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Structural characterization of a sulfated polysaccharide from *Gracilariopsis lemaneiformis* and its potentiation of cisplatin antitumor activity in Colon-26 carcinoma tumor-bearing mice by inducing ferroptosis

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Ferroptosis, a form of regulated cell death caused by iron-mediated lipid peroxidation, has become a potential strategy to overcome drug resistance and improve the efficacy of traditional cancer treatments. In this study, we investigated whether treatment with the combination of *Gracilariopsis lemaneiformis* polysaccharides and cisplatin (CP) potentiated the antitumor activity in a Colon-26 carcinoma tumor-bearing mouse model by ferroptosis activation. The *G. lemaneiformis* polysaccharide GP90 was mainly composed of (1→3) linked 4-O-sulfate-β-D-galactose and (1→4) linked 3,6-anhydro-α-L-galactose with a molecular weight of 12.45 kDa. Compared with the CP group, the combination of GP90 and CP significantly suppressed tumor growth. Based on the transcriptomic and metabolomic analyses of tumor tissue, GP90 enhanced the antitumor effect of CP by promoting ferroptosis and regulating ferroptosis-related metabolic pathways. Moreover, the accumulation of 4-hydroxy-2-nonenal (4-HNE) and down-regulation of the expression of solute carrier family 7 member 11 (SLC7A11) and glutathione peroxidase 4 (Gpx4) were verified by immunohistochemistry staining. Finally, gene set enrichment analysis showed that positive immunoregulatory pathways were significantly enriched in the GP90 and CP combination group. Our results indicate that GP90 potentiates chemotherapy sensitivity by targeting the transferrin receptor and SLC7A11/Gpx4 pathway to induce ferroptosis, which might be a useful therapeutic target in colorectal cancer patients.

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1. Introduction

Colorectal cancer (CRC) is the third most common malignancy and the second leading cause of cancer-related death,¹ with nearly 1.93 million new cases and 0.94 million deaths in 2020 worldwide.² Cisplatin (CP) is one of the most effective cytotoxic agents in CRC chemotherapy; however, its therapeutic efficacy is limited by chemoresistance and detrimental side effects.³ Hence, there remains a need for effective therapeutic

approaches to improve the CP sensitivity in CRC. Recent scientific evidence has shown that CP induces ferroptosis and apoptosis in CRC cells and that the combination of CP and the ferroptosis inducer erastin is effective in enhancing the antitumor activity of drugs.⁴ Ferroptosis, a new mode of cell death caused by iron-dependence and lipid peroxidation, plays a beneficial role in improving cancer treatment, tumor chemoresistance, and antitumor immunotherapy. Ferroptosis agonists can promote the accumulation of intracellular reactive oxygen species (ROS), reduce the transferrin content, or inhibit cystine/glutamate antiporter System Xc-mediated cystine transport, which leads to ferroptosis in drug-resistant tumor cells.⁵ In addition, ferroptosis may reshape the tumor ecology and stimulate the immune microenvironment, thereby inhibiting tumor growth and progression.⁶ Thus, strategies to induce ferroptosis have been proposed to improve the outcomes of current cancer therapy.

Gracilariopsis lemaneiformis is an edible red alga of economic importance that is not only directly consumed but can also be processed into dietary supplements, cosmetics, and

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pharmaceuticals, among others. *G. lemaneiformis* contains high levels of polysaccharides, phycobiliproteins, pigments, and other nutritional components, whereas polysaccharides are its main active components.⁷ *G. lemaneiformis* polysaccharides (GPs) exert potential antitumor activity without obvious toxicity. Shi *et al.* reported that the three polysaccharides extracted from *G. lemaneiformis* inhibited the proliferation of MCF-7, HepG-2, and HeLa cells *in vitro*.⁸ Additionally, *in vivo* antitumor tests showed that soluble GPs significantly inhibited mouse sarcoma S180 tumor growth and improved the activity of splenic lymphocytes and natural killer cells.⁹ Moreover, GPs alleviated dextran sulfate sodium-induced colitis in mice by remodelling the gut microbiota and intestinal metabolites, which reduces the risk of CRC.¹⁰ Thus, GPs not only directly inhibit tumor growth, but also interfere with tumor progression and can be used for tumor prevention and adjuvant therapy. Du *et al.* found that *Lycium barbarum* polysaccharide effectively prevented the proliferation and promoted the ferroptosis of breast cancer cells, mainly by the System Xc- and glutathione peroxidase 4 (Gpx4) pathway.¹¹ Studies have also shown that *Pinus massoniana* pollen polysaccharide treated colon cells exhibit electron-lucent cytoplasm and nuclei as well as cytoplasmic vacuolization, which are characteristic of apoptosis caused by ferroptosis.¹² The question remains whether the auxiliary antitumor activity of GPs is related to the induction of ferroptosis.

In this study, we investigated the synergistic antitumor effect of GPs in combination with CP and determined whether the underlying mechanism is related to ferroptosis induction.

2. Materials and methods

2.1. Materials

Dried *G. lemaneiformis* was provided by Qingdao Haiyouyou Trading Co., Ltd (Shandong, China), the origin being Qingdao. Colon-26 (C26) colon carcinoma cell lines were obtained from the American Type Culture Collection (ATCC, Manassas, VA, USA). Dulbecco's modified Eagle medium (DMEM), fetal bovine serum (FBS), penicillin-streptomycin antibiotics, and phosphate-buffered saline (PBS) were purchased from Invitrogen-Gibco (Grand Island, NY, USA). All chemical reagents were of analytical grade.

2.2. Preparation of GP90

Dried *G. lemaneiformis* powder was obtained by crushing and passing through an 80-mesh screen. Then the powder was washed three times with anhydrous ethanol, and soaked overnight in anhydrous ethanol at 4 °C. After filtration, the filtered residue was dried at 70 °C for 2 h and then extracted with 35 times the volume of 0.05 M citric acid solution (v/w, mL g⁻¹) at 100 °C. After the reaction for 2 h, the supernatant was collected by centrifugation at 10 000 rpm for 20 min, and the pH was adjusted to 7.0 with 1 M NaOH solution and concentrated. The concentrate was recovered, and anhydrous ethanol was added until the concentration of the final system alcohol was

30% for precipitating the fraction, followed by a continued regulation of the supernatant with anhydrous ethanol to final alcohol concentrations of 60% and 90% to obtain GP90. The precipitated GP90 was solubilized with distilled water, deproteinized by the Sevag method (chloroform : butanol = 5 : 1, v/v), concentrated, dialyzed, and lyophilized.

2.3. Structural characterization of GP90

The homogeneity and molecular weight were measured with an Agilent 1260 Infinity size exclusion chromatography and gel permeation chromatography system equipped with an Agilent RID-7115A refractive index detector (Agilent Technologies, Palo Alto, CA, USA) and a gel-filtration chromatography column (TSK-gel G3000PWXL column, 7.8 × 300 mm i.d., 7 µm; Tosoh Bioscience LLC, Tokyo, Japan). Dextran standards (1, 5, 12, 50, 80 and 270 kDa) were used for calibration. Elution was performed with 0.02 M ammonium acetate solution at a rate of 0.6 mL min⁻¹ and maintained at 35 ± 1 °C.

According to a previously reported high-performance liquid chromatography (HPLC) method,¹³ freeze-dried GP90 (5 mg) was hydrolysed with 4 M trifluoroacetic acid (TFA, 1 mL) for 4 h at 120 °C, and excess TFA was removed by rotary evaporation with methanol. Subsequently, the products or monosaccharide standards were derivatized with 3-methyl-1-phenyl-2-pyrazolin-5-one (PMP, 0.5 M) at 70 °C for 1.5 h. Then the reaction mixture was neutralized and excess PMP was removed using chloroform. The monosaccharide was analyzed using HPLC (Agilent-1260; Agilent Technologies) equipped with a YMC-Pack ODS-AQ column (250 × 4.6 mm, I.D. S-5 µm, 12 nm) and eluted with 50 mM ammonium acetate solution and acetonitrile at a ratio of 73 : 17 (v/v).

Fourier transform infrared spectroscopy (FTIR) for GP90 was carried out using 400 FT-IR spectrometers (Shimadzu, Japan) at wavelengths ranging from 4000 to 400 cm⁻¹.

GP was prepared with D₂O to a concentration of 30 mg mL⁻¹. The nuclear magnetic resonance (NMR) spectra of GP90 were analyzed using a 700 MHz NMR spectrometer (Bruker Daltonik GmbH, Leipzig, Germany), and the 1D- and 2D-NMR spectra were recorded.

2.4. Animal experimental design

BALB/c mice (8 weeks old, 20–24 g, male) were obtained from SiPeiFu (Beijing Biotechnology Co., Ltd, Beijing, China) and housed in a room under a 12 h light-dark cycle and maintained at 22 ± 2 °C. Animal experiments were performed following the guidelines of the Animal Ethics Committee of Guangdong Medical University (GDY1902062).

C26 colorectal models were prepared as described by Zhang *et al.*¹⁴ Briefly, C26 colon cells were cultured in DMEM containing 10% FBS, 1% glutamine, 1 mM sodium pyruvate, and 1% streptomycin/penicillin at 37 °C and 5% CO₂. Then the cells were collected in a 0.25% trypsin-EDTA solution and resuspended in sterile PBS before subcutaneous injection into mice (1 × 10⁶ cells per mouse). On day 1 after the C26 cell injection, all mice were randomly divided into four groups: model, GP90, CP and CP + GP90.¹⁵ Mice in the GP90 and CP + GP90 groups



were gavaged with 200 mg kg⁻¹ GP90 daily, whereas mice in the other two groups were gavaged with 0.2 mL sterile saline alone daily. In CP and CP + GP90 groups, the mice were treated with a combination of GP90 and intraperitoneal CP chemotherapy (3 mg kg⁻¹) starting from day 1 and every 4 days thereafter. During the study, the tumor volume was measured every 3 days based on the formula: $V = 0.5x^2y$, where x is the shortest and y is the largest superficial diameter. On day 16 after treatment, the mice in all groups were sacrificed by cervical dislocation. Spleen and tumor tissues were collected and stored at -80 °C for subsequent analyses.

2.5. RNA isolation, sequencing, and bioinformatics analysis

Total RNA from tumor samples was extracted using the Trizol reagent (Invitrogen, Carlsbad, CA, USA) based on the manufacturer's protocol, and the RNA quality was assessed using RNase-free agarose gel electrophoresis using an Agilent 2100 Bioanalyzer (Agilent Technologies). After the extraction of total RNA, eukaryotic mRNA was enriched using oligo (dT) beads, while rRNA was removed using the Ribo-Zero™ Magnetic Kit (Epicentre, Madison, WI, USA). Then the enriched mRNA was fragmented into short fragments with fragmentation buffer and reverse transcribed into cDNA using random primers. The cDNA fragments were purified using the QiaQuick PCR extraction Kit (Qiagen, Venlo, the Netherlands), end-repaired, polyadenylated, and ligated to Illumina sequencing adapters. The ligation products were separated by size by agarose gel electrophoresis, amplified by PCR, and sequenced using the Illumina HiSeq2500 (Gene Denovo Biotechnology Co., Guangzhou, China). The RNA-sequence data reported in this study was archived in the NCBI SRA database with the accession number PRJNA945228 and PRJNA945586.

The differential expression analysis of different RNAs in each group was performed using the DESeq2 software. The genes with a false discovery rate parameter below 0.05 and an absolute fold change ≥ 2 were considered differentially expressed genes (DEGs). All DEGs were mapped to Gene Ontology (GO) terms in the GO database (<https://www.geneontology.org/>), and then the number of genes per term was calculated. The GO terms that were significantly enriched in DEGs were defined using the hypergeometric test. The Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway enrichment analysis identified metabolic pathways or signal transduction pathways that were significantly enriched in DEGs compared to the genome-wide background. The calculated p -value was corrected for the false discovery rate (FDR) with an FDR of ≤ 0.05 as the threshold. GO terms and KEGG pathways that met this condition were defined as significantly enriched in DEGs.

2.6. Metabolite analysis

Tumor tissues (100 mg) were grounded with liquid nitrogen and resuspended in 500 μ L prechilled 80% methanol by vortexing. The homogenate was incubated on ice for 5 min and then centrifuged at 15 000g, 4 °C for 20 min. The supernatant was taken and diluted with LC-mass spectrometry (MS) grade water

to a final methanol concentration of 53%, and then centrifuged at 15 000g, 4 °C for 20 min. Finally, the supernatant was injected into an LC-tandem MS (MS/MS) system for analysis.

Ultra-performance LC (UPLC)-MS/MS analysis was performed using the Vanquish UPLC system (Thermo Fisher Scientific, Dreieich, Germany) coupled with the Orbitrap Q Exactive™ HF-X mass spectrometer (Thermo Fisher Scientific) at Gene Denovo Co., Ltd. Samples were injected into a Hypesil gold column (100 \times 2.1 mm, 1.9 μ m) at a flow rate of 0.2 mL min⁻¹. The eluents for the positive polarity mode were eluent A (0.1% formic acid in water) and eluent B (methanol), and the eluents for the negative polarity mode were eluent A (5 mM ammonium acetate, pH 9.0) and eluent B (methanol). The elution parameters were set as follows: 2% B, 1.5 min; 2–100% B, 12.0 min; 100% B, 14.0 min; 100–2% B, 14.1 min; and 2% B, 17.0 min. A Q Exactive™ HF-X mass spectrometer was operated using a spray voltage of 3.2 kV, a capillary temperature of 320 °C, a sheath gas flow rate of 40 arb, and an aux gas flow rate of 10 arb.

Raw data files generated by UPLC-MS/MS were processed using the Compound Discoverer 3.1 (Thermo Fisher Scientific) for peak alignment, peak extraction, and quantitation for each metabolite. Then peak intensities were normalized to the total spectral intensity and matched against mzCloud (<https://www.mzcloud.org/>), mz Vault, and the Mass List database to obtain accurate qualitative and relative quantitative results. Statistical analyses were performed using Python (Python 2.7.6 version), R (R version R-3.4.3), and the CentOS (CentOS release 6.6) statistical software.

Orthogonal projections to latent structure-discriminant analysis (OPLS-DA) were performed in metaX. Metabolites with variable importance in projection values >1 and a p -value of the t -test <0.05 were considered to be differentially expressed metabolites (DEMs). The functions of DEMs and metabolic pathways were analyzed using the KEGG database. The P -Value <0.05 was considered statistically significant.

2.7. Quantitative PCR

The gene expression was validated based on the transcriptomic data using quantitative PCR (qPCR). Total RNA was prepared using the Trizol reagent. HiScript II Q RT SuperMix was used for the qPCR of complementary DNA. Quantitative PCR was performed with the AceQ qPCR SYBR Green Master Mix (Vazyme, Nanjing, China) on the TL988 device (TianLong Group, Guangdong, China). The amplifying conditions for cDNA were as follows: denaturation at 95 °C for 90 s, 40 cycles of 95 °C for 5 s, and 60 °C for 15 s, followed by 72 °C for 20 s. The 2^{- $\Delta\Delta$ Ct} method was applied to calculate changes in the mRNA expression levels of candidate genes. The primer sequences are listed in Table 1.

2.8. Immunohistochemistry staining

Tumor tissues were weighed, fixed in 10% formalin, and embedded in paraffin. Briefly, after deparaffinization and blocking in 3% hydrogen peroxide, the sections were treated with citric sodium buffer (1 M, pH 6.0) and then treated by



Table 1 Sequences of qPCR primers

Gene	Forward sequence	Reverse sequence
ApoE	CTGAACCGCTTCTGGGATTA	GTTGCGTAGATCCTCCATGT
Tfrc	CAGTCATCAGGGTTGCCTAATA	CTGGGCTCCTACTACACATAAC
Bnip3	TCCAGCCTCCGTCTCTATTT	CGACTTGACCAATCCCATATCC
β -Actin	CTGAGTCTCCCTTGGATCTTTG	AGGCAGGTGAACTGTATG

microwave for antigen unmasking. Sections were blocked and then incubated with anti-4-hydroxy-2-nonenal (4-HNE, 1:1600 dilution; Bioss Antibodies, Massachusetts, USA), anti-solute carrier family 7 member 11 (SLC7A11, 1:1600 dilution; Cell Signaling Technology [CST], Shanghai, China), and anti-GPX4 (1:1600 dilution; CST) primary antibodies at 4 °C overnight. The next day, sections were incubated with horseradish peroxidase labelled secondary antibody (1:5000 dilution; CST) for 1 h at room temperature and then rinsed three times with PBS. The sections were coloured with DAB, counterstained with hematoxylin, and dehydrated. All images were viewed using the FilterMax F5 fluorescence microplate reader (Molecular Devices, Silicon Valley, CA, USA).

2.9. Statistical analyses

All data are expressed as the mean \pm standard deviation (SD). Differences were analyzed by the one-way analysis of variance using the IBM SPSS 22.0 software. $P < 0.05$ was considered statistically significant.

3. Results

3.1. Structural characterization of GP90

The contents of carbohydrates and sulfate in GP90 were $88.29 \pm 8.24\%$ and $22.93 \pm 0.94\%$, respectively. A single and sharp peak was observed in the chromatogram of GP90 (Fig. 1A) with an average molecular weight of 12.45 kDa. GP90 was mainly composed of galactose (97.95%; Fig. 1B), which indicated that GP90 was a homogeneous polysaccharide. As shown in Fig. 1C, the FTIR spectrum of GP90 showed typical characteristic peaks of polysaccharides. The broad absorption at 3402.43 cm^{-1} and sharp absorption at 2954.95 cm^{-1} were O–H and C–H stretching vibrations, respectively. The strong absorption peak at 1220.94 cm^{-1} and the sharp absorption peak at 848.68 cm^{-1} corresponded to the asymmetric stretching vibrations of S=O and C–O–S, respectively, which indicates the presence of sulfate.¹⁶ Peaks between 1000 and 1200 cm^{-1} may have the presence of C–O–H and C–O–C of galactose.¹⁷ The peaks at 925.83 and 848.68 cm^{-1} indicated the presence of 3,6-anhydro-galactopyranose and axial sulfate ester at C4 of D-galactopyranose (G4S), respectively.¹⁸ The low-intensity peak at 802.39 cm^{-1} was thought to be the sulfate group at C-2 of 3,6-anhydro-galactopyranose.¹⁹ The results revealed that GP90 was composed of galactose and anhydro-galactose.

The 1D NMR (^1H , ^{13}C , and DEPT-135) and 2D NMR (heteronuclear singular quantum correlation spectroscopy [HSQC],

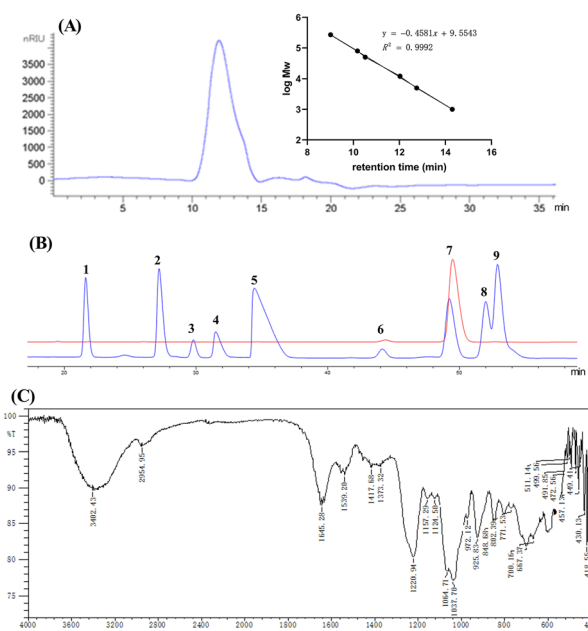


Fig. 1 Molecular weight distribution (A), monosaccharide composition (B), and FT-IR spectra of GP90 (C). The HPLC chromatograms of monosaccharide standards (blue line) and GP90 (red line). Peaks 1 = Man, 2 = Rib, 3 = Rha, 4 = GlcA, 5 = GalA, 6 = Glc, 7 = Gal, 8 = Xyl and 9 = Ara.

heteronuclear multiple correlation spectroscopy [HMBC], and correlation spectroscopy [COSY]) were performed to analyze the structural configurations of GP90. As shown in the ^1H NMR spectrum of GP90 (Fig. 2A), a typical proton of the anomeric carbon of 3,6-anhydrogalactose (DA) at δ 5.02 and of galactose-4-sulfate (G4S) at δ 4.54 are characteristic of κ -carrageenan.^{17,19} The ratio of G4S/DA was 2.34, which was calculated based on the H-4 proton of G4S and the H-1 signal of DA.²⁰ The signals at δ 102.27 and δ 94.34 (Fig. 2B) were ascribed to C-1 of G4S and DA, respectively.²¹ From the DEPT-135 spectrum (Fig. 2C), the intense inverted peak at δ 60.73 was the C-6 of D4S, whereas the other inverted peak at δ 69.40 was the C-6 of DA. The assignment of GP90 was analyzed with 2D NMR. The ^1H and ^{13}C chemical shifts of GP90 are summarized in Table 2. The HSQC spectrum depicts the correlations of anomeric carbons of G4S with protons at δ 102.27/4.54 and δ 94.34/5.02 for DA (Fig. 2D). From the COSY spectrum of GP90 (Fig. 2E), the coupling correlation between H-1 (δ 4.54) and H-2 (δ 3.42), H-3 (δ 3.73) and H-4 (δ 4.59), and H-5 (δ 3.72) and H-6 (δ 3.71) of G4S were found. For DA, H-1/H-2 was at δ 5.02/4.05, H-2/H-3 was at δ 4.05/4.43, and H-5/H-6 was



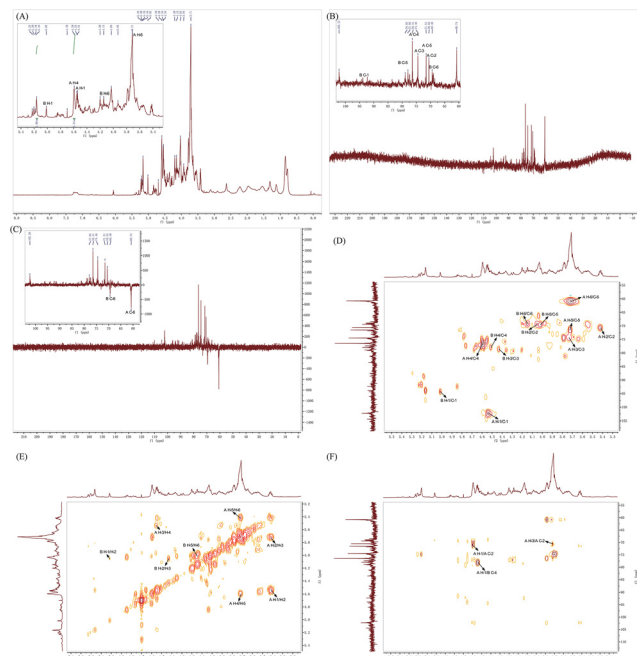


Fig. 2 The structural characterization of GP90 (A) ^1H NMR spectrum, (B) ^{13}C NMR spectrum, (C) DEPT135 spectrum, (D) HSQC spectrum, (E) COSY spectrum and (F) HMBC spectrum. DEPT: distortionless enhancement by polarization transfer; COSY: homonuclear chemical shift correlation spectroscopy; HSQC: heteronuclear singular quantum correlation spectroscopy; and HMBC: heteronuclear multiple correlation spectroscopy.

Table 2 ^1H and ^{13}C NMR chemical shifts of GP90

Sugar residues	$^1\text{H}/^{13}\text{C}$ chemical shift (ppm)					
A (1→3) linked 4-O-sulfate-β-D-galactose	4.54	3.42	3.73	4.59	3.72	3.71
B (1→4) linked 3,6-anhydro-α-L-galactose	102.27	70.55	74.45	76.41	71.51	60.73
	5.02	4.05	4.43	4.51	4.02	4.15
	94.34	69.54	78.67	77.97	69.54	69.40

at 4.02/4.15. The linkage sites of G4S and DA in GP90 were further confirmed from the HMBC NMR spectrum shown in Fig. 2F. The cross-peak at δ 4.54/77.91 (A H-1/B C-4) revealed that the O-1 of residue A was linked to the C-4 of residue B. The results of NMR spectroscopy also confirmed the presence of (1→3) linked 4-O-sulfate-β-D-galactose and (1→4) linked 3,6-anhydro-α-L-galactose. This similar structure of sulfated galactans has been found in other red algae such as *Gracilari caudate*, *Gracilariopsis persica*, *Ahnfeltiopsis flabelliformis*, and *Acanthophora spicifera*.^{22–25}

3.2. The antitumor effect of GP90

The C26 tumor-bearing mice were treated with GP90 for 15 days and were sacrificed on day 16. Compared with the model group, tumor growth was significantly inhibited in GP90-treated mice starting from day 10 after the tumor challenge (Fig. 3A). The average tumor weight in the GP90 group was decreased by 24.01% compared to that in the model group

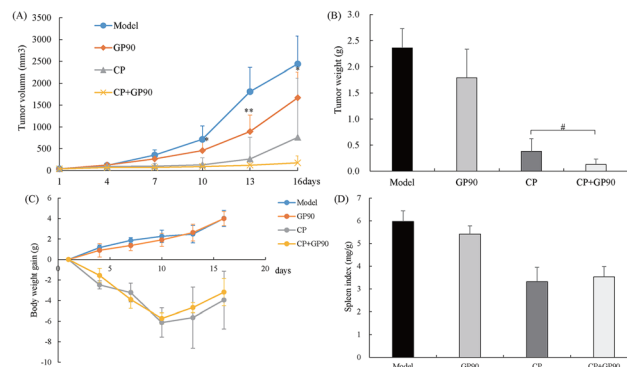


Fig. 3 GP90 exerted antitumor efficacy in the C26 carcinoma tumor-bearing mice. (A) The tumor growth curve was reported using the tumor volume one time per 3 days. (B) The average tumor weight for each group on the last day (i.e., day 16). (C) The average weight gain of mice every three days. Results are expressed as the mean \pm SD ($n = 6$). * $p < 0.05$, ** $p < 0.01$, the model group vs. the GP90 group; # $p < 0.05$, the CP group vs. the CP + GP90 group.

(Fig. 3B). In addition, the tumor growth of CP-treated mice was obviously inhibited, and the tumor inhibition rate was significantly increased to 94.55% when combined with GP90 (Fig. 3B). During the experiment, GP90 treatment had no significant effect on the mouse body weight and spleen index (Fig. 3C and 3D), suggesting that GP90 inhibited tumor growth without exerting general toxicity to the host. These results indicate that GP90 exerts potential antitumor effects and increases the sensitivity of tumor cells to CP.

3.3. Antitumor effect of the GP90 and CP combination treatment is associated with ferroptosis

To elucidate the potential mechanism of the combined antitumor activity of GP90 and CP, the transcriptome and metabolome were used to analyze the changes in the tumor tissue. A total of 22 322 genes were annotated in the tumor tissue from the CP + GP90 group, of which there were 13 DEGs (five up-regulated and eight downregulated) compared with the CP group (Fig. 4A). GO enrichment results showed that the DEGs were significantly enriched in biological processes such as localization, immune system response, and signalling; molecular functions such as binding, catalytic activity, and transporter activity; and significant enrichment in cellular components such as membrane, protein-containing complexes and the extracellular region (Fig. 4B). KEGG pathway terms enriched by DEGs were mainly involved in nitrogen metabolism, ferroptosis, and cholesterol metabolism (Fig. 4C). qPCR analysis of ferroptosis-related genes apolipoprotein (ApoE), transferrin receptor (Tfrc) and B-cell lymphoma 2-interacting protein 3 (Bnip3) in tumor tissues was performed to validate the RNA-seq results. Compared with the CP group, the relative mRNA expression of ApoE and Tfrc was up-regulated, whereas the relative mRNA expression of Bnip3 was down-regulated in the CP + GP90 group (Fig. 4D). Considering the DEGs and KEGG pathways involved in ferroptosis, we speculated that GP90



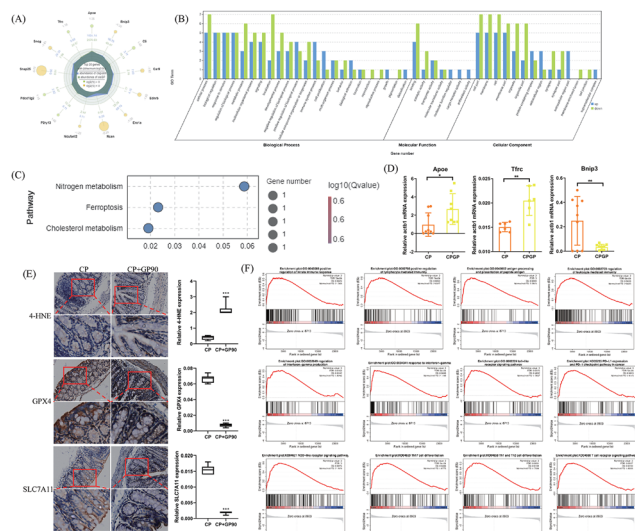


Fig. 4 Transcriptomic analysis of the effect of GP90 on enhancing the CP sensitivity in the colon cancer tissue. (A) The radar chart of DEGs in CP vs. the CP + GP90 group. (B) Gene ontology analysis. (C) The most enriched KEGG pathways. (D) Quantification of ferroptosis-related genes by qPCR in the tumor tissue. (E) Immunohistochemistry staining for 4-HNE, SLC7A11, and Gpx4 in the tumor tissue. The original magnification of H & E is at 400 \times , and the width of the bar corresponds to 100 μ m. (F) The gene set enrichment analysis (GSEA) of tumor tissue transcriptome data exhibited that C26 mice treated with the CP + GP90 combination therapy were significantly enriched in immune-related signalling pathways compared to mice treated with CP. The data are shown from at least four independent experiments. Results were accepted at $|NES| > 1$, $NOM\ p < 0.05$, $FDR\ q < 0.25$. Data are expressed as the mean \pm SD; * $p < 0.05$, ** $p < 0.01$, and *** $p < 0.001$ compared with the CP group.

could enhance the CP treatment sensitivity by inducing ferroptosis in tumor cells. The protein expression of ferroptosis-associated biomarkers including 4-HNE, Gpx4 and SLC7A11 in tumor tissues was identified by immunohistochemistry staining. Compared to the CP group, the relative expression of 4-HNE was clearly increased in the CP + GP90 group, whereas the relative expression of Gpx4 and SLC7A11 was sharply decreased (Fig. 4E). The gene set enrichment analysis (GSEA) results demonstrated various enriched immune- and tumor-related pathways in the GP + GP90 pathways, such as the positive regulation of the innate immune response, positive regulation of lymphocyte mediated immunity, antigen processing and presentation of peptide antigen, positive regulation of interferon- γ (IFN- γ) production, response to IFN- γ , Toll-like receptor signalling pathway, programmed death-ligand 1 (PD-L1) expression and programmed cell death protein 1 (PD-1) checkpoint pathway in cancer, nucleotide oligomerization domain (NOD)-like receptor signalling pathway, T helper 17 (Th17) cell differentiation, Th1 and Th2 cell differentiation and T cell receptor signalling (Fig. 4F).

The OPLS-DA plots for the CP and CP + GP90 groups show clustering in the respective regions (Fig. 5A). Compared with the CP group, there were 43 DAMs induced by the CP + GP90 group, and the volcano data show the distribution of the

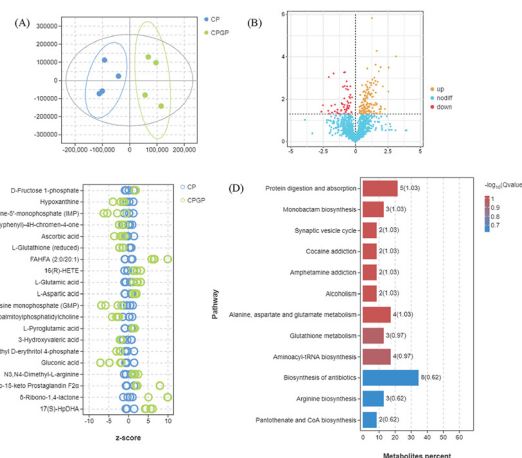


Fig. 5 GP90 combined with cisplatin altered metabolic profiling in tumor tissue. (A) A volcano plot of differentially accumulated metabolites (DAMs), (B) the OPLS-DA score plot, (C) the Z-score plot of top 20 DAMs, and (D) the KEGG enrichment bar graph in the tumor tissue between the CP and CP + GP90 groups. Data are shown from four independent experiments.

DEGs (Fig. 5B). Among the 43 DAMs, 27 genes were upregulated, and 16 genes were downregulated. These DAMs were mainly L-glutathione (reduced), L-glutamic acid, ascorbic acid, 17(S)-HpDHA, and 16(R)-HETE (Fig. 5C), which induced ferroptosis in tumor cells. The KEGG enrichment and pathway analysis showed that after GP90 treatment, the glutamate metabolism, glutathione metabolism, pantothenate and coenzyme A (CoA) biosynthesis were mainly regulated, which may be related to induced-ferroptosis of GP90 (Fig. 5D).

4. Discussion

CRC remains one of the most common malignancies worldwide. Among various chemotherapy drugs, CP is one of the most effective and widely used anticancer drugs for CRC. CRC initially responds to CP, followed by tumor recurrence after further treatment due to resistance, but does not affect CP-induced ferroptosis. Ferroptosis is a potentially novel strategy to address the problem of CP resistance. Studies have shown that CP combined with a ferroptosis inducer has a cumulative effect on its antitumor activity.²⁶ Polysaccharides have garnered great research interest in recent years because of their anticancer activity or can improve the efficacy of conventional chemotherapeutic drugs with fewer side effects.²⁷ The anticancer activity of some polysaccharides is related to the regulation of ferroptosis. Zhai *et al.* demonstrated that red ginseng polysaccharide induces ferroptosis effects on breast and lung cancer cells by targeting Gpx4.²⁸ Du *et al.* found that the anticancer activity of *L. barbarum* polysaccharide is related to the induction of ferroptosis in breast cancer cells.¹¹

GPs induce transcriptional alternations and modulate lung cancer cell viability, morphology, and apoptosis.²⁹ In our study, it was found that GP90 can significantly decrease the



colon tumor volume after 13 days of the animal experiment. Previous studies have shown that the higher antitumor activities of polysaccharides may be related to their sulfate groups, which can selectively act as ligands on the cancer cell lines, forming molecular interactions that induce apoptosis.^{30,31} GP90 has a relatively small molecular weight of 12.45 kDa and high sulfate content, accounting for 22.93%. It has been found that the polysaccharide activity increases with decreasing relative molecular weight in a certain molecular weight range.⁷ A similar report showed that low molecular weight GPs (average molecular weight of 20.96 kDa) could effectively inhibit the growth of transplanted S180 tumors.⁹ Smaller molecular GP fragments are tolerant to oral, gastric, and small intestinal conditions and reach the colon as a whole, are more easily metabolized by intestinal microbes, and have a better effect on cellular responses.³²

GPs can significantly regulate the expression of apoptosis and cell cycle-related genes *via* the death receptor-mediated apoptosis pathway and the p53 pathway in A549 cell lines.²⁹ Fan *et al.* showed that acidic GPs significantly suppressed the proliferation of tumor in ICR mice transplanted with H22 hepatoma cells by increasing both specific and non-specific cellular immune responses.³³ In this study it has been found for the first time that GPs combined with CP could induce ferroptosis in the tumor tissue and sensitize the antitumor effect of CP in C26 tumor bearing-mice (Fig. 6). Transcriptomic results from tumor tissues showed that GP90 regulated the differential expression of genes related to ferroptosis in mice with CP chemotherapy. The combination treatment significantly increased the protein expression of Tfrc, which is a potential biomarker for colon adenocarcinoma initiation and progression and drug targets.³⁴ Dysregulated iron metabolism in CRC patients is highly correlated with their poor prognosis, and the iron transporter protein Tfrc mediates iron transport, promotes Tfrc mRNA stability, increases ROS production and iron death sensitivity, and enhances the host immune response to cancer.³⁵ In addition, Apoe expression associated with cholesterol metabolism is elevated, with similar results in adipocytes treated with ferric

ammonium citrate resulting in an increased Apoe expression. Excess levels of cholesterol can also induce ferroptosis by elevating oxidative stress responses.³⁶ When cells become cancerous, the interaction between inflammation and tumors enhances ROS production, and the depletion of the mitotic receptor Bnip3, a key mitochondrial autophagy receptor in cells, promotes increased ROS levels and cell death.³⁷ Thus, Bnip3-depleted cells in GP90 combined with CP groups may contribute to tumor cell ferroptosis.

Based on these results, the ferroptosis-related proteins were validated, and an increase in the 4-HNE expression, and a decreased expression in Gpx4 and SLC7A11 were found in the CP + GP90 group. The final product of malondialdehyde, 4-HNE, produced by lipid peroxidation, may form covalent adducts with biomacromolecules, thereby reducing the membrane integrity and cross-linking, and inactivating proteins, and ultimately promoting cell membrane rupture and ferroptosis.³⁸ Recently, a growing number of studies have found that the induction of ferroptosis by increasing reactive oxygen species, decreasing intracellular levels of the antioxidant glutathione (GSH), or inactivating Gpx4 or SLC7A11 in CRC cells can facilitate the clinical treatment of CRC.³⁹ Gpx4, a key upstream regulator of ferroptosis, is the only glutathione peroxidase used for intracellular lipid peroxidation reduction. If GSH is depleted or Gpx4 is inactivated, intracellular ferrous ions induce ferroptosis by breaking down phospholipid hydroperoxide leading to lipid peroxidation, which can be lethal to tumor cells.⁴⁰ Ferroptosis can be induced when lipid peroxide accumulates excessively, this process is tightly regulated by SLC7A11, known as System Xc-, a key component of the cysteine-glutamate antiporter.⁴¹ Either knockdown of the SLC7A11 gene or inhibition of its activity increased ROS levels and decreased cysteine and glutathione levels, subsequently impairing the viability of colorectal cancer stem cells.⁴² Thus, inhibition of the SLC7A11 expression usually leads to GSH depletion, which induces ferroptosis. These results are consistent with the decreased glutathione (reduced) and ascorbic acid and increased L-glutamic acid in the tumor tissue of GP90 combined with CP treatment. The KEGG enrichment analysis of the metabolomics of colon cancer tissues similarly showed that the combination of GP90 and CP affected the glutamate metabolism, glutathione metabolism, and pantothenate and CoA biosynthesis. Additionally, the related lipid metabolites such as 16(R)-HETE and 17(S)-HpDHA were increased. The increased DHA effectively activates ferroptosis-mediated tumor killing by promoting ROS accumulation, lipid peroxidation, and protein oxidation.⁴³ Inhibition of System Xc- has been reported to decrease the levels of CoA, which is synthesized from cysteine *via* the pantothenate pathway and plays a role in many metabolic pathways, particularly lipid metabolism, and can affect the sensitivity to ferroptosis.⁴⁴ This finding indicates that GP90 improves the CP treatment sensitivity by activating transferrin and reducing Gpx4- and SLC7A11-induced ferroptosis.

The onset of tumor ferroptosis and the resulting immune response trigger immunogenic cell death, as well as promote dendritic cell maturation and T cell infiltration. CD8⁺ T cells

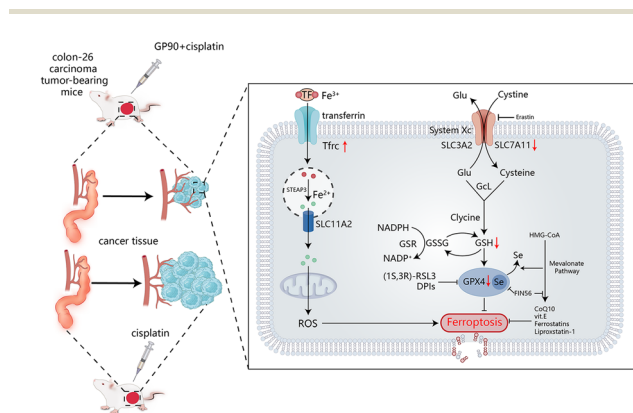


Fig. 6 GP90 combined with cisplatin potentiates the antitumor activity *via* inducing ferroptosis targeting the Tfrc and SLC7A11/Gpx4 pathway.



can synergistically enhance T cell-regulated antitumor immune activity and promote the ferroptosis of tumor cells by downregulating IFN- γ release from SLC3A2 and SLC7A11, in conjunction with the immune checkpoint inhibitor PD-L1.³⁸ Therefore, from GSEA, GP90 combined with CP could positively regulate the innate immune response, promote the production and response of IFN- γ , and assist the immune checkpoint inhibitor PD-L1, which contribute to tumor cell death. Additionally, the Gpx4-deficient regulatory T cells elevate lipid peroxidation that facilitates Th17 responses and the promotion of antitumor immunity.⁴⁵ We have found that the combination of GP90 and CP promotes the Th17, Th1, and Th2 cell differentiation in response to the reduction in Gpx4. GP90 activated the innate immune response in combination with CP, and also involved the NOD-like receptor signalling pathway, Toll-like receptor signalling pathway, and T cell receptor signalling pathway, and plays a role in ferroptosis by regulating oxidative stress.⁴⁶ Further study will address the effect of GP90 on the immune function in CP-treated C26 carcinoma tumor-bearing mice.

5. Conclusions

In summary, we found that GP90 enhanced the antitumor effect of CP by promoting ferroptosis, which might be *via* targeting Tfr and the SLC7A11/Gpx4 pathway. Furthermore, the combination treatment of GP90 and CP potentiated the cancer immunotherapy, which may be related to the NOD-like receptor, Toll-like receptor, T cell receptor, PD-L1 expression, and PD-1 checkpoint pathway. Overall, GP90 is a novel ferroptosis inducer and can be used as a functional food supplement for colon patients to promote chemotherapy sensitivity.

Author contributions

The authors confirm the contribution to the paper as follows: Bingna Cai: conceptualization, data curation, formal analysis, writing – original draft preparation, funding acquisition, review and editing. Lianxiang Luo: methodology and validation. Xiaodan Chen: investigation and validation. Xiangtan Zhao: investigation and validation. Jiak He: visualization. Hua Chen: formal analysis and validation. Peng Wan: visualization and data curation. Deke Chen: data curation. Jianyu Pan: supervision, funding acquisition, review and editing. All authors have read and agreed to the published version of the manuscript.

Conflicts of interest

There are no conflicts to declare.

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