The Long Noncoding RNA Linc01833 Enhances Lung Adenocarcinoma Progression via MiR-519e-3p/S100A4 Axis

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Yuan Zhang^{1,*}
Wenhua Li^{1,*}
Zongxiang Lin¹
Jingfeng Hu¹
Jingpu Wang¹
Yukai Ren¹
BoChong Wei¹
Yuxia Fan²
Yang Yang¹

¹Department of Thoracic Surgery, The First Affiliated Hospital, Zhengzhou University, Zhengzhou 450052, People's Republic of China; ²Department of Thyroid Surgery, The First Affiliated Hospital of Zhengzhou University, Zhengzhou 450052, People's Republic of China

*These authors contributed equally to this work.

Introduction: Lung cancer (LC) is among the most prevalent malignancies worldwide, with extremely high morbidity and mortality rates. Mounting evidence has suggested that the abnormally expressed long noncoding RNA (lncRNA) in lung cancer tissues may play vital roles in tumor progression. In the present research, we aimed to examine the functions and underlying mechanism of linc01833 in lung adenocarcinoma (LUAD).

Methods: qRT-PCR was employed to determine transfection efficiency. CCK-8, transwell invasion assay, Western blotting analysis and qRT-PCR were used to detect proliferation as well as migration of different LUAD cell lines, and were also applied to determine the changes during epithelial-mesenchymal transformation (EMT). Afterwards, bioinformatics and dual-luciferase reporter assay were utilized to explore and to identify the potential corresponding targets of linc01833 and miR-519e-3p.

Results: Linc01833 OE can significantly improve proliferation as well as invasion ability of LC cells and promote the EMT process. Dual-luciferase reporter assay demonstrated that linc01833 could directly bind to miR-519e-3p, thereby inhibiting its expression. Further experiments showed that S100A4 was a direct target of miR-519e-3p. Rescue assay demonstrated that linc01833 acted on the miR-519e-3p/S100A4 axis.

Conclusion: We verified the mechanism of linc01833 in promoting infiltration and metastasis in LUAD. To be specific, linc01833 can function as a competitive endogenous RNA (ceRNA) to adsorb miR-519e-3p through a sponge and regulate S100A4 in lung cancer, thereby being involved in LUAD progression. Collectively, our research provides new insights towards the in-depth understanding of LC progression mechanisms.

Keywords: lung cancer, linc01833, miR-519e-3p, S100A4, EMT

Introduction

Lung cancer (LC) is among the most prevalent malignancies globally due to its high morbidity and mortality rates. Among all types of LC, the incidence of lung adenocarcinoma (LUAD) has exceeded that of lung squamous cell carcinoma (LUSC), showing an increasing trend year by year. Up to now, LUAD accounts for nearly 40–50% of all types of LC. Due to the aggressiveness and metastasis of malignant cancer cells, poor prognosis is common among patients.¹ The 5-year survival rate of LC patients is only about 18%, without any significant breakthrough or improvement.²

Mounting attention has been paid to long noncoding RNA (lncRNA) in tumor research due to the accumulative evidence indicating the critical involvement of

Correspondence: Yuxia Fan
The First Affiliated Hospital of Zhengzhou University, No. 1 East Jianshe Road, Zhengzhou 450052, People's Republic of China
Email fanyuxia1981@126.com

Yang Yang
The First Affiliated Hospital, Zhengzhou University, No. 1 East Jianshe Road, Zhengzhou 450052, People's Republic of China
Email fccyangl@zzu.edu.cn

lncRNA in tumor development. lncRNA can regulate the pathophysiological process of a tumor through various pathways. At present, evidence shows the abnormal expression of lncRNA in diverse types of tumor tissues, with significant effects on tumor proliferation and metastasis. For instance, lncRNA MALAT1 can enhance the proliferation and metastasis of epithelial ovarian cancer cells,³ but it suppresses breast cancer development.⁴ Additionally, lncRNA FAL1 can accelerate the proliferation and metastasis of liver cancer cells by functioning as a ceRNA through competitive binding with miR-1236,⁵ which also promotes progression of colorectal cancer (CRC) by modulating the miR-637/NUPR1 pathway.⁶ lncRNA MEG3 is capable of suppressing proliferation and invasion of gastric cancer (GC) cells through the p53 signaling pathway, which is considered as a novel therapeutic target for GC.⁷ Similarly, lncRNA MEG3 can also inhibit metastasis as well as invasion in GC by regulating RNA-21, bringing new hope of long-term survival for GC patients.⁸ Despite a further understanding of the functional mechanisms of lncRNAs by conducting extensive studies on different lncRNAs, some lncRNAs have not been explored, therefore, their roles in tumorigenesis and tumor development remain unknown. Of note, linc01833 is a target that deserves further study. Our previous research showed that linc01833 was highly expressed in LUAD tissues, which enhanced metastasis and invasion of LC. Nevertheless, the functions of linc01833 in LUAD and its related mechanism have not been extensively reported.

The present study attempted to determine the impact of linc01833 on LUAD progression and to explore its underlying mechanism. For the first time, our experimental results confirmed that linc01833 was capable of enhancing proliferation and invasion in LUAD via the modulation of miR-519e-3p/S100A4, thereby being involved in LUAD tumorigenesis and progression.

Materials and Methods

Data Collection and Bioinformatics Analysis

The Cancer Genome Atlas (TCGA) database was used to extract relevant data for cluster analysis. linc01833 expression was assessed in non-small cell lung cancer (NSCLC) and normal tissue, followed by construction of a boxplot for intuitive observation and comparison. Afterwards, we analyzed and compared the overall

survival (OS) of LC patients with different expression levels of linc01833.

Cell Culture

NSCLC cell lines, A549 and HCC4006, were obtained from Shanghai Cell Bank of Chinese Academy of Sciences, which were maintained in DMEM (Invitrogen, Carlsbad, CA), containing 10% FBS (Gibco, MA, USA) and 100 µg/mL penicillin (Invitrogen) at 37 °C in a humidified incubator with 5% CO₂.

Transfection

The linc01833 overexpression (OE) vector, S100A4 OE vector and universal negative control (NC) were provided by Genescript (Shanghai, China). The 3'-untranslated regions (UTR) in the recognition and binding sites of wild type (WT) or mutant type (mut) of miR-519e-3p in S100A4 were amplified, followed by cloning into pGL3 vector to form pGL3-WT-S100A4-3'-UTR or pGL3-mut-S100A4-3'-UTR. Lipofectamine 2000 (Invitrogen) was adopted for cell transfection, followed by further experiments after incubation.

Cell Proliferation Assay

We measured and evaluated the cell proliferation at different time points using CCK-8 kit. Specifically, all transfected cells were inoculated in a 96-well plate, followed by addition of CCK-8 reagent every 24 h to ensure that CCK-8 was always maintained at a stable target concentration. After continuous incubation of cells in standard environment, the optical density values at 450 nm wavelength were measured at 0, 24, 48, 72, and 96 h by spectrophotometer, followed by measurement of cell proliferation rate. The assays were conducted in triplicate and repeated three times or more to ensure the reliability of results.

RNA Extraction as Well as qRT-PCR

TRIzol reagent (Invitrogen) was utilized to isolate total RNA from NSCLC cells, which was transcribed into cDNA by Prime Script RT kit (Takara, Osaka, Japan), followed by cDNA amplification by qRT-PCR utilizing SYBR Green master mix (Thermo Fisher). Relative expression of mRNA or lncRNA was calculated using the 2^{-ΔΔCT} method, followed by normalization to β-actin levels. Each sample was divided into three groups and analyzed repeatedly.

Transwell Invasion Assay

QCM-EC Matrix cell invasion test (Merck, Darmstadt, Germany) was employed to determine cell invasion

capacity. Specifically, the transfected cells suspended in FBS-free medium were added into matrix gel (BD Biosciences, San Jose, CA) in the chamber. Complete medium was further supplemented to lower chambers. Cells were incubated in standard conditions until they crossed the ECM layer and adhered firmly to the base of the fixed membrane. The cells crossing the fixed membrane were fixed with methanol, subjected to H & E staining, observed and counted under microscope at 10× magnification at three randomly selected views. The assay was repeated three times.

Dual-Luciferase Reporter (DLR) Assay

In this study, the dual-luciferase detection kit (Promega, Madison, USA) was used for relevant experimental studies and to analyze the dual-luciferase reporter activity. Specifically, A549 and HCC4006 cells were co-transfected with linc01833 WT, linc01833 mut, miR-519e-3p, S100A4 WT, S100A4 mut, or NC siRNA. Afterwards, cells were digested, dissolved, and centrifuged to collect supernatant, followed by measurement of renilla and firefly luciferase activity by microplate photometer (BioTeck, Winooski, VT, USA). All the experiments were performed in triplicate, followed by calculation of the average value to ensure the reliability of the experimental results.

Western Blot (WB) Analysis

Total protein was extracted from A549 and HCC4006 cells, lysed in RIPA buffer (Roche, Branchburg, NJ) containing protease inhibitor. Protein sample was subjected to SDS-PAGE, transferred to PVDF membrane, blocked with 5% BSA, rinsed with PBS, followed by reaction with primary antibody (anti-E-cadherin, anti-cytokeratin, anti-vimentin, anti-S100A4 and anti-GAPDH) overnight at 4°C. After rewashing, membranes were reacted with HRP-conjugated secondary antibody. Immunoreactivity was observed using Odyssey machine (Roche Diagnostics) with ECL detection reagent (Amersham Biosciences). GAPDH was used as an internal control.

Immunofluorescence Analysis

The slides were soaked and washed three times with phosphate buffer saline (PBS). After fixing with 4% paraformaldehyde for 15 min, the slides were soaked with PBS again. After draining PBS with absorbent paper, the slides were sealed with 5% goat serum at room temperature for 30 min. The slides were incubated with primary

antibodies against cytokeratin (1:400 dilution) and vimentin (1:200 dilution) at 4°C overnight, followed by incubation with anti-rabbit secondary antibodies at room temperature for 30 min. Finally, the liquid on the slide was dried using absorbent paper and the slides were sealed with the sealing liquid. The slides were observed and photographed under a fluorescence microscope.

Data Analysis

SPSS version 21.0 (SPSS, Chicago, IL) was used for data statistics and analysis. In brief, categorical data were displayed by frequency and percentage, and continuous data were shown as means ± SD. One-way ANOVA or Student's *t*-test was utilized for statistical analysis if appropriate. Pearson correlation was used to analyze the correlation between different gene expressions. Kaplan-Meier method and Log rank test were used to analyze the survival curve. *P*<0.05 suggested statistical significance.

Results

Biological Characteristics of Linc01833 and Its Relationship with Clinical Features

The purpose of this experiment was to examine the biological effects and mechanisms of linc01833 in LUAD. Further mining of the TCGA database uncovered increased linc01833 expression in LUAD tissues (Figure 1A). Survival analysis demonstrated an association of linc01833 expression with OS in LUAD patients (Figure 1B). qRT-PCR assay further revealed significantly higher expression of linc01833 in LUAD tissue than the adjacent normal tissue (Figure 1C).

Biological Role of Linc01833 in LUAD

To confirm the role of linc01833 in LUAD, linc01833 was overexpressed in two LUAD cell lines A549 and HCC4006, followed by measurement of the transfection efficiency (Figure 2A). CCK-8 assay revealed that linc01833 OE could significantly promote LUAD cell proliferation (Figure 2B and C). In consideration of the important role of EMT in cell metastasis, EMT markers were examined at mRNA and protein levels. WB showed that after OE of linc01833, the expression of cytokeratin and E-cadherin was significantly downregulated in A549 and HCC4006 cells, while that of vimentin was significantly upregulated, suggesting that OE of linc01833 promoted the EMT process of LUAD cells (Figure 2D). Similarly, qRT-PCR analysis showed consistent outcomes in A549 and HCC4006 cell

Figure 1 Mining the TCGA database uncovered the significant overexpression of linc01833 in LUAD ($P < 0.001$) - (A). Survival analysis demonstrated that the expression of linc01833 was related to OS in LUAD patients ($P < 0.001$) (B). qRT-PCR showed significantly increased level of linc01833 in LUAD tissue (cancer) than adjacent tissue (control). *** $P < 0.001$ (C).

Figure 2 Transfection efficiency of linc01833 overexpression in A549 and HCC4006 cells verified by qRT-PCR. *** $P < 0.001$ (A). Overexpression of linc01833 could significantly promote LUAD cell proliferation. *** $P < 0.001$ (B, C). Western blotting analysis indicated that linc01833 overexpression significantly enhanced EMT process in LUAD cells (D). qRT-PCR analysis confirmed that linc01833 overexpression significantly enhanced EMT process in LUAD cells. *** $P < 0.001$ (E, F). Immunofluorescence confirmed that linc01833 overexpression inhibited the expression of cytokeratin while promoting that of vimentin (G, H). Transwell invasion assays showed that overexpression of linc01833 significantly promoted migration of LUAD cells. *** $P < 0.001$ (I, J).

lines (Figure 2E and F). Immunofluorescence subsequently showed significant downregulation of cytokeratin expression but significant upregulation of vimentin expression (Figure 2G and H), which was consistent with the above

results. The migration ability of LUAD cells was assessed by transwell invasion assays, showing that OE of linc01833 could significantly improve the migration ability of LUAD (Figure 2I and J). Our experiments confirmed that linc01833

can significantly enhance proliferation, migration and metastasis of LUAD cells, which was also involved in the EMT process of LUAD cells.

MiR-519e-3p is a Direct Target of Linc01833

Bioinformatics software was applied to predict and analyze the possible binding miRNA targets of linc01833, aiming to explore its potential mechanism. This showed that miR-519e-3p had the potential to bind with linc01833, and its binding sequence was shown (Figure 3A). Subsequently, we used the DLR assay to determine whether miR-519e-3p could directly bind to linc01833 in A549 and HCC4006 cell lines. Consequently, miR-519e-3p can directly bind with wt linc01833, while there was insignificant difference in mut linc01833 group (Figure 3B and C). Moreover, miR-519e-3p expression was significantly inhibited in linc01833 OE group, indicating that linc01833 could negatively modulate miR-519e-3p level (Figure 3D).

Linc01833 Promotes LUAD via Negative Modulation of MiR-519e-3p

qRT-PCR analysis revealed that miR-519e-3p showed a significantly lower expression trend in LUAD tissue in comparison with adjacent tissue (control) (Figure 4A-a). Increased linc01833 expression in LUAD tissue is generally accompanied by decreased miR-519e-3p expression, and vice versa (Figure 4A-b). In addition, CCK-8 assay demonstrated that miR-519e-3p OE was capable of suppressing LUAD cell proliferation (Figure 4A-c-d). The functions of miR-519e-3p in LUAD were tested by WB (Figure 4A-e) and qRT-PCR (Figure 4A-f-g). The results showed that miR-519e-3p OE led to significantly increased expression of cytokeratin and E-cadherin, but significantly decreased expression of vimentin in LUAD cells, indicating that miR-519e-3p OE was capable of inhibiting the EMT process in LUAD cells (Figure 4A-h-i). Therefore, we believed that miR-519e-3p exerted an inhibitory effect on the

Figure 3 Potential binding sequences of linc01833 and miR-519e-3p (A). The dual-luciferase reporter assay suggested that miR-519e-3p was a direct target of linc01833. ***P<0.001 (B, C). Overexpression of linc01833 could suppress the level of miR-519e-3p. **P<0.01; ***P<0.001 (D).

Figure 4 (A): qRT-PCR analysis found significantly lower miR-519e-3p expression in LUAD, with negative association with linc01833 expression in LUAD. *** $P < 0.001$ (A-a-b). miR-519e-3p overexpression could significantly inhibit LUAD cell proliferation. *** $P < 0.001$ (A-c-d). Western blotting analysis proved that miR-519e-3p overexpression significantly inhibited EMT progression in LUAD (A-e). qRT-PCR analysis found that miR-519e-3p overexpression significantly inhibited EMT progression in LUAD. *** $P < 0.001$ (A-f-g). Transwell invasion assays showed that miR-519e-3p overexpression suppressed migration of LUAD cells. *** $P < 0.001$ (A-h-i). **(B):** miR-519e-3p overexpression reversed the proliferation of linc01833 overexpression in A549 and HCC4006 cells. *** $P < 0.001$ (B-a-b). Western blotting analysis proved that miR-519e-3p overexpression reversed the outcomes of linc01833 on EMT in LUAD cells (B-c). qRT-PCR analysis found that miR-519e-3p overexpression reversed the outcomes of linc01833 on EMT in LUAD cells. *** $P < 0.001$ (B-d-e). Transwell invasion assays revealed that miR-519e-3p overexpression reversed the effect of linc01833 on migration capacity of LUAD cells. *** $P < 0.001$ (B-f-g).

progression of LC. To further understand the correlation of linc01833 with miR-519e-3p, reverse assays were conducted. MiR-519e-3p OE could reverse the proliferative effect of linc01833 OE on A549 and HCC4006 cells (Figure 4B-a-b). Compared with linc01833 OE group, miR-519e-3p OE can upregulate the levels of cytokeratin and E-cadherin in LUAD cells, while it can downregulate the expression of vimentin (Figure 4B-c-e). Transwell invasion assay showed similar results (Figure 4B-f-g). The reverse assays revealed that miR-519e-3p OE reversed the effect of linc01833 on proliferation, migration and metastasis of LUAD cells. Therefore, our results suggested that linc01833 promoted LUAD development via negative modulation of miR-519e-3p.

The S100A4 is a Direct Target of MiR-519e-3p

For further investigation of the mechanism underlying the roles of miR-519e-3p on the biological behavior in LUAD, bioinformatics software was applied for prediction and analysis. A potential binding sequence of S100A4 and miR-519e-3p was found (Figure 5A). S100A4 might be a possible target of miR-519e-3p, which was further confirmed by DLR assay (Figure 5B and C). Further investigation found a negative correlation of miR-519e-3p expression with S100A4 expression in LC tissues (Figure 5D). WB and qRT-PCR analysis demonstrated that miR-519e-3p OE could significantly downregulate S100A4 expression, indicating the negative regulatory effects of miR-519e-3p on S100A4

Figure 5 Potential binding sequences of miR-519e-3p and S100A4 (A). Dual-luciferase reporter assay found S100A4 as a direct target of miR-519e-3p. ***P<0.001 (B, C). S100A4 expression was negatively correlated with miR-519e-3p expression in lung cancer tissues (D). Western blotting and qRT-PCR demonstrated that miR-519e-3p overexpression can significantly decrease S100A4 expression. ***P<0.001 (E, F). Western blotting and qRT-PCR revealed that linc01833 overexpression could significantly enhance the expression of S100A4. ***P<0.001 (G, H).

expression (Figure 5E and F). Linc01833 OE can significantly increase the expression of S100A4, either at protein or mRNA level, suggesting that linc01833 can positively regulate the expression of S100A4 (Figure 5G and H).

The Biological Role of Linc01833/MiR-519e-3p/S100A4 Axis in LUAD

Rescue assays were performed to examine the functions of the linc01833/miR-519e-3p/S100A4 axis in invasion and metastasis of LUAD. Compared with the co-transfection group, OE of S100A4 can reverse the effects of linc01833 and miR-519e-3p on LUAD cell proliferation (Figure 6A and B). We also detected the decreased expression of cytokeratin and E-cadherin, and increased vimentin level (Figure 6C–E). These outcomes collectively suggested that S100A4 OE can reverse the effects of OE of linc01833 and miR-519e-3p on EMT and proliferation in LUAD cells.

Discussion

An estimated 1.59 million people die of LC globally every year. Moreover, both incidence and mortality of LC are the highest among all types of malignant tumors, thus attracting more and more attention.^{9,10} LC is mostly characterized by relatively insidious onset and relatively rapid progression. Despite continuous advances in therapeutic concepts and technology, the 5-year survival rate of LC patients is still not high due to invasion and metastasis of LC. Thus, related research on the progress of LC metastasis has always been a hot spot.

LncRNA generally contains over 200 nucleotides (nt), generally without protein-coding function. LncRNAs can be classified according to different structures, including Sense lncRNA, Intronic lncRNA, Bidirectional lncRNA, Antisense lncRNA, Intergenic lncRNA (lincRNA), etc.^{11,12} The expression levels of some lncRNAs can be significantly changed during the development of malignant tumors, which are closely associated with diverse

Figure 6 Overexpression of S100A4 rescued the effect of linc01833 and miR-519e-3p on LUAD cell proliferation. *** $P < 0.001$ (A, B). Western blotting showed that overexpression of S100A4 could reverse the effects of linc01833 and miR-519e-3p on EMT process in LUAD cells (C). qRT-PCR analysis found that overexpression of S100A4 could reverse the effects of linc01833 and miR-519e-3p on EMT process in LUAD cells. ** $P < 0.01$; *** $P < 0.001$ (D, E).

biological processes and may participate in carcinogenesis and tumor development. LncRNAs are likely to be used as diagnostic and therapeutic markers in tumors.¹³ For instance, lncRNABORG can mediate the relapse and metastasis of breast cancer.¹⁴ By regulating SATB2, LncRNA SATB2-AS can affect the microenvironment of tumor immune cell growth and assist to decrease invasion and metastasis of CRC.¹⁵ However, there have been no relevant studies on linc01833 so far, thus, we explored its effect and mechanism in LC in our study.

Unlike lncRNA, miRNAs (with 18–25 nt in length) can bind with and degrade 3'UTR of target mRNA. Additionally, miRNA also has the ability to regulate coding and noncoding transcriptomes. Therefore, investigation into the regulatory mechanism of the lncRNA-miRNA-mRNA network can provide more comprehensive and in-depth understanding of pathogenesis as well as metastatic mechanisms of tumors, aiming to find possible therapeutic targets. At present, such networks have been widely applied to examine multiple kinds of malignancies.^{16–18} For instance, lncRNA H1 can function as a competitive ceRNA for modulating EMT and metastasis of bladder cancer via regulating miR-29b-3p expression.¹⁹ lncRANSNHG7 upregulates GALNT1 through sponge adsorption of miR-216b to enhance proliferation and liver metastasis in CRC.²⁰ In this study, bioinformatics was applied to reveal that miR-519e-3p is a possible target of linc01833. Combined with software prediction analysis, S100A4 is revealed as a possible target of miR-519e-3p. Afterwards, these results were also verified by DLR assay, showing a direct correlation between linc01833/miR-519e-3p/S100A4. S100A4 is a vital member of the calcium-binding protein S100 family, comprising 101 amino acids. Current studies have shown that S100A4 is involved in diverse physiological activities, including cell movement and adhesion, extracellular matrix reconstruction, tumor angiogenesis, etc.^{21–24} Meanwhile, studies have found that it is closely related to many malignant tumors, such as pancreatic cancer, CRC, bladder cancer, breast cancer, GC, prostate cancer, and is significantly expressed in these malignant tumors. Additionally, S100A4 is capable of modulating apoptosis and proliferation, promoting the EMT process of tumor cells, which also has a close association with invasion, lymph node involvement, and distant metastasis of various tumors (including NSCLC).^{25–34} In this study, we constructed the corresponding relationship of linc01833/miR-519e-3p/S100A4, shedding new light on the mechanism of LUAD progression.

Herein, we revealed the high expression of linc01833 in LUAD tissue and for the first time demonstrated that OE of linc01833 could significantly promote proliferation and invasion in LUAD cells. WB and qRT-PCR analysis were used to detect EMT-related factors, showing that OE of linc01833 also significantly enhanced the EMT process of LC cells. In addition, linc01833 OE can significantly inhibit miR-519e-3p expression. Combining the outcomes from DLR assay, linc01833 was likely to inhibit the expression of miR-519e-3p by sponge adsorption of ceRNA. We also verified that miR-519e-3p OE can inhibit proliferation and the EMT process of LC cells, and can also inhibit S100A4 expression. However, the reverse assays suggest that miR-519e-3p can partially reverse the biological effects of OE of linc01833. OE of S100A4 can significantly enhance proliferation and invasion capacity of LUAD cells and promote their EMT process, which is consistent with the existing literature on the biological role of S100A4.^{30,34} It can also reverse the combined effect of both OE of linc01833 and miR-519e-3p on LC cells.

Conclusion

In conclusion, we demonstrated that lncRNA linc01833 could attenuate its negative regulation of S100A4 and promote the infiltration and metastasis of LUAD through sponge adsorption of miR-519e-3p. Meanwhile, miR-519e-3p can inhibit the progression of LUAD through targeted inhibition of S100A4. Taken together, our study has supplemented understanding of the network mechanism of LUAD. However, further investigations are required on the interaction of linc01833/miR-519e-3p/S100A4 axis in the biological network of LC.

Data Sharing Statement

The datasets used and analyzed during the current study are available from the corresponding author on reasonable request.

Author Contributions

All authors made substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data; took part in drafting the manuscript or revising it critically for important intellectual content; agreed manuscript submission to the Journal; gave final approval of the version to be published; and agreed to be accountable for all aspects of the work. Yuan Zhang and Wenhua Li should be considered joint first author. Yuxia Fan, PhD

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Disclosure

The authors declare no conflicts of interest or financial ties to disclose.

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