Cancer-Secreted Exosomal MiR-620 Inhibits ESCC Aerobic Glycolysis via FOXM1/HER2 Pathway and Promotes Metastasis

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Background: Esophageal squamous cell carcinoma (ESCC) is a leading cause of cancer death worldwide. MicroRNAs (miRNAs) have been reported to regulate cell functions through exosomes. Through the Gene Expression Omnibus (GEO) database, miR-620 was selected as a serum miRNA highly expressed in ESCC, but its detailed role in ESCC has not been explored. Tumor-secreted miRNAs have been reported to promote cancer metastasis through reprogramming the aerobic glycolysis of lung fibroblasts. Therefore, we intended to verify whether exosomal miR-620 secreted in ESCC cells may regulate the aerobic glycolysis of lung fibroblasts.

Methods: The effect of miR-620 on the aerobic glycolysis of ESCC cells was firstly verified through bioinformatics prediction and mechanism assays. Exosomes secreted from ESCC cells was detected, and the influence of exosomal miR-620 in regulating the aerobic glycolysis of lung fibroblasts was then verified both in vitro and in vivo.

Results: MiR-620 inhibited ESCC malignancy and suppressed the aerobic glycolysis of ESCC cells via targeting Forkhead box M1 (FOXM1) and human epidermal growth factor receptor 2 (HER2). Moreover, exosomal miR-620 was highly secreted in ESCC and could regulate HFL1 aerobic glycolysis via FOXM1/HER2 signaling. Furthermore, exosomal miR-620 could promote ESCC metastasis by reprogramming the aerobic glycolysis of lung fibroblasts (HFL1).

Conclusion: Exosomal miR-620 secreted by ESCC cells inhibited the aerobic glycolysis via FOXM1/HER2 axis and promoted cancer metastasis.

Keywords: miR-620, exosomes, esophageal squamous cell carcinoma, aerobic glycolysis, metastasis
INTRODUCTION

Esophageal cancer is one of the leading causes of cancer-related death worldwide, and half of the cases of the disease occur in China (1). As one of the predominant histological subtypes of esophageal cancer, esophageal squamous cell carcinoma (ESCC) is characterized by late-stage diagnosis, metastasis, therapy resistance, and frequent recurrence, which results in challenging clinical management and effective therapeutic options (2). Therefore, it is urgently needed to explore more targeted biomarkers for ESCC treatment.

MicroRNAs (miRNAs) are small noncoding RNAs of 22 nucleotides, which have increasingly been recognized as potent posttranscriptional regulators of gene expression (3). They have been reported to function posttranscriptionally by usually base-pairing to the mRNA 3′-untranslated regions (3′ UTR) (4). Exosomes, which are one of the smallest extracellular vesicles released from cells, have been shown to carry different nucleic acids, including miRNAs (5, 6). More and more evidences have demonstrated that miRNAs can regulate cell functions not only between cells of the same origin but also to distant target cells (7, 8). In this study, we applied the GEO database (https://www.ncbi.nlm.nih.gov/geo/) (id: GSE122497) to obtain the serum microRNA profile of ESCC samples. Among them, many miRNAs have been demonstrated to play important parts in cancer development, especially in ESCC. Genes regulated by antitumor miR−145−3p, for example, were closely associated with the molecular pathogenesis of ESCC (9); miR-10a, miR-22, miR-100, miR-148b, miR-223, miR-133a, and miR-127-3p have all been discovered to be with significant high expression in the serum of patients with ESCC (10). Combined with relevant literature, we further searched for the miRNAs that have been reported to be associated with aerobic glycolysis, and finally a serum miRNA-miR-620 was sifted out as our study subject. MiR-620 has been reported to participate in cancer progression. For example, miR-620 promotes TGF-β-induced proliferation of airway smooth muscle cell (11), participates in the regulation of breast cancer-secreted IGF2 instigates fibroblasts and bone marrow-derived vascular progenitor cells to promote cancer progression (21). However, related research in ESCC has not been documented, which leads to our study. In this study, lung fibroblast HFL1 was cocultured with exosomal miR-620 to construct HFL1/Exo cells, and a series of mechanism experiments were conducted to explore the impact of exosomal miR-620 on lung fibroblast aerobic glycolysis and further on the progression of ESCC.

In a word, we aim to verify the upregulation expression of miR-620 in ESCC cells and further explored the effects of miR-620 on lung fibroblast aerobic glycolysis in ESCC.

MATERIALS AND METHODS

Ethical Statement

Relevant animal experiments were carried out in accordance with the approval of the ethics committee of the First Affiliated Hospital of Soochow University. All experiments are taken in accordance with the standard biosecurity and safety procedures of the First Affiliated Hospital of Soochow University.

Cell Lines

Human normal esophageal epithelial Het-1A cell line and human ESCC cell lines (ECA-109 and KYSE450) were obtained from American Type Culture Collection (ATCC, Manassas, VA, USA). The luciferase-labeled ECA-109-luc and KYSE450-luc cells were obtained from Shanghai Chenyishiye Co., Ltd (Shanghai, China). The above cells were cultivated in Dulbecco’s modified Eagle’s medium (DMEM) (MD207-050, Gibco-BRL/Invitrogen, Waltham, MA, USA) supplemented with 10% fetal bovine serum (FBS; 16000-044, Gibco, USA) and 100× penicillin/streptomycin solution (15140122, Gibco-BRL/Invitrogen, USA). Human embryonic lung fibroblast HFL1 cell line was obtained from BeNa Culture Collection (China) and cultured with 90% F-12K plus 10% FBS.

Real-Time Quantitative RT-PCR

In line with the instruction of TRIzol reagent (T9108, Takara, Japan), the isolation of total RNA samples was extracted in ECA-109 and KYSE450 cells. Synthesis of complementary DNA (cDNA) for miRNAs was carried out using TaqMan™ MicroRNA Reverse Transcription Kit (TaKaRa, Kusatsu, Japan), and the cDNA for mRNAs was synthesized by PrimeScript RT reagent Kit (TaKaRa, Japan). RT-qPCR reaction was achieved with qRT-PCR Kit (QR0100-1KT, Sigma-Aldrich, St. Louis, MO, USA) followed by 2−ΔΔCt method. In relevant assays, the reference gene was β-actin while the reference gene for miRNA quantification was U6 and miR-16 (in exosome experiments). Related sequence information is included in Supplementary Table S1.
**Cell Transfection**

Specific shRNAs targeting FOXM1 (sh1/2/3-FOXM1) and HER2 (sh1/2/3-HER2), together with their negative control shRNA (sh-NC) were provided by Ribobio Co., Ltd. (Guangzhou, China) to silence FOXM1 expression. For the overexpression of FOXM1 and HER2, the whole sequences were synthesized and subcloned into pcDNA3.1 vector to construct pcDNA3.1-FOXM1 and pcDNA3.1-HER2, with pcDNA3.1 empty vector as the negative control (NC). For the overexpression or silencing of miR-620, miR-620 mimics or miR-620 inhibitor was respectively used, with mimics-NC or inhibitor-NC as internal controls. Transfections were conducted with Lipofectamine 2000 (11668019, Invitrogen, USA) in line with the supplier’s protocols. After incubation for 48 h, cells were collected for subsequent experiments.

**Cell Counting Kit-8 Assay**

Cell viability was evaluated with the application of Cell Counting Kit-8 (CCK-8) according to the manufacturer’s guidelines. Transfected ECA-109 or KYSE450 cells were seeded to 96-well plates and then placed in an incubator with 5% CO2 at 37°C for 2000 (11668019, Invitrogen, USA) in line with the supplier’s protocols. Transfections were conducted with Lipofectamine 2000 (11668019, Invitrogen, USA) in line with the supplier’s protocols. After incubation for 48 h, cells were collected for subsequent experiments.

**Wound-Healing Assay**

ECA-109 and KYSE450 cells were seeded into the 6-well plates and then cultivated at 37°C with 5% CO2 for confluence. When the degree of cell confluence was above 90%, a pipet tip was used to scrape the cells as vertically as possible and then the cells were washed with PBS for three times. The scratched cells were removed, the serum-free medium was added, and then the cells were placed into an incubator at 37°C with 5% CO2. Finally, the absorbance at 450 nm after 24, 48, or 72 h was measured with the microplate reader to evaluate the viability of indicated cells.

**Transwell Assays**

ESCC cells were planted on the top of 24-well Transwell chambers coated with Matrigel for invasion assay or without Matrigel for migration assay. The lower chambers were loaded with complete medium. Twenty-four hours later, cells in the upper layer were removed by a cotton swab and the bottom of the chamber was fixed in methanol solution for 15 min. Crystal violet was adopted to stain the membranes for 10 min, and the invaded or migrated cells were observed and counted under a microscope (10 × 10).

**Western Blot Assay**

When the cell confluence was above 80%, the protein extracts were collected from ESCC cell lines using protein extraction kit (PROTTOT-1KT, Sigma-Aldrich, USA) and RIPA buffer (KGP701, KeyGEN BioTECH, Nanjing, China). After being separated through sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE; P0670-250ml, Beyotime Biotech, Shanghai, China), proteins were transferred to polyvinylidene fluoride (PVDF) membranes and cultured in 5% skim milk. The membranes were cultivated with primary antibodies over night at 4°C, followed by being cultivated with secondary antibody for 1 h. After washing in TBST, the secondary antibodies were added and finally assayed by ECL substrate.

The primary antibodies were obtained from Abcam (UK) and listed as follows: anti-E-cadherin (ab40772), anti-Vimentin (ab92547), anti-CD63 (ab1318), anti-CD81, anti-HSP70 and anti-FOXM1 (ab245309), anti-GM130, with anti-β-Actin (ab8227) as the internal control. Experiment was conducted three times.

**RNA Pull-Down Assay**

With the application of Pierce™ Streptavidin Magnetic Beads (Thermo Fisher, USA), RNA pull-down assay was carried out in ECA-109 cells. Experimental groups were divided into Input Antisense (control), HER2 Promoter Sense, and HER2 Promoter (mut) Sense groups. Biotinylated HER2 probes were incubated with cell extracts and streptavidin magnetic beads and then RNAs were purified with TRIzol reagent. The enrichment of FOXM1 was analyzed by Western blot. Likewise, biotinylated FOXM1 probes were incubated with cell extracts and streptavidin magnetic beads, and, finally, the enrichment of miR-620 was examined by qRT-PCR. The experiment was independently conducted in triplicate.

**Luciferase Reporter Assay**

For luciferase reporter assay, FOXM1 3′UTR possessing wild-type and mutant miR-620 binding sites were subcloned into pmirGLO luciferase vectors to obtain pmirGLO-FOXM1 (3′UTR) and pmirGLO-FOXM1 (3′UTR-Mut). MiR-620 mimics or mimics NC were cotransfected with pmirGLO-FOXM1 (3′UTR) and pmirGLO-FOXM1 (3′UTR-Mut) into NSCLC cells. After 48-h transfection, cells were extracted and the luciferase activities were analyzed utilizing the luciferase reporter assay system (Promega).

As for FOXM1 and HER2 promoter luciferase reporter assay, the HER2 promoter was constructed into pGL3 luciferase vectors and then cotransfected with pcDNA3.1 or pcDNA3.1-FOX1 into 293T cells. The luciferase activities were analyzed utilizing the luciferase reporter assay system (Promega).

**Exosome Uptake Experiment**

By referring to related document, we conducted the exosome uptake experiment (22). A total of 20 ml of culture media (1 × 107 (7) cells) were collected on ice, centrifuged at 800×g for 10 min, and then centrifuged at 12,000×g for 30 min to remove the cellular debris. Exosomes were centrifuged at 100,000×g for 2 h in a SW32 rotor (Beckman Coulter) for separating from the supernatant. The exosome pellet was washed once in a large volume of PBS and then resuspended in 100 μl of PBS. Finally, exosomes were then identified by transmission electron microscope (TEM) to observe the morphologies of exosomes secreted from miR-620 (labeled as miR-620-Exo).

In addition, the exosomes were marked by PKH67 Fluorescent Cell linker kits according to previous guidance (23). Briefly, ECA-109 and KYSE450 cells were seeded into 24-
MiR-620 Inhibits the Aerobic Glycolysis of ESCC Cells

Energy metabolism reprogramming has become an emerging hotspot in cancer research. In most of the cancers, even with abundant oxygen, cancer cells will promote the uptake and absorption of nutrients needed by newborn cells through aerobic glycolysis and reduce oxidative phosphorylation of mitochondria (16). Combining with the information above, we decided to explore whether miR-620 may regulate the aerobic glycolysis of ESCC cells. With the application of glucose determination kit, lactic acid determination kit and pyruvate kinase activity assay kit, it was verified that the glucose consumption, total lactate protein, and relative PK catalytic activity were reduced by miR-620 overexpression (Figures 2A–C). It was then measured through Seahorse XF Extracellular Flux Analyzers that the ECAR/OCR ratio was declined in ESCC cells transfected with miR-620 mimics (Figure 2D). By comparison, we analyzed the effect of miR-620 knockdown on the aerobic glycolysis of ESCC cells, and opposite results were exhibited (Supplementary Figures S2A–D). The above findings suggested that miR-620 inhibited the aerobic glycolysis of ESCC cells. For further verification, we treated ESCC cells with 50 or 100 μM 2-DG (the glycolytic inhibitor) for 48 h (24), and it was then discovered that the glucose consumption, total lactate protein, relative PK catalytic activity, and the ECAR/OCR ratio were all
reduced upon 2-DG treatment (Figures 2E–H), which demonstrated the effective inhibition of 2-DG on ESCC aerobic glycolysis. Moreover, Supplementary Figures S2E–H shows that adding 2-DG could reverse the promoted aerobic glycolysis caused by miR-620 inhibitor, which further manifested that miR-620 regulated the aerobic glycolysis of ESCC cells.

**MiR-620 Regulates HER2 Expression Through FOXM1**

After we confirmed that miR-620 could regulate the aerobic glycolysis of ESCC cells, we continued to probe into the specific molecular mechanism. Through the GEO database (GSE75241), we selected the top 30 genes upregulated in ESCC, and then potential mRNAs which could bind to miR-620 were selected through starBase database. As a result, 8 candidates were sifted out, which were ITM2C, FNDC38, CD276, LAMC2, LHFPL2, LAMC1, PARVB, and FOXM1 (Supplementary Figures S3A, B). Through qRT-PCR detection, it was found that only FOXM1 was significantly reduced by miR-620 overexpression in ECA-109 cells (Supplementary Figure S3C). Therefore, we chose FOXM1 for further investigations. The result of RNA pulldown assay with qRT-PCR showed that the enrichment of miR-620 was enhanced in the wild-type Bio-FOXM1 (3′ UTR) group rather than in the mutant group (Figure 3A). Subsequently, it was observed through luciferase reporter assay that the luciferase activity of HEK293T cells transfected with miR-620 mimics was declined in the pmirGLO-FOXM1 (3′ UTR) group while on obvious change was seen in the corresponding mutant group (Figure 3B). We utilized 50 mM α-amanitin to treat ESCC cells, and then it was discovered that the upregulation of miR-620 inhibited the stability of FOXM1 mRNA in ECA-109 cells (Figure 3C). The above findings manifested that miR-620 combined with FOXM1 3′UTR region to decline FOXM1 mRNA stability in ESCC cells. In order to figure out the downstream target gene of FOXM1, KEGG (https://www.genome.jp/kegg/) was applied and potential 6 genes related to aerobic glycolysis were detected (Supplementary Figure S3D), and then it was verified through hTFtarget database (http://bioinfo.life.hust.edu.cn/hTFtarget#!/) that only HER2 was forecast to be the target gene of FOXM1 (Supplementary Figure S3E). Therefore, HER2 was chosen for the next-step researches. It was shown from qRT-PCR assay that HER2 expression was suppressed upon sh1/2-FOXM1 transfection in ECA-109 cells, as evidenced by qRT-PCR assay (Figure 3D). In addition, FOXM1 expression was enhanced by transfecting
pcDNA3.1-FOXM1 in ESCC cells (Supplementary Figure S4A), and then the result of luciferase reporter assay manifested that HEK293T cells displayed an elevated luciferase activity in pGL3-HER2 promoter cotransfected with pcDNA3.1-FOXM1, while the corresponding mutant group exhibited no obvious change (Figure 3E). Through RNA pulldown assay with Western blot, it was shown that FOXM1 was pulled down by the HER2 promoter sense (Figure 3F). Finally, qRT-PCR data showed that the addition of pcDNA3.1-FOXM1 could normalize the reduced HER2 expression caused by miR-620 overexpression in ECA-109 cells (Figure 3G). To conclude, FOXM1 combined with the HER2 promoter so as to transcriptionally activate HER2 expression in ESCC cells.

FOXM1 Promotes the Aerobic Glycolysis of ESCC Cells via Upregulating HER2

In this part, we tried to verify the regulatory mechanism of FOXM1 and HER2 on the aerobic glycolysis of ESCC cells. As shown in Figure 4A–D, the aerobic glycolysis of ESCC cells was suppressed after FOXM1 was knocked down, as the glucose consumption, total lactate protein, relative PK catalytic activity, and the ECAR/OCR ratio were all declined. Same results were observed in the sh1/2-HER2 transfection group in ESCC cells (Figures 4E–H). By comparison, the effect of FOXM1 and HER2 overexpression on the aerobic glycolysis of ESCC cells was respectively verified in Supplementary Figures S4B–J, which further manifested that both FOXM1 and HER2 could effectively promote the aerobic glycolysis of ESCC cells. After that, experimental groups were divided into sh-NC, sh1-FOXM1, sh1-FOXM1+pcDNA3.1, and sh1-FOXM1+pcDNA3.1-HER2 for rescue assays. As shown in Figure J, the glucose consumption of ESCC cells was inhibited by FOXM1 silencing, while such effect was offset by the cotransfection of pcDNA3.1-HER2. Same opposite results were observed in the total lactate protein, relative PK catalytic activity, and the ECAR/OCR ratio of ESCC cells in different transfection groups (Figures 4K–M). In conclusion, FOXM1 promoted the aerobic glycolysis of ESCC cells via upregulating HER2 expression.

Exosomal miR-620 Is Highly Secreted in ESCC

According to the qRT-PCR data in Fig. S5A, the expression level of miR-620 in the supernatant of ESCC cells was lower than that of normal Het-1A cell, whereas the intracellular expression of miR-620 in ESCC cells was higher than that in normal Het-1A cell. Previous study has reported that miRNAs can be secreted to the extracellular environment by membrane-enclosed vesicles (such as exosomes) (6). Therefore, we tried to explore whether miR-620 may play its regulatory role through exosomes. At first, we observed through electron microscope that the exosomes secreted from miR-620-overexpressed ESCC cells (named miR-620-Exo) was 40–100 nm in diameter with oval-shaped vesicles, which conformed to the basic features of exosomes (Figure 5A;
Supplementary Figure S5B). Meanwhile, the expression of exosome-specific marker proteins (CD63, CD81, and HSP70) was detected in ESCC cell-derived exosomes instead of the exosome-negative marker GM130 (Figure 5B), and qRT-PCR data showed that miR-620 was highly expressed in the miR-620-Exo group (Figure 5C). In order to examine whether the receptor cells could accept the exosomes secreted from the donor cells as well as the exosomal miR-620, we constructed Cy3-miR-620 vectors in the exosomes secreted from ESCC cells (ECA-109/NC-Exo and KYSE450/NC-Exo) and the exosomes secreted from miR-620-overexpressed ESCC cells (ECA-109/ miR-620-Exo and KYSE450/miR-620-Exo), as previously described (23). During Transwell assays, the donor ESCC cells containing Cy3-miR-620 were cultured for 24 h and then collected and seeded into the apical chamber of Transwell, and the receptor lung fibroblast cells (HFL1) cells were placed into the basolateral chamber. Through fluorescence microscope, it was observed that most of the receptor cells displayed red fluorescence, indicating that miR-620 could be transmitted through exosomes (Figure 5D). With the application of PKH67 Green Fluorescent Cell Linker Kit, it was further manifested that PKH67-marked exosomes were absorbed and...
internalized by ESCC cells (Supplementary Figure S5C), and miR-620-Exo could enhance the expression of miR-620 in the receptor HFL1 cells (Figure 5E). Moreover, it was shown that miR-620 expression remained basically unchanged compared with the control group after we treated exosomes with RNase A, while the addition of Triton X-100 declined miR-620 expression in the exosomes, which proved that miR-620 was located in the membrane (Supplementary Figure S5D).

FIGURE 4 | FOXM1 promotes the aerobic glycolysis of ESCC cells via up-regulating HER2. (A–I) The impact of sh-FOXM1 and sh-HER2 transfection on the aerobic glycolysis of ESCC cells was respectively evaluated including the measurement of glucose consumption, total lactate protein, relative PK catalytic activity, and the ECAR/OCR ratio. (J–M) Experimental groups were divided into sh-NC, sh1-FOXM1, sh1-FOXM1+pcDNA3.1, and sh1-FOXM1+pcDNA3.1-HER2 groups, and glucose consumption, total lactate protein, relative PK catalytic activity, and the ECAR/OCR ratio were respectively examined by qRT-PCR in different groups. **p < 0.01.
Exosomal miR-620 Inhibits the Aerobic Glycolysis of Lung Fibroblasts

An increasing number of evidence has proven that miRNA can regulate cell function by transferring exosomes to adjacent or distant cells (7, 8). For example, it has been reported that breast cancer-secreted miR-122 reprograms glucose metabolism to promote metastasis (20); meanwhile, it has been illustrated that cancer cell-secreted IGF2 instigates fibroblasts and bone marrow-derived vascular progenitor cells to promote cancer progression (21). Therefore, we wanted to explore whether exosomal miR-620 may regulate the aerobic glycolysis of lung fibroblast cells (HFL1). HFL1 cells absorbing the exosomes were renamed HFL1/Exo, and then it was discovered from qRT-PCR that the expression of FOXM1 and HER2 was decreased in HFL1 cells receiving ECA-109/miR-620-Exo and KYSE450/miR-620-Exo (Figures 6A, B). In addition, it was observed in Figures 6C–F that the glucose consumption, total lactate protein, relative PK catalytic activity, as well as the ECAR/OCR ratio were all decreased (Figures 7B–E), indicating that exosomal miR-620 could inhibit the aerobic glycolysis of HFL1 cells.

Exosomal miR-620 Inhibits the Aerobic Glycolysis of Mouse Lung Fibroblasts to Promote ESCC Metastasis

In order to verify whether ESCC cell-secreted miR-620 may regulate the aerobic glycolysis of premetastatic mouse lung fibroblasts (MLF), related in vivo were conducted. qRT-PCR results show that FOXM1 and HER2 expression was reduced in MLF cells receiving ECA-109/miR-620-Exo and KYSE450/miR-620-Exo (Figure 7A). Moreover, it was discovered that after MLF received exosomal miR-620, the glucose consumption, total lactate protein, relative PK catalytic activity, as well as the ECAR/OCR ratio were all decreased (Figures 7B–E), indicating that exosomal miR-620 could inhibit the aerobic glycolysis of MLF.

Furthermore, we investigated the effect of exosomal miR-620 on ESCC metastasis. First of all, the mice were pretreated with exosomes secreted from different ESCC cells for 6 weeks. Luciferase-labeled ESCC cells (ECA-109-luc and KYSE450-luc) were then injected intracardiac, with exosome treatment for another 3 weeks. As shown on Figure 7F, qRT-PCR data manifested that the expression level of miR-620 in mouse serum was higher in ESCC cells accepting exosomal miR-620 (ECA-109/miR-620-Exo and KYSE450/miR-620-Exo) than in tumor-transplanted groups (ECA-109-luc tumor and KYSE450-luc tumor). Subsequently, it was analyzed that the expression of FOXM1 and HER2 was reduced in MLF receiving ECA-109/miR-620-Exo and KYSE450/miR-620-Exo (Figure 7G). Moreover, it was found that the relative MLF luciferase level was higher in mice treated with miR-620-Exo, which indicated that ESCC metastasis was more significant in mice with higher miR-620 expression in serum (Figure 7H). The number of pulmonary metastatic nodules was further tested to be enhanced in MLF receiving ECA-109/miR-620-Exo and...
KYSE450/miR-620-Exo (Figure 7I). To conclude, exosomal miR-620 inhibited the aerobic glycolysis of MLF and thus promoted ESCC metastasis.

**MiR-620 Inhibits the Aerobic Glycolysis of Primary Tumors and Promotes ESCC Metastasis**

To determine if the primary tumor-secreted miR-620 enhances metastasis, orthotopic mammary xenografts were constructed. It was shown that after miR-620 was upregulated in KYSE450-luc cells, the xenograft tumors displayed a slower growth tendency as well as a lighter tumor weight at the 28th day (Figures 8A, B). It was then observed through qRT-PCR that the expression of FOXM1 and HER2 in KYSE450-luc cells was reduced upon miR-620 overexpression (Figure 8C). In addition, the aerobic glycolysis of KYSE450-luc cells was discovered to be suppressed by miR-620 overexpression, as the glucose consumption, total lactate protein, relative PK catalytic activity, and the ECAR/OCR ratio of KYSE450-luc cells were all declined (Figures 8D–G). To identify the effect of miR-620 on the ESCC metastasis, miR-620 was overexpressed in KYSE450-luc cells, which turned out that miR-620 expression was enhanced in mouse serum (Figure 8H). It was then observed that the expression of FOXM1 and HER2 in KYSE450-luc cells was reduced in MLF after miR-620 was overexpressed (Figures 8I, J), and the relative MLF luciferase level was higher in the xenograft tumors treated with enhanced miR-620 expression (Figure 8K). The above findings showed that miR-620 inhibited the aerobic glycolysis of primary tumors while promoting ESCC metastasis.

**DISCUSSION**

ESCC is a common malignant tumor of the digestive system worldwide, especially in China. It is featured by advanced diagnosis and poor prognosis as a result of limited and ineffective early detection methods at an early stage (25). Through the GEO database (id: GSE122497), we found miR-620 in the serum miRNAs with obviously high expression. As miR-620 has been reported to exert an important role in many cancers (26, 27) but not in ESCC, we chose to verify its upregulation pattern in ESCC cells. In line with functional assays, miR-620 was verified to be an oncogene that inhibits cell proliferation, migration, invasion, and EMT process in ESCC.

Energy metabolism reprogramming is an emerging hotspot in cancer research. Altered aerobic glycolysis represents a well-recognized characteristic of cancer cell energy metabolism, termed as the Warburg effect. Even in the presence of abundant oxygen, a majority of tumor cells produce substantial amounts of energy through a high glycolytic metabolism (28). It has been documented that cancer-secreted miRNA can reprogram aerobic glycolysis of lung fibroblasts to promote metastasis in breast cancer (20), but such study has not been conducted in ESCC. In this study, we verified that miR-620...
FIGURE 7 | Exosomal miR-620 inhibits the aerobic glycolysis of lung fibroblasts and promotes ESCC metastasis. (A) The expression of FOXM1 and HER2 in MLF cells accepting different cancer-secreted exosomal miR-620 was detected by qRT-PCR. (B–E) qRT-PCR was utilized to detect the glucose consumption, total lactate protein, relative PK catalytic activity, and the ECAR/OCR ratio in MLF accepting different cancer-secreted exosomal miR-620. (F) Luciferase-labeled ESCC cells (ECA-109-luc and KYSE450-luc) were constructed, and qRT-PCR was used to assess the expression level of miR-620 in mouse serum in ESCC cells accepting exosomal miR-620 (ECA-109/miR-620-Exo and KYSE450/miR-620-Exo) in comparison with tumor-transplanted groups (ECA-109-luc tumor and KYSE450-luc tumor). (G) qRT-PCR was used to detect the expression of FOXM1 and HER2 in MLF treated with ECA-109/miR-620-Exo and KYSE450/miR-620-Exo. (H) qRT-PCR was used to detect the relative MLF luciferase level in mice treated with miR-620-Exo. (I) The metastasis of mice was observed through HE staining, and the number of pulmonary metastatic nodules was measured in different groups. *p < 0.05, **p < 0.01.
overexpression and the addition of 2-DG exerted consistent outcomes on ESCC cells, suggesting that miR-620 could regulate aerobic glycolysis in ESCC. Next, we further explored the specific molecular mechanisms of miR-620 in the regulation of aerobic glycolysis. Through bioinformatics prediction, FOXM1 was selected and verified to be the downstream target of miR-620, and miR-620 could bind to FOXM1 3′ UTR to reduce FOXM1 mRNA stability. In addition, FOXM1 was verified to bind to the HER2 promoter and transcriptionally activated HER2 in ESCC. Through a series of rescue assays, it was proved that FOXM1 could regulate aerobic glycolysis in ESCC through HER2. The above data suggested the miR-620/FOXM1/HER2 axis in regulating the process of aerobic glycolysis.

It has been reported that miRNAs can be secreted into the extracellular environment by extracellular vesicles, such as exosomes (6). Exosomes are effective carriers for the intercellular material transfer of material (29), and the function of exosomal miRNAs as well as miRNA dysregulation in cancer-associated fibroblasts has been reported (19). In this study, through exosome extraction and uptake experiments, we discovered that exosomal miR-620 was highly secreted in ESCC cells. Also, it was found that human embryonic lung fibroblast HFL1 cells could absorb and internalize exosomes, which was renamed as HFL1/Exo for the follow-up studies.

More and more evidences have demonstrated that miRNAs can regulate cell function by transferring exosomes to the neighboring...
or distant cells (7, 8). It has been reported that breast cancer-secreted miR-122 reprograms lung fibroblast aerobic glycolysis and thus promotes metastasis (20), and it was discovered that cancer cell-secreted IGF2 instigates fibroblasts and bone marrow-derived vascular progenitor cells to promote cancer progression (21). In this study, we verified through mechanism assays that exosomal miR-620 downregulated aerobic glycolysis of HFL1. Finally, in vivo experiments were carried out and demonstrated that ESCC-secreted miR-620 could promote reprogram aerobic glycolysis in lung fibroblasts and thus promote metastasis in ESCC. A graphical abstract has been provided for better understanding.

In summary, our study suggests that exosomal miR-620 secreted by ESCC cells regulates HFL1 aerobic glycolysis via FOXM1/HER2 signaling and promotes ESCC metastasis by reprogramming the aerobic glycolysis of lung fibroblasts.

Our study has some limitations; for example, we chose to study miR-620 at the beginning of our study and then found through experiments that it may exert functions through exosomes. There are many other potential miRNAs in the exosomes of ESCC, but we did not conduct related sequencing for them to better select the target miRNA, which needs further sequencing test in the future to help us complete our research. In addition, human epidermal growth factor receptor 2-positive (HER2+) breast cancer makes up about 20% of all invasive breast cancers, and Herceptin (trastuzumab), known as a human monoclonal antibody that interferes with the HER2 receptor, is currently the only FDA-approved therapeutic antibody for HER2-positive breast cancer (30) (31). In our study, HER2 was identified to be the target of FOXM1, which further participated in the miR-620/FOXM1/HER2 axis to regulate ESCC aerobic glycolysis. As the ESCC cell lines used in our study did not involve HER2 positivity, we did not focus on exploring the potential mechanism of Herceptin here. On the other hand, as we utilized lung fibroblasts in mechanism assays, we will try to utilize lung fibroblasts of esophageal origin to further complete our investigations on the aerobic glycolysis of ESCC. The extraction of exosomes using ultracentrifugation is controversial as it cannot exclude the contamination of other particles (such as microvesicles), while there are also reports proving that exosomes were extracted through this way (32) (33). We will consider this important matter and try to apply other approaches to extract exosomes in future researches. All in all, we hope that this study may provide a novel insight into therapeutic options for future ESCC treatment.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material. Further inquiries can be directed to the corresponding author.

**ETHICS STATEMENT**

All the relevant animal experiments were carried out in accordance with the Animal Care and Use Committee guidelines of the First Affiliated Hospital of Soochow University. Written informed consent was obtained from the owners for the participation of their animals in this study.

**AUTHOR CONTRIBUTIONS**

FL and YZ conceived and designed the study. YW and HL performed the experiments. SL and BP analyzed the data. LS and YX prepared all the figures. DJ wrote the paper. All authors read and approved the final manuscript.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fonc.2022.756109/full#supplementary-material

**Supplementary Figure 1 |** MiR-620 regulates the phenotype of ESCC cells. (A) With the application of GEO database (GSE122497), miR-620 was selected through large-scale serum microRNA profiling as a significantly up-regulated miRNA. (B) CCK-8 assay was carried out to assess the effect of miR-620 silencing on the proliferation of ESCC cells. (C) Wounding healing assay was conducted to evaluate the migration of ESCC cells upon miR-620 silencing. (D) The migration as well as invasion of ESCC cells transfected with miR-620 inhibitor was evaluated through Transwell assay. (E) The phenotype of ESCC cells upon miR-620 knockdown was observed through microscope. (F) Western blot was utilized to detect the expression of the EMT-related proteins (E-cadherin and Vimentin) upon miR-620 knockdown. **P < 0.01.

**Supplementary Figure 2 |** MiR-620 regulates the aerobic glycolysis of ESCC cells. (A) Glucose determination kit was applied to examine the glucose determination of miR-620 inhibitor-transfected ESCC cells. (B) Lactic acid determination kit was used to detect lactic acid content in ESCC cells upon miR-620 silencing. (C) Pyruvate kinase activity assay kit was applied to measure the PK content in miR-620 inhibitor-transfected ESCC cells. (D) The ECAR/OCR ratio in miR-620 inhibitor-transfected ESCC cells was tested by Seahorse XF Extracellular Flux Analyzer. (E-H) ESCC cells were treated with 50 or 100μM 2-DG (the glycolytic inhibitor) for 48h, and the glucose consumption, total lactate protein, PK catalytic activity and the ECAR/OCR ratio were respectively examined via qRT-PCR in different groups (inhibitor-NC, miR-620 inhibitor, miR-620 inhibitor+NC and miR-620 inhibitor+2-DG). **P < 0.01.

**Supplementary Figure 3 |** MiR-620 regulates HER2 expression through FOXM1. (A, B) Through GEO database (GSE75241), the top 30 genes up-regulated in ESCC were selected, and potential miRNAs which could bind to miR-620 were selected through starBase database. (C) The expression of 8 candidates (ITM2C, FNDC3B, CD278, LAMC2, LHFPL2, LAMC1, PARVB and FOXM1) was measured by qRT-PCR in ESCC cells transfected with miR-620 mimics. (D) KEGG database was applied to
Supplementary Figure 4 | FOXM1 regulates the aerobic glycolysis of ESCC cells through HER2. (A) FOXM1 expression was enhanced in ESCC cells by the transfection of pcDNA3.1-FOXM1. (B–E) qRT-PCR was utilized to test the glucose consumption, total lactate protein, relative PK catalytic activity and the ECAR/OCR ratio in FOXM1-overexpressed ESCC cells. (F–J). HER2 expression was enhanced in ESCC cells, and then qRT-PCR was utilized to test the glucose consumption, total lactate protein, relative PK catalytic activity and the ECAR/OCR ratio in HER2-overexpressed ESCC cells. *P < 0.01.

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