

Clinical Values of miR-23a-3p in Oral Lichen Planus and Its Role in Keratinocyte Proliferation and Inflammatory Response

Jian Wang^{1,*}
Mingyan Hu^{2,*}
Leilei Li²

¹Department of Stomatology, Dongying Hospital of Traditional Chinese Medicine, Dongying, Shandong, 257000, People's Republic of China; ²Department of Stomatology, Dongying People's Hospital, Dongying, Shandong, 257091, People's Republic of China

*These authors contributed equally to this work

Purpose: Oral lichen planus (OLP) is a chronic inflammatory disease occurring in the oral cavity, and several miRNAs have been identified to be involved in the disease progression and malignant transformation. This study investigated the expression changes of miR-23a-3p in OLP patients, and further explored its functional role in keratinocyte cell proliferation and inflammatory response.

Patients and Methods: Fifty buccal mucosal tissue samples were collected from OLP patients. HaCaT cells were cultured with lipopolysaccharides (LPS) to mimic the condition of OLP in vitro. RNA extraction and quantitative real-time PCR (qRT-PCR) were used for the measurement of miR-23a-3p levels. The cell viability and inflammation were detected by using cell counting kit-8 (CCK-8) and enzyme-linked immunosorbent assay (ELISA). The target gene of miR-23a-3p was verified by using luciferase reporter assay.

Results: Compared with the control group, miR-23a-3p was significantly downregulated in the buccal mucosal tissues of OLP patients, and a remarkably decreased level of miR-23a-3p was detected in patients with erosive OLP. ROC curve demonstrated the diagnostic value of miR-23a-3p for OLP with the AUC of 0.908, it can also distinguish erosive OLP from the non-erosive ones. MiR-23a-3p level was negatively associated with RAE (reticular, atrophic, erosive) score in OLP patients ($r = -0.790$, $P < 0.001$). The in vitro experiments indicated that overexpression of miR-23a-3p reversed the promotive effect of LPS on HaCaT cell proliferation and reduced the protein levels of TNF- α and IL-6. The cyclin D1 (CCND1) was a direct target gene of miR-23a-3p, it was overexpressed in OLP cell models.

Conclusion: MiR-23a-3p was at the low expression in OLP patients and showed close association with the disease severity. Overexpression of miR-23a-3p might inhibit keratinocyte proliferation and inflammatory response via targeting CCND1.

Keywords: oral lichen planus, miR-23a-3p, keratinocyte cell, CCND1

Introduction

Oral lichen planus (OLP) is a chronic inflammatory disease occurring in the oral cavity, affecting approximately 2% of the population, especially middle-aged women.¹ OLP has a malignant potency, and according to the World Health Organization, approximately 1.63% of patients initially diagnosed with OLP can develop into oral squamous carcinomas.² Therefore, the early diagnosis and treatment of OLP are crucial to prevent disease deterioration. However, the etiology and pathogenesis of OLP are still not well understood. Keratinocytes are a major cell type in human skin, which plays an important role in normal skin metabolism.³

Correspondence: Leilei Li
Department of Stomatology, Dongying People's Hospital, No. 317, South Dongcheng Road, Dongying, 257091, Shandong, People's Republic of China
Tel/Fax +86-546-8901234
Email lileileidongying@163.com

Studies have shown that the degeneration of basal keratinocytes plays an important regulatory role in the development and exacerbation of OLP.⁴

MicroRNAs (miRNAs) are a group of small non-coding RNAs, which regulate the target gene expression through binding to their target messenger RNAs (mRNAs).⁵ Specific miRNAs have been reported to be aberrantly expressed in various human diseases, and are involved in the development and pathogenesis of diseases.⁶ In previous studies, miRNAs are reported to be involved in inflammation, metabolism, and development.^{7,8} In the field of OLP, several miRNAs have been identified to be differentially expressed in OLP patients, and play a role in the disease progression and malignant transformation.^{9,10} For example, miR-4731 is identified to be low expressed in OLP patients, furthermore, the low level of miR-4731 also shows close association with the occurrence and survival outcome of oral squamous cell carcinomas (OSCCs).¹¹ In addition, several miRNAs have been detected to be aberrantly expressed in OLP cases, such as miR-155, miR-27b, and so on.^{12,13} MiR-23a-3p is a member of the miR-23a-27a-24-2 cluster, it has been suggested to exert its biological functions in different cancers through regulating cellular activation, cell motility and immune cell infiltration.^{14,15} In a study reported by Chen et al, miR-23a-3p was identified to be downregulated in the tissues of oral squamous cell carcinomas (OSCCs) patients, and overexpression of miR-23a-3p was suggested to suppress tumor proliferation and invasion of OSCC.¹⁶ Additionally, the dysregulation of miR-23a-3p was also detected in OLP patients through next-generation sequencing.¹⁷ However, the functions of miR-23a-3p in OLP remain unclear.

In the present study, the expression changes of miR-23a-3p were verified in OLP patients using qRT-PCR, and its clinical role was investigated. Additionally, we further explored the functional role of miR-23a-3p in keratinocyte cells behaviors.

Patients and Methods

Study Subjects

This study included 50 OLP patients, who were diagnosed with OLP in Dongying People's Hospital from May 2017 to March 2019. The OLP patients were diagnosed according to the clinicopathologically modified WHO diagnostic criteria of OLP.¹⁸ The exclusion criteria were as follows: cases with the age below 18 or above 70 years old;

pregnant women; patients who suffered from oral lesions adjacent to metal crowns or amalgam fillings; individuals with other detectable oral lesions or systemic diseases, or received treatment 3 months before the sample collection. Patients receiving any medication that can cause lichenoid reactions such as antihypertensive drugs and oral hypoglycemics were also excluded from the study. Besides, another 45 healthy individuals were recruited as a control group, who were detected to have no detectable oral lesions or systemic diseases. Control buccal mucosal tissues were collected during wisdom tooth extraction or orthognathic surgery. The OLP and control groups had no significant difference in age and gender. The clinical characteristics were recorded in Table 1. Before treatment, buccal mucosal tissue samples were collected from each participant.

Table 1 Clinical Characteristics of the Subjects

Variables	Healthy Group (n = 45)	OLP (n = 50)	P value
Age, years	49.02±13.00	48.52±12.33	0.847
Gender			0.694
Male	11	14	
Female	34	36	
Smoking			0.837
No	35	38	
Yes	10	12	
Drinking			0.783
No	37	40	
Yes	8	10	
Position			
Single site	–	23	
Buccal mucosa	–	18	
Tongue	–	4	
Lips	–	1	
Multiple sites	–	27	
Clinical classification			
Erosive type	–	31	
Reticular striation	–	15	
Atrophic type	–	2	
Patch type	–	2	

The Ethics Committee of Dongying People's Hospital approved all procedures of the present study, and written informed consent was obtained from each individual. This study was conducted in accordance with the Declaration of Helsinki.

Cell Culture and Transfection

The human immortalized skin keratinocyte cell line, HaCaT, was purchased from American Type Culture Collection (Manassas, VA, USA), and cultured in DMEM supplemented with 10% FBS (Thermo Fisher Scientific, Waltham, MA, USA) at 37°C in a humidified atmosphere containing 5% CO₂. To mimic the condition of OLP in vitro, HaCaT cells were cultured with 10 µg/mL of LPS for 48h according to the previous evidence.¹⁹ In brief, HaCaT cells were plated into a six-well plate at the density of 5×10^4 /well and maintained for 12 h. Then, the cells were cultured for another 48h in the absence of LPS (10 µg/mL).

To regulate the expression levels of miR-23a-3p, transfection expression was performed when the cells reached the growth stage. Lipofectamine 2000 (Invitrogen, Carlsbad, CA, USA) was used to transfect miR-23a-3p mimic, miR-23a-3p inhibitor, or the negative control (miR-NC) into cells. After transfection, qRT-PCR was used to evaluate the transfection efficiency.

RNA Extraction and Quantitative Real-Time PCR (qRT-PCR)

Total RNA was extracted by using TRIzol reagent, then was reversed into complementary cDNA by the miRNA cDNA Synthesis Kit (CWBiotech) according to the manufacturer's instructions. Finally, the expression levels of miR-23a-3p were measured by using miRNA qPCR Assay Kit (CWBiotech) in ABI7300 real-time PCR system (Applied Biosystems; Thermo Fisher Scientific, Inc). The following thermocycling conditions were used for the PCR: initial denaturation at 94°C for 2 min, followed by 40 cycles of 94°C for 20 sec, 60°C for 34 sec. In the reaction, U6 was used as the internal reference gene, and the relative expression of miR-23a-3p was calculated using the $2^{-\Delta\Delta C_t}$ method. The primer sequences were as follows: miR-23a-3p (5'-CCAGGAACCCCTCCTTACTC-3' and 5'-TCTAGGGATGGTCCGAAGGA-3'); U6 (5'-CTCGCTTCGGCAGCACATATACT-3' and 5'-ACGCTT CACGAATTTGCGTGTC-3').

CCK-8 Assay

The change of the cell viability was measured by using (cell counting kit-8) CCK-8 reagent (Dojindo, Japan). Cells were first seeded in 6-well plates with a density of 4×10^5 . Before detection, 10 µL of reagent was added to the well plate. After incubation in the incubator for 1 h, the absorbance at 490 nm was detected using a Bio-Rad iMark plate reader (Bio-Rad Laboratories, Inc). The detection was performed every 24h, and three consecutive tests were performed to assess the relative viability of HaCaT cells.

Enzyme-Linked Immunosorbent Assay (ELISA)

The protein levels of inflammatory cytokines were examined by using the ELISA kits (Boster Biotechnology Company, Wuhan, China) according to the manufacturer's protocols.

Luciferase Reporter Assay

The candidate target gene of miR-23a-3p was predicted by using TargetScan Release 7.0 (<http://targetscan.org/>), and then was verified through the luciferase reporter assay. The 3'-untranslated region (UTR) of CCND1 was cloned into the luciferase reporter vector psiCHECK-2 (Promega Corporation) according to the manufacturer's instruction. Briefly, 500 ng of each reporter construct [wild-type (WT) or mutant 3'-UTR of CCND1] and miR-23a-3p mimic, miR-23a-3p inhibitor or miR-NC were co-transfected into pre-confluent (60-70%) HaCaT cells using Lipofectamine 2000 (Invitrogen, USA) for 48 h at 37°C. The relative luciferase activity was measured by the Dual-Luciferase Reporter System (Promega Corporation, USA) according to the instructions of the manufacturer. Renilla fluorescence activity was identified as the internal reference.

Statistical Analysis

All data analysis was carried out using the SPSS version 18.0 software (SPSS Inc., Chicago, IL) and GraphPad Prism 5.0 software (GraphPad Software, Inc., USA). The differences between groups were analyzed by Student's *t*-test or one-way ANOVA analysis. Pearson's analysis was used for correlation analysis. The statistical outlier limits were calculated (greater than two standard deviations from the mean), and no significant outliers were detected. The receiver operating characteristic (ROC) curve was used to detect the diagnostic value of miR-23a-3p. The differences were considered significant when $P < 0.05$.

Results

Clinical Characteristics of the Study

Subjects

As shown in Table 1, there were 11 males and 34 females in the healthy group, with a mean age of 49.02 ± 13.00 years old. The mean age of the OLP patients was 48.52 ± 12.33 years, including 14 males and 36 females. The statistical analysis results indicated that there was no significant difference in age, gender, smoking, and drinking status between the control and OLP groups (all $P > 0.05$). In addition, the position and clinical classification of OLP patients were recorded in Table 1.

MiR-23a-3p is Downregulated in OLP

Patients

The expression levels of miR-23a-3p were detected by qRT-PCR in buccal mucosal tissues of OLP patients. As shown in Figure 1A, miR-23a-3p was significantly lower in OLP patients than that in healthy controls ($P < 0.001$). Additionally, all OLP patients recruited in the present study were divided into the non-erosive group ($n = 29$) and erosive group ($n = 21$). It was noted that miR-23a-3p was down-regulated in erosive OLP patients compared with non-erosive cases (Figure 1B, $P < 0.001$).

Diagnostic Value of miR-23a-3p in OLP

Patients

Considering the dysregulation of miR-23a-3p in OLP patients, a ROC curve was constructed to calculate the diagnostic value of miR-23a-3p for OLP. According to the levels of miR-23a-3p in OLP patients and healthy controls, the ROC curve was constructed. As shown in

Figure 2A, the area under the curve (AUC) was 0.908, with the sensitivity of 84.0% and specificity of 86.7% at the cutoff value of 0.820. It suggested that miR-23a-3p can distinguish OLP patients from healthy controls. Additionally, according to the levels of miR-23a-3p in non-erosive and erosive OLP patients, a ROC curve was also constructed. The results showed that the AUC was 0.831, with a sensitivity of 71.4% and specificity of 86.7% at the cutoff value of 0.550 (Figure 2B). Erosive OLP is featured by more evident oral mucosal damage, it is generally considered to be a more severe form than non-erosive OLP. In terms of the low level of miR-23a-3p in erosive OLP cases, it was concluded that miR-23a-3p might be associated with the severity of OLP.

Correlation of miR-23a-3p with RAE Scores in OLP Patients

RAE (reticular, atrophic, erosive) score is well known to evaluate the severity of OLP patients. In the current study, we further explored the association of miR-23a-3p level with RAE scores in OLP patients (Figure 3). The results indicated that miR-23a-3p level was negatively associated with RAE score in OLP patients ($r = -0.790$, $P < 0.001$), suggesting miR-23a-3p was associated with the severity of OLP.

Overexpression of miR-23a-3p Inhibits HaCaT Proliferation

To investigate the role of miR-23a-3p in vitro, HaCaT cells were treated with LPS to mimic the OLP condition. It was found that after LPS treatment, the miR-23a-3p levels were downregulated significantly in HaCaT cells compared with the control group, which was consistent with the results observed in buccal mucosal tissues of OLP patients (Figure 4A). After cell transfection, the level of

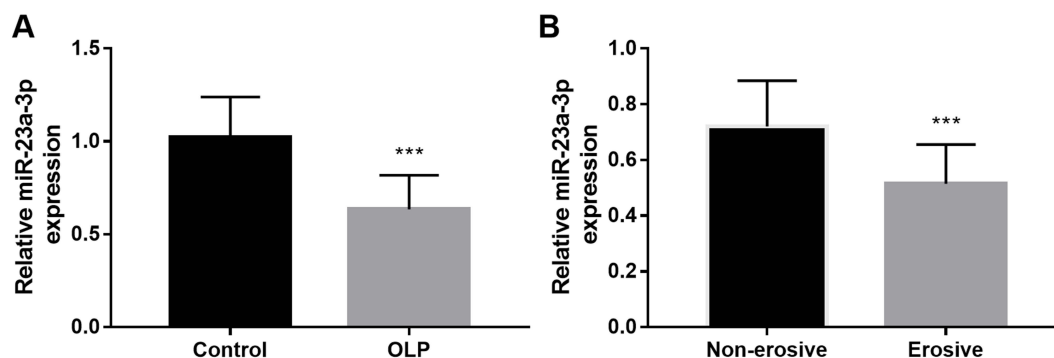


Figure 1 The expression levels of miR-23a-3p in OLP patients. (A) MiR-23a-3p was significantly lower in OLP patients than that in healthy controls. (B) MiR-23a-3p was down-regulated in erosive OLP patients compared with non-erosive cases. *** $P < 0.001$.

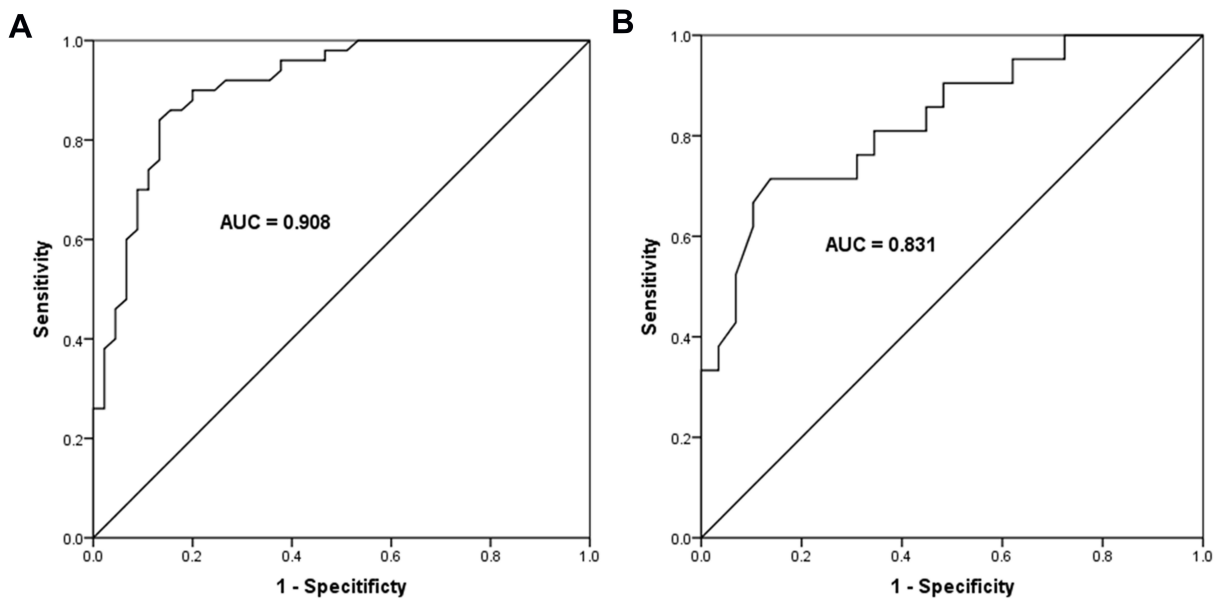


Figure 2 Diagnostic value of miR-23a-3p in OLP patients. **(A)** According to the levels of miR-23a-3p in OLP patients and healthy controls, miR-23a-3p was suggested to distinguish OLP patients from healthy controls, with the AUC of 0.908, the sensitivity of 84.0% and specificity of 86.7% at the cutoff value of 0.820. **(B)** According to the levels of miR-23a-3p in non-erosive and erosive OLP patients, miR-23a-3p could distinguish erosive OLP patients from non-erosive ones, with the AUC of 0.831, the sensitivity of 71.4% and specificity of 86.7% at the cutoff value of 0.550.

miR-23a-3p was increased significantly by miR-23a-3p mimic transfection, whereas miR-23a-3p inhibitor transfection reduced the level of miR-23a-3p (Figure 4A). The OD value in the CCK-8 assay suggested that LPS treatment inhibited the HaCaT cell viability compared with the untreated control group (Figure 4B). However, overexpression of miR-23a-3p reversed the inhibition of cell viability induced by LPS, whereas miR-23a-3p downregulation further inhibited the cell viability (Figure 4B). Moreover, the protein levels of inflammatory cytokines were further detected. It was found that LPS promoted

the release of TNF- α and IL-6, which was reversed by miR-23a-3p upregulation. However, downregulation of miR-23a-3p aggravated the release of TNF- α and IL-6 induced by LPS (Figure 4C).

CCND1 is the Target Gene of miR-23a-3p

According to the TargetScan analysis results, the complementary sequence of miR-23a-3p was found in the 3'-UTR of CCND1 (Figure 5A). Furthermore, the luciferase reporter assay results indicated that overexpression of miR-23a-3p inhibited the luciferase activity of WT 3'-UTR of CCND1 (Figure 5B). But for the luciferase activity of MUT 3'-UTR of CCND1, there was no significant difference among different groups. In addition, CCND1 was upregulated in HaCaT cells after LPS treatment, but miR-23a-3p mimic transfection reduced the level of CCND1, and miR-23a-3p downregulation further increased the level of miR-23a-3p (Figure 5C).

Discussion

MiRNAs are a class of non-coding RNAs composed of 19–25 nucleotides. MiRNAs can inhibit the target gene expression at the transcriptional levels through binding to the 3'-UTR of mRNA.¹³ The bioinformatic studies have suggested that one-third of known genes may be regulated by different miRNAs.²⁰ Many studies have reported that

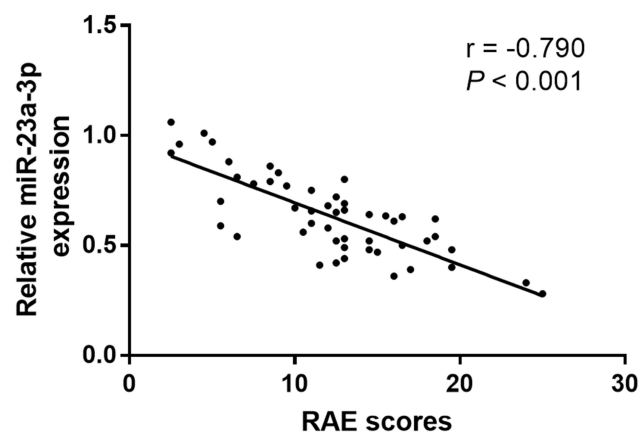


Figure 3 Correlation of miR-23a-3p with RAE scores in OLP patients. MiR-23a-3p level was negatively associated with RAE score in OLP patients ($r = -0.790$, $P < 0.001$).

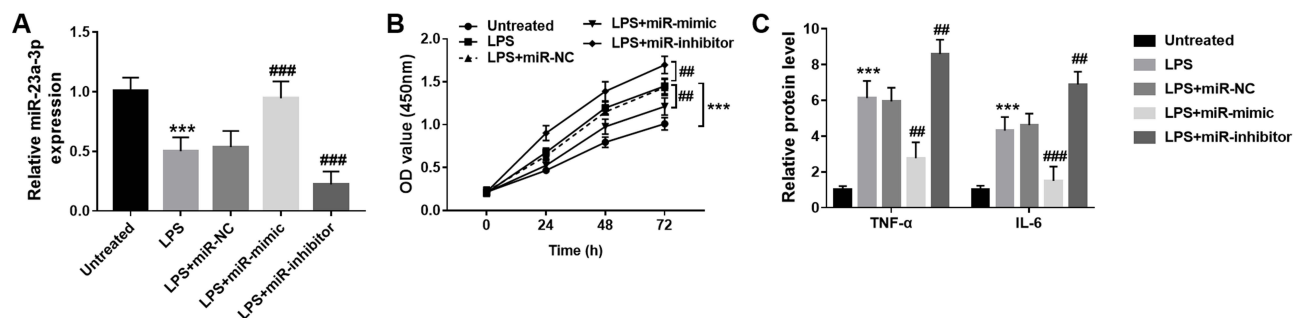


Figure 4 Overexpression of miR-23a-3p inhibits HaCaT proliferation and inflammatory response. (A) The expression level of miR-23a-3p was decreased by LPS treatment, which was reversed by miR-23a-3p mimic transfection. (B) LPS promoted the cell proliferation, which was reversed by miR-23a-3p mimic transfection. (C) LPS promoted the release of TNF- α and IL-6, which was reversed by miR-23a-3p upregulation. However, downregulation of miR-23a-3p aggravated the release of TNF- α and IL-6 induced by LPS. *** $P < 0.001$, compared with untreated group; ## $P < 0.01$, ### $P < 0.001$, compared with LPS group.

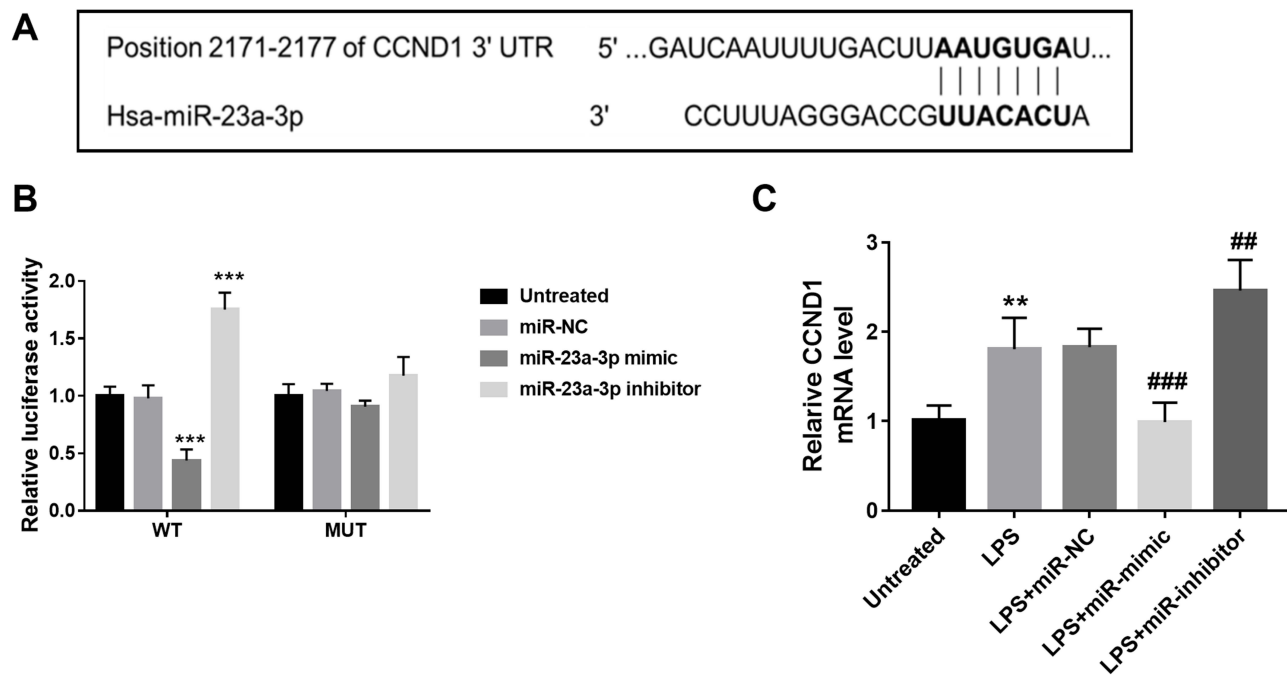


Figure 5 CCND1 was the target gene of miR-23a-3p. (A) According to the TargetScan analysis results, the complementary sequence of miR-23a-3p was found in the 3'-UTR of CCND1. (B) Overexpression of miR-23a-3p inhibited the luciferase activity of WT 3'-UTR of CCND1. But for the luciferase activity of MUT 3'-UTR of CCND1, there was no significant difference among different groups. (C) CCND1 was upregulated in HaCaT cells after LPS treatment, but miR-23a-3p mimic transfection reduced the level of CCND1, and miR-23a-3p downregulation further increased the level of miR-23a-3p. ** $P < 0.01$, *** $P < 0.001$ compared with untreated group. ## $P < 0.01$, ### $P < 0.001$, compared with LPS group.

miRNAs are involved in many important cellular processes, such as cell apoptosis, cell differentiation, proliferation, tumorigenesis and tumor inhibition.^{21,22} A large number of researches have proved that miRNA is an important epigenetic regulatory factor in the occurrence and progression of various human diseases.^{23,24}

Previous evidence has demonstrated that a variety of miRNAs are differentially expressed in OLP patients. For example, miR-146a and miR-155 were identified to be over-expressed in OLP lesions tissues.²⁵ Ma et al reported that approximately 70 miRNAs were identified to be differentially expressed in OLP patients compared with healthy controls

through microarray analysis.²⁶ In the present study, we confirmed that miR-23a-3p was down-regulated in the buccal mucosal tissues of OLP patients. And the ROC curve analysis results further indicated that miR-23a-3p expression can distinguish OLP patients from healthy controls. Consistent with the present results, a study by Chen et al reported the down-regulation of miR-23a-3p in the oral mucosa of OLP patients by using next-generation sequencing with DESeq and edgeR software algorithms, indicating its potential role in the etiology and pathogenesis of OLP, which supported our present results.¹⁷ All evidence indicated the potential role of miR-23a-3p in OLP.

Oral lichen planus is a chronic disease that can be controlled but not eliminated. An erosive form of OLP results in pain and ulceration, which may most likely transform or develop into malignant tumors.²⁷ In the current study, a total of 50 OLP patients were recruited, who were further divided into non-erosive group and erosive group. Then the qRT-PCR results indicated that miR-23a-3p was down-regulated in erosive OLP patients compared with non-erosive cases. Considering the expression changes of miR-23a-3p between erosive and non-erosive OLP patients, we speculated that miR-23a-3p might be associated with the severity of OLP. RAE score is well known to evaluate the severity of OLP patients. In the current study, we further explored the association of miR-23a-3p level with RAE scores in OLP patients, and a significantly negative association was detected between miR-23a-3p level and RAE scores. Additionally, according to the levels of miR-23a-3p in non-erosive and erosive OLP patients, a ROC curve was also constructed. The results indicated that miR-23a-3p can distinguish erosive OLP patients from non-erosive OLP patients. Erosive OLP is featured by more evident oral mucosal damage, it is generally considered to be a more severe form than non-erosive OLP. In terms of the low level of miR-23a-3p in erosive OLP cases, we presented that miR-23a-3p might be associated with the severity of OLP.

OLP is an immune disease, and the pathological process of OLP has been reported to be closely linked to basal keratinocytes degeneration.²⁸ HaCaT cells are a type of immortalized human skin keratinocytes, which are easily grown and passaged, thus HaCaT cells are suitable for the research of oral keratinocytes. Gram-negative bacterial lipopolysaccharide (LPS) has been widely used in previous studies to induce inflammatory mediators.²⁹ Accordingly, in the present study, HaCaT cells were treated with LPS to establish the in vitro inflammatory OLP model. Initially, we found that after LPS treatment, the miR-23a-3p levels were downregulated significantly in HaCaT cells, which was consistent with the results observed in buccal mucosal tissues of OLP patients. It has been indicated that cell proliferation of keratinocyte was promoted in OLP progression, the same result was also detected in vitro model of OLP, which were consistent with the results observed in LPS-treated HaCaT cells in the current study.^{30,31} Additionally, the gain and loss function experiments' results suggested that overexpression of miR-23a-3p reversed the effect of LPS on HaCaT cell proliferation. We concluded that miR-23a-3p might be involved in the pathogenesis of OLP through inhibiting keratinocyte proliferation.

OLP is an immune disease, and the pathological process of OLP has been reported to be closely linked to the release of

inflammatory cytokines.²⁸ In the OLP cell model, LPS promoted the release of TNF- α and IL-6, which was consistent with the previous evidence. Furthermore, the gain and loss function experiment results demonstrated that overexpression of miR-23a-3p inhibited the inflammatory response of keratinocyte cells. miR-23a-3p has been widely reported to be involved in the inflammatory response in various cell types and diseases. In the study of traumatic brain injury, miR-23a-3p is reported to alleviate the brain injury via inhibiting neuroinflammation.³² Downregulation of miR-23a-3p is also identified in psoriatic arthritis patients and enhanced secretion of inflammatory cytokines in synovial fibroblasts.³³ The present results indicated that overexpression of miR-23a-3p inhibited the keratinocyte cell inflammatory response during the progression of OLP, which was supported by the previous evidence.

The cyclin D1 (CCND1) gene, located on chromosome 11q13, is a proto-oncogenic positive regulator of the cell cycle, driving cells from G1 into S phase checkpoint.³⁴ Previous studies have indicated that the expression of cell cycle regulatory proteins is dysregulated in oral premalignant and malignant lesions.^{34,35} In the present study, the luciferase reporter assay results indicated that CCND1 was a direct target gene of miR-23a-3p, and CCND1 was upregulated in the LPS induced OLP cell model. The involvement of CCND1 in OLP has been widely reported in previous studies.^{36,37} In addition, an elevated level of CCND1 is identified by Yao et al in the epithelial cells of OLP patients by immunohistochemistry, and it was closely associated with the cell apoptosis index of the lymphocytes.³⁶ Besides, CCND1 is also detected to be at the high expression in the mucosa samples of OLP patients, and the aberrant expression is involved in the protective role of miR-138 in OLP.³⁸ It is known that CCND1 is a target of the WNT signaling pathway, which has been reported to have a significant role in malignant transformation of oral lesions such as OLP, and contribute to the chronic inflammatory disease.³⁹ Accordingly, we speculated that miR-23a-3p might inhibit keratinocyte proliferation and inflammatory response via targeting CCND1. However, the underlying mechanism is worthy of exploration. In the future, other studies involving the role of CCND1 as well as WNT signaling in OLP are needed to confirm our speculation.

Conclusion

In conclusion, the current study suggested that miR-23a-3p was downregulated in OLP patients, and might be associated with the disease severity. Overexpression of miR-23a-3p might inhibit keratinocyte proliferation via targeting CCND1. The

present results provide a basis for further study of the molecular mechanisms underlying the development and progression of OLP.

Ethics Statement

The Ethics Committee of Dongying People's Hospital approved all procedures of the present study, and written informed consent was obtained from each individual.

Disclosure

The authors report no conflicts of interest in this work.

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