Jarid1b promotes epidermal differentiation by mediating the repression of Ship1 and activation of the AKT/Ovol1 pathway

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Abstract
Objectives: Terminally differentiated stratified squamous epithelial cells play an important role in barrier protection of the skin. The integrity of epidermal cells is maintained by tight regulation of proliferation and differentiation. The aim of this study was to investigate the role of epigenetic regulator H3K4me3 and its demethylase Jarid1b in the control of epithelial cell differentiation.

Materials and methods: RT-qPCR, Western blotting and IHC were used to detect mRNA and protein levels. We analysed cell proliferation by CCK8 assay and cell migration by wound healing assay. ChIP was used to measure H3K4me3 enrichment. A chamber graft model was established for epidermal development.

Results: Our studies showed that H3K4me3 was decreased during epidermal differentiation. The H3K4me3 demethylase Jarid1b positively controlled epidermal cell differentiation in vitro and in vivo. Mechanistically, we found that Jarid1b substantially increased the expression of mesenchymal-epithelial transition (MET)-related genes, among which Ovol1 positively regulated differentiation gene expression. In addition, Ovol1 expression was repressed by PI3K-AKT pathway inhibitors and overexpression (O/E) of the PI3K-AKT pathway suppressor Ship1. Knockdown (KD) of Ship1 activated downstream PI3K-AKT pathway and enhanced Ovol1 expression in HaCaT. Importantly, we found that Jarid1b negatively regulated Ship1 expression, but not that of Pten, by directly binding to its promoter to modulate H3K4me3 enrichment.

Conclusion: Our results identify an essential role of Jarid1b in the regulation of the Ship1/AKT/Ovol1 pathway to promote epithelial cell differentiation.
INTRODUCTION

The integrity of stratified epithelia is maintained by tight regulation of proliferation and differentiation. Basal cell self-renewal maintains the cell population and gives rise to differentiated cells, which maintains tissue homeostasis and the skin barrier function to defend against environmental stresses and protect the body from dehydration. Aberrant keratinocyte differentiation is involved in many skin diseases, such as psoriasis, atopic dermatitis and skin cancers. Increasing evidence has documented that the switch to differentiation gene expression is epigenetically controlled and that histone modifications are involved in the cell fate commitment. Repressive H3K27me3 is enriched in non-epidermal and differentiation genes to sustain the stem state. Knocking out key subunits of the PRC2 complex, Ezh1/2, thereby inducing the ablation of H3K27me3 marks, leads to premature differentiation of keratinocytes and Merkel cells. H3K4me3 is usually found around transcription start sites as an active gene transcription marker. The transient poised state of H3K4me3 and H3K27me3 is resolved upon differentiation, with only one of the marks remaining at the promoter to stably activate or silence gene expression, which has been proposed to occur in embryonic stem cells (ESCs), induced pluripotent stem cells, some non-stem cell lines and cancer cells. Intriguingly, global loss of H3K4me3 accompanies the differentiation of ESCs to progenitor cells.

Lysine demethylase Jarid1 family members, including Jarid1a, Jarid1b, Jarid1c and Jarid1d, have been found to demethylate H3K4me3. The role of Jarid1b in cell differentiation has recently received increased attention. Jarid1b contributes to the myogenic, osteogenic and macrophage lineage specification of mesenchymal cells. Silencing of Jarid1b in ESCs impairs neural lineage commitment. An RNAi screen identified Jarid1b as a major regulator of haematopoietic stem cell differentiation. Jarid1b contributes to trophoblast stem cell differentiation by downregulating self-renewal genes via removing H3K4 methylation. Jarid1b has been reported to be highly expressed in various types of tumours. In luminal breast cancer, Jarid1b is a luminal lineage-driving oncogene. Leukaemia stem cell (LSC) differentiation is associated with downregulation of H3K4me3 profiles, and overexpression (O/E) of Jarid1b leads to LSC differentiation, reducing leukaemia stem cell oncogenic potential. While these studies provide insight into the role of H3K4me3 in cell fate decisions and that of Jarid1b in regulating the differentiation of a variety of cell types, it is unclear how Jarid1b contributes to stratified epithelium differentiation.

Therefore, to investigate the role of H3K4me3 demethylases in regulating stratified epithelium differentiation, we evaluated the global expression levels of H3K4me3 and its demethylases in the Jarid1 family. Our findings demonstrated that at the global level, H3K4me3 expression decreased upon keratinocyte differentiation and that both methyltransferases and demethylases of H3K4me3 were increased, suggesting a crucial role of demethylases during differentiation. Furthermore, overexpression of Jarid1b promoted keratinocyte differentiation, whereas depletion of Jarid1b delayed differentiation in vitro and impaired differentiation layer formation in reconstituted epidermis in vivo. Mechanistically, we demonstrate that Jarid1b regulates the key differentiation transcription factor Ovol1 by directly binding to the PI3K-AKT pathway suppressor Ship1, but not to Pten. These findings suggest an epigenetic role of Jarid1b in regulating keratinocyte differentiation.

MATERIALS AND METHODS

Animals

Nude mice (male, 6 weeks old, 20-25 g) were purchased from the Beijing Vital River Laboratory, China, and housed in the SPF-grade centre of the animal facility in the affiliated hospital of Qingdao University. All experimental protocols were approved by the ethics committee on animal care of the Affiliated Hospital of Qingdao University. All animal studies were performed in strict accordance with the approved protocol.

Cell culture and treatment with inhibitors

HaCaT cells were cultured in low-calcium growth medium. Briefly, calcium-restricted media were obtained from Gibco, and calcium in serum was removed by incubation with Chelex 100 (Bio-Rad 142-2842). Low-calcium growth medium and high-calcium differentiation medium were produced by adding CaCl₂ at 0.03 mmol/L and 2.8 mmol/L, respectively. Cells were treated with inhibitors (Selleck) at the indicated concentration for 24 hours.

Plasmids and transfection

Jarid1b and Ovol1 were subcloned into the cumate-inducible Cumate-pLenti-GIII-CMV-IRE5-puro-SV40-GFP vector. In this system, the repressed CymR target gene is expressed when cumate is present. Ship1 was subcloned into the pCDNA3-3Flag vector. The target shRNA sequence was chemically synthesized in the pLKO.1 vector, which is included in a Tables S1-S3. Lentivirus vectors were transfected into HEK293FT cells with the helper plasmids delta8.9142-2842). Target gene expression was induced via cumate treatment at the indicated concentration when necessary. The Ubi-Pten-3FLAG and control plasmids were purchased from GeneChem Company, China. Transient transfection was performed with Lipofectamine 3000 following the instructions of the manufacturer.

Western blot analysis

Cells were lysed in 2% SDS lysis buffer and sonicated. A total of 15 μg of protein was loaded after quantification (Pierce 23225). Then, the proteins were transferred to a 0.45 μm PVDF membrane. After 1 hour of blocking with 5% BSA, the membrane was incubated with the
primary antibody overnight at 4°C and then the secondary antibody at room temperature for 1 hour on the next day. Antibody information can be found in Tables S1-S3. After washing, the blots were developed with the Super Signal Pico substrate (Pierce Biotechnology).

2.5 | Real-time reverse-transcription PCR

Total RNA was isolated by using RNAiso Plus (Takara D9108) and reverse-transcribed using All-In-One RT MasterMix with the AccuRT Genomic DNA Removal Kit (Abm G492). Quantitative PCR amplification (Abm MasterMix-S) using a Roche LightCycler 480 was performed via initial denaturation at 95°C for 5 minutes, followed by 40 cycles of 95°C for 10 s and 60°C for 15 s. The sequences of primers can be found in Tables S1-S3. Relative quantification was performed using the 2-ΔΔCt method normalized to GAPDH.

2.6 | Chromatin immunoprecipitation (ChIP)

The ChIP procedure has been described previously.27,18 In brief, formaldehyde cross-linked HaCaT cells were sonicated for 200 cycles (25 seconds ON and 15 seconds OFF, 40% amplitude) with the 2 mm probe of VCX750 sonicator. Dynal Protein G magnetic beads (Thermo 10003D) were incubated with an antibody overnight and then with the sonicated samples for at least 16 hours at 4°C. The conjugated beads were washed with low-salt, high-salt, LiCl and TE buffers in turn for 45 minutes each. DNA was isolated with phenol/ chloroform/isoamyl alcohol after reverse cross-linking and RNase A and proteinase K treatment.

2.7 | Immunohistochemistry and immunofluorescence

For immunohistochemistry, antigen retrieval was performed with pH 6.0 citrate buffer for paraffin-embedded tissue sections. After removing endogenous peroxidase with 0.3% hydrogen peroxide and non-specific protein blocking reagent incubation, primary antibody incubation was conducted overnight at 4°C. The mean intensity was quantified by using Image-Pro Plus from at least three raw, single-channel greyscale images that were obtained under the same conditions.

For immunofluorescence, 4% PFA-fixed cells were blocked with blocking buffer for 1 hour and incubated with the primary antibody overnight at 4°C. After washing three times, the cells were incubated with the secondary antibody for 1 hour at room temperature, followed by Hoechst counterstaining.

2.8 | Wound healing assay

A total of 600,000 cells were seeded in six-well plate wells. The next day, scratches were introduced using 200 μL pipet tips to create scratches with a similar diameter. The cells were washed once with PBS to remove floating cells. Then, 2 mL of fresh media supplemented with 1% FBS and the indicated concentration of cумate was added. Images of the scratches were recorded at the indicated time points at the same positions. The wound areas were quantified with ImageJ software.

2.9 | Cell viability

For the CCK8 assay, cells were seeded at density of 1 × 10^4 per well in a 96-well plate. Following the instructions of the Cell Counting Kit-8 (Dojindo CK04), the attached cells were incubated with 110 μL of medium containing 10 μL of CCK-8 solutions for 30 minutes to 2 hours. The absorbance was measured at 450 nm.

A live-cell imaging system was used to evaluate cell viability upon Jarid1b knockdown (KD). A total of 5000 cells per well were transfected with control or Jarid1b shRNA under low- or high-calcium culture conditions, and images were obtained every 6 hours. Cell confluence was calculated to reflect cell proliferation.

2.10 | Preparation of skin cell suspensions and chamber grafting

Engraftments were performed as described previously.19,20 Neonatal mice were sacrificed within 24 hours, and the back skin was dissected. The skin was floated on 1 U/mL dispase II overnight at 4°C. The next morning, the epidermis and dermis were separated with sharp forceps. The dermis was digested with 0.1% collagenase type 1 (LS004196) and 0.005% DNase I (LS002058) at 37°C for 1 hour with agitation, and this step was repeated twice. Collagenase activities were quenched by DMEM containing 10% foetal bovine serum. Single dermal cells were filtered through a 40 μm cell strainer. HaCaT cells were trypsinized to obtain a single-cell suspension. All cell lines were counted. Equal numbers of HaCaT and dermal cells were combined at 2 × 10^6 cells/μL in a 500 μL volume. The combined cells were injected into a silicon chamber implanted in the back muscle fascia of anaesthetized nude mice (Beijing Vital Research Laboratory, China). The chambers were removed after 1 week. The reconstituted skins were collected after 27 days of implantation.

2.11 | Statistical analysis

To determine the significance of data obtained from human samples or cell culture assays, comparisons were performed using descriptive and inferential statistics accompanied by graphs from the Prism software program (GraphPad). In all column bar graphs, the mean value ± 1 standard deviation is presented. For all the statistics, the 0.05 level of confidence was accepted for statistical significance.

3 | RESULTS

3.1 | H3K4me3 expression decreases and Jarid1b expression increases during epidermal differentiation

The epidermis offers an ideal in vivo model for investigating cell differentiation due to the physical separation of differentiated suprabasal layers from basal layer cells. Immunohistochemistry
assays demonstrated that H3K4me3 expression was decreased (Figure 1A,B) in suprabasal cells compared with basal cells, while Jarid1b expression was increased (Figure 1C,D). Calcium-induced differentiation has been widely accepted as an in vitro model for investigating epidermal differentiation. HaCaT cells were cultured in low-calcium conditions to maintain an undifferentiated state and underwent differentiation upon high-calcium stimulation.21 The results showed that the expression of K10 and involucrin (IVL) increased with time in high-calcium conditions, indicating that differentiation was successfully induced in our system. We found that H3K4me3 decreased upon differentiation (Figure 1E). Next, we examined the gene expression of methyltransferases and demethylases related to H3K4me3. We found that both methyltransferase and demethylase gene expression increased (Figure 1F and S1), indicating that the Jarid1 family is responsible for the global downregulation of H3K4me3 upon differentiation. Jarid1b displayed the greatest increase among all four members of the Jarid1 family (Jarid1d is not shown due to a high Ct value). Western blotting confirmed the increase in Jarid1b (Figure 1E). In addition, we compared Jarid1 gene expression levels in human primary keratinocytes based on the GEO database (GSE21413) and found that Jarid1b expression was comparatively higher than that of the other two genes (Tables S1-S3).

3.2 | Jarid1b deregulates HaCaT cell differentiation

Next, we silenced Jarid1b to explore its function in HaCaT cell differentiation. Jarid1b knockdown was confirmed at the mRNA and protein levels (Figure 2A). The mRNA and protein expression of K10 and IVL in Jarid1b knockdown cells was lower than that in the control cells at the indicated time points (Figure 2B,C), suggesting that Jarid1b silencing delayed HaCaT cell differentiation.

3.3 | Overexpression of Jarid1b enhances HaCaT cell differentiation and inhibits proliferation

A stable, inducible Jarid1b-overexpressing HaCaT cell line was produced by lentivirus infection followed by selection with 2 mg/mL puromycin to obtain a pure cell population. Jarid1b was overexpressed upon cumate treatment at the indicated concentration. K10 and IVL expression was upregulated with Jarid1b overexpression (Figure 2D and S2). In addition, the CCK8 assay showed that cell proliferation decreased upon cumate treatment compared with the control cells (Figure 2E), whereas knockdown of Jarid1b increased cell proliferation (Figure 2F), suggesting that the induction of Jarid1b may play a significant role in terminal differentiation.

3.4 | Jarid1b silencing impairs terminal differentiation in reconstituted skin

To further understand the role of Jarid1b in vivo, HaCaT cells and mouse dermal fibroblasts were grafted onto nude mice.19 After 27 days, the reconstituted skin was analysed by histological and immunohistochemical assays. First, we confirmed that in the reconstituted skin, Jarid1b expression was depleted compared with the control (Figure S3). H&E staining showed that the control cell differentiated into well-structured epithelia, whereas the Jarid1b-silenced epidermal
layers were thinner (Figure 3A,B). Moreover, staining for the differentiation marker K10 showed that the intensity was much lower and that the layer was thinner upon Jarid1b knockdown (Figure 3A,C,D), demonstrating substantial impairment of terminal differentiation. We also noted that all of the reconstituted epidermis consisted of one non-K10-stained basal cell layer (Figure 3A), though HaCaT cells retained K10 expression in the low-calcium culture in vitro (Figure 1E), which showed that the transplantation assay was a reliable and sensitive model and that essential regulation by stoma interactions is needed for epidermis formation in vivo. This native epidermis-resembling model strongly suggests a crucial role of Jarid1b in terminal differentiation.

### 3.5 Jarid1b controls MET gene expression

To determine whether Jarid1b inhibits FaDu cell migration, a scratch wound healing assay was performed. Initially, both groups exhibited almost the same wound size at 0 hour. However, in the Jarid1b-overexpressing group, the scratch wound area was still significantly larger than that in the control at 24 and 48 hours. Compared with the control, the cell migration rate was substantially decreased ($P < 0.05$) upon Jarid1b overexpression (Figure 4A). We speculated that Jarid1b could be involved in the epithelial-mesenchymal transition (EMT) process, as Jarid1b contributes to cell migration and terminal differentiation. We found that the expression of mesenchymal-epithelial transition (MET) genes, including Klf4, Foxa1, Ovol1 and Ovol2, was enhanced or suppressed upon Jarid1b overexpression or knockdown, respectively (Figure 4B,C). Western blot assays confirmed that Jarid1b overexpression resulted in elevated expression of Ovol1, whereas Jarid1b knockdown led to reduced Ovol1 expression (Figure 4D,E).

Recent studies have suggested that Ovol1 contributes to epidermal differentiation. Our results demonstrated that the expression of K10 and IVL was upregulated by Ovol1 overexpression and downregulated upon knockdown of Ovol1 (Figure 4F).

### 3.6 Ovol1 is controlled by the Jarid1b-Ship1-PI3K/AKT pathway

Our previous study demonstrated that the PI3K-AKT pathway promoted FaDu cell differentiation,$^{17}$ which raised the possibility that Ovol1 could be involved in this process. Here, our results showed that Ovol1 expression was inhibited upon treatment with the PI3K inhibitor LY294002 and the AKT inhibitor perifosine, suggesting that PI3K-AKT signalling plays an important role
**FIGURE 3** Depletion of Jarid1b impaired suprabasal layer formation. A, Representative images of H&E and K10 immunohistochemical staining in HaCaT reconstituted skin. B, Full-thickness analysis of H&E staining in the epidermis. C, K10 staining thickness and (D) intensity analysis (#P < 0.01 shRNA group vs control at each time point, Student’s t test)

**FIGURE 4** Jarid1b controlled MET gene expression. A, Representative images of wound healing scratches upon cumatate-induced Jarid1b overexpression at the indicated time points and quantification of the gap area (#P < 0.01 Jarid1b overexpression group vs control at each time point, Student’s t test). B, Jarid1b overexpression and (C) knockdown regulated the expression of the MET gene (Klf4, Foxa1, Ovol1 and Ovol2). D, Jarid1b overexpression or (E) knockdown enhanced or inhibited Ovol1 protein expression, respectively. Western blot analysis of the expression of keratinocyte differentiation markers (K10 and IVL) upon Ovol1 (F) overexpression and (G) knockdown.
in the control of Ovol1 expression (Figure 5A,B). We also found that inhibition of PI3K/AKT signalling repressed the expression of the differentiation genes K10 and IVL (Figure 5A,B and S4a), which was consistent with our previous study in cancer cells. Jarid1b is a demethylase that functions as an eraser of H3K4me3, leading to transcription repression. Thus, there is less of a possibility that Jarid1b directly regulates Ovol1 transcription. We hypothesized that Jarid1b activated PI3K/AKT/Ovol1 by controlling suppressors of this pathway and that Ship1 could be one such potential candidate. Our results showed that Ship1 expression was repressed by Jarid1b overexpression and enhanced by Jarid1b depletion (Figure 5C,D). Furthermore, Ovol1 expression was decreased upon Ship1 overexpression and increased upon Ship1 knockdown (Figure 5E,F). We also overexpressed another important suppressor, Pten. Ectopic expression of Pten in HaCaT cells increased K10 expression (Figures S4B,C), indicating the specificity of Jarid1b-Ship1 in the regulation of keratinocyte differentiation.

3.7 | Jarid1b directly binds to the SHIP1 gene promoter

Next, we examined whether another PI3K/AKT pathway suppressor, Pten, was regulated by Jarid1b. Unfortunately, there were no obvious changes in Pten upon Jarid1b overexpression (Figure 6A), suggesting that there is a specific mechanism by which Jarid1b regulates Ship1. We analysed the sequence of the SHIP1 gene and found the consensus binding sequence (GCACA/C) of Jarid1b near its promoter. We designed primers containing the binding motif around the TSS gene to perform ChIP assays. Our results demonstrated that Jarid1b and H3K4me3 were enriched at the TSS of the SHIP1 gene promoter (Figure 6B). Furthermore, overexpression of Jarid1b resulted in a reduction in H3K4me3 enrichment (Figure 6C). We next conducted a Ship1 rescue assay in Jarid1b-overexpressing cells. The results showed that Ship1 overexpression could enhance cell proliferation, which was inhibited by Jarid1b knockdown (Figure 6D). In addition, Ovol1 overexpression attenuated the stimulation of cell proliferation by Jarid1b knockdown (Figure 6E). Taken together, our results demonstrated that Jarid1b directly binds the SHIP1 gene by effectively activating the PI3K-AKT-Ovol1 pathway, resulting in keratinocyte differentiation.

4 | DISCUSSION

Genome-wide analyses have revealed that H3K4me3 is enriched at transcription start sites, which is regarded as an active histone
The global H3K4 methylation state is involved in the ESC fate decision during development. Global loss of H3K4me3 has been described during ESC differentiation. Our results showed that global H3K4me3 expression was decreased during differentiation in vitro and in vivo, which suggested that progenitor cell commitment is accompanied by H3K4me3 loss at the global level. Interestingly, LSC differentiation is also associated with stem maintenance gene repression via loss of H3K4me3, but not H3K79me2. Next, we explored the roles of H3K4-specific demethylases and methyltransferases in the global loss of H3K4me3. Our results showed that the levels of both types of enzymes increased during calcium-induced differentiation, indicating that only demethylases are responsible for the global loss of H3K4me3. The greatest increase was observed for Jarid1b among the four increased demethylases; Jarid1b has been reported to negatively regulate LSC stem potential and leukaemogenesis in murine and human MLL-rearranged AML cells. A global reduction in H3K4me3 levels may be a common feature in stem cell specification, where stem gene silencing results from the removal of H3K4me3 by demethylases.

Next, our study demonstrated that the histone demethylase Jarid1b is a positive regulator of epidermal differentiation. This finding is consistent with the essential role of Jarid1b in ESC differentiation. Jarid1b is indispensable in the neural commitment of ESCs, though a minor effect on self-renewal was observed. Knockdown of Jarid1b resulted in delayed ESC differentiation upon LIF depletion, which was also consistent with our results in vitro and in vivo. In addition, Jarid1b plays an important role in myogenic lineage specification from mesenchymal cells by regulating the enrichment of H3K4me3 at the Runx2 gene promoter. Here, we showed that Jarid1b controls Ship1 gene expression by directly binding to its promoter.

Our previous study demonstrated that Jarid1b overexpression is positively associated with hypopharyngeal squamous cell carcinoma (SCC) differentiation and that forced expression of Jarid1b leads to FaDu cell differentiation. Jarid1b has been reported to show elevated expression in various types of tumours, including breast, prostate and bladder tumours, and has been proposed as an oncogene. Controversially, Jarid1b has been reported as an epigenetic regulator with tumour-suppressive potential in melanoma. A slow-cycling melanoma cell subpopulation was shown to be characterized by high Jarid1b expression. In leukaemia, high expression of Jarid1b promotes cell differentiation, showing tumour-suppressive potential rather than acting as an oncogene. Collectively, the available results indicate that the contribution of Jarid1b to tumours may be context-dependent. Here, we demonstrated that Jarid1b contributed to epidermal cell differentiation. Considering these findings together with our previous results in FaDu cancer cells, we believe that Jarid1b may be a tumour suppressor rather than an oncogene, at least in squamous cell cancer carcinoma. For SCC differentiation, we usually have to use adjacent normal tissue as a control because the initial undifferentiated state has elapsed. Our study may provide a good example for characterizing gene function in SCC differentiation.
From our results, we assume that SCC and epithelial cells may evolutionarily share common mechanisms in promoting cell differentiation. Jarid1b represents a barrier to ESC EMT and the reprogramming of differentiated cells by regulating mesenchymal master regulators,32 whereas in human lung and colon cancer cells, overexpression of Jarid1b promotes the EMT process via upregulating the expression of Zeb1 and Zeb2, which are targeted by microRNA-200.33 Our results showed that Jarid1b overexpression impaired cell migration in a wound healing assay and promoted the expression of MET genes, among which Ovol1 has been proven to be a key molecule in epithelial differentiation. Ovol1 can repress Ovol234 and its own transcription.35 Ovol1 knockout mice exhibit skin barrier impairment,34,36 suggesting the important role of Ovol1 in epidermal differentiation. Filaggrin and involucrin levels are reduced upon Ovol1 knockout in vivo,36 which is consistent with our results regarding the regulation of Inv and K10. In skin melanoma, high expression of Ovol1 prevents tumour invasion and predicts a better prognosis than that associated with low expression.37 In cutaneous and oral SCC, the expression of Ovol1 markedly decreases in the invasive portion compared with the in situ state, implying a role of Ovol1 in potentiating differentiation.38,39 Despite the importance in cell differentiation, the regulation of Ovol1 expression is largely unknown. NGN3 negatively regulates the transcription of Ovol1 in an E-box-dependent fashion in the rodent pancreas.40 In SCC cell lines, IRF6 binds to the Ovol1 gene promoter and positively regulates Ovol1 expression.41 Li et al.42 showed that the mOvol1 promoter is activated by the LEF1/β-catenin complex. Our previous research has shown that β-catenin is regulated by the Jarid1b-β‐catenin pathway. Thus, we speculated that this pathway may also be involved in the regulation of Ovol1 expression. As expected, we found that Ovol1 expression was positively controlled by the Jarid1b-P3K-AKT pathway. Interestingly, Jarid1b only controls the AKT pathway suppressor Ship1, and not Pten, by directly binding to its promoter, which suggests that there may be specific mechanisms to trigger distinct downstream biological effects.

Together, our studies clearly show the crucial role of Jarid1b in promoting epidermal cell differentiation (Figure 6F). As the observation of roles of Jarid1b in controlling cell differentiation and tumorigenesis remains controversial, a clear understanding of its roles in the regulation of differentiation in a tissue-dependent system will be essential and beneficial for determining their functional involvement in SCC and, further, for developing proper therapeutic strategies in the future. In summary, our studies addressed a representative epithelial tissue, the skin, and demonstrated a clear role of Jarid1b-mediated repression of Ship1 and activation of the AKT/Ovol1 pathway in enhancing the epithelial cell differentiation state.

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CONFLICT OF INTEREST

The authors declare no conflict of interest.

AUTHOR CONTRIBUTIONS

ZJ, JY, YT, FT and AX designed the experiments and analysed the data; SX, NY, LZ, ZL, HY, LQ, ZS, CT and ZJ performed the research; and ZJ wrote the paper. All authors read and approved the final manuscript.

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SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section at the end of the article.

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