Naringenin suppresses BEAS-2B-derived extracellular vesicular cargoes disorder caused by cigarette smoke extract thereby inhibiting M1 macrophage polarization

Zhen Chen, Hao Wu, Weiyang Fan, Jiashuo Zhang, Yue Yao, Weiwei Su, Yonggang Wang and Peibo Li*

Guangdong Engineering and Technology Research Center for Quality and Efficacy Re-evaluation of Post-marketed Traditional Chinese Medicine (TCM), State Key Laboratory of Biocontrol, Guangdong Provincial Key Laboratory of Plant Resources, School of Life Sciences, Sun Yat-sen University, Guangzhou, China

Extracellular vesicles (EVs)-mediated epithelium-macrophage crosstalk has been proved to maintain lung homeostasis in cigarette smoke-induced lung diseases such as chronic obstructive pulmonary disease (COPD). In our previous study, we found that EVs derived from cigarette smoke extract (CSE) treated BEAS-2B promoted M1 macrophage polarization, which probably accelerated the development of inflammatory responses. Naringenin has been proved to suppress M1 macrophage polarization, but whether naringenin regulates macrophage polarization mediated by EVs has not been reported. In this study, we firstly found that EVs derived from naringenin and CSE co-treated BEAS-2B significantly inhibited the expression of CD86 and CD80 and the secretion of tumor necrosis factor (TNF)-α, interleukin (IL)-6, IL-1β, inducible nitric oxide synthase (iNOS), and IL-12 in macrophage induced by EVs derived from CSE-treated BEAS-2B. Further research revealed that naringenin downregulated BEAS-2B-derived EVs miR-21-3p which targeted phosphatase and tensin homolog deleted on chromosome ten/protein kinase B (PTEN/AKT) cascade in macrophages and then suppressed M1 macrophage polarization. Subsequent proteomics suggested that naringenin decreased BEAS-2B-derived EVs poly ADP-ribose polymerase (PARP)1 expression thereby suppressing M1 macrophage polarization probably. Our study provides novel pharmacological references for the mechanism of naringenin in the treatment of cigarette smoke-induced lung inflammatory diseases.

KEYWORDS
cigarette smoke, naringenin, exosome, macrophage polarization, epithelium-macrophage crosstalk
Introduction

Long-term exposure to cigarette smoke predisposes to and accelerates the development of inflammation in chronic lung diseases such as chronic obstructive pulmonary disease (COPD) and asthma (1). Extracellular vesicles (EVs), novel intercellular communication tools, contain bioactive molecules such as nucleic acids, proteins, and lipids (2). Recently, EVs-mediated epithelium-macrophage crosstalk was proved to maintain lung homeostasis, whereas aberrant crosstalk probably led to immune imbalance and inflammatory response in lung diseases (3). Excessive accumulation of M1 macrophages at inflammatory sites can lead to exacerbated tissue damage and more severe inflammatory storms (4). In our previous study, we found that cigarette smoke extract (CSE) upregulated the human bronchial epithelial cell line (BEAS-2B)-derived EVs miR-21-3p and miR-27b-3p thereby promoting M1 macrophage polarization (5).

Naringenin, a natural flavone, is abundantly present in citrus fruits and has shown strongly anti-inflammatory activities in multiple in vitro and in vivo studies of lung diseases such as COPD, asthma, lung fibrosis, and acute lung injury (6). Furthermore, Karuppagounder et al. (7) found that naringenin could suppress M1 macrophage polarization thereby ameliorating skin inflammation in atopic dermatitis mice. However, whether naringenin can affect the extracellular vesicular cargoes profiles derived from airway epithelium and then regulate macrophage polarization has not been reported.

Consequently, in this study, we investigated whether naringenin could downregulate the EVs miR-21-3p or miR-27b-3p levels derived from airway epithelium thereby regulating M1 macrophage polarization. Moreover, considering that contained proteins were also critical for EVs to exert biological functions, we employed proteomics to explore whether naringenin altered protein expression profiles of EVs and then regulated M1 macrophage polarization. Our findings provided more comprehensive pharmacological evidence for naringenin in the treatment of cigarette smoke-induced lung diseases.

Materials and methods

Chemicals and reagents

The BEAS-2B cells were obtained from iCell (Shanghai, China), and the murine alveolar macrophage cell line (MH-S) was purchased from Procell (Wuhan, China). The murine leukemia macrophage cell line (RAW264.7) and the human myeloid leukemia mononuclear cell line (THP-1) were obtained from EK-Bioscience (Shanghai, China). High glucose dulbecco’s modified eagle medium (DMEM) was acquired from HyClone (Logan, UT, USA). Roswell park memorial institute (RPMI) 1640 medium was purchased from Gibco (Grand Island, NY, USA). Fetal bovine serum was purchased from Corning (Auckland, New Zealand) and penicillin/streptomycin was obtained from Gibco. Naringenin, dimethyl sulfoxide (DMSO), and phorbol ester (PMA) were obtained from Sigma-Aldrich (St. Louis, MO, USA). Specific phosphatase and tensin homolog deleted on chromosome ten (PTEN) inhibitor (SF1670) and poly ADP-ribose polymerase (PARP)1 inhibitor (AG14361) were acquired from MCE (Monmouth Junction, NJ, USA).

CSE preparation

CSE was prepared as previously reported (8). Briefly, the smoke of three commercial cigarettes (Cocopalm, Guangzhou, China) was pumped into 5 ml warm medium by a bt100-2j peristaltic pump (LongerPump, Baoding, China). Before administration to cells, the CSE solution is needed to be filtered through 0.22-μm-pore-size sterilizing filters. The solution was diluted 10-fold, after which the absorbance was measured at 320 nm. If the absorbance approaches 0.8, the resultant CSE solution was regarded as 100% CSE, which was diluted with DMEM medium before being used.

Cell culture and treatment

BEAS-2B cells were cultured in high glucose DMEM plus 10% fetal bovine serum and 1% penicillin/streptomycin at 37°C and 5% CO2. THP-1 monocytes were treated with 100 ng/ml PMA for 48 h and differentiated into M0 macrophages on day 3. RAW264.7, MH-S, and THP-1 macrophages were cultured in RPMI 1640 medium plus 10% fetal bovine serum and 1% penicillin/streptomycin at 37°C and 5% CO2. When 80-90% confluence was reached, BEAS-2B cells were incubated with EVs-free DMEM (100,000×g overnight to remove the EVs in fetal bovine serum) at 37°C and 5% CO2 for 24 h. Then, the supernatants were harvested to isolate EVs.

EVs extraction

EVs were isolated as described previously (9). Briefly, the supernatants were centrifuged at 300×g for 2,000×g for 10 min, respectively. Next, the supernatants were centrifuged at 10,000×g for 30 min. Then, we filtered the harvested supernatants with 0.22-μm filters. Subsequently, ultracentrifuge (Beckman, Brea, CA, USA) was utilized for the centrifugation of supernatants at 100,000×g for 70 min. Finally, the required pellets were resuspended with PBS before being kept at -80°C.

The abbreviations of each group of EVs are as follows: Control-EVs: BEAS-2B cells were treated without any specific treatment, and supernatants were harvested after 24 h for EVs...
isolation; CSE-EVs: the cells were treated with 7.5% CSE and supernatants were harvested for EVs isolation; CSE+NRG-EVs: the cells were co-treated with 7.5% CSE and 100 μM naringenin and supernatants were harvested for EVs isolation; CSE+DMSO-EVs: the cells were co-treated with 7.5% CSE and 0.1% DMSO (vehicle of naringenin) and supernatants were harvested for EVs isolation.

**EVs identification**

Isolated EVs were imaged utilizing the JEM-1400Flash transmission electron microscopy (TEM) (JEOL Ltd., Japan) at a voltage of 120 kV. The size distributions of EVs were determined by nanoparticle tracking analysis (NTA) using a NanoSight-NS300 (Malvern Instrument, Malvern, UK). Total protein concentrations of EVs were measured using BCA Protein Assay Kits (Beyotime, Nanjing, China) according to the manufacturer’s protocols. Marker proteins of EVs were detected by western blotting. ALIX (1:1000, ab275377, Abcam), TSG101 (1:1000, ab133586, Abcam), CD9 (1:2000, ab50249, Abcam) were used.

**High-performance liquid chromatography with tandem mass spectrometry**

Naringenin in CSE+NRG-EVs was detected by HPLC-MS/MS using a 1200SL HPLC-6410 QQQ liquid chromatograph-mass spectrometer (Agilent, Santa Clara, CA, USA). EVs samples were prepared in acetonitrile. Naringenin at a concentration of 200 ng/ml was prepared as a reference standard. The chromatographic separation was achieved on an Agilent Poroshell 120 EC-C18 column (3.0 × 30 mm, 2.7 μm) at 45°C. Mobile phase: 0.1% formic acid, with a flow rate of 400 μl/min. Mass spectrometric analysis was performed on the negative ion multiple reaction monitoring mode, with Capillary 4000 V, Drying Gas 10 L/min, Neb Pressure 25 psi, Gas Temp 350°C. For naringenin, the collision energy was 12 eV and the multiple reaction monitoring modes with a precursor to product qualifier transition m/z was 270.9/150.7.

**Detection of uptake of EVs by macrophages**

EVs extracted from the supernatants of BEAS-2B cells were labeled with PKH67 green fluorescent (Sigma-Aldrich) according to the experimental procedures described by the previous study (10). Briefly, 400 μl EVs were mixed with 1.6 ml Diluent C and 8 μl PKH67 dye for 5 min at room temperature, and 1% bovine serum albumin (BSA) was added to terminate the dyeing process. PKH67-labeled EVs were recollected at 100,000×g for 70 min at 4°C. PKH67-labeled EVs or the PKH67-PBS control were separately incubated for 12 h with macrophages seeded on confocal dishes. Nuclei of macrophages were stained with 5 μg/ml DAPI (Sigma-Aldrich) for 10 min at room temperature and then observed under an SP8X laser scanning confocal microscope (Leica, Wetzlar, Germany).

**Flow cytometry**

The polarization-related surface markers of RAW264.7, MH-S, and THP-1 macrophages were detected by flow cytometry. In our previous study (5), we found that 10 μg/ml CSE-EVs could significantly promote M1 polarization of RAW264.7 macrophages, thus we treated macrophages (5×10⁴ cells/cm²) with 10 μg/ml EVs in EVs-free RPMI 1640 mediums for 24 h in this study. For positive control, 2 μM SF1670 and 1 μM AG14361 were used to inhibit PTEN and PAPR1 expression of THP-1 cells, respectively. Then, the cells were collected and stained with the following antibodies on ice for 20 min in the dark. APC anti-mouse CD86, APC anti-mouse CD80, APC anti-human CD11b, PE anti-human CD86, and PE anti-human CD80 (Biolegend, San Diego, CA, USA) were used in this procedure. After being washed twice with PBS, cells were fixed with fixation buffer and then analyzed by flow cytometry (CytoFLEX, Beckman).

**Enzyme-linked immunosorbent assay**

Cell supernatants were collected from RAW264.7 and THP-1 macrophages and centrifuged at 3000 rpm at 4°C for 20 min. The concentrations of tumor necrosis factor (TNF)-α, interleukin (IL)-6, IL-1β, inducible nitric oxide synthase (iNOS), and IL-12 were determined with ELISA Assay Kits (MeiMian, Wuhan, China) according to the manufacturer’s instructions. All samples were measured at 450 nm wavelength by an Epoch-2 microplate reader (BioTek, Winooski, VT, USA).

**RNA isolation and quantitative real-time PCR**

Total RNA from THP-1 macrophages was extracted using TRIzol (Invitrogen, Carlsbad, NY, USA). Total EVs RNA was isolated using Cell Culture Media Exosome Purification Kits (Norgen Biotek, Canada) following the manufacturer’s instructions. The RNA quality and quantity were determined using the NanoDrop 2000c (Thermo Fisher Scientific, MA, USA). Real-time PCR was performed with GoTaq® qPCR Master Mix (Promega, Madison, WI, USA) using a LightCycler 480 System (Roche, Mannheim, Germany). The
primer sequences are listed in Supplementary 1. The relative expression of the target mRNA was normalized to GAPDH and the target miRNA was normalized to U6. All relative changes in gene expression were analyzed by the 2-ΔΔCT method.

**Western blotting**

THP-1 macrophages were seeded in 6-well plates at 5×10^4 cells/cm^2 and treated with 10 μg/mL EVs for 24 h. Then, they were lysed by RIPA buffer containing protease inhibitors (Beyotime, Shanghai, China). Protein lysates were collected after centrifuging at 16,000 rpm for 30 min and the concentration was determined by BCA Protein Assay Kits. Then, proteins were loaded on 10% SDS-polyacrylamide gels and transferred to polyvinylidene difluoride membranes. The membranes were blocked with 5% fat-free milk for 2 h at room temperature and incubated with specific antibodies overnight at 4°C. PTEN (1:1000, 9188T, CST), AKT (1:1000, 4691T, CST), p-AKT (1:1000, 4060T, CST), and GAPDH (1:10000, ab8245, Abcam) were used in this procedure. After being washed, the membranes were incubated with corresponding HRP-conjugated secondary antibodies at room temperature for 1 h. The bands were visualized by a chemiluminescence imaging system (Tanon, Shanghai, China). ImageJ software (NIH, Bethesda, MD, USA) was used to quantify the integrated density of bands.

**Protein filter-aided sample preparation of EVs**

EVs were isolated with SDT buffer (4% SDS, 100 mM Tris-HCl, PH 7.6) and the proteins of EVs were quantified with the BCA Protein Assay Kit. Two hundred μg of proteins for each sample were reduced with 50 mM dithiothreitol and repeated ultrafiltration using a UA buffer (8 M Urea, 150 mM Tris-HCl, pH 8.5). Then 100 μl iodoacetamide was added to block reduced cysteine residues and the filters were washed with 100 μl UA buffer three times and then 100 μl 25 mM NH_4HCO_3 buffer twice. Finally, the protein suspensions were digested with 4 μg trypsin in NH_4HCO_3 buffer overnight at 37°C, and the resulting peptides were collected as a filtrate.

**Nanoscale liquid chromatography coupled to tandem mass spectrometry**

Samples were analyzed on a nanoElute (Bruker, Bremen, Germany) coupled to a timsTOF Pro (Bruker, Bremen, Germany) equipped with a CaptiveSpray source. Peptides were separated on an analytical column, 1.6μm C18 beads with a packed emitter tip (IonOpticks, Australia). The column was equilibrated using 4 column volumes before loading the sample in 100% buffer A (99.9% MilliQ water, 0.1% FA). Samples were separated at 300 nl/min using a linear gradient and the timsTOF Pro was operated in PASEF mode. The raw data were analyzed using MaxQuant software version 1.6.14.0. The cutoff of the false discovery rate for peptide and protein identification was set to 0.01. Protein abundance was calculated based on the normalized spectral protein intensity. Proteins which Fold change > 2 or < 0.5 and p-value < 0.05 were considered to be differentially expressed proteins.

**Bioinformatics analysis**

Bioinformatics analyses for differentially expressed EVs proteins in this study were performed using the OmicStudio tools at https://www.omicstudio.cn/tool. The heat map was performed using the Advanced Heatmap Plots tool to visualize all of the differentially expressed proteins. The annotation from Gene Ontology (GO) terms to proteins was completed by Blast2GO Command-Line. Then further improvement of annotation and connection between GO terms was carried out by ANNEX. Pathway analysis was performed using the KEGG database. Fisher’s Exact Test was used to identify the significantly enriched GO terms and pathways by comparing the number of differentially expressed proteins and total proteins.

**Statistical analysis**

Experiment results were expressed as the mean ± SD of at least 3 independent experiments and statistical analyses were performed using GraphPad Prism 8 software. Differences between groups was assessed by unpaired two-tailed Student’s t-test for comparison of two groups and one-way analysis of variance (ANOVA) followed by Dunnett’s test for multiple groups. A p < 0.05 was considered the minimum level of statistical significance.

**Results**

The characterization of BEAS-2B-derived EVs

EVs were isolated from the supernatants of BEAS-2B cells. TEM, NTA, and western blotting were carried out to identify the EVs. The morphology of EVs was visualized using TEM. We observed that the purified EVs have cup or spherical shapes with membrane vesicle structures (Figure 1A). NTA analysis showed mere peaks of EVs at approximately 120 nm with mean diameters within 150 nm (Figure 1B). The concentrations of EVs were approximately ranging from 5.7 × 10^8 to 6.8 × 10^8 particles/ml. Western blotting illustrated that ALIX, TSG101, and CD9 were enriched in the EVs pool but not in the BEAS-2B cells pool (Figure 1C).
CSE+NRG-EVs almost did not contain naringenin

To clarify whether CSE+NRG-EVs contain naringenin, the content of naringenin in CSE+NRG-EVs was determined by HPLC-MS/MS. As shown in Figure 2, compared with the naringenin reference standard, the peak height of CSE+NRG-EVs was extremely low, indicating that naringenin was almost absent in CSE+NRG-EVs.
CSE+NRG-EVs suppressed M1 polarization of RAW264.7 macrophages

RAW264.7 macrophage was considered a typical cell line for studying macrophage function, thus we firstly utilized RAW264.7 macrophages to investigate the role of CSE+NRG-EVs on macrophage polarization. To clarify whether BEAS-2B-derived EVs can be internalized by RAW264.7 macrophages, PKH67-labeled EVs were incubated with RAW264.7 cells for 12 h. Subsequent confocal microscope imaging showed that the green fluorescence dyed EVs were dotted around the blue fluorescence dyed nuclei of macrophages, suggesting that BEAS-2B-derived EVs could be concentrated in the cytoplasm of RAW264.7 macrophages (Figure 3A).

Flow cytometry and ELISA were performed to investigate the role of CSE+NRG-EVs on M1 macrophage polarization of RAW264.7 macrophages. In contrast to CSE-EVs, the percentages of M1 phenotype (CD86+ and CD80+) cells were found to be decreased by the treatment of CSE+NRG-EVs (Figure 3B). Furthermore, we observed that the secretion of M1-associated cytokines including TNF-α, IL-6, IL-1β, iNOS, and IL-12 in the CSE+NRG-EVs group was significantly reduced in comparison with that in the CSE-EVs group (Figure 3C). These data suggested that CSE+NRG-EVs inhibited M1 polarization of RAW264.7 macrophages induced by CSE-EVs.

CSE+NRG-EVs suppressed M1 polarization of MH-S macrophages

Alveolar macrophages, important immune cells in the lung, are mainly responsible for eliminating invaded pathogens and timely adjusting the immune status to alleviate tissue damage (11). To further investigate the role of CSE+NRG-EVs on M1 macrophage polarization, we analyzed M1-associated markers of MH-S macrophages by flow cytometry. Confocal microscope imaging confirmed that BEAS-2B-derived EVs were taken up by MH-S macrophages (Figure 4A). The flow cytometry results showed that CSE+NRG-EVs markedly decreased the ratios of CD86+ and CD80+ macrophages, indicating that the M1 polarization of MH-S macrophages was inhibited by the treatment of CSE+NRG-EVs (Figure 4B). These findings suggest that CSE+NRG-EVs exerted an inhibitory effect on the M1 polarization of MH-S macrophages induced by CSE-EVs.

CSE+NRG-EVs suppressed M1 polarization of THP-1 macrophages

Monocytes are a major source of resident macrophages in the lung, especially in the initial phase of inflammation response (12). To clarify if the inhibition of M1 macrophage polarization was also observed in the human macrophage cell line. We incubated the THP-1 macrophages with BEAS-2B-derived EVs and examined the levels of M1-associated macrophage markers. THP-1 monocytes presented as rounded cells and differentiated into spindle-shaped or irregular polygon M0 macrophages by PMA treatment (Supplementary 2). Similarly, we observed the internalization of BEAS-2B-derived EVs by THP-1 macrophages utilizing confocal microscopy (Figure 5A).

The percentage of polarization-related surface markers of macrophages was detected by flow cytometry (Figure 5B). High positivity for CD11b demonstrated that THP-1 monocytes were efficiently differentiated into M0 macrophages in response to PMA. Furthermore, CSE-EVs significantly increased the CD11b+CD86+ and CD11b+CD80+ cells percentage of THP-1 macrophages, instead, CSE+NRG-EVs markedly reduced the percentages of CD11b+CD86+ and CD11b+CD80+ cells. The concentration of M1-associated cytokines of THP-1 macrophages in the supernatants was measured by ELISA (Figure 5C). CSE+NRG-EVs also reversed the secretion of TNF-α, IL-1β, IL-6, iNOS, and IL-12 induced by CSE-EVs in THP-1 macrophages, as the same in RAW264.7 macrophages. Together these results suggested that CSE+NRG-EVs also played an inhibitory role in the M1 polarization of THP-1 macrophages induced by CSE-EVs.

Naringenin downregulated EVs miR-21-3p which targeted PTEN/AKT pathway thereby suppressing M1 macrophage polarization

In our previous study, we found that CSE promoted M1 macrophage polarization via upregulated EVs miR-21-3p and miR-27b-3p derived from BEAS-2B cells (5). Thus, we speculated that naringenin suppressed M1 macrophage polarization via downregulating these miRNAs in this study. QRT-PCR indicated the miR-21-3p level, but not the miR-27b-3p level, in CSE+NRG-EVs was markedly decreased compared with CSE-EVs (Figure 6A).

Zhu et al. (13) performed dual-luciferase reporter assays and affirmed that the miR-21-3p targeted the conserved sequence of PTEN 3′-UTR in human cancerous liver cells, which was predicted by TargetScan and PicTar databases. Furthermore, Hu et al. (14) found that transfection of miR-21-3p inhibitors increased the PTEN level in human microvascular endothelial cells. PTEN is a classical suppressor that antagonizes the phosphatidylinositol 3-kinase/protein kinase B (PI3K/AKT) signaling pathway (15). In this study, we examined whether CSE+NRG-EVs targeted the PTEN/AKT pathway of macrophages. Western blotting and qRT-PCR confirmed that the PTEN protein and mRNA levels remarkably decreased in THP-1 macrophages with CSE-EVs treatment, while...
significantly increasing in the CSE+NRG-EVs group (vs. CSE-EVs group) (Figures 6B, C). Correspondingly, phosphorylation of AKT/total AKT (p-AKT/AKT) protein level was upregulated by CSE-EVs treatment, while reversed by CSE+NRG-EVs treatment (Figure 6D). These findings suggested that naringenin could downregulate BEAS-2B-derived EVs miR-21-3p, thereby increasing the PTEN level and inhibiting the activation of AKT of macrophages.
We then clarified the relation between PTEN and the inhibitory effect of CSE+NRG-EVs on M1 macrophage polarization. Flow cytometry analysis revealed that the ratio of the CD11b+CD80+ cells in the CSE+NRG-EVs in combination with SF1670 group was significantly higher than in the CSE-EVs group but lower than in the CSE-EVs group (Figure 6E). These results indicated that inhibition of PTEN expression attenuated but did not completely abolish the inhibitory effect of CSE+NRG-EVs on M1 macrophage polarization. Altogether, the outcomes suggested that naringenin could downregulate BEAS-2B-derived EVs miR-21-3p targeting PTEN/AKT cascade in macrophages, which was one of the mechanisms of suppressing M1 macrophage polarization.
Naringenin downregulating EVs PARP1 thereby suppressing M1 macrophage polarization probably

Considering that proteins are also essential cargoes in EVs, we performed EVs proteomics to further investigate whether naringenin altered protein profiles of EVs thereby inhibiting M1 macrophage polarization. By nano-LC-MS/MS, 1001 proteins were identified, 208 of which were selected as significantly differential expressions with Fold change > 2 or < 0.5 and p-value < 0.05 in CSE-EVs compared to Control-EVs, while 251 differentially expressed proteins were identified in

**FIGURE 5**
CSE+NRG-EVs inhibited M1 macrophage polarization of THP-1 macrophages induced by CSE-EVs. (A) Uptake of BEAS-2B-derived EVs by THP-1 macrophages detected by confocal microscopy. Scale bar = 20 μm. (B) Flow cytometry analysis of the percentages of CD86+ and CD80+ THP-1 macrophages treated with different groups of EVs (n = 3). (C) ELISA was performed to measure the secretion levels of TNF-α, IL-6, IL-1β, iNOS, and IL-12 of THP-1 macrophages (n = 6). **p < 0.01 vs. Control-EVs group. #p < 0.05, ##p < 0.01 vs. CSE-EVs group.
CSE+NRG-EVs compared to CSE-EVs. The heat map illustrated that 91 proteins were significantly reversed by the treatment of naringenin in EVs, including 84 proteins upregulated in CSE-EVs and downregulated in CSE+NRG-EVs, and 7 proteins with the opposite trend (Figure 7A). The original proteomic data were shown in Supplementary 3.

For the 91 reversed proteins, functional enrichment analysis was performed for the three GO domains “biological process”, “cellular component”, “molecular function” and KEGG signaling pathways. Concerning the biological process, the reversed proteins were involved in the “cell adhesion”, the “extracellular matrix organization”, the “negative regulation of macrophage-derived foam cell differentiation”, and the “positive regulation of gene expression”. Proteins of the cellular component showed association with the “plasma membrane”, the “extracellular exosome”, the “cytoplasm”, the “cell surface”, and the “endocytic vesicle lumen”. Finally, the biological process revealed that the reversed proteins represented the “protein binding”, the “antioxidant activity”, the “zinc ion binding”, the “enzyme binding”, the “protein homodimerization activity” and so on (Figure 7B). KEGG pathways were mainly enriched in “the focal adhesion”, the “ECM-receptor interaction”, the “VEGF signaling pathway”, the “PI3K/AKT signaling pathway”, and so on (Figure 7C). Enrichment analysis suggested that CSE+NRG-EVs suppressed M1 macrophage polarization might involve these biological processes and signaling pathways.

**FIGURE 6**
Naringenin downregulated BEAS-2B-derived EVs miR-21-3p which targeted tensin homolog deleted on chromosome ten/protein kinase B (PTEN/AKT) cascade thereby suppressing M1 macrophage polarization. (A) Quantitative real-time PCR (qRT-PCR) was performed to detect the expression levels of miR-21-3p and miR-27b-3p in EVs (n = 5). (B) Expression of PTEN was quantified using western blot analysis (n = 6). (C) The mRNA level of PTEN was determined using qRT-PCR analysis (n = 6). (D) Western blotting was utilized for the p-AKT/AKT level (n = 6). (E) The CD11b+CD80+ THP-1 macrophages treated by EVs or SF1670 (PTEN inhibitor) were indicated by flow cytometry (n = 3). **p < 0.01 vs. Control-EVs group. *p < 0.05, **p < 0.01 vs. CSE-EVs group.
Existing evidence showed that PARP1 promoted a high glucose-induced phenotype shift in macrophages toward an inflammatory M1 phenotype (16). And in our proteomics, PARP1 was found to be remarkably upregulated in CSE-EVs (vs. Control-EVs, p = 0.0016) but downregulated in CSE+NRG-EVs (vs. CSE-EVs, p = 0.0123). Thus, we investigated whether EVs PARP1 was associated with the promotion effects of CSE-EVs on M1 macrophage polarization in the present study. Flow cytometry analysis indicated that the percentage of CD11b+CD80+ cells in the CSE-EVs in combination with the AG14361 group was significantly lower than in the CSE-EVs group but higher than in the CSE+NRG-EVs group (Figure 7D). These outcomes indicated that inhibition of PARP1 expression attenuated but did not eliminate the promoting effect of CSE-EVs on M1 macrophage polarization, suggesting that naringenin could downregulate BEAS-2B-derived EVs PARP1, which was another mechanism of suppressing M1 macrophage polarization probably.

Discussions

Epitheliums are the main barrier cells against the invasion of pathogens and macrophages are the major immune cells in the lung that determine the resolution of inflammation (17, 18). In particular, macrophage polarization involves and influences the orientation and the magnitude of inflammatory responses (19). A great number of publications have shown that both epitheliums and macrophages play critical roles in cigarette smoke-induced lung diseases such as COPD (20–23). However, the mechanism of EVs-mediated epithelium-macrophage crosstalk has not been elucidated and their roles in cigarette smoke-induced lung diseases are few studied. In our previous study, we found that CSE-treated BEAS-2B-derived EVs promoted M1 macrophage polarization (5).

Inhibition of M1 macrophage polarization seems to be a feasible strategy for the treatment of inflammatory diseases (24). Accessible evidence (7, 25) has proved that naringenin could suppress M1 macrophage polarization to attenuate...
inflammation, but whether naringenin could alter epithelium-derived extracellular vesicular cargoes thereby regulating macrophage polarization has not been reported. In this study, we firstly found that naringenin and CSE co-treated BEAS-2B-derived EVs significantly inhibited the expression of CD86 and CD80 and the secretion of TNF-α, IL-6, IL-1β, iNOS, and IL-12 in macrophage. These outcomes indicated that naringenin suppressed M1 macrophage polarization induced by CSE-treated BEAS-2B-derived EVs via altering their cargoes.

Accumulating studies showed that PTEN was involved in the process of M1 macrophage polarization. In this study, we found that naringenin downregulated EVs miR-21-3p to activate PTEN in macrophages. And inhibiting PTEN expression attenuated the inhibitory effect of naringenin and CSE co-treated BEAS-2B-derived EVs on M1 macrophage polarization. Furthermore, PTEN could inhibit the AKT activation and the PI3K/AKT pathway possessed a low p-value and a high rich factor in the KEGG enrichment. Liu et al. (26) found that mesenchymal stem cell-derived EVs inhibited PTEN/AKT pathway, thereby suppressing the inflammatory response in diabetic mice. Kai et al. (27) proved that regulatory T lymphocytes ameliorated intracerebral hemorrhage-induced inflammatory injury by suppressing M1 macrophage polarization through the IL-10/GSK3β/PTEN axis. Together these findings indicated that naringenin downregulated BEAS-2B-derived EVs miR-21-3p which targeted the PTEN/AKT cascade of macrophages and then suppressed M1 macrophage polarization.

EVs can influence the function of recipient cells by delivering information through their contained abundant proteins (28). In the present study, we performed proteomic analysis and found that naringenin decreased EVs PARP1 expression. Further research revealed that inhibition of PARP1 attenuated the promotion effect of CSE-treated BEAS-2B-derived EVs on M1 macrophage polarization. Clinical studies showed that PARP1 activity and mRNA level were significantly higher in COPD patients compared to non-smokers, which were positively correlated with the progression of COPD (29). Choudhuri et al. (30) found that inhibiting PARP1 activity or knocking out PARP1 gene could significantly suppress M1 macrophage polarization. These outcomes suggested that naringenin downregulated BEAS-2B-derived EVs PARP1 and then suppressed M1 macrophage polarization probably.

In conclusion, we found that naringenin suppressed BEAS-2B-derived extracellular vesicular cargoes disorder caused by CSE thereby inhibiting M1 macrophage polarization. The underlying mechanisms included that naringenin downregulated BEAS-2B-derived EVs miR-21-3p which targeted PTEN/AKT cascade and naringenin decreased BEAS-2B-derived EVs PARP1 expression, and then suppressed M1 macrophage polarization (Figure 8). Indeed, further studies are needed to address several unanswered questions. It is still not clear whether naringenin regulates macrophage polarization to attenuate lung inflammation in vivo via altering extracellular vesicular cargoes derived from airway epithelium. Our results also warrant further investigation to
discover whether naringenin regulates macrophage polarization via altering the other cargoes such as long non-coding RNAs, circular RNAs, and lipids. Furthermore, a limitation of our study is that the macrophage cell lines but not human primary macrophages were used. Therefore, experiments using human primary CD14 positive peripheral blood mononuclear cell-derived macrophages will be worth conducting to demonstrate M1 macrophage polarization in our future investigations. In a word, our study provides novel scientific bases for clarifying the mechanism of naringenin in the treatment of cigarette smoke-induced lung inflammatory diseases.

Data availability statement

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found below: http://www.proteomexchange.org/, PXD033545.

Author contributions

PL and WS provided the concept and designed the research. PL and YW supervised the research. ZC, HW, WF, JZ, and YY performed the experiments and collected data. ZC and HW analyzed the data. ZC and PL wrote the manuscript. All authors have read and approved the final manuscript. All authors contributed to the article and approved the submitted version.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fimmu.2022.930476/full#supplementary-material

SUPPLEMENTARY 1

The list of primer sequences used in this study.

SUPPLEMENTARY 2

Differentiation induction of THP-1 cells. (A) Rounded THP-1 monocytes differentiated into (B) spindle-shaped or irregular polygon M0 macrophages with the treatment of phorbol ester. Scale bar = 200 μm.

SUPPLEMENTARY 3

The original proteomic data of EVs in this study.

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