IL-13 regulates human nasal epithelial cell differentiation via H3K4me3 modification

Introduction: Epigenetic regulation has been shown to play an important role in the development of inflammatory diseases, including chronic rhinosinusitis and nasal polyps. The latter are characterized by epithelial mis-differentiation and infiltration of inflammatory cytokines. H3K4me3 has been shown to be involved in regulating lineage commitment. However, the underlying mechanisms, especially in human nasal epithelial cells (HNEpC), remain underexplored. The objective of this study was to investigate the role of H3K4me3 in HNEpC differentiation treated with the Th2 cytokine IL-13.

Patients and methods: The expression levels of mRNA and proteins were investigated using reverse transcription-polymerase chain reaction (RT-PCR) assays and Western blot in nasal polyp tissues and human nasal epithelial cells respectively. We measured these levels of H3K4me3, MLL1 and targeted genes compared with control subjects.

Results: We demonstrate that expression of H3K4me3 and its methyltransferase MLL1 was significantly upregulated in IL-13-treated HNEpC. This elevation was also observed in nasal polyps. Expression of cilia-related transcription factors FOXJ1 and DNAI2 decreased, while goblet cell-derived genes CLCA1 and MUC5a increased upon IL-13 treatment. Mechanistically, knockdown of MLL1 restored expression of these four genes induced by IL-13.

Conclusion: These findings suggest that H3K4me3 is a critical regulator in control of nasal epithelial cell differentiation. MLL1 may be a potential therapeutic target for nasal inflammatory diseases.

Keywords: IL-13, H3K4me3 modification, nasal epithelial cell, differentiation

Introduction

The nasal epithelium possesses a pseudo-stratified structure, with specialized cell types including goblet cells, ciliated or non-ciliated columnar cells, and basal cells. This epithelium plays an important role in protecting the airway from infection, inflammation, and physical injury.1 Epithelial remodeling is induced upon damage, which is characterized by accumulation of pseudocyst formations, lack of collagen, and excessive inflammatory infiltrations, resulting in potentially irreversible structural changes.2 Epithelial cells undergo migration, proliferation, and differentiation in response to environment stimuli. The process is highly organized and is regulated by diverse growth factors and cytokines. Cytokines play essential roles in mediating allergic inflammation. Chronic rhinosinusitis (CRS) is a prevalent condition causing poor quality of life. CRS is divided into two subtypes: CRS with or without nasal polyps (CRSwNP or CRSsNP). CRSwNP displays epithelial barrier dysfunction with ciliary impairment and excessive mucus secretion.3 CRSwNP is characterized by a Th2 inflammatory pattern with high expression...
of type 2 cytokines IL-4, IL-5, and IL-13. CRSsNP merely expresses these biomarkers, but with high levels of INF-γ, characterized by predominance of Th1 cell.4−6 In addition, some CRS patients express a neutrophilic type of inflammation via Th1 and/or Th17 cells.7 Type 2 cytokines are induced by epithelium cells, which can activate type 2 innate lymphoid cells such as mast cells. These effector cells also produce type 2 cytokines, contributing to tissue remodeling. Nasal polyps display epithelial mucociliary dysfunction with excessive mucus secretion and cilia disappearance.8 Although the cause of nasal polyps is not clear, accumulated evidence suggests that cytokines play crucial roles. Previous studies suggest that IL-13 leads to stasis of sinonasal mucus production and cilia dysfunction,9 resulting in persistent inflammation. In human bronchial epithelial cells, IL-13 induced goblet cell hyperplasia and ciliated cell loss.10,11 The underlying mechanisms have yet to be fully explored.

Epigenetic regulation has been found to be involved in a number of inflammatory disorders.12−15 Previous studies16 have shown that, in nasal polyps, transcription and protein expression levels of HDCA2 are increased. Treatment with histone deacetylase inhibitor and histone modifications suppressed myofibroblast differentiation and altered extracellular matrix production.16,17 HDAC inhibitors suppressed inflammation via induction of FoxP3+ regulatory T-cells that also have relevance to asthma.18,19 In addition, in nasal polyp fibroblasts, H3 lysine27 acetylation (H3K27Ac) was highly expressed, suggesting that histone modifications regulated development of nasal polyps.20 It is noteworthy that increased active histone markers, including H3-K9 acetylation and H3-K4 trimethylation across the IL-4 and IFN-γ loci, were observed with Th1 or Th2 cell lineage commitment.21−23 It has been found that MLL1 can regulate the development of Th2 reactions by H3K4me3 modification through stabilizing expression of GATA3.24 MLL1 can also influence Th1 cell proliferation via regulating IL-12 responsiveness,25 suggesting that MLL1 has played a key role of regulating cellular inflammatory processes.

In the present study, we observed elevated H3K4me3 expression in the nasal polyps. We investigated the function of the histone methyltransferase MLL1 in human nasal epithelial cells (HNEpC) upon IL-13 treatment. We found that H3K4me3 may play an important role in the mis-differentiation of nasal epithelium in inflammatory disorders.

Patients and methods

Subject collection

Nasal polyp tissues and normal inferior turbinate tissues were collected from 16 patients with CRSwNP undergoing functional endoscopic sinus surgery from the Department of Otorhinolaryngology, Affiliated Hospital of Qingdao University, China. All tissues were used immediately and/or snap-frozen at −80°C. The study was approved by the local ethics committee and the regulatory authorities of China. Written informed consent was obtained from all subjects before sample collection. Patients with an established immunodeficiency and pregnancy were excluded from the study. None of the patients had allergy, asthma, or aspirin sensitivity and treated with corticosteroids for at least 1 month before surgery.

Cell culture and treatment with IL-13

HNEpCs were purchased from PromoCell (Atlanta, GA, USA) and cultured in 1640 medium (Thermo Fisher Scientific, Waltham, MA, USA), supplemented with 10% fetal bovine serum and 1% penicillin–streptomycin (Thermo Fisher Scientific) in 5% CO2 atmosphere at 37°C. The cells were stimulated with Recombinant Human IL-13 (2#00-13, Escherichia coli; Peprotech, Rocky Hill, NJ, USA) at a final concentration of 5, 10, 50, 100, and 200 ng/mL for indicated time.

Plasmids and transfection

MLL1 shRNA plasmids were obtained from GeneChem Company (Montreal, QC, Canada). The targeted sequences were 5′-GATTATGACCTCCAATTTAA-3′ and 5′-GCACTGTTAACCATTCCACTT-3′. The plasmids were transfected with helper vectors, pDelta8.9 and pVSV-G, into HEK293FT cells. After 48 h of transfection, the medium was collected and centrifuged at 50,000 × g for 3 h. The pellets were resuspended in PBS. Lentivirus was transduced into the HNEpC. To obtain a stable and pure MLL1-knockdown cell population, we performed selection with 2 µg/mL of puromycin after 48 h of transfection. It usually takes 2 days for all the control cells to die. After selection, we collected cells and examined the efficiency of transfection through real-time polymerase chain reaction (PCR) and Western blot.

Western blot analysis

To obtain cell and tissue proteins, samples were processed with 2% sodium dodecyl sulfate (SDS) lysis buffer and sonicated to break up DNA. Lysates were boiled for 10 min at 98°C. Then, the samples were measured by BCA Protein Assay Kit (Beyotime Institute of Biotechnology, Shanghai, China), and 20 µg of total protein was loaded. Transferred polyvinylidene fluoride membranes were incubated with primary antibodies against H3K4me3 (1:1000; #GC-263, PTMbiolabs, Chicago, IL, USA), MLL1 (1:1000; #14197, Cell Signaling Technology, Danvers, MA, USA), and β-tubulin (1:3000; Beyotime Institute of Biotechnology) overnight at 4°C, followed by incubation.
with secondary antibodies of anti-mouse IgG and anti-rabbit IgG, respectively (1:2000; CWbiotech, Beijing, China) for 1 h at room temperature. Western blot analyses were normalized to β-tubulin. The blots were developed with Super Signal Pico substrate (Pierce Biotechnology, Shanghai, China). Each immunoblot was repeated three times, with samples obtained from different experiments. The relative intensity of protein bands was measured with NIH image J software.

**RNA preparation and quantitative real-time qPCR**

Samples were stored at −80°C until homogenization and no more than 25 mg tissues were homogenized in Trizol. For quantitative real-time PCR, total RNA was extracted from HNEpC and tissues using RNAiso Plus (D9108; Takara Bio, Tokyo, Japan) following the instructions from the manufacturer. RNA quantity and purity were determined by Nanodrop spectrophotometer. GAPDH was used as an internal control. Reverse cloning of cDNA by 500 ng RNA was performed according to the manufacturer’s instructions. Real-time PCR was performed to determine the mRNA expression. In brief, real-time PCR was conducted using the Roche Lightcycler480 Real-time PCR System with SYBR green reagents. Quantifications were normalized to cycler480 Real-time PCR System with SYBR green reagents according to the manufacturer’s instructions. Real-time PCR was performed to determine the mRNA expression. In brief, real-time PCR was conducted using the Roche Lightcycler480 Real-time PCR System with SYBR green reagents. Quantifications were normalized to GAPDH. Relative gene expression was calculated using the 2^ΔΔCT method.

The primer sequences used for application were as follows:

- FOXJ1 forward: 5′-GTTCCTCCCGAGGCACCTTTGGA-3′ and reverse: 5′-CACCAGATCCACCTGTCCGG-3′;
- DNAI2: forward: 5′-GTTGAGGTCAGAGGTTGCG-3′ and reverse: 5′-GGATGAGGAGCACCAGTGG-3′; MLL5: forward: 5′-ACCGATGAAATTGCCCGGACCACGGG-3′ and reverse: 5′-TTGTACACCACCGTGCTG-3′;
- CLCA1: forward: 5′-TTGTTAAGGCACCGCTCTGTG-3′; MLL1: forward: 5′-TTTTAGAGGAGAAGCGACCGCC-3′ and reverse: 5′-AGGGTGATAGCTGTTTCGGC-3′;
- MUC5a: forward: 5′-ACCATCGGAATTCGGCAAC-3′ and reverse: 5′-TGTCTATGCCACTCTGTGGC-3′; SETD1A: forward: 5′-GGTGCCTACAGGTCCCGGC-3′ and reverse: 5′-GACGTGTTCATCAGCCTTCT-3′; and JARID1B: forward: 5′-CTATCTGCCAAATGGTGCG-3′ and reverse: 5′-TCTAACACTGCGACACGTCC-3′.

### Statistical analyses

Statistical analyses were done using descriptive and inferential statistics by GraphPad Prism software version 6.0 (GraphPad Software, Inc., La Jolla, CA, USA). Student's t-test and ANOVA test were performed to determine the statistical significance between two groups and among more than two groups, respectively. All in vitro experiments were done and repeated at least three times. In Figures 1–3, mean value ± 1 SD are presented. For all statistics, P=0.05 was considered to be statistically significant.

### Results

#### Increased expression of H3K4me3 and relative epithelial gene mRNA expression in nasal polyps

Pathological remodeling of nasal polyps is characterized by epithelial dysfunction. First, we collected nasal polyp tissues and inferior turbinate samples from the same side of nasal polyps patients undergoing polypectomy for the treatment of nasal obstruction. mRNA expression of FOXJ1, DNAI2, CLCA1, and MUC5a was examined. Expression of FOXJ1 and DNAI2, major cilia-related transcription factors, was decreased in nasal polyps compared to control, whereas that of CLCA1 and MUC5a, goblet cell-derived genes, was elevated (Figure 1A), suggesting mis-differentiation of epithelium. H3K4me3 expression was increased in nasal polyps samples compared with control (Figure 1B and 1C), suggesting that histone methylation may play an important role in metaplasia of nasal epithelia.

#### Elevation of H3K4me3 and MLL1 expression upon IL-13 treatment in HNEpC

To further understand the role of H3K4me3 in progression of nasal Th2 inflammatory diseases, we treated HNEpC cells with IL-13 at varying concentrations. H3K4me3 expression was elevated by IL-13 treatment (Figure 2A). Peak expression of H3K4me3 occurred at 10 ng/ mL concentration of IL-13 (Figure 2B). Thus, we chose 10 ng/mL concentration of IL-13 for further experiments. Next, we analyzed mRNA expression of FOXJ1, DNAI2, CLCA1, and MUC5a. IL-13 induced mRNA expression of CLCA1 and MUC5a, but suppressed FOXJ1 and DNAI2.
Increased expression of H3K4me3 and relevant epithelial gene mRNA expression in nasal polyps.

Notes: (A) mRNA expression of FOXJ1, DNAI2, MUC5a, and CLCA1 compared with the control group. (B) H3K4me3 expression was measured in nasal polyps by immunoblotting and compared with the control. (C) Relative quantification of H3K4me3 intensity normalized by β-tubulin in tissues. Data are expressed as mean ± SD. *P < 0.05, Student’s t-test.

H3K4me3 is a dynamic and reversible process that is governed by histone methyltransferases and demethylases. We examined a variety of important genes regulating the methyl group of H3K4me3 by quantitative real-time PCR. The methyltransferase genes (MLL1, MLL2, MLL3, MLL4, MLL5, and SETD1A) were elevated and demethylase Jarid1b was decreased with IL-13 treatment (Figure 2D). Of these, MLL1 showed the greatest change. We validated IL-13-induced elevation of MLL1 by Western blot (Figure 2E). These results suggest that IL-13-induced H3K4me3 elevation was probably regulated by both increase in methyltransferase and reduction of demethylase. However, the methyltransferase MLL1 plays a predominant role.

Knockdown of MLL1 reversed IL-13-induced changes of gene expression

To evaluate whether MLL1 influences IL-13-induced metaplasia, we knocked down MLL1 with lentivirus and examined mRNA expression of FOXJ1, DNAI2, MUC5a, and CLCA1. First, we validated MLL1 shRNA efficiency. Compared with control, MLL1 mRNA and protein expression was significantly reduced upon MLL1 knockdown (Figure 3A and B), and this was proved (Figure 2F). As expected, H3K4me3 expression markedly declined with MLL1 reduction (Figure 3B). The expression of FOXJ1 and DNAI2 significantly increased, while that of CLCA1 and MUC5a decreased compared with control (Figure 3C). These data suggest that MLL1 knockdown reverses the alterations of hallmark genes in nasal epithelium induced by inflammation.

Discussion

To our knowledge, this study is the first to investigate the connection between H3K4me3 and HNEpC metaplasia by IL-13. We provide additional evidence in favor of histone methylation involvement in nasal inflammation.

Histone modifications and DNA methylation are crucial for sustaining distinct gene expression. Cho et al showed that in nasal polyps, expression of HDAC2 increased compared to normal nasal inferior turbinates. We found that expression of H3K4me3 increased in nasal polyps compared with controls.

Many studies have suggested that a variety of Th2 inflammatory cytokines are implicated in development and maintenance of nasal inflammation, such as three representative cytokines (IL-4, IL-5, and IL-13). It is widely accepted that increased IL-4 and IL-13 can upregulate eotaxin production in epithelial cells, and IL-5 plays pivotal role in the recruitment and survival of eosinophils, which is also positively associated with serum total IgE. Nasal polyp tissue significantly highly expressed IL-5 and IgE in
CRSwNP compared with CRSsNP patients, which may lead to nasal polyp formation at specific sites with mucosal inflammation. As for immunoglobulin production, total IgE has also often been highly expressed within polyp tissue and fluid, especially in eosinophil CRSwNP, which in turn can contribute to local inflammation.

H3K4 methylation has played an important role in regulating inflammatory gene transcription through different ways. For example, Li et al. found that SET7/9, H3K4 methyltransferase, was involved in gene expression of TNF-α via recruitment of NF-κB p65 to inflammatory gene promoters in inflammation and immunity. Enrichment of H3K4me3 could increase the expression of IFN-γ and IL-4 produced by Th17 cells through interrupting the balance between native CD4+ T-cell precursors and Th1, Th2, and Th17 T-helper cell subsets in asthma, and IL-4 stimulation could increase H3K4me3 at IgE locus in CL-01 and primary B cells, resulting in high expression of IgE.

IL-13 has been widely recognized as an important cytokine in this process. In our study, we found that expression of H3K4me3 and MLL1 showed higher levels in IL-13-induced HNEpCs. To our knowledge, this is the first study to show that H3K4me3 and MLL1 expression levels increased in nasal polyps. As for the status of H3K4me3 in CRSwNP, further studies should be applied according to diverse clusters, which would be informative for individual treatment.

Increased gene expression of CLCA1 and MUC5AC and eosinophilic infiltration is seen following instillation of IL-13...
into mouse airway,\(^\text{42}\) mediated by the JAK/STAT6 pathway.\(^\text{43}\) In nasal polyps, STAT6-positive cells were localized in epithelium, gland cells, and inflammatory cells, and expression of STAT6 in epithelium had significantly increased compared with the control, which was positively associated with the recruitment of eosinophils.\(^\text{44}\) In the airway epithelium, IL-13 treatment decreased FOXJ1 mRNA expression via binding of STAT6 to the FOXJ1 promoter.\(^\text{45,46}\) We found increased MUC5A and CLCA1 and decreased FOXJ1 and DNAI2 at the transcriptional level with IL-13 treatment in HNEpC, consistent with previous reports.\(^\text{58}\) Recently, H3K4me3, an activated histone marker, was shown to be important in inflammatory processes,\(^\text{47}\) especially with high expression of IL-13.\(^\text{48}\) Our results suggest that H3K4me3 expression increases upon IL-13 treatment, which is at least partially attributed to methyltransferase MLL1. We examined four genes following MLL1 knockdown. It appears that CLCA1 and MUC5A expression is positively regulated by MLL1, although H3K4me3 enrichment at the promoter of these two genes needs to be further investigated. Carson et al\(^\text{49}\) found that MLL1-dependent H3K4me3 modification could regulate macrophage proinflammatory responses, indicating that MLL1 could be a novel therapeutic target for inflammatory diseases. Regarding FOXJ1 and DNAI2, the underlying mechanism remains to be explored in future. The signaling pathway of H3K4me3 and MLL1 remains poorly understood. Some studies have found that histone acetylation regulates chronic inflammatory disorders induced by IL-13 via STAT6 signaling pathway.\(^\text{50}\) We suspect that H3K4me3 may be involved in the STAT6 pathway. In addition, some studies have shown that Th2 cytokines such as IL-13 induce the airway inflammatory environment via the mitogen-activated protein kinase (MAPK) pathway.\(^\text{51,52}\) Histone modification could affect the p38 MAPK pathway, suggesting that histone modifications affect inflammatory development induced by IL-13 through MAPK.\(^\text{53,54}\) The function of the H3K4me3 and MLL1 signaling network needs to be further investigated.

Meanwhile, some studies have forced the necessity to develop inhibitors of MLL1 methyltransferase activity. MM-401 was able to inhibit MLL1 activity by blocking MLL1–WDR5 interaction without affecting other MLL family histone methyltransferases,\(^\text{55–57}\) supporting that MLL1 could be the target of epigenetic therapy. MLL1 may be a potential target for CRSwNP via influencing the process of H3K4me3 modification, especially in Th2 cytokine-dominant patients.

Figure 3 Knockdown of MLL1 reversed IL-13-induced changes of gene expression in HNEpC.

Notes: (A) MLL1 mRNA and (B) protein expression were measured upon MLL1 silencing. (C) Relative mRNA expression of nasal epithelial hallmark genes upon MLL1 knockdown in IL-13-treated HNEpC. Data are expressed as mean ± SD. *P < 0.05; Student’s t-test.

Abbreviation: HNEpC, human nasal epithelial cells.
Disclosure
The authors report no conflicts of interest in this work.

References


