamberov ([yana2@pennmedicine.upenn.edu](mailto: yana2@pennmedicine.upenn.edu)).

Materials availability

This study did not generate new, unique reagents.

Data and code availability

Raw and processed snRNA-seq data files have been deposited in NCBI's the Gene Expression Omnibus⁶¹ and are accessible through GEO Series accession number GSE220977 (<https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE220977>). Any additional information required to reanalyze the data reported in this paper is available from the [lead contact](#) upon request.

EXPERIMENTAL MODEL AND STUDY PARTICIPANT DETAILS

Animals

*Tg(tetO-cre)1Jaw/J (tetO-Cre)*³² and *Gt(ROSA)26Sor^{tm1(DTA)Lky}/J (ROSA-DTA)*⁴⁸ mice were purchased from the Jackson Laboratory. *Tg(KRT5-rtTA)T2D6Sgkd/J (Krt5-rtTA)*³³ mice were provided by Dr. Sarah Millar (Icahn School of Medicine at Mount Sinai), *En1tm8.1Alj/J (En1^{fllox})*³¹ mice were provided by Dr. Alexandra Joyner (Memorial Sloan Kettering Cancer Center) and *Lgr6tm2.1(cre/ERT2)Cle/J (Lgr6-eGFP)*⁶² and *Gt(ROSA)26Sor^{tm14(CAG-tdTomato)Hze}/J (ROSA-tdTomato)*⁶³ mice were provided by Dr. Pantelis Rompolas (University of Pennsylvania). S100a4-CreERT2 mice were generated by Dr. Mayumi Ito (New York University) and provided under material transfer agreement. *Gt(ROSA)26Sortm4(ACTB-tdTomato,-EGFP)Luo/J (mTmG)*⁶⁴ mice were provided by Dr. George Cotsarelis (University of Pennsylvania). En1^{fllox}, Krt5-rtTA, and tetO-Cre mice were maintained on an FVB/N background (Charles River Laboratories) for more than 10 generations. ROSA-DTA mice were maintained on a C57BL/6J background (Jackson Laboratory) until they were bred to S100a4-CreERT2 mice, which were maintained on a mixed genetic background. Lgr6-eGFP mice were also maintained on a mixed genetic background. FVB/N and C57BL/6N wild-type mice were obtained from Charles River Laboratories and Taconic Biosciences, respectively. Both males and females were used for all experiments in this study.

For inducible En1 conditional knock-out (En1-cKO) experiments, male mice harboring the tetO-Cre, Krt5-rtTA, and homozygous En1^{fllox} alleles were mated to females homozygous for En1^{fllox} and carrying the Krt5-rtTA allele. Timed pregnant dams were continuously administered a doxycycline dosed diet (6 g/kg) *ad libitum* beginning at E16.5. Offspring were euthanized at either P2.5 or P28 via decapitation or CO₂ inhalation, respectively.

For embryonically induced S100a4 lineage tracing experiments, male S100a4-CreERT2/+ mice were mated to ROSA-tdTomato homozygous females. Timed pregnant females were dosed with tamoxifen in corn oil via oral gavage (3 mg/kg) at E15.5 and pups were euthanized at P2.5 for volar hindpaw collection. For perinatally induced S100a4 lineage tracing, male S100a4-CreERT2/+;mTmG/+ mice were mated to mTmG/+ or mTmG/mTmG females and pups were dosed orally with tamoxifen dissolved in corn oil (10 μl of 50mg/ml stock) at P1.5 and P2.5, coinciding with placode stage of IFP eccrine glands in the volar skin, and volar skin from the forelimb and hindlimb was harvested at P22.

For S100a4+ cell ablation experiments, male mice harboring at least one copy of the S100a4-CreERT2 and ROSA-DTA alleles were mated to females either homozygous for ROSA-DTA or heterozygous for both S100a4-CreERT2 and ROSA-DTA. Timed pregnant females were dosed with tamoxifen in corn oil via oral gavage (3 mg/kg) at both E14.5 and E15.5. Pregnant dams were euthanized at E19.5 (developmental equivalent of P0) and pups were harvested via endpoint caesarean section due to dystocia in the dams.

All mice used in this study were housed on a 12h light/dark cycle in a University Laboratory Animal Resources (ULAR) managed vivarium at the Perelman School of Medicine (PSOM). All procedures were performed in accordance with guidelines established by the National Institutes of Health and have been approved by the University of Pennsylvania PSOM Institutional Animal Care and Use Committee (protocol # 806105). Mice were housed under standard laboratory conditions and received food and water *ad libitum*.

Human tissue

Specimens from fetal (120 gestational days) human plantar foot skin were obtained from the Birth Defects Research Laboratory at the University of Washington (Uniform Biological Material Transfer Agreement #48286A) with ethics board approval and maternal written consent. Deidentified samples from adult human cheek skin (64 years old) were obtained from the Skin Biology and Disease Resource Center at the University of Pennsylvania with ethics board approval and written consent. This study was performed in accordance with ethical and legal guidelines of the University of Pennsylvania institutional review board.

METHOD DETAILS

Quantification of appendages in adult volar skin

Quantification of eccrine glands and hair follicles at P28 was performed in whole mount preparations of hindlimb volar skin as previously described.¹⁶ Briefly, the epidermis was separated from the underlying dermis by dispase digestion and fixed with 10% neutral buffered formalin. Hair follicle-associated sebaceous glands were stained with Oil Red O and appendages (eccrine glands and hair follicles) were stained with Nile Blue, and whole mounts were imaged on a Leica M680 dissection scope fitted with a Leica IC90E camera. All data reported in this manuscript represents the average counts from the right and left hindlimb volar skin of an individual mouse.

Isolation and purification of nuclei for snRNA-seq

En1-cKO and Control pups were euthanized at P2.5 and volar hindlimb skin (right and left) was rapidly dissected on ice and snap frozen on dry ice and stored at -80°C until nuclear isolation. Single nuclear suspensions were generated from the pooled volar hindlimb skin (right and left) from 5 mice of the same genotype, for a total of 10 pooled volar skins, as previously described with some modifications.²³ Pooled samples were dissociated in homogenization buffer using a Polytron homogenizer (Kinematica Inc.) followed by dounce homogenization (20 times with loose pestle; 10 times with tight pestle). Suspensions were filtered to remove debris and single nuclear suspensions were quantified via hemocytometer after staining with Trypan Blue, which stains all nuclei. For P2.5 volar skin, we obtained $\geq 2.1 \times 10^5$ nuclei/ml for each sample. Sucrose gradient centrifugation was not performed on single nuclear suspensions. Biological replicates were prepared and sequenced in two separate batches; one replicate of each genotype (En1-cKO: tetO-Cre/+; Krt5-rtTA/+; En1^{flox/flox}; Control: En1^{flox/flox}; Krt5-rtTA/+; En1^{flox/flox}) was processed and sequenced in the same batch to avoid autocorrelation of condition and batch effects.

snRNA-seq library preparation and sequencing

Single nuclei were co-encapsulated with barcoded beads (ChemGenes) and reverse transcription was performed as previously described.²³ Optimal PCR cycle number for library amplification was determined via qPCR using 6,000 beads, cDNA was tagged, and libraries were further amplified as previously described.²³ cDNA libraries were quantified via Qubit 3.0 (Invitrogen) and library quality was determined via Bioanalyzer (Agilent) prior to sequencing on an Illumina NextSeq 500 using the 75-cycle High Output v2 kit (Illumina). In total, six neonatal volar skin snRNA-seq libraries were sequenced in four sequencing runs. See open access snRNA-seq protocol for more details: <https://www.protocols.io/view/snucdrop-seq-protocol-n2bvjr36plk5/v2>.²³

In situ hybridization, immunofluorescence, RNAscope, and imaging

Volar skin was fixed in 4% PFA, cryoprotected using sucrose, and embedded in OCT (Tissue Tek) for cryo-sectioning at a thickness of 10–12 μ m. Sections were collected in 3–4 series for staining of adjacent sections. *In situ* hybridization was performed as previously described¹⁹ with custom anti-sense DIG-labeled riboprobes transcribed *in vitro* (Roche) from amplified cDNA (see Key Resources Table for primers). Immunofluorescence was performed for KRT14 (1:10,000, BioLegend 905303), EDAR (1:100, R&D Systems AF745), PDGFRA (1:100, R&D Systems AF1062), and GFP (1:1000, Abcam ab13970) after blocking tissue in PBT (0.2% Triton-X100 in PBS) with 10% serum. Secondary detection was performed with antibodies conjugated to Alexa Fluor⁴⁸⁸ (Invitrogen A-11039 or Abcam ab150129), Alexa Fluor⁵⁹⁴ (Jackson ImmunoResearch 711-585-152), or Alexa Fluor⁶⁴⁷ (Jackson ImmunoResearch 711-605-152) and samples were counterstained with 4',6-diamidino-2-phenylindole (DAPI, Sigma D9542).

FFPE adult and fetal human tissue was processed and sectioned by SBDR Core A. RNAscope was performed on 5 μ m paraffin sections after standard target retrieval for *S100A4* (ACD Bio #422071), *PPIB* (positive control; ACD Bio #313901), and *DapB* (negative control; ACD Bio #310043) using an RNAscope 2.5 HD-RED assay kit (ACD Bio #322350). Subsequently, immunofluorescence for KRT14 was performed as described above and samples were counterstained with DAPI.

Samples were imaged on a Leica DM5500B microscope equipped with Leica DFC 500 (bright field) and Leica DFC 360X (fluorescence) cameras. Images were processed using FIJI software.⁶⁵

Scoring of footpad eccrine glands at E19.5

S100a4 Ablated and Control samples were harvested as described above and whole hindlimbs were cryo-sectioned at 10 μ m and three adjacent series of sections were collected. The first series was stained with hematoxylin and eosin (H&E, Sigma Aldrich) for scoring. The second and third series were used for *in situ* hybridization for *En1* and *S100a4*, respectively, to confirm efficient ablation

of *S100a4*+ cells. Scoring of E19.5 Hematoxylin and eosin-stained sections was performed by counting all nascent eccrine glands in the footpads throughout the entire series of sections. Eccrine counts were then normalized to the number of sections scored to yield an eccrine density estimate.

RNA extraction and qRT-PCR

FVB/N and C57BL6/N neonatal pups were euthanized at P2.5 and volar hindpaw skin was harvested and snap frozen as described above. Biological replicates are comprised of left and right hindpaw skin pooled from 3 mice. Pooled skin samples were dissociated using a Polytron homogenizer and total RNA was isolated using TRIzol (Life Technologies) extraction followed by clean up and on-column DNase I treatment using a RNeasy Mini Kit (Qiagen). cDNA was reverse transcribed using SuperScript III (Thermo Fisher) following the manufacturer's instructions. qRT-PCR was performed in biological triplicate and technical quadruplicate using Power SYBR PCR master mix (Thermo Fisher) for *Dkk4* and *En1*. *Rpl13a* served as a reference gene for normalizing Ct values. Each data point reported in Figure 2L represents the mean of technical replicates for a sample. Primers used for qRT-PCR are found in Table S4.

QUANTIFICATION AND STATISTICAL ANALYSIS

snRNA-seq data pre-processing

Paired-end snRNA-seq reads were processed using publicly available the Drop-seq Tools v1.12 software⁶⁶ with modifications described previously.²³ A digital expression matrix was generated by assembling a list of UMIs in each gene (as rows) within each cell (as columns), and UMIs within ED = 1 were merged.

snRNA-seq cluster identification and marker gene analysis

Digital expression matrices for each sample were loaded into Seurat v4.0.2⁶⁷ and merged. UMI counts were normalized by scaling by library size, multiplying by 10,000, and transforming to log scale. Genes were filtered out if they were expressed in <10 nuclei and nuclei were filtered out if they contained a high proportion of UMIs mapped to mitochondrial genes (≥ 0.05), fewer than 300 or more than 6000 detected genes, resulting in 45,370 nuclei in the final filtered dataset.

Prior to clustering, individual sample datasets were integrated using the Harmony algorithm,²⁵ the top 2000 variable genes were identified using the *FindVariableFeatures* function in Seurat with the VST selection method, and expression of these variable genes was scaled and centered for principal components analysis. Based on the cumulative standard deviations of each principal component (PC), visualized by the function *ElbowPlot* in Seurat, we selected the first 40 PCs for two-dimensional uniform manifold approximation and projection (UMAP) implemented by Seurat *RunUMAP* with default parameters. Initial clustering on the full dataset (all nuclei, merged conditions) identified 23 clusters with the resolution parameter for *FindClusters* set to 0.7. Based on the expression pattern of well-established marker genes in this dataset, we assigned 25,859 nuclei to dermal cells (~57% of our data), 9,070 nuclei to epidermal cells (~20% of our data), and 10,441 nuclei to various other cell types (see Figure S2). Differential expression analysis was performed to identify marker genes for each of these 23 initial clusters using Seurat's *FindAllMarkers* function with Wilcoxon test. Differentially upregulated genes that were expressed in at least 25% of nuclei with a log fold change (LFC) ≥ 0.58 were considered marker genes (Data S1A). Dermal cluster specific marker genes were identified using the same approach, but with only dermal clusters of nuclei compared to each other (Data S1C).

Subclustering, pseudobulk differential expression, trajectory inference, and CellChat analysis

Nuclei from clusters determined to be epidermal (original clusters 2, 9, 10, 14, and 15) or dermal (original clusters 0, 1, 3, 4, 5, 8, 11, 20) were subsetted, the top 2000 variable genes were identified, and PCA was performed as above. The top 40 PCs were selected for clustering analysis with a resolution of 0.4 for epidermal nuclei and 0.7 for dermal nuclei, leading to the identification of 9 epidermal subclusters and 11 dermal subclusters. Of these 9 epidermal subclusters, 6 are basal (6,799 nuclei) and 3 of these subclusters are suprabasal (2,271 nuclei). Marker genes of each subcluster were identified as described above (Data S1B and S1C) and gene ontology (GO) enrichment analysis was performed on these marker gene lists using the *compareCluster* function from the Cluster Profiler R package.⁶⁸

We performed pseudo-bulk analysis to identify differentially expressed transcripts between Control and En1-cKO placode nuclei (Epi3). Pseudobulk samples were created using the Seurat function *PseudobulkExpression* on sample counts for each subcluster using the aggregate pseudo-bulk method. Differential expression analysis was performed on these pseudo-bulk samples for Epi3 using DESeq2⁶⁹ with a likelihood ratio test. Genes were deemed significantly differentially expressed between Control and En1-cKO pseudo-bulk samples if p adjusted < 0.01 and LFC ≥ 0.58 .

Trajectory inference of the subclustered nuclei was performed using Slingshot⁷⁰ on either the subsetted epidermal or dermal PCA reduction (merged conditions). Epi0 was supplied as the starting cluster to the *slingshot* function for the epidermal trajectory inference; for dermal trajectory inference, subcluster Derm11 was provided as the end cluster. The inferred lineages were then mapped onto UMAP embeddings for visualization using the *embedCurves* Slingshot function. Differential progression analysis was performed on the epidermal lineage via Kolmogorov-Smirnov test of the lineage pseudotime distributions between Control and En1-cKO nuclei.

CellChat⁴⁸ was performed on Control nuclei from the dermal and epidermal subclusters that were determined to be involved in the relevant inferred lineages using the default CellChat mouse database. Average gene expression per cluster was computed using the "truncatedMean" method (trim=0.1) within CellChat's *computeCommunProb* function. All significant receptor-ligand interactions

($p < 0.05$) identified between any pair of clusters are summarized as a chord diagram (Figure 4C). Complete results are provided as a supplemental data file (Data S2).

Statistical analysis of phenotypic data and qRT-PCR expression

Statistical parameters, including sample sizes, are reported in figures, figure legends, tables, and/or corresponding results text. A Kruskal-Wallis test followed by a *post-hoc* Dunn's test with Benjamini-Hochberg correction for multiple comparisons was performed on P28 appendage counts for En1-cKO phenotyping ($\alpha = 0.05$). A Mann-Whitney-Wilcoxon test was used for pairwise comparisons of qRT-PCR data ($\alpha = 0.05$). All statistical analyses were performed in R⁷¹.

Supplemental information

**Sweat gland development requires an eccrine
dermal niche and couples two epidermal programs**

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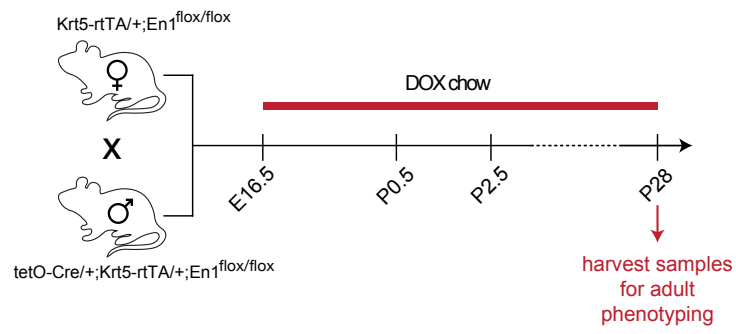
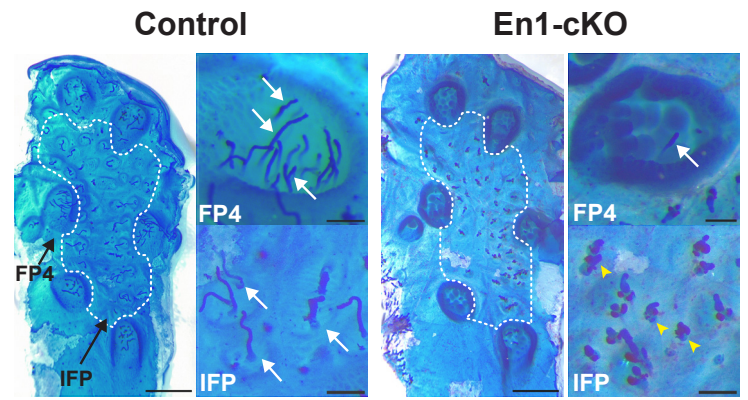
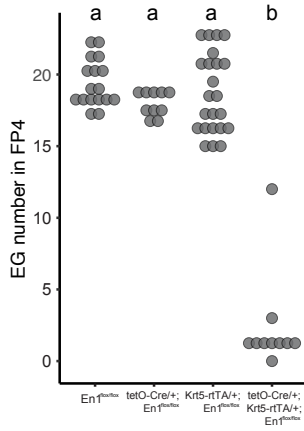
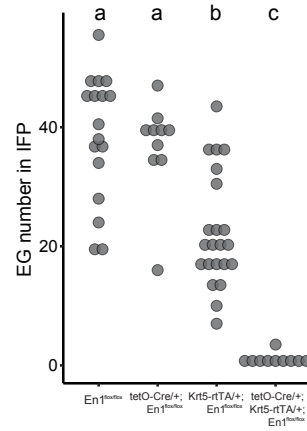
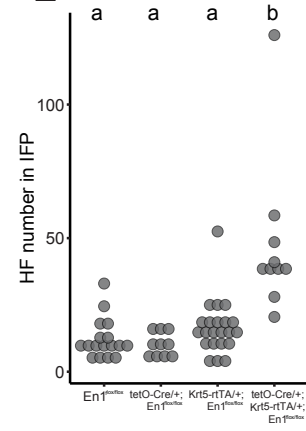
FIGURE S1**A****B****C****D****E**

Figure S1. Appendage counts in Control vs. En1-cKO volar skin split by genotype, related to Figure 1

A) Mating scheme to generate animals, dosing schedule, and time points for analyses performed in this study. All En1-cKO and Control mice analyzed in this study were administered doxycycline (DOX) starting on embryonic day (E) 16.5, when eccrine placodes first form in the footpads, until day of harvest. Volar hindpaw skin was harvested on post-natal day (P) 28 for adult phenotyping. B) Representative whole-mount preparations of P28 Control (left) and En1-cKO (right) hindlimb, volar skin. All appendages are stained with Nile blue (blue color) and hair follicle-associated sebaceous glands are stained with Oil Red O (red color). Footpad 4 is labeled and indicated by a black arrow. White dotted line outlines the interfootpad region (IFP). Scale bars in whole-mount images represent 1mm. Insets show higher magnification images of footpad 4 (FP4; top), and of the interfootpad regions (IFP bottom) of Control and En1-cKO skin. Eccrine gland (white arrow). Hair follicle (yellow arrowhead). Inset scale bars represent 200 μ m. C) Plots of eccrine gland (EG) number in footpad 4 (FP4) showing samples split according to the specific genotypes that comprise Control and En1-cKO groups. D) Quantification of eccrine gland number in the interfootpad (IFP) from Figure 1E with samples split by genotypes that comprise Control and En1-cKO groups. E) Quantification of hair follicle (HF) number in the interfootpad region from Figure 1F with samples split by genotypes that comprise control and En1-cKO groups. Dots in quantification plots (C-D) represent the average of counts from left and right volar skin of one mouse, which constitutes a biological replicate. N = number of mice. Sample sizes are as follows: En1^{flox/flox} n=17; tetO-Cre/+;En1^{flox/flox} n = 10; Krt5-rtTA/+;En1^{flox/flox} n = 23; tetO-Cre/+;Krt5-rtTA/+;En1^{flox/flox} n = 10. Letter annotations on all plots (C-D) indicate significant pairwise differences found via Kruskal-Wallis followed by Dunn *post-hoc* testing (p<0.05). Groups annotated with the same letter are not significantly different from each other. Statistics reported in Tables S1-S3.

FIGURE S2

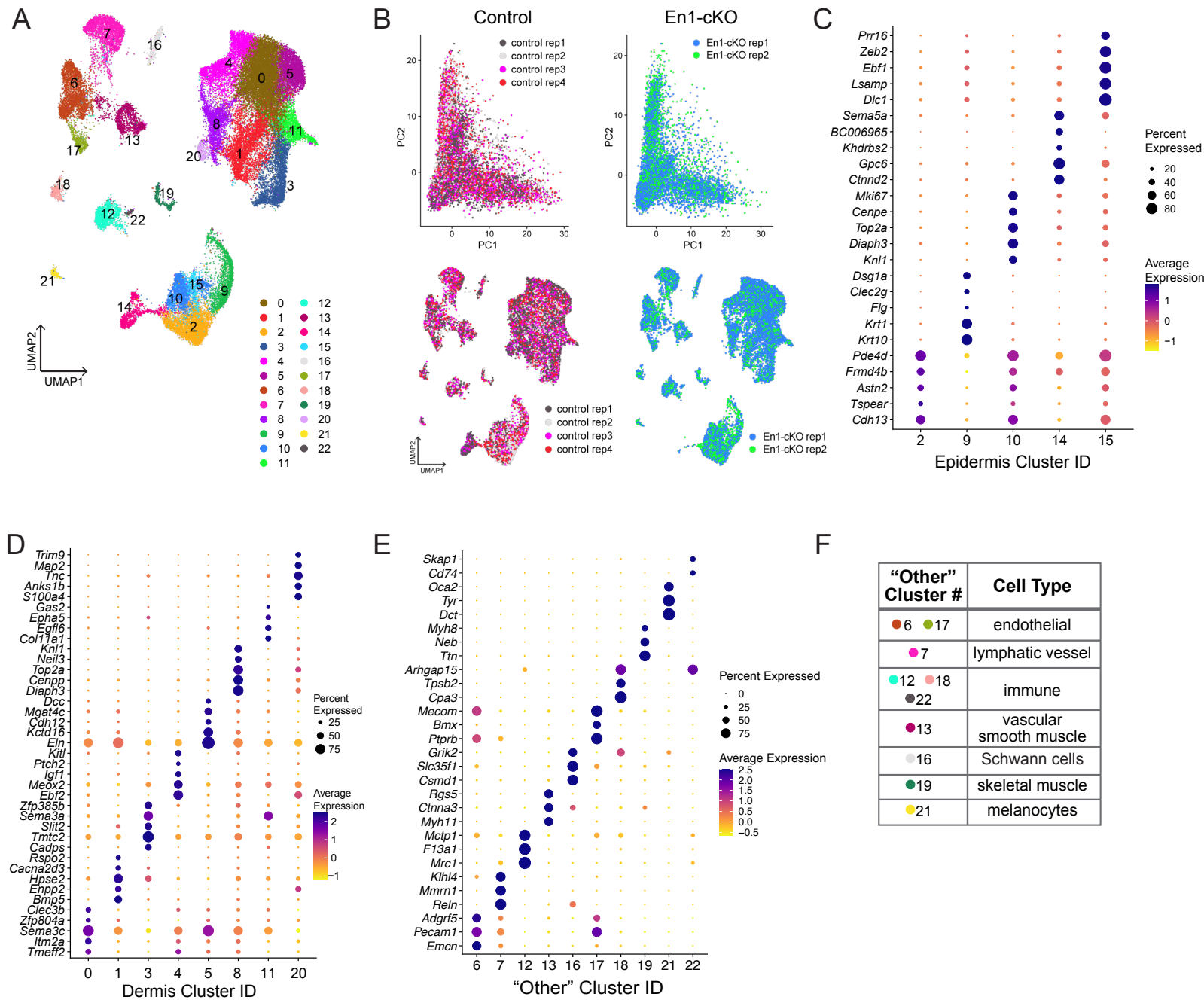


Figure S2. Clustering details for all nuclei from merged conditions, related to Figure 1, 3, and STAR Methods

A) UMAP projection from main Figure 1D colored by cluster identity. B) Plot of first two principal components (top) and UMAP (bottom) of Harmony integrated snRNA-seq data. Data from control (left) and En1-cKO (right) samples are shown separately. Points are colored according to replicate. C-E) Dot plots showing the expression of the top 5 marker genes identified for epidermal (C), dermal (D), and non-epidermal or dermal clusters (E; labeled “other” in main Figure 1). Dot color indicates average relative expression level for a cluster and dot size indicates the percent of nuclei in that cluster expressing the marker gene. F) Summary of cell type annotation for non-epidermal or dermal (“other”) clusters based on marker gene analysis.

FIGURE S3

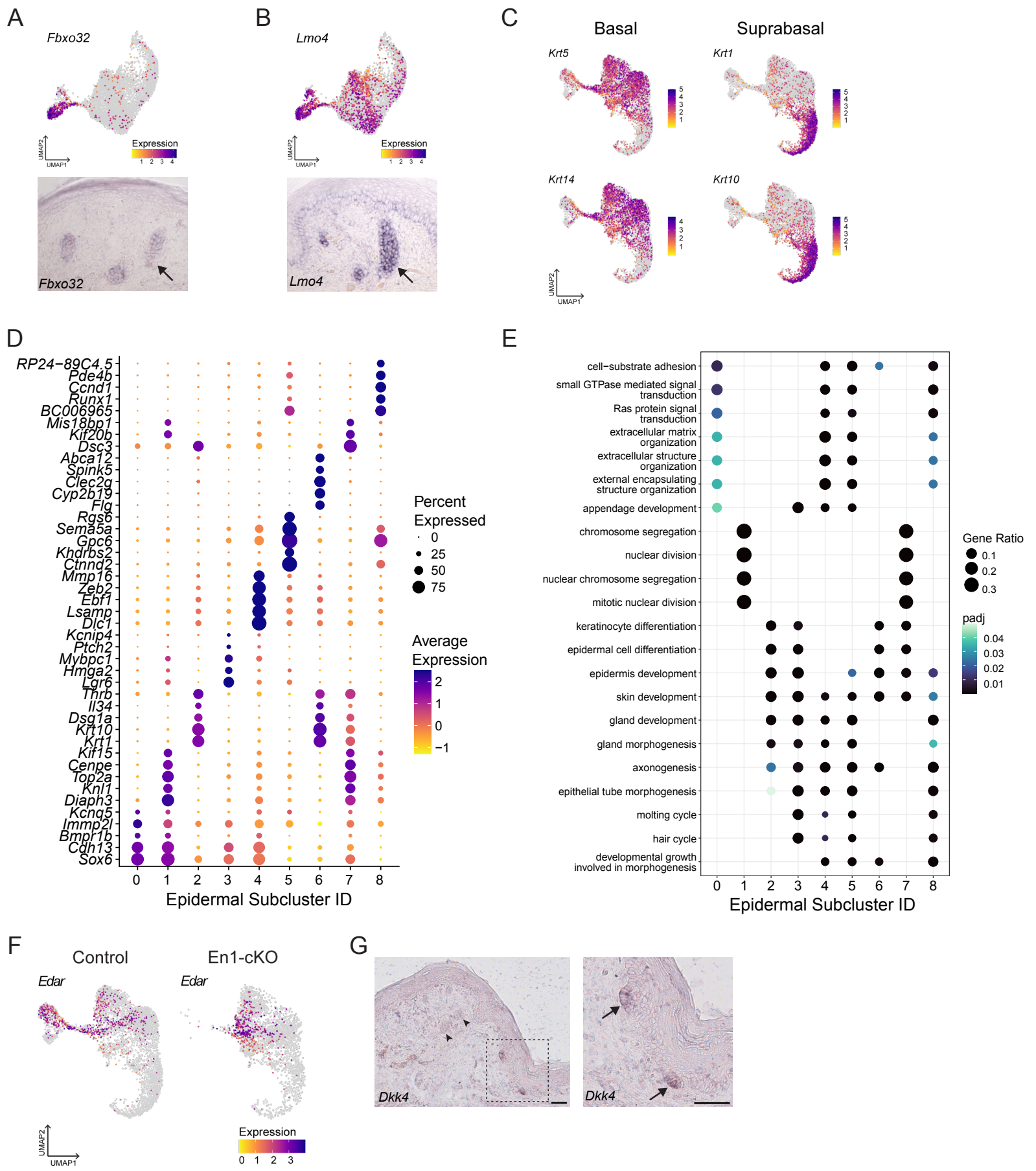


Figure S3. Epidermal subcluster marker genes and enrichment by gene ontology (GO), related to Figure 2

A) Feature plot (top) for *Fbxo32*, a specific marker gene of cluster 14. *In situ* hybridization for *Fbxo32* (bottom) in the volar limb in the footpad region at P2.5. B) Feature plot (top) for *Lmo4*, an additional marker of cluster 14. *In situ* hybridization for *Lmo4* (bottom) in the volar limb in the footpad region at P2.5. Arrows indicate nascent eccrine glands (A,B). C) Feature plots for known markers of basal keratinocytes (*Krt5*, *Krt14*) and suprabasal keratinocytes (*Krt1*, *Krt10*) showing normalized gene expression projected onto the UMAP embedding for subclustered epidermal nuclei. D) Dot plot depicting expression of the top 5 marker genes identified for each epidermal subcluster. Dot color indicates average relative expression level for the subcluster and dot size indicates the percent of nuclei in that subcluster expressing the marker gene. E) Dot plot depicting the top 4 results from a GO overrepresentation analysis conducted on significant differentially expressed marker genes for each epidermal subcluster (LFC ≥ 0.58 , p adj. < 0.001). Dot color represents adjusted p value for GO term enrichment and dot size represents the gene ratio. F) Feature plot showing normalized expression of *Edar*, a known marker of ectodermal appendages, for Control (left) and En1-cKO (right) subclustered epidermal nuclei. G) *In situ* hybridization for *Dkk4* in wildtype FVB/N volar hindpaw skin at P2.5. Lower magnification image (left) includes both nascent eccrine glands in the footpad (*Dkk4* negative; arrowheads) as well as eccrine placodes at the edge of the interfootpad space (*Dkk4* positive). Image to right shows eccrine placodes, indicated by arrows, at higher magnification. Scale bars represent 50 μ m.

FIGURE S4

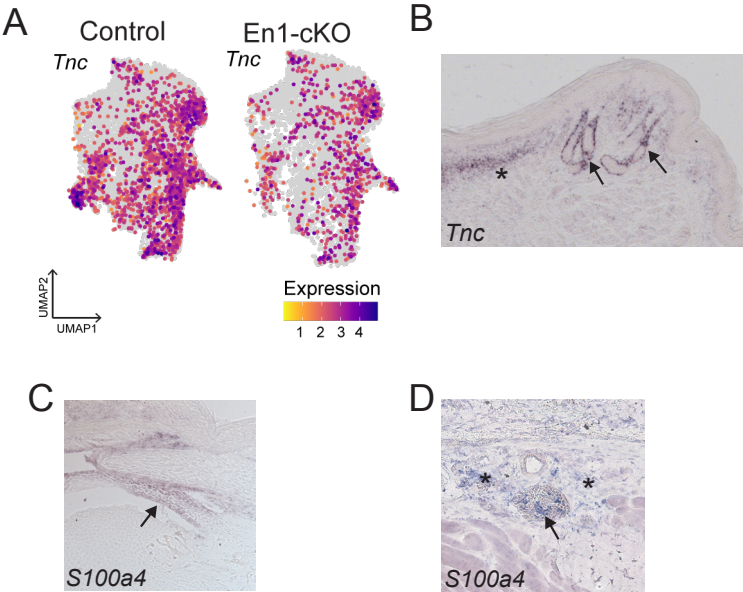


Figure S4. Additional validation of Cluster 20 markers, related to Figure 3 and Figure 4.

A) Feature plots show relative expression of *Tnc*, a marker gene of cluster 20, mapped onto UMAP of dermal nuclei from Control (left) and En1-cKO (right) samples. B) *In situ* hybridization for *Tnc* (purple) in the footpad region of the volar skin of a P2.5 FVB/N mouse. Nascent eccrine glands (arrows), as well as non-eccrine associated dermis (asterisk). C) *S100a4 in situ* hybridization signal (purple) in connective tissue of En1-cKO foot at P2.5 serves as internal positive control staining (same sample as shown in Figure 3E, right). D) *In situ* hybridization in the adult foot (P28) detects *S100a4* in a nerve (arrow) and connective tissue (asterisk) deep to the footpad. This serves as internal positive control for the mature eccrine gland *S100a4 in situ* shown in Figure 4A. Scale bars represent 50µm.

FIGURE S5

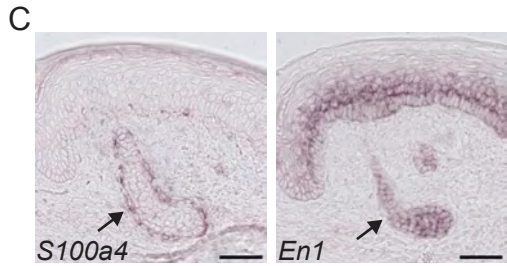
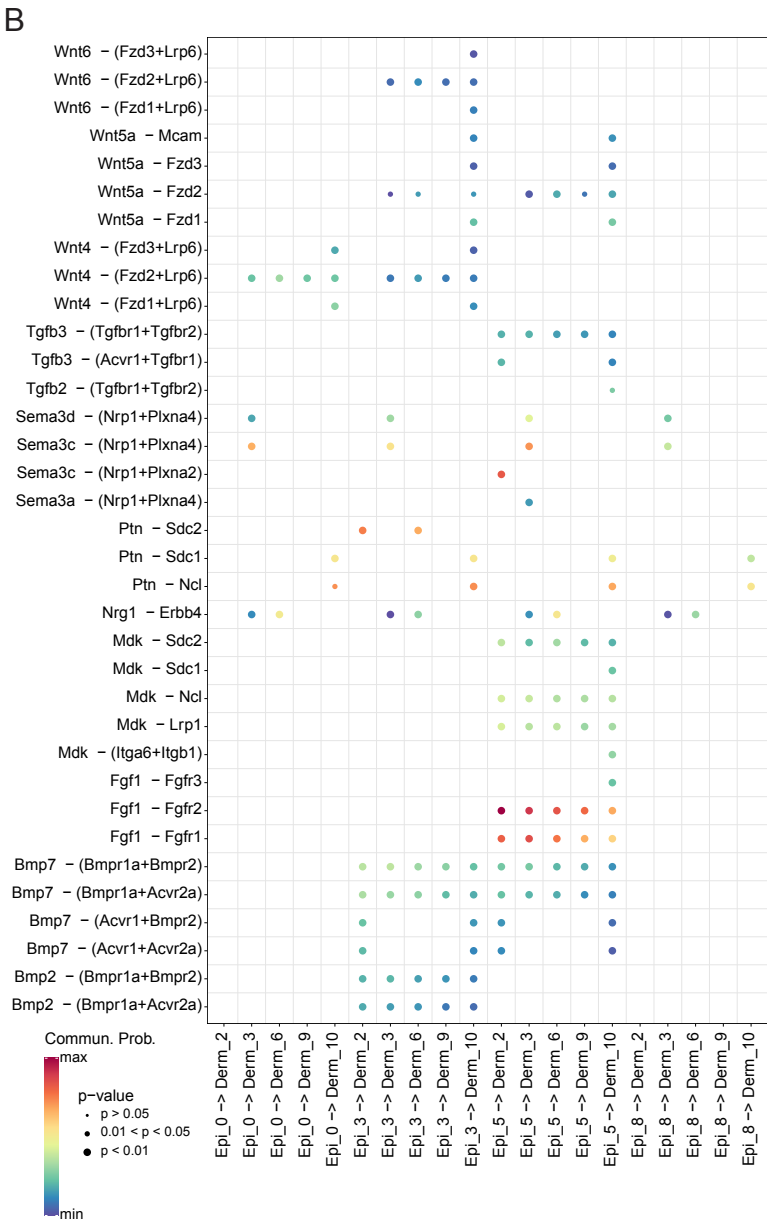
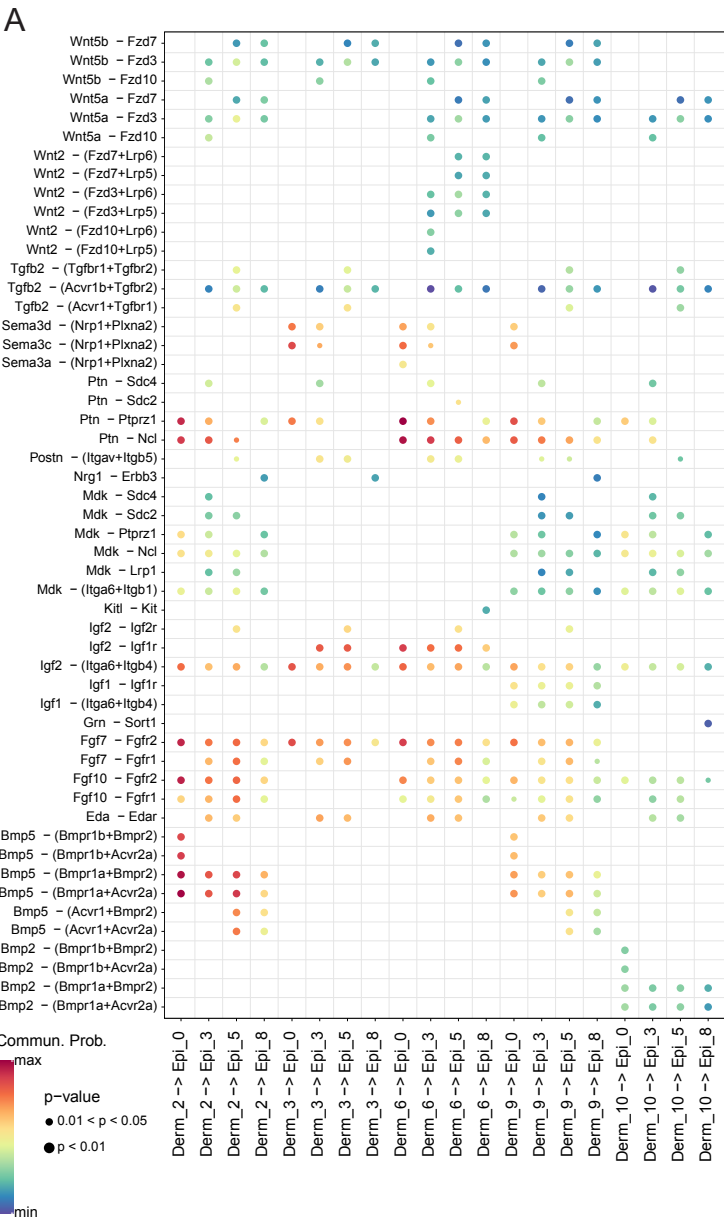


Figure S5. Inferred receptor-ligand interactions between dermal and epidermal subclusters and *S100a4* expression in volar skin, related to Figure 5

A,B) Dot plots representing the communication probability for all significant receptor-ligand pair interactions inferred by CellChat ($p < 0.05$) showing signaling from dermis to epidermis (A) and epidermis to dermis (B). Dot color reflects communication probability and dot size indicates p value.

C) *In situ* hybridization for *S100a4* (purple) and *En1* (purple) shows adjacent sections of Ablated volar skin with retained nascent glands. Sections shown are from the same Ablated sample as those shown in Figure 5C (bottom).

Table S1. Results of statistical analysis of the effect of genotype on footpad 4 (FP4) eccrine gland number in adult mice, related to Figure 1

Kruskal-Wallis FP4 ~ genotype			
chi-squared = 26.869, df = 3, p-value = 6.271e-06			
FP EG Dunn test results			
Comparison	Z	p.unadj	p.adj (BH)
creCtrl - floxCtrl	-1.38	0.17	0.25
creCtrl - ko	3.23	1.25E-03	2.49E-03
floxCtrl - ko	5.00	5.77E-07	3.46E-06
creCtrl - rttaCtrl	-0.53	0.60	0.60
floxCtrl - rttaCtrl	1.09	0.28	0.33
ko - rttaCtrl	-4.34	1.43E-05	4.30E-05

Results of Kruskal-Wallis test followed by *post-hoc* Dunn's test with Benjamini-Hochberg (BH) correction related to Figure S1A. Genotype codes: "floxCtrl" refers to En1^{flox/flox}; "creCtrl" refers to tetO-Cre/+; En1^{flox/flox}; "rttaCtrl" refers to Krt5-rtTA/+; En1^{flox/flox}; "ko" refers to tetO-Cre/+;Krt5-rtTA/+;En1^{flox/flox}. Alpha = 0.05.

Table S2. Results of statistical analysis of the effect of genotype on interfootpad (IFP) eccrine gland (EG) number in adult mice, related to Figure 1

Kruskal-Wallis IFP EG ~ genotype			
chi-squared = 38.202, df = 3, p-value = 2.561e-08			
IFP EG Dunn test results			
Comparison	Z	p.unadj	p.adj (BH)
creCtrl - floxCtrl	-0.41	0.68	0.68
creCtrl - ko	4.67	3.00E-06	9.00E-06
floxCtrl - ko	5.65	1.56E-08	9.36E-08
creCtrl - rttaCtrl	2.49	1.26E-02	1.51E-02
floxCtrl - rttaCtrl	3.46	5.48E-04	1.10E-03
ko - rttaCtrl	-2.98	2.86E-03	4.29E-03

Results of Kruskal-Wallis test followed by *post-hoc* Dunn's test with Benjamini-Hochberg (BH) correction related to Figure S1B. Genotype codes: "floxCtrl" refers to En1^{flox/flox}; "creCtrl" refers to tetO-Cre/+; En1^{flox/flox}; "rttaCtrl" refers to Krt5-rtTA/+; En1^{flox/flox}; "ko" refers to tetO-Cre/+;Krt5-rtTA/+; En1^{flox/flox}. Alpha = 0.05.

Table S3. Results of Kruskal-Wallis followed by *post-hoc* Dunn's test on the effect of genotype on interfootpad (IFP) hair follicle (HF) number in adult mice, related to Figure 1

Kruskal-Wallis IFP HF ~ genotype			
chi-squared = 25.635, df = 3, p-value = 1.137e-05			
IFP HF Dunn test results			
Comparison	Z	p.unadj	p.adj (BH)
creCtrl - floxCtrl	-0.52	0.60	0.60
creCtrl - ko	-4.46	8.15E-06	2.44E-05
floxCtrl - ko	-4.48	7.41E-06	4.45E-05
creCtrl - rttaCtrl	-1.77	0.08	0.12
floxCtrl - rttaCtrl	-1.44	0.15	0.18
ko - rttaCtrl	3.47	5.29E-04	1.06E-03

Results of Kruskal-Wallis test followed by *post-hoc* Dunn's test with Benjamini-Hochberg (BH) correction related to Figure S1C. Genotype codes: "floxCtrl" refers to $En1^{flox/flox}$; "creCtrl" refers to $tetO-Cre/+$; $En1^{flox/flox}$; "rttaCtrl" refers to $Krt5-rtTA/+$; $En1^{flox/flox}$; "ko" refers to $tetO-Cre/+; Krt5-rtTA/+; En1^{flox/flox}$. Alpha = 0.05.