KMT2D regulates p63 target enhancers to coordinate epithelial homeostasis

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Epithelial tissues rely on a highly coordinated balance between self-renewal, proliferation, and differentiation, disruption of which may drive carcinogenesis. The epigenetic regulator KMT2D (MLL4) is one of the most frequently mutated genes in all cancers, particularly epithelial cancers, yet its normal function in these tissues is unknown. Here, we identify a novel role for KMT2D in coordinating this fine balance, as depletion of KMT2D from undifferentiated epidermal keratinocytes results in reduced proliferation, premature spurious activation of terminal differentiation genes, and disorganized epidermal stratification. Genome-wide, KMT2D interacts with p63 and is enriched at its target enhancers. Depletion of KMT2D results in a broad loss of enhancer histone modifications H3 Lys 4 (H3K4) monomethylation (H3K4me1) and H3K27 acetylation (H3K27ac) as well as reduced expression of p63 target genes, including key genes involved in epithelial development and adhesion. Together, these results reveal a critical role for KMT2D in the control of epithelial enhancers and p63 target gene expression, including the requirement of KMT2D for the maintenance of epithelial progenitor gene expression and the coordination of proper terminal differentiation.

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Proper regulation of epigenetic information through the precise control of gene enhancer and transcriptional networks is essential for establishing and maintaining cell fate and identity [Sur and Taipale 2016]. Disruption of these pathways critical for differentiation can lead to a loss of proliferative control, ultimately driving carcinogenesis [Gonda and Ramsay 2015]. In line with this, the gene encoding the chromatin modifier and master enhancer regulator KMT2D (MLL4) is among the most frequently mutated genes in human cancer [Kandoth et al. 2013; Ford and Dingwall 2015; Rao and Dou 2015; Sze and Shilatifard 2016]. There is an exceptionally high prevalence of KMT2D mutations in self-renewing epithelial tissues ranging from the lung and esophagus to the skin and bladder, with the highest frequencies seen in keratinocyte cancers (cutaneous squamous cell carcinoma and basal cell carcinoma), the two most common forms of cancer worldwide [http://cancer.sanger.ac.uk/cosmic; Cerami et al. 2012; Gao et al. 2013].

Functionally, KMT2D and the related protein KMT2C (MLL3) are the major histone methyltransferases that deposit histone H3 Lys 4 (H3K4) monomethylation (H3K4me1), which poises enhancers for activation and transcription factor binding [Herz et al. 2012; Hu et al. 2013; Lee et al. 2013]. Furthermore, in addition to its catalytic role in H3K4me1 deposition, KMT2D is also required for the binding of the histone acetyltransferase complex CBP/p300, which acetylates H3K27 (H3K27ac) at enhancers, resulting in enhancer activation during cell identity gene induction and changes in cellular fate [Wang et al. 2016, 2017; Lai et al. 2017]. In mice, complete deletion of Kmt2d is embryonic-lethal [Lee et al. 2013]. Tissue-specific deletion studies have demonstrated that KMT2D is essential for the normal development and
function of enhancers during adipogenesis and myogenesis (Lee et al. 2013), cardiac development (Ang et al. 2016), and B-cell lymphocyte development (Ortega-Molina et al. 2015; Zhang et al. 2015). Despite the high incidence of KMT2D mutations in the epidermis and established roles in enhancer function and developmental programs in other tissues, little is understood about KMT2D function in normal epithelial homeostasis. Given the poor understanding yet potential importance of KMT2D in transcriptional enhancer function, we examined the transcriptional consequences of KMT2D loss in keratinocytes.

Results
To study KMT2D in the epithelium, we reduced KMT2D levels in primary neonatal human epidermal keratinocytes (NHEKs) and spontaneously immortalized human epidermal keratinocytes (HaCaTs) using shRNAs directed against KMT2D (shKMT2D) compared with scrambled control shRNA (shSC). RNA sequencing (RNA-seq) carried out in both the shRNA-treated NHEKs and HaCaTs demonstrated a significant reduction in KMT2D expression (Fig. 1A). This was confirmed by using three distinct shKMT2D hairpins that displayed a consistent twofold to threefold reduction in KMT2D expression using RT-qPCR, Western blotting, and immunofluorescence (IF) (Fig. 1B; Supplemental Fig. S1A,B). In addition, proliferative growth of shKMT2D keratinocytes was reduced by KMT2D loss compared with shSC keratinocytes (Fig. 1C), consistent with previous reports in other cell types (Issaeva et al. 2007; Guo et al. 2013). To confirm that the reduced proliferative capacity did not reflect a simple global stress response, we examined levels of the senescence-associated tumor suppressor p16/CDKN2A and the stress and senescence-associated transcription factor phospho-NF-κB p65 (Ser536) and did not observe any differences between either shSC or shKMT2D keratinocytes (Supplemental Fig. S1C).
of KMT2D depletion in keratinocytes. We performed three biological replicates of RNA-seq from NHEKs treated with shKMT2D. Strikingly, we found a broad reduction in the expression of genes critical for epithelial development and differentiation as well as adherence to the epidermal basement membrane (Fig. 1D). These encode proteins involved in epithelial development, migration, and differentiation [CYP27B1, GRHL2, PDFGB, FRKCH, RARG, Sox7, VDR, and WNT10A] (Rollman et al. 2003; Lee et al. 2009; Hara et al. 2011; Bikle 2012; Chen et al. 2012; Xu et al. 2017), the establishment and maintenance of cell polarity [DLG3, DLG5, and WNT7A] (Rodriguez-Boulan and Macara 2014); and cellular adhesion and the extracellular matrix organization [CLDN11, CNTN1, EMP2, IL4R, LOXL1, LOXL2, and SERPINE1] (Ferone et al. 2015; Petridou et al. 2017) and include disease-associated markers of interfollicular basal epidermal stem cells essential for the function of the epidermal basement membrane [COL4A2, ITGA3, ITGA5, ITGB2, ITGB4, LAMA5, and LAMB2] (representative genes are shown in Fig. 1D with corresponding adjusted \(P\)-values; a full list is in Supplemental Table S1A; Ko and Marinovich 2010, Simpson et al. 2011). Similarly, we also performed three RNA-seq replicates in the HaCaTs treated with shSC and shKMT2D. Overlap of all reduced genes (with \(P\)-value <0.1) from both sets of keratinocytes demonstrated a significant overlap in the transcriptional alterations observed with shKMT2D treatment (134 overlapping genes; \(P = 3.57 \times 10^{-76}\)) [Supplemental Fig. S1D]. Gene ontology (GO) analysis was carried out on the common 134 genes and showed enrichment of genes related to cellular growth, extracellular matrix organization, and adhesion, including key genes involved in epithelial development, differentiation, and morphogenesis. Notably, the vitamin D signaling pathway, a key cell signaling pathway involved in epithelial differentiation, was also among the most enriched pathways [Fig. 1E; Supplemental Table S2A; Bikle 2012]. Visualization of the keratinocytes by IF confirmed the loss of protein expression of representative genes such as RARG and VDR [Fig. 1F]. Collectively, these results show that KMT2D reduction drives a broad loss of key genes typically expressed by epidermal stem and progenitor cells of the basal layer as well as genes involved in proper epidermal differentiation, together suggesting a role for KMT2D in maintaining the hallmark gene expression program and proliferative capacity characteristic of epithelial progenitors.

To gain further insight into the function of KMT2D in the epidermis and the mechanism underlying keratinocyte gene expression, we mapped KMT2D binding genome-wide using chromatin immunoprecipitation (ChIP) combined with high-throughput sequencing (ChIP-seq) [full mapping information is in Supplemental Table S3A]. Consistent with its role in gene regulation at enhancers, a total of 1887 KMT2D ChIP-seq binding sites were analyzed by DiffBind [http://bioconductor.org/packages/release/bioc/vignettes/DiffBind/inst/doc/DiffBind.pdf] and the HOMER suite, among which half of the KMT2D peaks (52%) bound to intergenic regions, while introns (and likely intronic enhancers) made up the majority of the remaining binding sites (40%) [Fig. 2A]. Analysis of the gene nearest to each peak showed that 76% of peaks were associated with protein-coding genes, while the majority of the others occurred near noncoding RNAs (ncRNAs, 22%) [Fig. 2B].

We evaluated transcription factor-binding motifs that were enriched at KMT2D-binding regions. HOMER motif analysis of all KMT2D peaks identified p63, considered the master regulator of epithelial development and differentiation (Romano et al. 2012), as the most highly enriched transcription factor motif at KMT2D-binding sites [Fig. 2C]. To examine the overlap of KMT2D peaks with p63-binding sites, we intersected the gene nearest to our KMT2D peaks with 1751 previously published consensus p63-binding site-associated genes [McDade et al. 2012; Kouwenhoven et al. 2015]. We verified that KMT2D peaks indeed overlapped directly with p63-binding sites in shSC keratinocytes using ChIP-seq. Although our p63 antibody identified fewer total binding sites than observed in previous reports [McDade et al. 2012; Kouwenhoven et al. 2015], our data suggested that p63 and KMT2D bound to overlapping sites across the genome, as 49% of p63 peaks contained a directly overlapping KMT2D peak, while another 13% of p63 peaks were within 100 kb of a KMT2D peak. Overlapping peaks included known p63 target genes such as MINK1, SERINC2, and TP63 itself [Fig. 2E–G].

We next wanted to test for a potential interaction between KMT2D and p63. Importantly, chromatin co-immunoprecipitation (co-IP) experiments in NHEKs were consistent with the ChIP-seq data in showing that KMT2D interacts with p63 on chromatin [Supplemental Fig. S2A]. In line with data showing that the ΔNp63 isoform of p63 is the predominant form of p63 active in the epidermis [Truong et al. 2006], using a ΔNp63-specific antibody, we determined that KMT2D interacts with the α, β, and γ isoforms of ΔNp63, suggesting that this interaction is not mediated by the C-terminal sterile α motif (SAM) or the transactivation (TA) domains absent in the ΔN forms of p63 [Supplemental Fig. S2B]. Formaldehyde cross-linking followed by immunoprecipitation and mass spectrometry (IP-MS) further confirmed that p63 and KMT2D interact on chromatin [Fig. 2H,I]. Intriguingly, these data also suggest that this interaction may in fact be mediated by a shared interacting partner of the >200 that we identify here for the first time [Fig. 2H,J; Supplemental Fig. S3A–C; Supplemental Table S4].

To test the effects of KMT2D loss, we also performed KMT2D ChIP-seq on the shKMT2D keratinocytes. Hierarchical clustering showed that both shSC and shKMT2D samples clustered with their respective replicate data set [Fig. 3A]. According to DiffBind [http://bioconductor.org/packages/release/bioc/vignettes/DiffBind/inst/doc/DiffBind.pdf], which uses a stringent peak-based approach, of the 1887 total KMT2D peaks, 198 were identified as having at least a twofold loss of KMT2D peak size [Fig. 3B [heat map], C [black line in metaplot]]. In addition, in order to capture more of the KMT2D changes in enrichment that we observed in our primary data with shKMT2D
treatment, we also used a non-peak-based approach by comparing the tag density (AUC) alterations in ChIP-seq enrichment with shKMT2D treatment. This approach identified 2271 regions of at least a twofold decrease in KMT2D enrichment in shKMT2D keratinocytes in comparison with shSC keratinocytes (Supplemental Table S3A). In contrast, just 43 regions across the genome demonstrated a twofold or greater increase in KMT2D enrichment (Supplemental Table S3B). Consistent with KMT2D’s interaction with p63 and binding at the same sites on chromatin, motif analysis of KMT2D-reduced enrichment regions in shKMT2D keratinocytes once again showed that p63 was the most highly enriched transcription factor motif (Fig. 3D). In contrast, regions that gained KMT2D were enriched for transcription factors associated with epidermal differentiation, including CEBPB and AP1-associated factors BATF and Fra1 (Fig. 3E; Maytin and Habener 1998; Eckert et al. 2013; Watt 2016).

p63 has a critical role in regulating dynamic enhancers, including superenhancers critical for cell identity and cell fate specification, during epidermal development and differentiation (Kouwenhoven et al. 2015; Cavazza et al. 2016). Hence, we explored overlaps between established keratinocyte p63-binding sites and target genes (McDade et al. 2012; Kouwenhoven et al. 2015) and genes near KMT2D-bound regions, focusing on genes that lost >25% enrichment with shKMT2D across both replicates. Here, the intersection of p63-binding sites and KMT2D-depleted genes was highly significant \( P = 9.75 \times 10^{-127} \) (Fig. 3F; Supplemental Table S3C). GO analysis of these 362 “KMT2D-depleted/p63 target” genes included pathways similar to above—hemidesmosome assembly, cell and adherens junctions, and focal adhesion (Fig. 3G; Supplemental Table S2B)—again including genes involved in epidermal development, stemness, polarity, and adhesion (CNTN1, CTTNB1P1, DLG5, ITGA3, ITGB4, LAMA5, LAMC1, PRKCH, RARG, SERPINE1, SOX7, VDR, WNT7A, and WNT10A), many of which are known to be p63-regulated (Carroll et al. 2006; Kommagani et al. 2006; McDade et al. 2012; Ferone et al. 2015; Kouwenhoven et al. 2015). Together, these data demonstrate that in keratinocytes, KMT2D is enriched at p63 targets and maintains the expression of many key genes involved in epithelial homeostasis (including pathways involved in epidermal commitment and stratification such as the vitamin D and retinoid pathways) as well as numerous genes involved in regulating cell junctions and adhesion to the epidermal basement membrane.
KMT2D is a histone methyltransferase, thus, we assayed relevant histone modifications at key genes using ChIP-seq for key enhancer chromatin modifications, H3K4me1 (catalyzed by KMT2D), and H3K27ac (KMT2D recruits the acetyltransferase CBP/p300) [Wang et al. 2016, 2017; Lai et al. 2017]. We performed two replicates for each histone modification, and, as expected, H3K4me1 and H3K27ac clustered separately (Supplemental Fig. S4A). We identified the top 5% depleted regions (and the nearest associated gene) in each H3K27ac and H3K4me1 ChIP-seq replicate [Supplemental Table S5]. Consistent with both the RNA-seq and KMT2D-binding data, the top 5% regions of loss of both H3K4me1 and H3K27ac were enriched for GO categories involved in focal adhesion, adherens junctions, and cell migration and proliferation [Supplemental Table S5]. Overlap of the KMT2D-depleted/p63 target genes that lost gene expression by RNA-seq with the genes that lost enhancer histone modification enrichment identified 82 genes (Fig. 4A, Supplemental Table S6A). The most highly significant GO categories enriched within these 82 genes involved development and differentiation, consistent with both our current understanding of the role of p63 in stratified epithelium (and KMT2D in other tissues, such as skeletal muscle and fat) [Fig. 4B; Supplemental Table S2C] and our other results demonstrating the effects of KMT2D depletion on the expression of key genes involved in the epidermal basement membrane [such as COL4A2, ITGA3, ITGB4, and LAMA5] and epithelial development and differentiation [CTNNBIP1, Dlg5, Ehd1, Ehd2, EMP2, Prkch, RARG, Vdr, and Serpine1] (Fig. 4C, D; Supplemental Fig. S4B–F; Supplemental Table S6A). In addition, there were 20 KMT2D-depleted/p63 target genes significantly losing expression that did not lose enhancer histone modifications [at least within the top 5% loss for either H3K4me1 or H3K27ac] [Supplemental Table S6B]. These genes were also involved in development and adhesion and included ITGA3, Krt15, IL1B, and SPARC, and Tjap1 [Supplemental Table S6B]. These latter observations are consistent with recent evidence suggesting that KMT2D depletion can lead to extensive gene expression changes that are independent of its...
histone methyltransferase catalytic activity (Dorighi et al. 2017). Collectively, these results indicate that KMT2D can drive the expression of p63 target genes broadly involved in multiple aspects of epithelial development and differentiation through both catalytic histone methylation activity and other catalytically independent mechanisms at enhancers.

We then examined the genes that were up-regulated upon shKMT2D treatment of keratinocytes. GO analysis of the common genes that exhibited a significant increase ($P < 0.1$) in the shKMT2D keratinocytes identified a striking enrichment of genes involved in keratinocyte differentiation, with the GO category “cornification” displaying a $>40$-fold enrichment (Fig. 5A). These genes included markers of early epidermal stratification (such as $KRT10$) as well as canonical transcription factors involved in epidermal differentiation and stratification ($KL4$, $GRHL1$, $GRHL3$, and $ZNF750$) (Fig. 5B, Supplemental Table S1B). Other genes significantly increased in shKMT2D keratinocytes included genes such as $KRT16$ and $KRT17$, which are typically associated with hyperproliferative conditions such as squamous cell carcinoma and psoriasis (Fig. 5B, Supplemental Fig. S5A). The combination of data showing a loss of expression of genes involved in the basal epidermal stem cell layer and the progenitor state along with the spurious up-regulation of genes associated with keratinocyte differentiation suggests that in undifferentiated keratinocytes, KMT2D expression actively coordinates with p63 to maintain epidermal progenitor gene expression and thus prevent terminal differentiation.

To address how these chromatin and gene expression alterations may affect human skin, we used three-dimensional (3D) human organotypic skin cultures (Simpson et al. 2010). These cultures allow for the analysis of epidermal genes and proteins during differentiation in a more human and architecturally relevant context than standard submerged cultures (Simpson et al. 2010). While shSC-treated keratinocytes displayed a relatively normal-appearing stratified epidermis, shKMT2D-treated keratinocytes appeared highly disorganized, with cells showing a loss of polarity, irregular morphologies, and premature random cornification (Fig. 5C) consistent with many of the observed gene expression changes. IF of these sections confirmed decreased levels of KMT2D expression (Fig. 5D). Staining for other markers associated with

Figure 4. KMT2D depletion provokes a loss of enhancer histone modifications and p63 target genes. (A) The intersection of the KMT2D-depleted/p63 target genes that lose gene expression with those that lose enhancer histone modifications (H3K4me1 or H3K27ac) in shKMT2D keratinocytes identified 82 genes. (B) GO analysis of these 82 KMT2D-depleted/p63 target genes that lose enhancer histone modifications shows enrichment for genes broadly involved in development and differentiation. ChIP-seq tracks from representative genes that lose KMT2D enrichment at p63-bound regions as well as enhancer histone modifications. Examples include $VDR$ (C) and $LAMAS$ (D).
differentiation, such as filaggrin and loricrin, demonstrated disorganized stratification dynamics with premature and elevated expression of these proteins as well as a notable reduction in the expression of keratin 5, a marker of the epidermal progenitor cells of the basal epidermis (Fig. 5E,F). In order to determine whether a loss of p63 target gene expression was solely responsible for the premature activation of differentiation-associated genes, we overexpressed p63 in the shKMT2D keratinocytes. While significant overexpression of p63 was achieved, it was unable to reverse the gene expression changes observed in the KMT2D-depleted keratinocytes [Supplemental Fig. S5B]. These results were not entirely surprising given KMT2D’s critical role in enhancer, transcription, and transcription factor-binding dynamics. They potentially suggest that p63 may require sufficient levels of KMT2D in order to fully coordinate target gene expression and that the effects of KMT2D on epithelial homeostasis and stratification dynamics are more broad than simply acting solely through p63.

Discussion

An abundance of evidence has shown that p63 is critical for limb development and epidermal specification (Yang et al. 1998, 1999; Mills et al. 1999), and mutations in TP63 are associated with several developmental disorders that display epithelial abnormalities (Rinne et al. 2007). Over time, p63 has come to be known as the master regulatory transcription factor of epithelial tissues, playing key roles at multiple levels of development and differentiation (Truong et al. 2006; Soares and Zhou 2017). Excitingly, here we present evidence that KMT2D, a
major enhancer histone methylation enzyme and one of the most frequently mutated genes in epithelial cancers, is a critical regulator of p63-dependent gene expression programs and epithelial homeostasis. Our data indicate that KMT2D interacts with p63 on chromatin and at a broad array of enhancers involved in epithelial development, adhesion, and differentiation. Furthermore, we also identified numerous potential new interacting proteins that may play a role in mediating the KMT2D–p63 interaction. In turn, depletion of KMT2D in human keratinocytes leads to extensive alterations in normal epidermal transcriptional programs—characterized by a broad loss of expression of genes typically expressed in the self-renewing basal epidermal stem cell compartment, such as basal integrin and laminin genes—as well as key regulators of proper epidermal stratification, including the retinoid and vitamin D signaling pathways (Lee et al. 2009; Bikle 2012). In support of our findings, p63 has been shown to regulate many of the same genes that we observed here to be regulated by KMT2D, including those involved in adhesion and vitamin D signaling (Carroll et al. 2006; Kommagani et al. 2006, 2009). In response to the loss of the expression of these genes, genes involved in keratinocyte differentiation and proliferation (KRT1, KRT10, KRT6, KRT16, and KRT17) and transcription factors that promote terminal differentiation (KLF4, GRHL1, GRHL3, and ZNF750) are turned on prematurely.

As one of the largest organs in the body, the skin serves as a major barrier to the external environment. Similar to other epithelial surfaces, in order to maintain homeostasis, the epidermis undergoes regular highly coordinated self-renewal that involves dramatic alterations in gene transcription and cellular architecture (Avgustinova and Benitah 2016). This self-renewal ability is maintained in the interfollicular epidermis by stem cells in the basal layer that are typically slow cycling but with a high proliferative potential and self-renewing capacity, serving to ultimately supply the more differentiated suprabasal epidermis (Blanpain and Fuchs 2014; Watt 2016). Our data indicate that KMT2D plays a critical role in epithelial homeostasis by maintaining the adhesion and proliferative capacity of the epidermal basement membrane through its interaction with p63 to regulate critical enhancers for genes involved in both the maintenance of the basal epidermis and the proper coordination of epithelial stratification.

Similar to p63, while KMT2D may be critical for epithelial stem cell maintenance and gene expression, our data indicate that it may also play key roles in coordinating later steps in terminal differentiation, as does p63 (Truong et al. 2006). Indeed, given its effects on genes also involved in the promotion of epidermal differentiation, such as the vitamin D and retinoid signaling pathways, it is likely KMT2D participates in the coordination of all stages of epidermal commitment and differentiation. Consistent with this model, IF staining of normal skin sections for KMT2D [from both our laboratory and publicly available data sets] shows the presence of KMT2D in both basal epidermal stem cells and the nuclei of cells nearing terminal differentiation (Supplemental Fig. S6A,B). This observation may offer insight into why KMT2D has been shown to serve as both an oncogene and a tumor suppressor in cancer, depending on the cellular context (Ford and Dingwall 2015; Ortega-Molina et al. 2015; Zhang et al. 2015; Dawkins et al. 2016; Toska et al. 2017). Here we observed that KMT2D depletion reduces keratinocyte proliferation in undifferentiated keratinocytes, which is consistent with reports that KMT2D deficiency slows the proliferation of both breast and colon cancer cell lines (Mo et al. 2006; Guo et al. 2013; Kim et al. 2014) and correlates with improved outcomes in pancreatic cancer (Dawkins et al. 2016). In contrast, our data suggesting a role for KMT2D in regulating genes involved in vitamin D and retinoid signaling would suggest a potential tumor-suppressive role (Bikle 2014; Uray et al. 2016). Indeed, deciphering these potential oncogenic versus tumor-suppressive functions of KMT2D in epithelial cancers remains a critical question and an important direction for future studies.

Materials and methods

Keratinocyte cultures

NHEKs were isolated from deidentified discarded neonatal human foreskin obtained by Core B of the Penn Skin Biology and Diseases and Resource-based Center. Foreskin was incubated for 12 h at 4 °C in 2.4 U/mL dispase II. Sterile forceps were used to separate the underlying dermis. The epidermal sheet was transferred to a 60-mm tissue culture plate, incubated in 0.25% trypsin for 10 min at 37 °C, and then neutralized with 1 mL of fetal bovine serum (FBS). Sterile forceps were used to scrape the epidermal sheet against the dish to dissociate cells. The suspension was passed through a 40-µm strainer and then centrifuged at 200g for 5 min. The cell pellet was resuspended in 5 mL of medium (see below). HaCaTs were obtained from AddexBio. Both were cultured in a 50:50 mix of 1x keratinocyte–SFM supplemented with human recombinant epidermal growth factor 1–53 and bovine pituitary extract combined with medium 154 supplemented with human keratinocyte growth supplement and 1% 10,000 U/mL penicillin–streptomycin [all from Thermo Fisher Scientific]. For differentiation experiments, NHEKs were cultured in medium containing 1.22 mM calcium chloride for 48 h and then harvested.

Viral transduction

Lentiviral particles expressing scrambled control or KMT2D (MLL4)-specific shRNAs (D4, D5, and D9) were produced using the pLKO.1-Puro viral backbone plasmid generously provided by the Xianxin Hua laboratory at the University of Pennsylvania. Three KMT2D shRNAs were used: KMT2D shRNA D4 (CCGG CCCACCTGAATCATCACCTTTCTCGAGAAAGGTGATGACCCACCTGAATCATCACCTTTCTCGAGAAAGGTGATGATGATA TCCAGGTTG), KMT2D shRNA D5 (CCGGCCTGAATTGAAA CAACAGTCTTCTCGAGAAGACGTGTTGTTCAATTCA), and KMT2D shRNA D9 (CCGGCGAATGGGAAAGAACAAGCTGTTGTTCAATTCA). For the viral transductions, keratinocytes were infected with virus at 60% confluence on 10-cm² plates for 16 h in the presence of polybrene. Forty-eight hours following infections, cells underwent selection with puromycin to obtain completely puromycin-resistant cell populations. For the p63 overexpression studies, NHEKs at ~60% confluence were transfected with 10 µg of pcDNA3.1
ΔNp63α-Flag (Addgene, no. 26979) and 10 µL of Lipofectamine 2000 diluted in 1 mL of OptiMEM overnight (~16 h). Cells were placed in fresh medium, and RNA was harvested 48 h following the initial transfection.

Growth curve measurement

Two-hundred-thousand cells were seeded on a 950-mm² surface area (one well of a six-well plate) on day 0. Cell number was measured every 2 d with a Countess automated cell counter (Life Technologies) following standard procedures and default parameter settings, after which 200,000 cells were plated back for the next count.

RT-qPCR

RNA was extracted using RNeasy kit (Qiagen) following the manufacturer’s instructions. cDNA was obtained by High-Capacity RNA-to-DNA kit (Thermo Fisher) and then analyzed by standard qPCR methods on a 7900HT Fast real-time PCR (ABI). Primer sequences are available on request. RT-qPCR data analysis was performed by first obtaining the normalized QT values (normalized to 18S ribosomal RNA). The average and standard deviations of biological replicates were obtained and assessed for significance using an unpaired t-test (http://www.graphpad.com). For all experiments, three asterisks indicate \( P < 0.01 \), two asterisks indicate \( P < 0.05 \), one asterisk indicates \( P < 0.1 \), and “NS” indicates nonsignificance \( (P > 0.1) \).

RNA-seq

RNA was extracted using RNeasy kit (Qiagen, catalog no. 74014) following the manufacturer’s instructions. All RNA-seq libraries were prepared using the NEBNext polyA mRNA magnetic isolation module followed by NEBNext Ultra Directional RNA library preparation kit for Illumina (both from New England Biolabs). Library quality was checked by Agilent BioAnalyzer 2100 (Agilent), and libraries were quantified using KAPA library quantification kits (KAPA Biosystems). Libraries were then sequenced using a NextSeq500 platform (75-base-pair [bp] single-end reads) for Illumina. All RNA-seq was aligned using RNA STAR under default settings to Homo sapiens University of California at Santa Cruz (UCSC) hg19 (RefSeq and Gene core annotations). FPKM (fragments per kilobase per million mapped fragments) generation and differential expression analysis were performed using DESeq2. NHEK and HaCaT RNA-seq each consisted of three independent biological replicates.

IF

Cultured keratinocytes were fixed in 4% PFA in PBS for 30 min at room temperature. Cells were washed twice with PBS and permeabilized with 0.5% Triton X-100 in PBS for 10 min. After washing twice, cells were blocked in 10% BSA in PBS for 1 h at room temperature. Cells were incubated with primary antibodies in 5% BSA in PBS supplemented with 0.1% Tween 20 (PBST) overnight at 4°C. Antibodies are listed below. Next, cells were washed four times with PBST for 10 min each followed by incubation with fluorophore-conjugated secondary antibody in 5% BSA in PBST for 1 h at room temperature. Cells were then washed three times in PBST and once with PBS and incubated with 1 µg/mL DAPI for 5 min. The cells were then washed twice with PBS and mounted with ProLong Gold (Invitrogen). For the 3D organotypic skin cultures and normal human skin sections, following exposure to xylene and ethanol, sections were treated with target-unmasking fluid to deparaffinize the tissues. Sections were incubated in primary antibody overnight following blocking for 2 h in BlockAid (Thermo Fisher Scientific). Following secondary antibody incubation and washes, the sections were incubated with 1 µg/mL DAPI for 5 min, washed twice with PBS, and mounted with ProLong Gold with DAPI (Invitrogen). The slides were observed and imaged using a Nikon Eclipse microscope.

Antibodies

 Antibodies included anti-Histone H3 (Abcam, ab1791), anti-Histone H3 (monomethyl K4) (Abcam, ab8895), anti-Histone H3 (acetyl K27) (Abcam, ab4729), anti-ΔNp63 (HPA035977) (Sigma-Aldrich), anti-p63 α (D2K8X) (Cell Signaling, no. 13109), anti-p63 (4A4) (Santa Cruz Biotechnology, sc-8431), anti-ΔNp63 (Biolegend, no. 619001), anti-cytokeratin 16 antibody (EP1615Y) (Abcam, ab76416), anti-vitamin D receptor antibody (ChIP-grade) (Abcam, ab3508), and RARY antibody (G-1) (Santa Cruz Biotechnology, sc-7387).

MS analysis and protein identification

Flag-tagged KMT2D protein was transfected (Life Technologies, no. 11668019) and expressed in HEK293T cells, and Flag-tagged ΔNp63α was generated by in vitro translation (Thermo, no. 88881). Subsequently, both were separately subjected to immunoprecipitation with Flag antibody-conjugated protein G Dynabeads (Life Technologies, no. 10004D). Following stringent washes, equal amounts of lysates of undifferentiated or differentiated NHEKs were lysed for 1 h at 4°C in modified RIPA buffer containing 150 mM NaCl, 1% NP-40, 50 mM Tris-Cl (pH 8.0), and 1% SDS supplemented with protease inhibitors, or lysates of undifferentiated cross-linked NHEKs (cross-linked/lysed/sonicated as above) were added. After overnight co-IP, samples were washed thoroughly using a buffer containing 20 mM Tris (pH 8.0), 137 mM NaCl, 1 mM MgCl₂, 1 mM CaCl₂, 1% NP-40, and protease inhibitors. Beads were resuspended in 50 µL of 50 Mm ammonium bicarbonate (NH₄HCO₃), and samples were reduced with 0.5 µL of 1 M DTT for 30 min at 55°C and then cooled to room temperature. Samples were then alkylated using 2.5 µL of 500 mM iodoacetamide for 30 min at room temperature and protected from light. Following this, 1.5 µg of trypsin (Promega REVLV511C) was added to each sample and allowed to digest overnight. The next day, samples were subjected to 80°C for 10 min to stop trypsin digestion. Tubes were placed on a magnet, and peptides were transferred to a new tube. Beads were washed twice with 50% acetonitrile and transferred to the same tube. Finally, samples were dried on a SpeedVac and desalted by stage tip-ping. After in-bead digestion, samples were analyzed using nanoliquid chromatography coupled online to tandem MS [nLC-MS/MS]. Peptides were loaded onto a 75-µm ID × 17-cm in-house-packed 3-µm column (ReproSil-Pur C18-AQ) using an Easy-nLC-1000 (Thermo Scientific). Peptides were eluted over a 75-min gradient with 3%–35% solvent B [A: 0.1% formic acid, B: 80% acetonitrile, 0.1% formic acid] with a flow rate of 300 nL/min. The nLC was coupled to an Orbitrap Fusion (Thermo Scientific) operating in data-dependent mode (DDA). The spray voltage was set at 2.3 kV, and the capillary temperature was set at 275°C. Full MS scans were performed within the range of 450–1200 m/z at a resolution of 120,000 (at 200 m/z). Data-dependent MS/MS was performed in the ion trap using top-speed mode set to 3 sec; signals with charge state ≥2+ with an intensity >10,000 counts were automatically selected for MS/MS fragmentation performed using high-energy collision dissociation (HCD) with normalized collision energy of 27. MS raw files were analyzed using MaxQuant.
Illumina sequencing, all data were demultiplexed from the raw reads using NextSeq500 platform (75-bp, single-end reads) (Illumina). After (KAPA Biosystems). Libraries were then sequenced using a libraries were quantified using KAPA library quantification kits. Quality was checked by Agilent BioAnalyzer 2100 (Agilent), and libraries were grown using NEBNext Ultra DNA library making sequencing libraries using NEBNext Ultra DNA library preparation kit for Illumina (New England Biolabs). Library quality was checked by Agilent BioAnalyzer 2100 (Agilent), and libraries were quantified using KAPA library quantification kits (KAPA Biosystems). Libraries were then sequenced using a NextSeq500 platform (75-bp, single-end reads) (Illumina). After sequencing, all data were demultiplexed from the raw reads using Illumina’s BCL2FASTQ from BaseSpace. Fastq files were aligned to H. sapiens UCSC hg19. Tracks were visualized using the UCSC genome browser as described further below.

**ChIP-seq bioinformatics analyses**

Raw reads were mapped to the reference human genome assembly GRCCh37 (hg19) using Bowtie2 local alignment (Langmead and Salzberg 2012). The alignment result was further processed by SAMtools to remove low-quality alignments (a minimum mapping quality of 10), duplicate reads, reads mapped to the mitochondria, unplaced (chrUn) and random (random) contigs, and reads mapped to the ENCODE blacklisted genomic regions. MACS2 was used to identify ChIP-seq enriched regions (or peaks) (Zhang et al. 2008), requiring peak calling FDR <0.01. Distribution of resulting peaks in different genomic regions and motif analysis were conducted using the Homer suite (‘annotatedPeaks -size given’ and ‘findMotifsGenome’) (Heinz et al. 2010). The gene target for each peak was identified as the gene with the transcription start site (TSS) nearest to the peak. Genome-wide tag density was computed using BETools and UCSC command line utilities to generate tracks for visualization. Each individual track was normalized to 1 million reads. For each ChIP sample, the corresponding H3 (for H3K27ac and H3K4me1) or input control (for KMT2D and p300) was subtracted from the track using deepTools. Differential binding sites in shSC and shKMT2D conditions were identified by DiffBind Bioconductor package (default parameters). Heat maps were also generated by DiffBind. For KMT2D differential binding, we retrieved all sites with twofold differences for downstream analysis. Metaplots showing average ChIP signal near differential binding sites were generated using deepTools. Furthermore, to include more differential binding sites that were not identified due to the stringency of DiffBind, a statistical test-based differential analysis was performed. Tag density (AUC) was computed from each normalized input-subtracted KMT2D sample using bwtool. For each merged KMT2D peak region, we used Student’s t-test to examine whether its tag density was significantly different between shSC and scKMT2D samples. To identify the top 5% regions of histone modification loss, sequence tags for two replicates of H3K27ac and H3K4me1 ChIP-seq were aligned to the NCBI version 37 assembly of the human genome (hg19) using Bowtie version 1.1.1 (parameters -m 1 –best). Peaks were called for the two replicates using MACS2 callpeak, with matching whole-histone subunit H3 ChIP-seq as a sonication efficiency control [parameters -s 75 -q 0.01]. The peak sets for both histone post-translational modifications (PTMs) were further filtered to use a 0.1% FDR. Peaks were then assessed for losses in the shKMT2D keratinocytes, and those peaks with a quantitative loss in the 95th percentile were associated with their nearest gene by distance to the TSS.

**Statistics**

For Venn diagrams, hypergeometric testing was performed. For RT-qPCR and the growth curve measurements, the standard error of the mean of three replicates is presented. Additional statistical methods are listed in the “ChIP-seq Bioinformatics Analyses” section above.

**3D organotypic human skin cultures**

The 3D organotypic human skin cultures were performed as described previously (Simpson et al. 2010). Briefly, 12 T3 fibroblasts were grown in DMEM + 10% FBS. Cells were released from culture plates using 0.25% trypsin for 5 min at 37°C, resuspended in DMEM + 10% FBS, and counted using a hemacytometer to determine the volume needed to obtain 0.75 million to 1 million fibroblasts per organotypic culture. The required volume was centrifuged in a 50-mL sterile conical tube at 200g for 5 min, and the supernatant was removed. The fibroblast cell pellet was resuspended in 1/10 the final required volume (2 mL per culture) of 10× collagen resuspension buffer ([1.1 g of NaHCO3 plus 2.39 g of HEPES in 50 mL of 0.05 N NaOH] and held on ice. One-tenth the final volume of 10× DMEM (Sigma) was then added, and the cells were mixed by vigorous pipetting. Purified high-concentration rat tail collagen I (Corning) was added and diluted with sterile dH2O to a final concentration of 4 mg per milliliter of the final volume. NaOH (0.05 N) was added to a pH of ~7. The collagen–fibroblast slurry was mixed by inverting, and then 2 mL was pipetted into the upper chamber of a six-well transwell insert (Corning) placed within a deep-well six-well tissue culture plate (Corning). The fibroblast–collagen matrices were allowed to polymerize for 60 min at 37°C. Next, the matrices were submerged in DMEM + 10% FBS and placed overnight at 37°C. The next day, NHEKs were trypsinized, resuspended in DMEM + 10% FBS, counted to collect 1 million cells per culture, and centrifuged at 200g for 5 min, and the supernatant was discarded. The NHEK pellet was resuspended in E-medium supplemented with 5 ng/mL EGF (Sigma) to a volume of 2 mL per culture. The DMEM was removed from both the upper and lower chambers of the transwell plates containing the collagen–fibroblast matrices. Two milliliters of NHEKs (1 million cells) was seeded atop each matrix in the upper transwell chamber, and 14 mL of E-medium with 5 ng/mL EGF was added to the bottom chamber. The cultures were placed overnight at 37°C. The next day, the medium was aspirated from both the top and bottom chambers of the transwell. To place the NHEK monolayers at an air–liquid interface and induce stratification, 10 mL of E-medium (without EGF supplementation) was added only to the bottom chamber of the transwell, and the cultures were grown for up to 12 d at 37°C and fed 10 mL of E-medium every other day. To generate protein
lysates, the transwell apparatus was removed from the plate, and the organotypic culture was separated from the underlying matrix using sterile forceps. The culture was transferred into urea sample buffer (8 M urea, 1% SDS, 10% glycerol, 60 mM Tris, 5% β-mercaptoethanol at pH 6.8) and dissolved by vigorous pipetting using a 25-gauge needle and a 1-mL syringe. Organotypic cultures were prepared for routine histology by submerging the culture in 10% neutral-buffered formalin for 24–48 h.

Data availability
The relevant data sets generated during this study have been uploaded to NCBI Gene Expression Omnibus and are available under accession numbers GSE110475, GSE110453, GSE110467, GSM2991421–GSM2991485, and GSM2991609–GSM2991620.

GO analyses
All GO analyses were performed using PANTHER at http://geneontology.org/page/go-enrichment-analysis.

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