LETTER TO EDITOR

IncRNA187415.1 silence in BCAMs ameliorated breast cancer progression by blocking C/EBPβ-IncRNA187415.1-CISH axis and reversing pro-tumor characteristic of BCAMs

Dear Editor,

Recently, we demonstrated that IncRNA187415.1 knock-down in breast cancer-associated macrophages (BCAMs) could change their phenotype and function of BCAMs, benefit the BCAMs inducing breast cancer cells apoptosis and inhibit their invasion. IncRNA187415.1 maintained BCAMs function in local microenvironment by inhibiting CISH mRNA degradation; however, transcription factor C/EBPβ regulated transcription of IncRNA187415.1 via binding promoter region of –1500∼–2000 bp of IncRNA187415.1.

As our previous research about IncRNA profiles of BCAMs using IncRNAs microarray showed that IncRNA187415.1 was the most significantly changed gene. The data were further confirmed in BCAMs isolated from breast cancer bearing mice in vitro (Figures 1A and 1B). To further investigate the biological role of IncRNA187415.1, its expression in BCAMs was successfully silenced by siRNAs (Figure 1C). IncRNA187415.1 silence in BCAMs significantly inhibited the proliferation, migration, and invasion of 4T1 cells (Figures 1D and 1E). Conversely, IncRNA187415.1 silence in BCAMs resulted in a significantly increase of the apoptosis in 4T1 cells (Figure 1F). To further confirm the similar effects also exist in vivo, the jetPEI-Man/ si-IncRNA187415.1 complex was used to specifically downregulate the IncRNA187415.1 in BCAMs of breast cancer bearing mice. A dramatic reduction of tumor volumes and tumor weight were observed in si-IncRNA187415.1 group compared to si-NC (Figure 1G and 1H). These results clearly demonstrated that IncRNA187415.1 knockdown target on BCAMs significantly ameliorated breast cancer progression.

To determine the effect of IncRNA187415.1 on the phenotype and function of BCAMs, IncRNA187415.1 was silenced in BCAMs with siRNA. After IncRNA187415.1 silence, CD86 and MHCII were significantly increased, conversely, CD206 was downregulated, LPS-treated macrophages (pro-inflammatory type) and IL-4-treated macrophages (anti-inflammatory type) were used as positive or negative control, respectively (Figure 2A). The qPCR and ELISA also indicated that the expression of Tnfa and Il-1β mRNA as well as their proteins levels were also increased after IncRNA187415.1 silence, and the mRNA and protein expression of Il-4, Il -10 were decreased in si-IncRNA187415.1 group (Figures 2B and 2D). Furthermore, the protein level of Arginase 1 (Arg1) decreased and iNOS increased in si-IncRNA187415.1 group (Figure 3C). Taken together, these results suggested that IncRNA187415.1 silence could reprogram the phenotype of BCAMs.

As the results above showed that IncRNA187415.1 silence in BCAMs could benefit the shift of BCAMs phenotype and function from anti-inflammatory to pro-inflammatory. To further clarify the mechanism of IncRNA187415.1 maintaining BCAMs showing anti-inflammatory phenotype in local microenvironment, downstream target genes related to IncRNA187415.1 in BCAMs were analyzed by RNA microarray. The results indicated that CISH is the potential target gene (Log2FC = 8.07) (Figure S1A and Table S1). Previous data also indicate that CISH is related to the polarization of macrophages.2 To confirm whether IncRNA187415.1 could promote the expression of CISH, firstly, IncRNA187415.1 was silence in BCAMs, IncRNA187415.1 silence significantly reduced the expression of CISH mRNA and protein level (Figures 3A and 3B). To further confirm how IncRNA187415.1 regulated CISH, actinomycin D, an RNA polymerase inhibitor, was employed. After actinomycin D treatment, the half-life of CISH mRNA was significantly shortened in IncRNA187415.1 silence group compared with control (Figures 3C and 3D), which indicated that IncRNA187415.1 could be involved in transcription-related processes of CISH and facilitate the stability of CISH mRNA. To further confirm, the shift of BCAMs phenotype and function caused by CISH downregulation induced by...
**FIGURE 1** LncRNA187415.1 silence in BCAMs ameliorated breast cancer progression. (A) Volcano plot of LncRNAs gene profiles results in BCAMs. The expression of LncRNAs in macrophages and BCAMs was detected by LncRNAs microarray technology. Upregulation (red), downregulation (green), and no-significance (gray). (B) qPCR analysis for LncRNA187415.1 expression in BCAMs from breast tumors (n = 8). (C) siRNA knockdown efficiency. siRNAs (50 nM) were transiently transfected into BCAMs, after 48 h, BCAMs were harvested for qPCR analysis (n = 3). (D) Representative images of 4T1 are stained by Ki67 (green) and DAPI (blue). LncRNA187415.1-downregulated BCAMs were co-cultured with 4T1 for 48 h. The proliferation of 4T1 was evaluated by Ki-67/DAPI staining. Data were statistically analyzed based on three independent experiments. (E) Invasion and migration ability of 4T1 cells co-cultured with LncRNA187415.1 downregulated BCAMs. Data represent means ± SD from three independent experiments. (F) Apoptosis of 4T1 cells was analyzed by Annexin V and PI staining. Note that 4T1 cells were co-cultured with LncRNA187415.1-downregulated BCAMs for 48 h; the apoptosis of 4T1 cells was analyzed by flow cytometry (n = 3). (G) Tumor volume. Note that 4T1 tumor-bearing mice were given jetPEI-Man/si-NC complex or jetPEI-Man/siRNA2 complex. Tumor volume measurement was performed by digital caliper at the indicated days (n = 7). (H) Tumor weight. Tumors were resected at day 24, and tumor weight was measured (n = 7). All data are shown as means ± SD. *p < 0.01, **p < 0.001

Abbreviation: PM, peritoneal macrophage.
LncRNA187415.1 silence, overexpressed plasmid of CISH was transfected into the BCAMs following lncRNA187415.1 silence for 48 h, the results showed that expression of CD86 and MHC II decreased, conversely, CD206 increased (Figure 3E).

To clarify which molecule regulates lncRNA187415.1 in BCAMs, PROMO database (http://alggen.lsi.upc.es/home.html) was used to predict the transcription factors that bind to lncRNA187415.1 promoter region. C/EBPβ was a candidate transcription factor for lncRNA187415.1. To further investigate the regulation of C/EBPβ on lncRNA187415.1, Cebpb was silenced in BCAMs by siRNA (Figure S1B and S1D), the expression of lncRNA187415.1 and Cish was significantly decreased (Figure S1C and S1E),
FIGURE 3 LncRNA187415.1 activated by C/EBPβ maintained the pro-tumor characteristics of BCAMs by facilitating CISH mRNA stability. (A and B) qPCR analyzed Cish mRNA, and protein levels were detected by qPCR and Western blotting. Data are shown as means ± SD. (C and D) BCAMs with siRNA2 or si-NC transfected for 48 h, and then treated by 5 μg/ml actinomycin D for indicated time. The mRNA
stability was evaluated using mRNA decay curves, and protein level was analyzed by Western blotting. Data were analyzed based on three independent experiments. (E) BCAMs with siRNA2 or si-NC co-cultured for 48 h, and then CISH overexpressed plasmid was transfected for the other 48 h. Flow cytometry analysis for surface protein markers. (F) Luciferase assay of 293T cells after co-transfection of lncRNA187415.1 promoter plasmid (Lnc-pro-luc) and C/EBPβ overexpressing plasmid, n = 3. (G) Luciferase assay of 293T cells after co-transfection of lncRNA187415.1 promoter plasmids with different length of promoter and C/EBPβ overexpressing, n = 3. **p < 0.01, ***p < 0.001. pro-all: sequence -2 kb ∼ ±0 bp, including all potential binding sites; pro-1500: sequence -1.5 kb ∼ +0 bp, including five potential binding sites; pro-800: sequence -800 bp ∼ ±0 bp, including three potential binding sites; pro-500: sequence -500 bp ∼ ±0 bp, including one potential binding sites. All data were analyzed based on three independent experiments. **p < 0.01, ***p < 0.001.

which confirmed that C/EBPβ regulated the expression of lncRNA187415.1 as a transcript factor. Then the JASPER database (http://jaspar.genereg.net/) was used to predict eight sites that could bind to the lncRNA187415.1 promoter region as follows (Figure S1F), and sequential truncations of this lncRNA187415.1 luciferase reported were created to identify regions critical for lncRNA187415.1 transcription. Four luciferase reporter plasmids containing pro-all (including all the sites), pro-1500 (containing five sites), pro-800 (containing three sites), and pro-500 (containing one site) were constructed in order to determine the approximate binding region. The result indicated that the fewer binding sites included, the lower the luciferase activity was. Additionally, Only the pro-1500 of the transcription factor was activated and bound to more than 30% (C/EBPβ+Lnc-pro-luc/C/EBPβ EV+Lnc-pro-luc) (Figures 3F and 3G); therefore, the C/EBPβ bound to the lncRNA187415.1 promoter region from −1.5 kb to −2 kb, and ChIP-qPCR experiment was performed to verify the binding sites (Figure SIG).

Our data clearly demonstrated that C/EBPβ-LncRNA187415.1-CISH maintained the pro-tumor characteristics of BCAMs and contributed to breast cancer progression. LncRNA187415.1 silence in BCAMs inhibited breast cancer cells proliferation, migration, and invasion, promoted apoptosis, and ameliorated breast cancer progression. Our results shed a light on immunotherapy of breast cancer targeting BCAMs.

CONFLICT OF INTEREST
The authors have declared that no conflict of interest exists.

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ETHICS APPROVAL AND CONSENT TO PARTICIPATE
All animal procedures were in compliance with the Guide for the Care and Use of Laboratory Animals (NIH, 76 FR 91 [May 11, 2011]) and were approved by the Animal Care and Use committee of Jiangsu University.

CONSENT FOR PUBLICATION
All authors approve to submit it as the present style.

DATA AVAILABILITY STATEMENT
All the data were shown in the manuscript and are available from the correspondence author upon request.

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REFERENCES
1. Zhou L, Tian Y, Guo F, et al. LincRNA-p21 knockdown reversed tumor-associated macrophages function by promoting MDM2 to antagonize p53 activation and alleviate breast cancer development. *Cancer Immunol Immunother*. 2020;69:835–846.
2. Chen YH, Hsieh SC, Chen WY, et al. Spontaneous resolution of acute gouty arthritis is associated with rapid induction of the anti-inflammatory factors TGFbeta1, IL-10 and soluble TNF receptors and the intracellular cytokine negative regulators CIS and SOCS3. *Ann Rheum Dis*. 2011;70:1655–1663.

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