INTRODUCTION

Cervical cancer is a leading cause of cancer death in women worldwide. During the past decades, the application of new risk-screening techniques, such as The ThinPrep cytologic test (TCT), human papillomavirus (HPV) DNA detection, and high-risk HPV E6/E7 mRNA detection help to detect premalignant dysplasia and cervical cancer in the early stages in developed countries. The growing vaccination coverage also makes cervical cancer preventable. However, in low- and middle-income countries, many patients are diagnosed in advanced stages with limited therapeutic options.

Currently, radiation therapy remains an essential component for cervical cancer to achieve a cure. In the definitive setting, chemoradiation is recommended for stages IB3-IVA, and radiotherapy or chemoradiation is conditionally recommended for stages IA1-IB2 when medically inoperable. In addition, brachytherapy is strongly
Heatshock protein 90 (HSP90) is a group of inducible ATP-dependent molecular chaperones in response to cellular stress. Four isoforms of HSP90 have been identified in mammalian cells, including Hsp90α (encoded by HSP90AA1) and Hsp90β (HSP90AB1) in the cytosol, Grp94 in the endoplasmic reticulum, and Trap1 in mitochondria. Unlike the small HSPs (such as HS40s and HSP70s), which primarily support the degradation of irreversibly damaged proteins and reduce aggregation of misfolded proteins, HSP90 mainly enhances the stability and function of conformationally labile proteins after primary folding. Hsp90α and Hsp90β are highly homologous, with 85% sequence identity. Therefore, these 2 proteins share similar activities in regulating client proteins. A series of HSP90 client proteins is involved in oncogenic processes, such as receptor tyrosine kinases (vascular endothelial growth factor receptor [VEGFR], epidermal growth factor receptor [EGFR], and rb-b2 receptor tyrosine kinase 2 [ERBB2]), transcription factors (HIF-1α, BCL6, and OCT4), apoptosis-associated proteins (BCL2 and BIRC5), cell cycle regulatory proteins (Cyclin D1, CDK4, and WEE1) and signal transduction proteins (AKT, NF-κB, B-RAF, and c-MET). Therefore, HSP90 inhibition has been considered as a high-potential strategy for cancer therapy. Several HSP90 inhibitors have been developed for animal studies or have entered clinical trials for various refractory cancers, such as lung cancer cells, soft tissue sarcoma cells, pancreatic cancer, and cervical cancer. In the current study, we further explored the mechanisms of HSP90 in regulating the radiosensitivity of cervical cancer.

2 | MATERIALS AND METHODS

2.1 | Bioinformatic analysis

The expression profile of HSP90AA1 and HSP90AB1 in patients with primary cervical cancer and their association with survival outcomes, including progression-free survival (PFS) and overall survival (OS) were analyzed based on data extracted from The Cancer Genome Atlas Cervical Squamous Cell Carcinoma and Endocervical Adenocarcinoma (TCGA-CESC), using the UCSC Xena Browser. Kaplan-Meier survival curves were generated between patients with the top 25% and bottom 25% of gene expression.

2.2 | Gene set enrichment analysis (GSEA)

GSEA was conducted using software 4.1.0 from the Broad Institute. Patients with primary CESC in TCGA-CESC were separated into 2 groups by median HSP90AA1 or HSP90AB1 expression. Then, single-gene GSEA was conducted within the Hallmark gene sets. Gene set permutations were set to 1000 to obtain a normalized enrichment score (NES). Only the gene sets with NES >1, Nominal (NOM) P < .05, and a false discovery rate (FDR) q-value <0.25 were compared.

2.3 | Cell culture and treatment

Cervical cancer cell lines CaSki and SiHa were obtained from the American Type Culture Collection (ATCC, Manassas, VA, USA) and were cultured following the method introduced previously. X-ray resistant sublines (CaSki-RR and SiHa-RR) were generated following the strategy described previously. In brief, cells were cultured in 25-cm² culture flasks and were irradiated with a 2 Gy dose at a rate of 1.2 Gy/min using an RS 2000 Xray Irradiator (160 kV, 25 mA, 0.3 mm copper filters) (Rad Source Tech, Suwanee, GA, USA). When cells reached ~80% confluence, the cells were cultured in new dishes. When they reached ~50% confluence, they were irradiated again. The irradiation (IR) cycles were stopped when the total dose reached 60 Gy. Cells were allowed to recover for 2 wk before further experiments.

Lentiviral CD147 shRNAs were generated based on the pLKO.1-EGFP-puro plasmid, with the following shRNA sequence: #1, 5′-CCAGAATGACAAAGGCAAGAA-3′; #2, 5′-GCTACACATTGAGAACCTGAA-3′. Lentiviral particles carrying CD147 (NM_001728) with flag-tag, wild-type FBXO6 (NM_018438, FBXO6-WT) or mutant FBXO6 (p.Y241A and p.W242A, FBXO6-MT) were generated based on the pLV-IRESS-GFP-puro plasmid. Cells were infected with lentivirus at an MOI of 10. Cells were infected with harvested lentivirus for 48 h for transient infection. Cells with stable gene expression or knockdown were selected using 2 μg/mL puromycin for 1 wk. MG132 (a typical proteasome inhibitor), tansipimycin (17-AAG) and cycloheximide (CHX) (a protein synthesis inhibitor) were purchased from Sigma-Aldrich (Saint Louis, MO, USA). 17-AAG treatment was performed depending on experimental readouts.

2.4 | Western blotting and immunoprecipitation (IP)

Total protein from cellular samples was extracted using a cold radiomunoprecipitation assay lysis buffer (RIPA) buffer (20 mM Tris [pH7.5], 150 mM NaCl, 1% Triton X-100, sodium pyrophosphate, β-glycerophosphate, EDTA, Na3VO4, leupeptin and protease inhibitors) (P0013, Beyotime, Shanghai, China) buffer. The cell lysates were cleared by centrifugation at 100 000 × g for 30 min at 4°C to generate the soluble samples. Then, protein concentration was quantified using a BCA Protein Assay Kit (Beyotime). RIPA-insoluble pellets were then extracted with urea buffer (7 M urea, 2 M thiourea, 4% CHAPS, 30 mM Tris, pH 8.5 + protease inhibitors). Then, the
samples were centrifuged at 100 000 × g for 30 min at 22°C. Next, 20 μg of protein samples were loaded into each lane of an SDS-PAGE gel, separated using electrophoresis, and then transferred to nitrocellulose (NC) membranes (Merck Millipore, Burlington, MA, USA). After blocking with 5% BSA, the membranes were incubated with primary antibodies and corresponding secondary antibodies. The protein bands were developed using BeyoECL Star chemiluminescence reagents (Beyotime).

An IP assay was conducted using the Pierce Co-Immunoprecipitation Kit (ThermoFisher Scientific), and following the manufacturer’s instruction. For each co-IP assay, 10 μg antibody was used. The eluted samples were subjected to western blotting.

The primary antibodies using included rabbit anti-HSP90α (1:1:1000, PA3-013, ThermoFisher Scientific) and rabbit anti-HSP90β (1:1:1000, PA3-012, ThermoFisher Scientific), mouse anti-FBXO6 (sc-134339, Santa Cruz Biotechnology, Dallas, TX, USA), mouse anti-CD147 (1:5000, 66443-1-Ig, Proteintech, Wuhan, China), rabbit anti-γ-H2AX (1:1000, ab2893, Abcam, Cambridge, USA) and mouse anti-β-actin (1:5000, 66009-1-lg, Proteintech).

2.5 Liquid chromatography-tandem mass spectrometry (LC-MS/MS)

LC-MS/MS assay was conducted to analyze the protein samples immunoprecipitated by anti-HSP90α or anti-HSP90β, following the strategy introduced in 1 previous study.19

2.6 Quantitative real-time reverse transcription-PCR (qRT-PCR) analysis

Total RNA was extracted from cellular samples using TRIzol reagent (Invitrogen, Carlsbad, CA, USA) and then reversely transcribed into cDNA using the PrimeScript RT Kit (Takara, Dalian, Liaoning, China). qRT-PCR was performed using a SYBR Green kit (Takara). β-Actin mRNA was used as a reference gene for normalization. Relative gene expression was calculated using the 2^ΔΔCt method. Primer sequences are listed in Table S1.

2.7 Clonogenic assay

Cells were seeded into 24-well plates and were irradiated at defined doses (0, 2, 4, 6, or 8 Gy) after adhesion, using a RS 2000 X-ray Irradiator. After 14 d of incubation, cells were washed, fixed with methanol, and stained with 0.1% crystal violet. Colonies containing >50 cells were counted using microscope inspection. The plating efficiency (PE) of un-irradiated cells (0 Gy) was calculated by the formula: PE = number of colonies counted/number of cells plated. Surviving fraction (SF) of the irradiated cells was calculated using the formula: SF = number of colonies formed after treatment/number of cells seeded × PE. A linear-quadratic model was utilized to generate survival curves using the following equation:

\[ Y = \exp(-1 \times (A \times X + B \times X^2)) \]

in which Y is the fraction survival and X is the dose. A equals −1 times the initial slope, and the initial value of B equals −0.1 times the initial slope.

2.8 Mice and treatment

Animal procedures were approved by the Institutional Animal Care and Use Committee (IACUC) of the Fifth Medical Center of PLA General Hospital, Beijing, China. Nude mice (BALB/c-nude, 5-wk-old females) were purchased from Vital River Laboratories (Beijing, China). Mice were randomly assigned into 6 groups. Here, 2 × 10^6 cells (CaSki-RR, SiHa-RR, and the cell lines with stable knockdown or overexpression) suspended in a 1:1 mixture of culture medium and Matrigel Matrix (Corning, Glendale, AZ, USA) were subcutaneously injected into the lower back. For 17-AAG treatment groups, 17-AAG was administered 50 mg kg^-1 for 5 times per wk via ip injection after xenograft reached ~100 mm^3. For IR treatment groups, mice were anesthetized with 2% isoflurane before partial body IR and shielded with lead. Only tumors were then irradiated using the RS 2000 Xray Irradiator (Rad Source) irradiation source. A total dose of 20 or 30 Gy (10 Gy/ every 8 d) X-ray radiation was given, at a dose rate of 1.2 Gy/ min. Tumor size was measured every 4 d using calipers. Tumor volume (V) was calculated with the following formula: V = A/B^2/2, in which A is the long axis diameter and B is the short axis diameter. When tumor volume reached ~1000 mm^3 in the largest tumor group mice were sacrificed. Tumors were removed, photographed, and sectioned. Tumor proliferation was determined using Ki-67 immunohistochemistry (IHC) staining, while tumor cell apoptosis was assessed using terminal transferase-mediated dUTP nick end labeling (TUNEL) staining.

2.9 Immunohistochemistry (IHC) staining

Commercial human cervical cancer tissue arrays were purchased from Taibosi Biotechnology (Xian, China). IHC staining was performed with a Leica BOND-III, automatic IHC and ISH Stainer (Leica, Wetzlar, Germany). The following antibodies were used, including anti-HSP90α (1:50, PA3-013, ThermoFisher Scientific, Carlsbad, CA, USA), anti-HSP90β (1:50, 37-9400, ThermoFisher Scientific) and anti-Ki-67 (1:2000, 27309-1-AP, Proteintech). Protein expression scores were conducted by 2 experienced pathologists without authorship in this study, according to the criteria proposed by the Human Protein Atlas. In brief, the score is a combination and conversation of staining intensity and fraction of stained cells. Four staining scores were defined, including not detect, low, medium, and high (https://www.proteinatlas.org/about/help#4).21
2.10 | Immunofluorescence assay

The level of phosphorylated histone variant H2AX (γ-H2AX), which is a surrogate marker of DNA damage, is detected using immunofluorescence assay. In brief, CaSki-RR and SiHa-RR cells with FBOX6 overexpression alone or together with CD147 overexpression were plated onto culture slides (Corning Inc, Corning, NY, USA). When cells reached ~50% of confluence, cells were subjected to 17-AAG (20 nM/24 h for CaSki-RR and 50 nM/24 h for SiHa-RR) or control pre-treatment and then treated with 2 Gy irradiation. At 6 h later, the cells were fixed with buffered formaldehyde for 15 min and then permeabilized with 0.02% Triton. The cells were then blocked in 3% albumin in Dulbecco’s phosphate-buffered saline (DPBS) for 1 h. Then, the slides were incubated with a primary antibody against γ-H2AX (AP0687, Abclonal, Wuhan, China) in a 1:200 dilution in the 3% albumin solution overnight at 4°C. Then, the slides were thoroughly washed and incubated with goat anti-rabbit IgG (H&L) (Alexa Fluor 647) (ab150079, Abcam, Cambridge, UK) secondary antibody in a 1:500 concentration in albumin. The slides were then incubated in darkness for 1 h before washing. The slides were mounted using VECTASHIELD Antifade Mounting Medium with 4’,6-diamidino-2-phenylindole (DAPI). Then, fluorescence images were captured using an FV1000 confocal laser scanning biological microscope (Olympus, Tokyo, Japan).

2.11 | Statistical analysis

Statistical analysis was performed using GraphPad Prism 8.1.2. Quantitative data were reported as mean ± standard deviation (SD). Statistical analysis of the statistical difference between 2 groups was estimated based on the unpaired t test with Welch correction. One-way ANOVA with Tukey’s post hoc multiple-comparisons testing was performed. The difference between the survival curves was estimated by Log-rank (Mantel-Cox) test. P-values of .05 or less were determined as statistically significant. * and #, P < .05; ** and ##, P < .01; *** and ###, P < .001.

3 | RESULTS

3.1 | HSP90AA1 and HSP90AB1 upregulation was associated with poor survival of patients with cervical cancer

With RNA-seq data acquired from GTEx-fallopian tube (FT) (n = 5), GTEx-cervix (n = 13), and primary cervical cancer tissues from TCGA (n = 304), we examined the expression profile of HSP90AA1 and HSP90AB1. Results showed that HSP90AA1 expression was significantly upregulated in the tumor group compared with normal tissue groups (Figure 1A, left). In comparison, no significant alteration of HSP90AB1 mRNA expression was observed among these groups (Figure 1A, right). Then, K-M survival analysis was performed between the primary tumor cases with the top 25% and bottom 25% of HSP90AA1 and HSP90AB1 expression. Patients with the top quartile of HSP90AA1 expression had substantially shorter PFS, compared with the counterparts with the bottom quartile of gene expression (Figure 1B). However, no significant difference was observed in OS between the 2 groups (Figure 1C). High HSP90AB1 expression was associated with significantly shorter PFS and OS, compared with the counterparts with the bottom quartile of gene expression (Figure 1D,E). IHC staining showed that among 20 cases of cervical tumor tissues examined, 13/20 and 12/20 cases had medium to high level of HSP90α and HSP90β expression, respectively (Figure 1F–H).

To understand the functional role of HSP90 in cervical cancer, we conducted a GSEA assay between patients with high and low HSP90AA1 or HSP90AB1 expression in TCGA-CESC (Table S2 and S3). GSEA identified that the gene set of DNA repair was associated with higher HSP90AA1 or HSP90AB1 expression (bold front, Figure S1). As tumor cells with a strong DNA repair capacity are usually radioresistant and HSP90 inhibitor has been considered a promising radio-sensitizer, we decided to explore the underlying mechanisms. Radioresistant CaSki (CaSki-RR) and SiHa (SiHa-RR) cells were generated (Figure S2A–C). These 2 subclones had significantly elevated HSP90 expression at both mRNA and protein levels (Figure S2D,E).

3.2 | HSP90 binds to and stabilizes CD147 in radioresistant cervical cancer cells

To explore the regulatory mechanisms of HSP90 in cervical cancer, we explored their interacting proteins via sequential use of IP, SDS-PAGE, and LC-MS/MS. By cross-comparing the candidates with the potential interactors of HSP90α/β in BioGRID (https://thebiogrid.org/), we found that CD147 and FBOX6 are 2 high-potential candidates (Figure 2A). Co-IP assays confirmed the interaction among HSP90α/β, CD147, and FBOX6 in CaSki-RR (Figure 2B–D) and SiHa-RR (Figure 2E–G) cells. In CaSki-RR and SiHa-RR cells, CD147 expression was significantly upregulated at the protein level, but not the mRNA level, compared with parental cell lines (Figure S2F). As FBOX6 acts as a substrate recognition component of the Skp1-Cullin1-F-box protein (SCF) ubiquitin E3 ligase complex, we decided to explore the functional significance of the bindings in terms of CD147 protein stability. 17-AAG was applied to treat CaSki-RR and SiHa-RR cells. Interestingly, treatment with 17-AAG reduced total CD147 protein levels in a dose-dependent manner (Figure 2H,I). Then, CaSki-RR and SiHa-RR cell lysates were separated into RIPA-soluble and RIPA-insoluble urea parts. 17-AAG treatment reduced CD147 protein levels in both soluble and insoluble samples (Figure S3A) but did not alter CD147 mRNA expression (Figure S3B). Therefore, these findings implied that HSP90 interacts with and stabilizes CD147 in cervical cancer cells.
Inhibition of HSP90 sensitizes radioresistant cervical cancer cells to radiotherapy partially via CD147

Previous studies have shown that HSP90 inhibition might be a promising strategy to overcome radioresistance in multiple cancers, but the mechanisms were still not fully understood. As HSP90 can stabilize CD147 in cervical cancer cells, we explored whether CD147 expression contributed to radioresistance.

CaSki and SiHa cells were transiently infected lentivirus carrying CD147 shRNA (Figure S4A,C,D) or flag-CD147 (Figure S4B,E). Cells with CD147 knockdown had reduced colony formation when exposed to X-ray IR (Figure S4F,G). In contrast, cells with CD147 overexpression had enhanced colony formation after IR (Figure S4H,I).

Similar experiments were performed using CaSki-RR and SiHa-RR cells. These RR cells with stable CD147 knockdown (Figure 3A,C) exhibited reduced colony formation when exposed to X-ray IR (Figure 3E,F). In comparison, cells with CD147 overexpression (Figure 3B,D) exhibited enhanced colony formation after IR and were less responsive to 17-AAG (Figure 3G,H). In vivo assays showed that CD147 knockdown alone significantly reduced the growth of CaSki-RR and SiHa-RR cell-derived tumors and also enhanced IR-induced growth suppression (Figure 3I-K). CD147 overexpression reduced the radio-sensitizing effect of 17-AAG (Figure 3I-K). Immunofluorescence TUNEL staining and
IHC staining of Ki-67 revealed a higher level of TUNEL-positive cells, but a lower level of Ki-67-positive cells in CD147 knockdown and 17-AAG treatment groups compared with the IR alone group (Figure 3L). However, CD147 overexpression partly weakened the synergistic effect of 17-AAG and IR on inducing cell apoptosis (Figure 3L).

### 3.4 Inhibition of HSP90 enhances the polyubiquitination of CD147 that leads to proteasomal degradation

To understand the molecular mechanisms through which HSP90 stabilizes CD147 in radioresistant cervical cancer cells, we checked the protein levels of CD147 influenced by 17-AAG, with or without the presence of MG132. Results showed that treatment with MG132 prevented 17-AAG associated flag-CD147 degradation in both CaSKi and SiHa cells (Figure 4A,B) and reduced endogenous CD147 degradation in both CaSKi-RR and SiHa-RR cells (Figure 4C,D). Treatment with 17-AAG enhanced flag-CD147 polyubiquitination (Figure 4E,F). Treatment with MG132 led to accumulated CD147 polyubiquitination, which was further enhanced by HSP90 inhibition (Figure 4G,H). CHX chase assay confirmed that 17-AAG treatment significantly facilitated CD147 degradation in both CaSKi-RR and SiHa-RR cells (Figure 4I-L). These results suggested that CD147 was more susceptible to ubiquitin-mediated proteasomal degradation when HSP90 is functionally inhibited.
FBXO6 ubiquitates and degrades CD147

As endogenous protein interactions among FBXO6, HSP90, and CD147 were confirmed by co-IP in both CaSki-RR and SiHa-RR cells (Figure 2B-F), we then explored whether FBXO6 was involved in CD147 degradation. FBXO6 overexpression reduced the CD147 protein levels and, conversely, knockout of FBXO6 led to elevated CD147 protein levels in CaSki-RR and SiHa-RR cells with stable CD147 knockdown (A, C) or overexpression (B, D). E-H, Quantitation of clonogenic assay to examine the effects of X-ray IR (0, 2, 4, 6, and 8 Gy) on cell growth of CaSki-RR and SiHa-RR cells with CD147 knockdown (E, F) or CD147 overexpression alone or combined with 17-AAG treatment (20 nM/24 h for CaSki-RR and 50 nM/24 h for SiHa-RR) (G, H). *, Comparison between shNC and shCD147#1, and between blank and flag-CD147. #, Comparison between shNC and shCD147#2, and between 17-AAG and 17-AAG+flag-CD147. I-K, Representative images of TUNEL staining and IHC staining of Ki-67 in tumor sections from panel (I)
reduced the degradation of CD147 and canceled 17-AAG-induced CD147 degradation (Figure 5E-G). Overexpression of FBXO6-WT promoted the degradation of CD147, the effect of which was enhanced by 17-AAG treatment (Figure 5H-J). However, overexpression of FBXO6-MT (p.Y241A and p.W242A), a mutant lacking ubiquitin ligase activity,25 lessened the degradation of CD147 (Figure 5H-J). In addition, this mutant also counteracted 17-AAG-induced CD147 degradation (Figure 5H-J).

3.6 | FBXO6 degrades CD147, which consequently sensitizes radioresistant cells to IR

As we found that FBXO6 could promote CD147 degradation, we then studied whether it modulated radiosensitivity via CD147. FBXO6 overexpression significantly increased the sensitivity of CaSki-RR and SiHa-RR cells to IR (2 Gy), the effect of which was enhanced by 17-AAG treatment but was weakened by CD147 overexpression (Figure 6A-C). Moreover, western blot and immunofluorescence assays FBXO6 overexpression enhanced IR-induced expression of γH2AX and γH2AX foci (Figure 6D,E). Similarly, the effect was enhanced by 17-AAG treatment, but was weakened by CD147 overexpression (Figure 6D,E). Overall, these data indicated that FBXO6 sensitized radioresistant cervical cancer cells to IR predominantly by inducing destabilization of CD147 via proteasomal degradation. Using survival data from TCGA-CESC, we confirmed that patients with high FBXO6 mRNA expression had significantly better PFS compared with their counterparts with low FBXO6 expression (Figure 6F). However, this trend was not confirmed in OS (Figure 6G).

4 | DISCUSSION

In this study, we confirmed that HSP90 upregulation at the protein level was associated with enhanced radioresistance of cervical...
FIGURE 5 FBXO6 ubiquitates and degrades CD147. A-D, Representative images (A, C) and quantitation (B, D) of the effect of FBXO6 overexpression (A, B) or knockdown (C, D) on CD147 levels. E-G, CHX was chase assay performed to compare CD147 protein stability in CaSki-RR and SiHa-RR cells after FBXO6 knockdown, with or without 17-AAG treatment. Cells were infected for transient FBXO6 knockdown. 48 h later, cells were treated with 17-AAG (100 nM for CaSki-RR and or 250 nM for SiHa-RR) or blank control for 4 h, followed by 5 μg/mL CHX treatment for the indicated time. H-J, CHX was chase assay performed to compare CD147 protein stability in CaSki-RR and SiHa-RR cells with FBXO6 overexpression (FBXO6-WT or FBXO6-MT) alone or in combination with 17-AAG treatment. Cells were first infected for transient FBXO6 overexpression. At 48 h later, cells were treated with 17-AAG (100 nM for CaSki-RR and or 250 nM for SiHa-RR) or blank control for 4 h, followed by 5 μg/mL CHX treatment for the indicated time. CD147 amount was determined by density analysis (F, G, I, J) of western blotting images (E, H). Data are representative of 3 independent biological experiments. Data were reported as the mean ± SD from 3 technical replicates.
SONG et al. cancer. Inhibiting HSP90 might be a promising strategy to conquer radioresistance in multiple cancers.\textsuperscript{13,26,27} Functionally, HSP90 can enhance the stability and function of conformationally labile proteins after primary folding, thereby modulating the functions of the binding proteins.\textsuperscript{8} Via inhibiting HSP90, its binding proteins might have accelerated degradation. For example, WT-EGFR is a client protein of HSP90 in cancer cells. Treatment with HSP90 inhibitors geldanamycin and AT13387 facilitated the degradation of WT-EGFR, thereby sensitizing EGFR-dependent cancer cells to chemotherapy and radiotherapy.\textsuperscript{28} Ganetespib, a selective inhibitor of HSP90, can act as a radio-sensitizer by reducing IR-induced HIF-1\textalpha upregulation in pancreatic cancer cells.\textsuperscript{13} Furthermore, 17\textsuperscript{-}AAG can reduce the binding between Hsp90\textbeta and its client protein MAST1 and promote proteasomal degradation of Hsp90\textbeta, thereby enhancing cisplatin

\textbf{FIGURE 6} FBXO6 degrades CD147, which consequently sensitizes radioresistant cells to IR. A-C, Representative images (A) and quantitation (B, C) of clonogenic assay to examine the effects of X-ray IR (2 Gy) on the growth of CaSki-RR with SiHa-RR cells with transient \textit{FBXO6} overexpression alone or in combination with 17\textsuperscript{-}AAG pre-treatment (20 nM/24 h for CaSki-RR and 50 nM/24 h for SiHa-RR) or \textit{CD147} overexpression. D, Western blotting assay (D) of FBXO6, γ-H2AX and CD147 expression in cells treated as in panel (A), 10 h after X-ray IR (2 Gy). E, Immunofluorescence assay of γ-H2AX foci formation in cells treated as in panel (A), 6 h after X-ray IR (2 Gy). F, G, K-M, survival analysis was performed to compare the difference in PFS (F) and OS (G) between primary cervical cancer cases with the top 25% and bottom 25% of \textit{FBXO6} mRNA expression. Data are representative of 3 independent biological experiments. Data are reported as the mean ± SD from 3 technical replicates.
sensitivity in human cancers. These mechanisms help explain why HSP90 inhibitors show antitumor activity.

Generally, HSP90 forms a clamp structure after homodimerization, anchored by dimerization of the C terminals. Then, the N-terminal domains act as a jaw, which is responsible for ATP hydrolysis and co-chaperones binding. Co-chaperones (such as CDC37, FKBP51, and TAHI) may help to recruit certain client proteins to HSP90 and contribute to regulating HSP90 conformational switch by inhibiting or enhancing ATP hydrolysis. These structural features determined that HSP90 has different client proteins under different conditions. By performing IP and LC-MS/MS in combination, we found that HSP90 interacted with CD147 and FBXO6. CD147 (also known as EMMPRIN or Basigin) is a ubiquitously expressed glycoprotein that is commonly overexpressed in a series of tumors and participates in cancer development. The N-terminal domain of CD147 has 3 glycosylation sites. Therefore, CD147 has non-glycosylated, low-glycosylated, and high-glycosylated forms, ranging from 27 kDa to ~65 kDa. High-glycosylated CD147 (ranging from 46 to 65 kDa) is the mature and active form on the transmembrane, with a long-term half-life. Previous studies confirmed that CD147 upregulation contributed to enhanced cell proliferation and reduced the chemosensitivity and radiosensitivity of cervical cancer. In this study, we confirmed that 17-AAG treatment decreased CD147 expression, but had little influence on FBXO6 expression. In addition, 17-AAG-induced radio-sensitizing effects were partly abrogated by CD147 overexpression both in vitro and in vivo. These findings suggested that HSP90 inhibitor 17-AAG exerted radio-sensitizing effects partly by reducing CD147 protein levels in cervical cancer cells.

FBXO6 is a critical substrate adapter of the evolutionarily SCF complex that mediates the ubiquitination of glycoproteins. Its F-box associated (FBA) domains are required for glycoprotein recognition. By performing CHX chase assays, we confirmed that FBXO6 could mediate CD147 degradation via the proteasomal pathway, the effect of which was enhanced by 17-AAG treatment. Therefore, we inferred that HSP90 can enhance CD147 stability by reducing FBXO6-mediated proteasomal degradation. The role of FBXO6 is paradoxical in cancers. In non-small-cell lung cancer, it decreases expression and phosphorylation of Chk1, thereby increasing the sensitivity of the cancer cells to cisplatin-based chemotherapy. However, FBXO6 promoted ovarian cancer cell proliferation, migration, and invasion partially by inducing RNAseT2 degradation. Therefore, FBXO6 might have cancer-specific functions, depending on its downstream substrates. In the current study, we further explored the effect of FBXO6 on the radiosensitivity of radioresistant cervical cancer cells. Results showed that enforced FBXO6 overexpression sensitized CaSk-R and SiHa-RR cells to IR. These effects were enhanced by 17-AAG treatment, but were weakened by CD147 overexpression. Survival analysis further confirmed the association between high FBXO6 expression and favorable PFS among patients with cervical cancer.

Although FBXO6 was identified as a functional E3 ubiquitin ligase facilitating CD147 degradation, the specific binding site and the type of ubiquitination chain remain unexplored. Traditionally, FBXO6 acts as a substrate recognition component of the SCF component. One recent study found that it interacted with the IAD domain of IFN-regulatory factor 3 (IRF3) through its FBA domain to induce ubiquitination and degradation without the involvement of SCE. Considering the promising radio-sensitizing effects of FBXO6 overexpression and CD147 inhibition in cervical cancer, it is meaningful to explore the detailed mechanisms of FBXO6-mediated CD147 degradation in the future.

In conclusion, this study found that HSP90 (both HSP90α and HSP90β) bound to FBXO6 and CD147 and reduced FBXO6-mediated CD147 polyubiquitination through the proteasomal pathway, thereby promoting radioresistance of cervical cancer cells. These mechanisms further explain why the HSP90 inhibitor exerts radio-sensitizing effects.

ACKNOWLEDGMENTS

None.

CONFLICTS OF INTEREST

None of the authors have conflicts of interest to declare.

DATA AVAILABILITY STATEMENT

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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**SUPPORTING INFORMATION**

Additional supporting information may be found in the online version of the article at the publisher’s website.

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