Long noncoding RNA ILF3-AS1 promotes cell proliferation, migration, and invasion via negatively regulating miR-200b/a/429 in melanoma

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Introduction

Melanoma is the most malignant skin cancer, which account for most of skin-cancer-related deaths. Long noncoding RNA (lncRNA) is a class of noncoding RNAs with crucial roles in many cancers. However, the roles of lncRNAs in melanoma have not been well studied. In the present study, using public available data and clinical tissues samples, we found that lncRNA ILF3-AS1 is up-regulated in melanoma tissues and cell lines, and correlated with poor prognosis of melanoma patients. Functional experiments showed that knockdown of ILF3-AS1 inhibits melanoma cell proliferation, migration, and invasion. Mechanically, we found that ILF3-AS1 interacts with EZH2, promotes the binding of EZH2 to the miR-200b/a/429 promoter, and represses miR-200b/a/429 expression. The expression of ILF3-AS1 is negatively correlated with that of miR-200b/a/429 in melanoma tissues. Moreover, inhibition of miR-200b/a/429 abrogates the biological roles of ILF3-AS1 knockdown on melanoma cell proliferation, migration, and invasion. In conclusion, these results demonstrate that melanoma-upregulated lncRNA ILF3-AS1 promotes cell proliferation, migration, and invasion via negatively regulating miR-200b/a/429, and imply that ILF3-AS1 may be a potential prognostic biomarker and therapeutic target for melanoma.
dysregulated in cancers [22-25]. The miR-200 family (miR-200s) is well known to function as tumor suppressors via inhibiting tumorigenesis and progression in many cancers [26,27]. miR-200s contains two clusters, with miR-200b, miR-200a, and miR-429 as the first cluster, and miR-200c and miR-141 as the second cluster. miR-200s are frequently dysregulated in many cancers, including melanoma [28,29]. But the factors contributing to the dysregulation of miR-200s in melanoma are still unknown.

In the present study, using a previously reported RNA sequencing results of melanoma [30], we found that IncRNA ILF3-AS1 (XLOC_013222, Refseq NR_024333.1) is up-regulated in melanoma. We further detected ILF3-AS1 expression in public available database and our own cohort of melanomas. Moreover, the biological roles and action mechanisms of ILF3-AS1 in melanoma are also investigated.

Materials and methods

Tissues samples

A total of 37 benign nevi, 60 primary melanomas, 25 metastatic melanomas, 13 breast tissues, 12 breast cancer tissues, 8 metastatic breast cancer tissues, 11 lung tissues, 9 non-small cell lung cancer (NSCLC) tissues, and 6 metastatic NSCLC tissues were obtained from patients who underwent surgery in the 253rd Hospital of PLA (Hohhot, Inner Mongolia, China) with signed informed consent. All fresh tissues samples were immediately frozen in liquid nitrogen and stored at −80°C until use. Pathological diagnosis of all tissues samples was confirmed by two independent pathologists. The Review Board of the 253rd Hospital of PLA reviewed and approved the present study.

Cell culture and transfection

The human epidermal melanocyte HEMa-LP was obtained from Invitrogen (Carlsbad, U.S.A.) and cultured in Medium 254 and Human Melanocyte Growth Supplement-2 (Invitrogen). The human melanoma cell lines SK-MEL-2, SK-MEL-28, and A375 were obtained from American Type Culture Collection (ATCC). SK-MEL-2 and SK-MEL-28 cells were cultured in Eagle’s Minimum Essential Medium. A375 cells were cultured in Dulbecco’s Modified Eagle’s Medium. All cells were maintained in medium supplemented with 10% fetal bovine serum (Invitrogen) in an atmosphere containing 5% CO₂ at 37°C.

miR-200a, miR-200b, and miR-429 inhibitors were obtained from GenePharma (Shanghai, China). All plasmids and miRNAs inhibitors were transfected using Lipofectamine 3000 (Invitrogen) following the manufacturer’s protocol.

RNA extraction and quantitative real-time polymerase chain reaction (qRT-PCR)

Total RNA was isolated from tissues samples and cells using TRIzol (Invitrogen) following the manufacturer’s protocol. The RNA was treated with DNase I to remove genomic DNA. Reverse transcription was performed using the M-MLV Reverse Transcriptase (Invitrogen) following the manufacturer’s protocol. Quantitative real-time polymerase chain reaction (qRT-PCR) was performed on ABI StepOnePlus system (Applied Biosystems, Foster City, CA, U.S.A.). The expression of ILF3-AS1 was measured using SYBR® Premix Ex Taq™II (Takara, Dalian, China) and normalized to GAPDH. The expression of miRNAs was measured using TaqMan MicroRNA Assays (Applied Biosystems) following the manufacturer’s protocol. The primer sequences used were as follows: 5’-TAAAGCCCATGTCCTTCC-3’ (forward) and 5’-TTCTGTGCTTTCTGTGCTC-3’ (reverse) for ILF3-AS1; 5’-CTCTTTGTGCCCTTTCTT-3’ (forward) and 5’-ATGGCTTTCGATCTCCTAT-3’ (reverse) for HEIH; 5’-GGAGGCGAGTCCTCAAAAT-3’ (forward) and 5’-GGCTGTGATCATCTCCGTAGG-3’ (reverse) for GAPDH. The relative expression of RNAs was calculated using the comparative C_t method.

Vectors and stable cell lines construction

Two independent cDNA oligonucleotides specifically targeting ILF3-AS1 (sh-ILF3-AS1-1 and sh-ILF3-AS1-2) were synthesized by GenePharma (Shanghai, China) and inserted into the shRNA expression vector pGPH1/Neo. The shRNAs target sites were as follows: for sh-ILF3-AS1-1: GCCTGTGATTCCAGACCTTC; for sh-ILF3-AS1-2: GCCTGTGATCCACAACGGTG. The shRNAs were transfected into A375 and SK-MEL-2 cells, and selected with neomycin (1000 µg/ml) for 4 weeks.

ILF3-AS1 full-length transcript was PCR amplified with the Phusion Flash High-Fidelity PCR Master Mix (Thermo Fisher, Waltham, MA, U.S.A.) and subcloned into the Hind III and BamH I sites of pcDNA3.1 (Invitrogen) or pSPT19 (Roche, Mannheim, Germany), named pcDNA3.1-ILF3-AS1 or pSPT19-ILF3-AS1 respectively. The primers used were as follows: 5’-CCCAAAGCCTTATTCTACCGCGCTGG-3’ (forward)
and 5'-CGGGATCCGACACGGAAACAGGAGGATTTA-3' (reverse). IncRNA HEIH overexpression vector pcDNA3.1-HEIH was constructed as previously described [31].

Cell proliferation assays
Glo cell viability assays and ethynyl deoxyuridine (EdU) incorporation assays were performed to assess cell proliferation. For Glo cell viability assays, 3000 indicated melanoma cells were seeded in 96-well plates. The luminescence at each time point was acquired with the Cell Titer-Glo® Luminescent Cell Viability Assay (Promega, Madison, WI, U.S.A.) according to the manufacturer's protocol. Cell viability was assessed using the proliferation curves plotted with the luminescence values. EdU incorporation assays were carried out using the EdU kit (Roche) according to the manufacturer's protocol. The results were acquired and quantified using the Zeiss photomicroscope (Carl Zeiss, Oberkochen, Germany) and Image-Pro plus 6.0 software.

Transwell migration and invasion assays
Transwell assays were carried out to assess cell migration and invasion capability. In brief, indicated melanoma cells were suspended in serum-free medium with 1 μg/ml Mitomycin C to inhibit cell proliferation, and seeded in the upper chamber of a 24-well transwell insert (8 μm pore size, Millipore, Bedford, MA, U.S.A.). For invasion assays, Coating Matrigel (BD Biosciences, San Jose, CA, U.S.A.) was plated in the upper chamber before the seeding of cells. The lower chamber was filled with complete medium. After being incubated for 24 h, the cells at the upper chamber were removed using cotton swabs, and the migratory or invasive cells at the bottom of the inserts were fixed by methanol, stained using Crystal Violet, photographed by Zeiss photomicroscope, and quantified by counting in five random fields.

RNA pull-down assays
ILF3-AS1 and its antisense RNA were in vitro transcribed and biotin-labeled from vector pSPT19-ILF3-AS1 using the Biotin RNA Labeling Mix (Roche) and T7 or SP6 RNA polymerase (Roche) respectively. Then 50 pmol of biotin-labeled RNA was mixed with 1 μg of protein extracts from A375 cells for 1 h at 4°C, followed by being incubated with Dynabeads Myone Streptavidin T1 (Invitrogen) for additional 1 h. The proteins binding to the Dynabeads were resolved in SDS buffer and separated by sodium dodecyl sulfate/polyacrylamide gel electrophoresis, followed by being transferred to nitrocellulose membrane. Then the membranes were incubated with antibodies for EZH2 (Millipore) or GAPDH (Cell Signaling Technology, Boston, MA, U.S.A.). After being washed, the membranes were incubated with fluorescence-labeled secondary antibodies, and detected using an Odyssey infrared scanner (LI-Cor, Lincoln, NE, U.S.A.).

RNA immunoprecipitation (RIP) assays
RNA immunoprecipitation (RIP) assays were performed using the EZ-Magna RIP™ RNA Binding Protein Immunoprecipitation Kit (Millipore) and EZH2 antibody (Millipore) following the manufacturer's protocol. Retrieved RNA was reverse transcribed and measured by qRT-PCR as above described.

Chromatin immunoprecipitation (ChIP) assays
Chromatin immunoprecipitation (ChIP) assays were performed using the EZ-Magna ChIP™ A/G Chromatin Immunoprecipitation Kit (Millipore) and EZH2 antibody (Millipore) or H3K27me3 antibody (Millipore) following the manufacturer's protocol. Retrieved DNA was measured using SYBR® Premix Ex Taq™ II (Takara) on ABI StepOnePlus system (Applied Biosystems). The primer sequences were as follows: 5'-CTCGCGTCACCGCTCAGTTG-3' (forward) and 5'-ACAATCCTGGTGCTCTGTG-3' (reverse) for miR-200b/a/429 promoter; 5'-GCTGGCCTGACTTTAC-3' (forward) and 5'-GAGTGGTGGTTGCGGGA-3' (reverse) for β-actin promoter.

Statistical analysis
Statistical analyses were implemented using the SPSS 18.0 software. Differences among the groups were estimated by Mann–Whitney U test, Log-rank test, Student’s t test, or Pearson's correlation analysis as indicated. P<0.05 was considered as statistically significant.
ILF3-AS1 is up-regulated in melanoma tissues and cell lines, and indicates poor prognosis of melanoma patients

To investigate the expression of ILF3-AS1 in melanoma, we first searched the MiTranscriptome database (www.mitranscriptome.org) [8], and found that ILF3-AS1 is increased in primary melanomas and metastatic melanomas compared with normal human melanocytes (NHMEs) (Figure 1A). Furthermore, we collected 37 benign nevi, 60 primary melanomas, and 25 metastatic melanomas, and measured ILF3-AS1 expression by qRT-PCR. Our results
showed that ILF3-AS1 is also significantly up-regulated in primary melanomas compared with benign nevi, and is further up-regulated in metastatic melanomas (Figure 1B). Analyzing the correlation between ILF3-AS1 expression and clinicopathological characteristics of these 60 primary melanomas, we noted that ILF3-AS1 is higher in melanomas at least 1 mm thick, which is more likely to metastasize and has more severity (T2/T3/T4 versus T1) (Figure 1C). To analyze the correlation between ILF3-AS1 expression and melanoma patients’ prognosis, we performed Kaplan–Meier survival analysis. As shown in Figure 1D, patients with higher ILF3-AS1 expression had a worse overall survival than those with lower ILF3-AS1 expression. These results showed that ILF3-AS1 is up-regulated in melanoma tissues and high ILF3-AS1 expression is correlated with metastatic characteristic and poor prognosis. Then we measured the expression of ILF3-AS1 in human epidermal melanocyte (HEMa-LP) and melanoma cell lines (SK-MEL-2, SK-MEL-28, and A375) by qRT-PCR. The results showed that ILF3-AS1 is significantly up-regulated in melanoma cell lines compared with epidermal melanocyte (Figure 1E).

To investigate the expression pattern of ILF3-AS1 in other tumors, we also collected 13 breast tissues, 12 breast cancer tissues, 8 metastatic breast cancer tissues, 11 lung tissues, 9 NSCLC tissues, and 6 metastatic NSCLC tissues. The results showed that ILF3-AS1 is also significantly up-regulated in breast cancer tissues and NSCLC tissues, compared with breast and lung tissues respectively (Figure 1F and G). Furthermore, ILF3-AS1 is also further up-regulated in metastatic breast cancer and NSCLC tissues (Figure 1F and G).

Knockdown of ILF3-AS1 inhibits melanoma cell proliferation

To explore the biological function of ILF3-AS1 in melanoma cells, we stably depleted ILF3-AS1 in A375 cells using two independent ILF3-AS1 specific shRNAs (Figure 2A). Glo cell viability assays revealed that knockdown of ILF3-AS1 by both shRNAs significantly inhibits A375 cell proliferation (Figure 2B). EdU incorporation assays also revealed that knockdown of ILF3-AS1 in A375 cells significantly decreases percentage of EdU positive cells (Figure 2C). To further confirm the effects of ILF3-AS1 on melanoma cell proliferation, we stably depleted ILF3-AS1 in SK-MEL-2 cells using two independent ILF3-AS1 specific shRNAs (Figure 2D). Glo cell viability assays revealed that knockdown of ILF3-AS1 by both shRNAs significantly inhibits SK-MEL-2 cell proliferation (Figure 2E). EdU incorporation assays also revealed that knockdown of ILF3-AS1 in SK-MEL-2 cells significantly decreases percentage of EdU positive cells (Figure 2F). These results demonstrated that knockdown of ILF3-AS1 inhibits melanoma cell proliferation.

Knockdown of ILF3-AS1 inhibits melanoma cell migration and invasion

To further explore the effects of ILF3-AS1 on melanoma cell migration and invasion, we performed transwell migration and invasion assays. The results revealed that the number of migrated A375 cells in ILF3-AS1 knockdown group is much fewer than that in the control group (Figure 3A). The number of invasive A375 cells in ILF3-AS1 knockdown group is also much fewer than that in the control group (Figure 3B). The effects of ILF3-AS1 on melanoma cell migration and invasion were further confirmed on SK-MEL-2 cells (Figure 3C and D). These results demonstrated that knockdown of ILF3-AS1 inhibits melanoma cell migration and invasion.

ILF3-AS1 negatively regulates the expression of miR-200b/a/429 via binding to EZH2

Increasing evidence have revealed that many IncRNAs bind and recruit EZH2 to target genes [32,33]. EZH2 is an important subunit of polycomb repressive complex 2 (PRC2), which increases H3K27me3 levels across the promoters of target genes and inhibits the expression of target genes [34]. To test whether ILF3-AS1 also functions in such a manner, we performed RNA pull-down assays using in vitro transcribed biotin-labeled ILF3-AS1. The results showed that ILF3-AS1 specifically binds to EZH2, but not GAPDH protein (Figure 4A). To further confirm the interaction between ILF3-AS1 and EZH2, RIP assays with EZH2 specific antibody were performed. As shown in Figure 4B, a significant enrichment of ILF3-AS1, but not GAPDH mRNA with EZH2 antibody was observed. IncRNA HEIH was used as positive control, which has been reported to associate with EZH2 [35]. miR-200a/429 is a well-known EZH2 target gene [36], which also has crucial roles in the proliferation, migration, and invasion of melanoma cells. To investigate whether ILF3-AS1 is involved in the transcriptional regulation of miR-200b/a/429 via interacting with EZH2, we performed ChIP assays using EZH2 and H3K27me3 specific antibodies in ILF3-AS1 stably depleted and control A375 cells. As shown in Figure 4C, knockdown of ILF3-AS1 decreased the binding of EZH2 and H3K27me3 levels across the miR-200b/a/429 promoter, but not the β-actin promoter. Furthermore, knockdown of ILF3-AS1 up-regulates miR-200b/a/429 expression in both A375 cells and SK-MEL-2 cells (Figure 4D and E). Collectively, these results demonstrated that ILF3-AS1 binds to EZH2, recruits EZH2 to the miR-200b/a/429 promoter, up-regulates H3K27me3 levels across the promoter, and represses the expression of miR-200b/a/429.
Figure 2. Knockdown of ILF3-AS1 inhibits melanoma cell proliferation

(A) The expression of ILF3-AS1 in ILF3-AS1 stably depleted and control A375 cells. (B) Glo cell viability assays in ILF3-AS1 stably depleted and control A375 cells. (C) EdU incorporation assays in ILF3-AS1 stably depleted and control A375 cells. The blue color indicates the nuclei, and the red color indicates EdU-positive nuclei; scale bar = 100 μm. (D) The expression of ILF3-AS1 in ILF3-AS1 stably depleted and control SK-MEL-2 cells. (E) Glo cell viability assays in ILF3-AS1 stably depleted and control SK-MEL-2 cells. (F) EdU incorporation assays in ILF3-AS1 stably depleted and control SK-MEL-2 cells. The blue color indicates the nuclei, and the red color indicates EdU-positive nuclei; scale bar = 100 μm. For all panels, data are represented as mean ± SD; **P<0.01, ***P<0.001 by Student’s t test.

The expression of ILF3-AS1 is negatively correlated with that of miR-200b/a/429 in melanoma tissues

To explore whether the regulation of miR-200b/a/429 by ILF3-AS1 also exists in vivo, we measured the correlation between the expression of ILF3-AS1 and miR-200b/a/429 in melanoma tissues. The results showed that the expression of ILF3-AS1 is negatively correlated with that of miR-200b, miR-200a, and miR-429 in melanoma tissues (Figure 5A–C), supporting the regulation of miR-200b/a/429 by ILF3-AS1 in vivo.

Inhibition of miR-200b/a/429 abrogates the repressing effects of proliferation, migration, and invasion caused by ILF3-AS1 knockdown

To explore whether the regulation of miR-200b/a/429 mediates the biological effects of ILF3-AS1 on melanoma cell proliferation, migration, and invasion, we inhibited miR-200b/a/429 expression by transfecting miR-200b/a/429 inhibitors in ILF3-AS1 stably depleted A375 cells (Figure 6A). Glo cell viability assays and EdU incorporation assays revealed that inhibition of miR-200b/a/429 abrogates the proliferation repression caused by ILF3-AS1 knockdown (Figure 6B and C). Furthermore, transwell migration and invasion assays revealed that inhibition of miR-200b/a/429 abrogates the migration and invasion repression caused by ILF3-AS1 knockdown (Figure 6D and E). These results demonstrated that the repressing effects of proliferation, migration, and invasion caused by ILF3-AS1 knockdown are dependent on miR-200b/a/429.
Enhanced expression of ILF3-AS1 promotes melanoma cell proliferation and invasion

A previous report showed that IncRNA HEIH also has oncogenic roles in melanoma via repressing miR-200b/a/429, which is consistent with the action mechanisms of ILF3-AS1 in melanoma. Therefore, we next investigated whether the oncogenic role of ILF3-AS1 is redundant with that of HEIH in melanoma. Glo cell viability assays revealed that enhanced expression of ILF3-AS1 significantly promotes A375 cell proliferation, and concurrent overexpression of ILF3-AS1 and HEIH slightly further promotes A375 cell proliferation (Figure 7A). Transwell invasion assays revealed that enhanced expression of ILF3-AS1 significantly promotes A375 cell invasion, and concurrent overexpression of ILF3-AS1 and HEIH slightly further promotes A375 cell invasion (Figure 7B). These results demonstrated that overexpression of ILF3-AS1 promotes melanoma cell proliferation and invasion, which is at least partially redundant with the roles of HEIH in melanoma.

Discussion

As one of the deadliest skin cancers, the incidence of melanoma is increasing faster than most of other solid cancers worldwide [37]. The tumorigenesis and progression of melanoma involve complex changes in many genes networks [38]. Among the critical molecules involved in the pathogenesis of melanoma, lncRNAs gradually attract human’s attentions for their important roles in melanoma [39]. For example, IncRNA SLNCR1 associates with poor melanoma survival and increases melanoma invasion via transcriptionally activating MMP9 [30]. SAMMSON functions as an oncogene in melanoma and silencing SAMMSON delivers effective antimelanoma therapeutic responses [12]. MALAT1 promotes melanoma growth and metastasis via sponging miR-22 [40]. BANCR promotes melanoma...
Figure 4. ILF3-AS1 negatively regulates miR-200b/a/429 expression via interacting with EZH2

(A) RNA pull-down assays followed by Western blot analysis revealed the specific enrichment of EZH2, but not GAPDH protein with \textit{in vitro} transcribed biotin-labeled ILF3-AS1 compared with antisense RNA (negative control). (B) RIP assays followed by qRT-PCR revealed the specific enrichment of ILF3-AS1, but not GAPDH mRNA with EZH2 antibody compared with nonspecific IgG (negative control). HEIH was used as positive control. (C) The specific binding of EZH2 and H3K27me3 levels across the \textit{miR-200b/a/429} promoter and the \textit{\(\beta\)-actin} promoter in ILF3-AS1 stably depleted and control A375 cells were measured by ChIP assays followed by qPCR. (D) The expression of miR-200b, miR-200a, and miR-429 in ILF3-AS1 stably depleted and control A375 cells was measured by qRT-PCR. (E) The expression of miR-200b, miR-200a, and miR-429 in ILF3-AS1 stably depleted and control SK-MEL-2 cells was measured by qRT-PCR. Data are represented as mean ± SD; *\(P<0.05\), **\(P<0.01\), ***\(P<0.001\) by Student's \(t\) test.
The correlation between ILF3-AS1 expression level and miR-200b (A), miR-200a (B), and miR-429 (C) expression levels in 60 melanoma tissues. P values were acquired by Pearson's correlation analysis.

Inhibition of miR-200b/a/429 abrogates the inhibitory effects of ILF3-AS1 knockdown on melanoma cell proliferation, migration, and invasion

The expression of miR-200b, miR-200a, and miR-429 in ILF3-AS1 stably depleted and control A375 cells transfected with miR-200b/a/429 inhibitors or control was measured by qRT-PCR. (B) Glo cell viability assays in ILF3-AS1 stably depleted and control A375 cells transfected with miR-200b/a/429 inhibitors or control. (C) EdU incorporation assays in ILF3-AS1 stably depleted and control A375 cells transfected with miR-200b/a/429 inhibitors or control. The blue color indicates the nuclei, and the red color indicates EdU-positive nuclei; scale bar = 100 μm. (D) Transwell migration assays in ILF3-AS1 stably depleted and control A375 cells transfected with miR-200b/a/429 inhibitors or control. Representative images are shown; scale bar = 100 μm. (E) Transwell invasion assays in ILF3-AS1 stably depleted and control A375 cells transfected with miR-200b/a/429 inhibitors or control. Representative images are shown; scale bar = 100 μm. For all panels, data are represented as mean ± SD; **P<0.01, ***P<0.001 by Student's t test.

proliferation via activating MAPK pathway [41]. Also in our previous study, we found that MHENCN promotes melanoma growth and metastasis via activating PI3K–Akt pathway [42]. In our another previous report, we found that PVT1 is up-regulated in melanoma tissues and serum of melanoma patients. Functional experiments demonstrated that PVT1 promotes melanoma cell proliferation and migration. These evidence further confirm the critical
roles of IncRNAs in melanoma. The amount of IncRNAs is much larger than that of protein-coding mRNAs [8], and many genes are involved in melanoma [43]. Thus, it is important to further reveal more functional IncRNAs in melanoma.

Searching the differently expressed IncRNAs in a previously reported RNA sequencing results of melanoma [30], we noted a novel IncRNA ILF3-AS1, which localizes at chromosome 19p13.2 and is up-regulated in melanoma tissues. The MiTranscriptome database also demonstrates that ILF3-AS1 is up-regulated in melanoma. However, the functions of ILF3-AS1 in cancers are unknown. In the present study, we further examined the expression, clinical significances, and biological roles of ILF3-AS1 in melanoma. We found that ILF3-AS1 is up-regulated in melanoma tissues and cell lines, and associates with metastatic characteristic and poor prognosis of melanoma patients. Functional assays demonstrated that silencing of ILF3-AS1 using two independent shRNAs both significantly inhibit melanoma cell proliferation, migration, and invasion. Overexpression of ILF3-AS1 promotes melanoma cell proliferation and invasion. These data reveal that as a previously unknown IncRNA, ILF3-AS1 is dysregulated and also has critical functions in melanoma. Our study enlarges our recognitions of IncRNAs and provides a novel prognostic biomarker and therapeutic target for melanoma.

miR-200s is known to have complex roles in many cancer processes, including proliferation, migration, invasion, drug resistance, epithelial–mesenchymal transition, and et al. [44–46]. miR-200s has also been reported to be down-regulated in invasive melanoma [28]. But the factors contributing to miR-200s dysregulation in melanoma are still unknown. In the present study, we found that ILF3-AS1 interacts with EZH2. EZH2 is a histone methyltransferase, catalyzes H3K27 trimethylation, and inhibits target genes expression. In the present study, we further found that through interacting with EZH2, ILF3-AS1 recruits EZH2 to the miR-200b/a/429 promoter, up-regulates H3K27me3 levels across the miR-200b/a/429 promoter, and inhibits miR-200b/a/429 expression. The expression of ILF3-AS1 is negatively correlated with that of miR-200b/a/429 in melanoma tissues. Silencing miR-200b/a/429 abrogates the repressive effects of ILF3-AS1 knockdown on melanoma cell proliferation, migration, and invasion. These data demonstrate that ILF3-AS1 epigenetically silences miR-200b/a/429 expression and the biological roles of ILF3-AS1 in melanoma are dependent on miR-200b/a/429.

In conclusion, our results identify a previously unknown IncRNA ILF3-AS1, which is up-regulated in melanoma, associates with poor prognosis of melanoma patients, and promotes melanoma cell proliferation, migration, and invasion. Mechanistically, ILF3-AS1 interacts with EZH2 and epigenetically silences miR-200b/a/429. Our data imply that ILF3-AS1 would be a potential prognostic biomarker and therapeutic target for melanoma.

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Author Contribution
X.C. and S.L. conceived and designed the study. S.L., X.Z., X.M., G.G., L.Y., D.Y., H.D., and W.S. performed the experiments. X.C. and S.L. analyzed the data and wrote the manuscript. All authors read and approved the manuscript.

Competing Interests
The authors declare that there are no competing interests associated with the manuscript.

Abbreviations
ChIP, chromatin immunoprecipitation; Edu, ethynyl deoxyuridine; ILF3-AS1, ILF3 antisense transcript 1; IncRNA, long non-coding RNA; NHME, normal human melanocyte; NSCLC, non-small cell lung cancer; PRC2, polycomb repressive complex 2; qRT-PCR, quantitative real-time polymerase chain reaction; RIP, RNA immunoprecipitation.

References
1 Rogers, H.W., Weinstock, M.A., Feldman, S.R. and Coldiron, B.M. (2015) Incidence estimate of nonmelanoma skin cancer (keratinocyte carcinomas) in the U.S. population, 2012. JAMA Dermatol. 151, 1081–1086
2 Siegel, R.L., Miller, K.D. and Jemal, A. (2015) Cancer statistics, 2015. CA-Cancer J. Clin. 65, 5–29
3 Little, E.G. and Eide, M.J. (2012) Update on the current state of melanoma incidence. Dermatol. Clin. 30, 355–361
4 Nikolau, V.A., Stratigos, A.J., Flaherty, K.T. and Tsao, H. (2012) Melanoma: new insights and new therapies. J. Invest. Dermatol. 132, 854–863
5 (2015) Melanoma research gathers momentum. Lancet 385, 2323
6 George, B.P., Abrahamse, H. and Hemmaragala, N.M. (2016) Caspase dependent apoptotic inhibition of melanoma and lung cancer cells by tropical Rubus extracts. Biomed. Pharmacother. 80, 193–199
7 Mustapha, N., Szczepanik, L.M., Ghedira, K., Hennebelle, T. and Chekir-Ghedira, L. (2015) Compounds isolated from the aerial part of Crataegus azarolus inhibit growth of B16F10 melanoma cells and exert a potent inhibition of the melanin synthesis. Biomed. Pharmacother. 69, 139–144
8 Iyer, M.K., Niknafs, Y.S., Malik, R., Singhal, U., Sahu, A., Hosono, Y. et al. (2015) The landscape of long noncoding RNAs in the human transcriptome. Nat. Genet. 47, 199–208
9 Muers, M. (2011) RNA: genome-wide views of long non-coding RNAs. Nat. Rev. Genet. 12, 742
10 Ponting, C.P., Oliver, P.L. and Reik, W. (2009) Evolution and functions of long noncoding RNAs. Cell 136, 629–641
11 Fatica, A. and Bozzoni, I. (2014) Long non-coding RNAs: new players in cell differentiation and development. Nat. Rev. Genet. 15, 7–21
12 Leucci, E., Vendramini, R., Spina, M., Laurent, F., Piers, M., Wouters, J. et al. (2016) Melanoma addiction to the long non-coding RNA SAMMSON. Nature 531, 518–522
13 Yuan, J.H., Yang, F., Wang, F., Ma, J.Z., Guo, Y.J., Tao, Q.F. et al. (2014) A long noncoding RNA activated by TGF-beta promotes the invasion-metastasis cascade in hepatocellular carcinoma. Cancer Cell 25, 666–681
14 Li, C., Wang, S., Xing, Z., Lin, A., Liang, K., Song, J. et al. (2017) A ROR1-HER3-IncRNA signalling axis modulates the Hippo-YAP pathway to regulate bone metastasis. Nat. Cell Biol. 19, 106–119
15 Liu, X., Xiao, Z.D., Han, L., Zhang, J., Lee, S.W., Wang, W. et al. (2016) LncRNA NBR2 engages a metabolic checkpoint by regulating AMPK under energy stress. Nat. Cell Biol. 18, 431–442
16 Ronnau, C.G., Verhaegh, G.W., Luna-Velez, M.V. and Schalken, J.A. (2014) Noncoding RNAs as novel biomarkers in prostate cancer. BioMed Res. Int. 2014, 591703
17 Zhu, X.T., Yuan, J.H., Zhu, T.T., Li, Y.Y. and Cheng, X.Y. (2016) Long noncoding RNA glypican 3 (GPC3) antisense transcript 1 promotes hepatocellular carcinoma progression via epigenetically activating GPC3. FEBS J. 283, 3739–3754
18 Lin, A., Li, C., Xing, Z., Hu, Q., Liang, K., Han, L. et al. (2016) The LINK-A lncRNA activates normoxic HIF1alpha signalling in triple-negative breast cancer. Nat. Cell Biol. 18, 213–224
19 Liu, C. and Lin, J. (2016) Long noncoding RNA ZEB1-AS1 acts as an oncogene in osteosarcoma by epigenetically activating ZEB1. Am. J. Transl. Res. 8, 4095–4105
20 Leucci, E., Coe, E.A., Marine, J.C. and Vance, K.W. (2016) The emerging role of long non-coding RNAs in cutaneous melanoma. Pigm. Cell Melanoma Res. 29, 619–626
21 Bartel, D.P. (2004) MicroRNAs: genomics, biogenesis, mechanism, and function. Cell 116, 281–297
22 Yuan, J.H., Yang, F., Chen, B.F., Lu, Z., Zhu, W.P. et al. (2011) The histone deacetylase 4 (HDAC4) family regulates the expression of microRNA-200a. PLoS Genet. 7, e1002109
23 Yoon, A.J., Wang, S., Chen, J., Robine, N., Philipeau, E., Oster, M.W. et al. (2014) Prognostic value of miR-375 and miR-214-3p in early stage oral squamous cell carcinoma. Am. J. Transl. Res. 6, 580–599
24 Bhattacharjee, S., Renganaath, K., Mehrotra, R. and Mehrotra, S. (2013) Combinatorial control of gene expression. BioMed Res. Int. 2013, 407263
25 Long, J.D., Sullivan, T.B., Humphrey, J., Logvinenko, T., Summerhayes, K.A., Kozinn, S. et al. (2015) A non-invasive miRNA based assay to detect bladder cancer in cell-free urine. Am. J. Transl. Res. 7, 2500–2509
26 Wellner, U., Schubert, J., Burk, U.C., Schmalhofer, O., Zhu, F., Sonntag, A. et al. (2009) The EMT-activator ZEB1 promotes tumorigenicity by repressing stromness-inhibiting microRNAs. Nat. Cell Biol. 11, 1487–1495
27 Gregory, P.A., Bert, A.G., Paterson, E.L., Barry, S.C., Tsykin, A., Farshid, G. et al. (2008) The miR-200 family and miR-205 regulate epithelial to mesenchymal transition by targeting ZEB1 and SIP1. Nat. Cell Biol. 10, 593–601

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van Kempen, L.C., van den Hurk, K., Lazar, V., Michiels, S., Winnepenninckx, V., Stas, M. et al. (2012) Loss of microRNA-200a and c, and microRNA-203 expression at the invasive front of primary cutaneous melanoma is associated with increased thickness and disease progression. *Virchows Arch.* **461**, 441–448

Zhang, H., Sun, Z., Li, Y., Fan, D. and Jiang, H. (2017) MicroRNA-200c binding to FN1 suppresses the proliferation, migration and invasion of gastric cancer cells. *Biomed. Pharmacother.* **88**, 285–292

Schmidt, K., Joyce, C.E., Buquicchio, F., Brown, A., Ritz, J., Distel, R.J. et al. (2016) The IncRNA SLNCR1 mediates melanoma invasion through a conserved SRA1-like region. *Cell Rep.* **15**, 2025–2037

Zhao, H., Xing, G., Wang, Y., Luo, Z., Liu, G. and Meng, H. (2017) Long noncoding RNA HEIH promotes melanoma cell proliferation, migration and invasion via inhibition of miR-200b/a/429. *Biosci. Rep.* **37**(3), doi:10.1042/bsr20170682

Wan, L., Sun, M., Liu, G.J., Wei, C., Zhang, E.B., Kong, R. et al. (2016) Long noncoding RNA PVT1 promotes non-small cell lung cancer cell proliferation through epigenetically regulating LATS2 expression. *Mol. Cancer Ther.* **15**, 1082–1094

Gupta, R.A., Shah, N., Wang, K.C., Kim, J., Horlings, H.M., Wong, D.J. et al. (2010) Long non-coding RNA HOTAIR reprograms chromatin state to promote cancer metastasis. *Nature* **464**, 1071–1076

Rossi, U.A., Gil-Cardeza, M.L., Villaverde, M.S., Finocchiaro, L.M. and Glikin, G.C. (2015) Interferon-beta gene transfer induces a strong cytotoxic bystander effect on melanoma cells. *Biomed. Pharmacother.* **72**, 44–51

Chen, X., Guo, W., Xu, X.J., Su, F., Wang, Y., Zhang, Y. et al. (2016) Melanoma long non-coding RNA signature predicts prognostic survival and directs clinical risk-specific treatments. *J. Dermatol. Sci.* **85**, 226–234

Luan, W., Li, L., Shi, Y., Bu, X., Xia, Y., Wang, J. et al. (2016) Long non-coding RNA MALAT1 acts as a competing endogenous RNA to promote malignant melanoma growth and metastasis by sponging miR-22. *Oncotarget* **7**, 63901–63912

Li, R., Zhang, L., Jia, L., Duan, Y., Li, Y., Bao, L. et al. (2014) Long non-coding RNA BANCR promotes proliferation in malignant melanoma by regulating MAPK pathway activation. *PLOS One* **9**, e90083

Chen, X., Dong, H., Liu, S., Yu, L., Yan, D., Yan, X. et al. (2017) Long non-coding RNA MHENCR promotes melanoma progression via regulating miR-425/489-mediated PI3K-Akt pathway. *Am. J. Transl. Res.* **9**, 90–102

Dror, S., Sander, L., Schwartz, H., Steinboim, D., Barzilai, A., Dishon, Y. et al. (2016) Melanoma miRNA trafficking controls tumour primary niche formation. *Nat. Cell Biol.* **18**, 1006–1017

Liu, S., Tetzstaif, M.T., Cui, R. and Xu, X. (2012) miR-200c inhibits melanoma progression and drug resistance through down-regulation of BMI-1. *Am. J. Pathol.* **181**, 1823–1835

Li, Q., Zhang, C., Chen, R., Xiong, H., Qiu, F., Liu, S. et al. (2016) Disrupting MALAT1/miR-200c sponge decreases invasion and migration in endometrioid endometrial carcinoma. *Cancer Lett.* **383**, 28–40

Zhang, L., Yang, F., Yuan, J.H., Yuan, S.X., Zhou, W.P., Huo, X.S. et al. (2013) Epigenetic activation of the MiR-200 family contributes to H19-mediated metastasis suppression in hepatocellular carcinoma. *Carcinogenesis* **34**, 577–586