Supercritical carbon dioxide-developed silk fibroin nanoplatform for smart colon cancer therapy

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Purpose: To deliver insoluble natural compounds into colon cancer cells in a controlled fashion.

Materials and methods: Curcumin (CM)–silk fibroin (SF) nanoparticles (NPs) were prepared by solution-enhanced dispersion by supercritical CO2 (SEDS) (20 MPa pressure, 1:2 CM:SF ratio, 1% concentration), and their physicochemical properties, intracellular uptake efficiency, in vitro anticancer effect, toxicity, and mechanisms were evaluated and analyzed.

Results: CM-SF NPs (<100 nm) with controllable particle size were prepared by SEDS. CM-SF NPs had a time-dependent intracellular uptake ability, which led to an improved inhibition effect on colon cancer cells. Interestingly, the anticancer effect of CM-SF NPs was improved, while the side effect on normal human colon mucosal epithelial cells was reduced by a concentration of ~10 μg/mL. The anticancer mechanism involves cell-cycle arrest in the G1/G0 and G2/M phases in association with inducing apoptotic cells.

Conclusion: The natural compound-loaded SF nanoplatform prepared by SEDS indicates promising colon cancer-therapy potential.

Keywords: supercritical, silk fibroin, nanoparticle, natural compounds, colon cancer

Introduction

Curcumin (CM) is a natural polyphenolic compound obtained from the root of Curcuma longa Linn (turmeric).1 It is reported that CM possesses multiple biomedical functions, such as antibacterial, antioxidant, anticancer, anti-inflammatory, anticarcinogenic, antispasmodic, and anticoagulant properties.2–4 Importantly, oral administration of CM 12 g/day exhibited little toxicity in clinical trials.5 However, clinical applications of insoluble CM face more challenging issues, such as low dissolution rate, low bioavailability, and controlled delivery.

Micronization is an effective strategy to overcome these drawbacks, such as with nanoprecipitation and spray-drying.6 However, these conventional micronization methods are always limited by organic solvent residuals and extreme processing conditions (such as high temperature). Owing to advantages such as mild process conditions (Tc =304.1 K, Pc =7.38 MPa), no organic solvent residues, and being environmentally benign, supercritical (sc) CO2 has been used in preparing nanoplatforms recently.7,8 Among several sc-CO2 techniques, solution-enhanced dispersion by sc-CO2 (SEDS) emerges as a novel nanotechnology due to its broad range in selection of biomaterials and drugs. In SEDS, a special coaxial nozzle is used to produce very small droplets and improve the mixing effect between compound solution and sc-CO2 flow, in order to promote mass transfer.9 Therefore, the surface morphology and particle size of the obtained particles can be controlled by parameters easily.10,11 Our group prepared CM
NPs using SEDS technology,\(^\text{12}\) and results showed that CM nanoparticles (NPs) exhibited higher solubility and dissolution than original CM. Also, the CM NPs obtained showed effective biomedical functions.\(^\text{13}\)

In order to control drug release, biomaterials are introduced to prepare polymeric nanoparticles.\(^\text{14–16}\) However, synthetic polymers may induce biocompatibility concerns. Natural silk is a type of fiber polymer which consists of glue-like silk sericin covering silk fibroin (SF).\(^\text{17}\) SF is an excellent biomaterial, due to its good biocompatibility, ready availability, and controllable degradability.\(^\text{18–20}\) A recent study showed that tumors were noticeably reduced after being injected with CM/5-fluorouracil (Fu)-entrapped SF NPs.\(^\text{21}\) However, the mechanism to deliver incorporated drugs from SF nanoplatforms in a controlled fashion needs further investigation. Furthermore, traditional technologies to prepare SF NPs would induce protein degradation or drug volume loss.

In this study, CM-SF NPs were prepared by SEDS and the physicochemical properties of the CM-SF NPs prepared were tested. Intracellular uptake activity was visualized to study the interactions between CM-SF NPs and cancer cells. The in vitro viability of colon cancer cells and normal human colon mucosal epithelial cells when treated with CM-SF NPs was evaluated. Cell-cycle arrest and apoptosis assays were performed to investigate the mechanism of the anticancer effect of CM-SF NPs.

**Materials and methods**

**Materials**

\(\text{CO}_2\) was from Hong Kong Specialty Gases (Hong Kong). Raw silk fibers of *Bombyx mori* were from the Jiangsu Wujing China Eastern Silk Market (Suzhou, China). Hexafluoroisopropanol (HFIP) was purchased from Huifengda Chemical (Jinan, China). Curcumin was from International Laboratory USA (San Francisco, CA, USA). Both human colorectal cancer cell line HCT116 and normal human colon mucosal epithelial cell line NCM460 were supplied from the cell bank of the Chinese Academy of Sciences (Shanghai, China).

**Preparation of curcumin–silk fibroin nanoparticles**

The procedure for preparing CM-SF NPs was based on our previous study,\(^\text{3}\) which is shown in Figure 1. Materials and procedure details are explained in the Supplementary material. Briefly, SF and CM were mixed and dissolved in hexafluoroisopropanol to the desired concentration (1%). After dissolving completely, the blend solution was ready for the SEDS process (Figure S1).

**Physicochemical characterization**

To assess surface morphology, the prepared samples were observed via field-emission scanning electron microscopy (SEM, JSM-6490; JEOL, Tokyo, Japan). For particle size and size distribution, the SEM images were analyzed with Nano Measurer software and about 500 particles were measured.\(^\text{9,13}\) Molecular structure change was tested using Fourier-transform infrared spectroscopy (FTIR, 1720 spectrophotometer; PerkinElmer, Waltham, MA, USA). Preparation details are provided in the Supplementary material.

**Solubility test**

Solubility testing was conducted in accordance with our previous studies, with minor modifications.\(^\text{9,13}\) Briefly, 10 mL PBS solution (pH 7.4) was added to dry CM-SF NPs and original CM (10 mg) powders and the solution was kept under mild agitation (60 rpm). Insoluble residues were filtered after 1 hour, and then the concentration of the samples was detected at 410 nm using ultraviolet spectrometry for calculating solubility (Figure S2).

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**Figure 1** Preparation process of CM-SF NPs via SEDS.

**Notes:** SF powder and CM powder were mixed and dissolved in HFIP solution, ready for the SEDS process. After complete removal of HFIP solution, CM-SF NPs were collected. Yellow curves and circles, curcumin; blue curves, silk fibroin; blue oval, HFIP.

**Abbreviations:** CM, curcumin; SF, silk fibroin; NPs, nanoparticles; SEDS, solution-enhanced dispersion by supercritical \(\text{CO}_2\); HFIP, hexafluoroisopropanol.
Cellular uptake study
All cell lines used in our research were purchased commercially from the cell bank of the Chinese Academy of Sciences (Shanghai, China). Procedures were based on our previous studies. First, 5x10^4 HCT116 cells were seeded onto cover slips in 24-well plates for cell attachment. Then, cells were treated with samples for 0.5 and 6 hours. After that, cells were washed with PBS and fixed with 4% formaldehyde for 1 minute. Secondly, formaldehyde was removed completely and 500 μL methanol (−20°C) was added, followed by 10 minutes’ permeabilization with 0.2% Triton X-100 in PBS and washed with pure PBS. Thirdly, cells were sealed with 10% bovine serum albumin. Fourthly, cell nuclei were stained for 30 minutes with 5 μL 4′,6-diamidino-2-phenylindole. Lastly, stained cells were observed using confocal laser-scanning microscopy (TCS SP8; Leica Microsystems, Wetzlar, Germany).

Cell-viability test
The viability of HCT116 and NCM460 cells treated with CM-SF NPs were evaluated with MTS assay (CellTiter 96; Promega, Fitchburg, WI, USA). Briefly, samples (CM-SF NPs, CM-dimethyl sulfoxide [DMSO], SF NPs, 5-Fu, and control) were sterilized for 30 minutes via ultraviolet light. HCT116 and NCM460 cells were cultivated in the 96-well plates (5x10^3 cells/well) for 24 hours. On the next day, the cell medium was replaced by the sterilized samples in DMEM. After 2–6 days of incubation, the culture medium was removed and cells were washed three times with PBS, then 4 hours of reaction was initiated with 200 μL MTS solution added. A microplate reader (model 680; Bio-Rad Laboratories Inc., Hercules, CA, USA) was used to measure absorbance values at 490 nm. Cell viability was calculated as:

\[
\text{Cell viability} (\%) = \left( \frac{\text{Absorbance of test cells}}{\text{Absorbance of control}} \right) \times 100
\]

where the OD_{490} values of cells treated with different groups corresponded to the absorbance of test cells and the OD_{490} values of untreated cells reflected the absorbance of control cells.

Cell-cycle and apoptosis measurement
HCT116 cells (10^6) were treated with CM-SF NPs, CM-DMSO, 5-Fu, and control, collected after removing supernatant, and then resuspended with PBS. Treated cells were fixed and kept at 4°C for 12 hours in ethanol (70%, −20°C). Afterward, cells were collected and resuspended in 500 μL PBS, followed by staining with 50 μL 7-ADD (7-amino-actinomycin D) and incubating at room temperature in the dark for 30 minutes. Finally, flow cytometry (FACSCanto II; BD Biosciences, San Jose, CA, USA) was used to test the stained samples.

For apoptosis measurement, cells were treated with CM-SF NPs, CM-DMSO, 5-Fu, and control for 24 hours at IC_{50}. All treated cells (10^6) were collected and washed twice in PBS. Ice-cold binding buffer (500 μL) was used to resuspend treated cell aggregates. The cell suspension was stained with 5 μL annexin V-phycoerythrin and 5 μL 7-ADD, then incubated in the dark for 5 minutes and tested by flow cytometry.

Statistical analysis
Each experiment was performed in triplicate, and all data are presented as means ± SD. Statistical analysis was performed using one-way analysis of variance with the level of statistical significance set at P<0.05.

Results
Physicochemical properties
Surface morphologies of samples are shown in Figure 2A and B. As can be seen, the irregular original CM powders showed uneven large particles (>1 μm), while the regular spherical CM-SF NPs were much smaller in size (<100 nm). Chemical structures of samples were compared (Figure 2C). Characteristic peaks of 1,648 (C=O stretching vibration), 1,531 (N–H stretching vibration), and 1,245 cm\(^{-1}\) (stretching vibration), and 1,260 cm\(^{-1}\) in original CM corresponded to 1,260, 1,155, 960, and 828 cm\(^{-1}\) in CM-SF NPs, respectively. Meanwhile, major characteristic peaks of 1,648 (C=O stretching vibration), 1,531 (N–H stretching vibration), and 1,245 cm\(^{-1}\) (N–H bending and C–N stretching vibrations) in SF corresponded to 1,648, 1,531, 1,260 cm\(^{-1}\) in the CM-SF NPs, respectively, from which it can be concluded that CM was successfully incorporated into the SF nanoplatform via SEDS. Importantly, the solubility of incorporated CM was greatly improved over the original CM (Figure 2D–F).

Cellular uptake activity
For studying the intracellular transport potential of CM-SF NPs, qualitative intracellular uptake assays were performed, and both 0.5- and 6-hour treated cells were chosen for observation. As shown in Figure 3A, there was no clear green fluorescence found in the control group or the CM-DMSO group at the 0.5-hour time point. Only slight green fluorescence was observed after 6 hours of treatment in the CM-DMSO group. By contrast, clear green fluorescence presented in the CM-SF NPs group at the 0.5-hour time point and its intensity increased greatly after 6 hours of treatment, which indicates...
that CM-SF NPs had time-dependent intracellular transportability and showed nuclei-targeting potential as well.\textsuperscript{13,22} Anticancer activity

The cell viability of HCT116 was measured in response to CM-SF NPs, CM-DMSO, SF NPs, 5-Fu, and control treatments for 48 hours (Figure 4A). The inhibitory effect of 5-Fu improved with increased concentration (0.5–20 \( \mu \)g/mL). However, neither the CM-SF NP nor CM-DMSO group showed any inhibitory effect when concentrations were <2 \( \mu \)g/mL, but when concentration reached 5 \( \mu \)g/mL, their anticancer effects sharply increased, maybe owing to the intrinsic property of CM. Meanwhile, CM-SF NPs had a stronger inhibitory effect (~68\%) than CM-DMSO (~42\%) group at 5 \( \mu \)g/mL, and the IC\textsubscript{50} value of CM-SF NPs group was lower compared to the CM-DMSO group (Table S1). CM-SF NPs presented superior anticancer potential (~94\%) compared to 5-Fu (~83\%) when the concentration was higher than 10 \( \mu \)g/mL. In contrast, cells treated with SF NPs retained about 90\% viability at all tested concentrations.
concentrations. In summary, CM-SF NPs prepared by SEDS exhibited superior anticancer potential at concentrations higher than 5 μg/mL.

**Cytotoxic activity**

It is necessary to evaluate the cytotoxicity of CM-SF NPs on normal human colon mucosal epithelial cells for potential biomedical applications. MTS results on NCM460 cells for concentrations of 0.1–200 μg/mL are shown in Figure 4B. As can be seen, 5-Fu exhibited toxic effects from 1 μg/mL, while both CM-SF NPs and CM-DMSO exhibited toxic effects at 10 μg/mL, and a significant difference in cell viability was observed (~58% in CM-SF NPs and ~14% in CM-DMSO), suggesting that the toxicity of CM incorporated...
into SF nanoplatforms was reduced at an effective concentration in the treatment of colon cancer.

Mechanism of anticancer effect

Cell-cycle arrest
To elucidate the mechanism of the inhibitory effect on colon cancer cells of CM-SF NPs, cells were treated for 24 hours at IC$_{50}$. Results of cell-cycle distributions are shown in Figure 5A and C. 5-Fu arrested high populations (~51%) of cells in the G$_2$/M phase, while both CM-DMSO and CM NPs presented high percentages of cells in the G$_0$/G$_1$ (~57% and ~51%, respectively) and G$_2$/M phases (~18% and ~19%, respectively) compared to the control group (~47% and ~12%, respectively). Therefore, the inhibitory mechanism on colon cancer cells of CM-SF NPs was presumably due to enhanced efficacy in cell-cycle arrest at the G$_0$/G$_1$ and G$_2$/M phases.

Induction of cell apoptosis
It is believed that cell apoptosis may represent a mechanism to counteract neoplastic development, which is important for cancer therapy. Reports on cancer therapy in recent years have revealed that CM can cause apoptosis signals in various cancers.$^{23-25}$ As shown in Figure 5B and D, cells treated with CM-SF NPs showed a higher percentage of apoptosis (~23%) than those treated with 5-Fu (~16%). This indicates that CM-SF NPs had better apoptotic cell-induction potential than 5-Fu. In summary, the mechanism of inhibitory effect on colon cancer cells of CM-SF NPs was mainly due to cell-cycle arrest in the G$_0$/G$_1$ and G$_2$/M phases in association with inducing apoptosis.

Discussion
CM is recognized as a potential chemotherapeutic drug. However, its clinical application is severely hindered by...
poor solubility, low bioavailability, lack of bioavailability to tumors, and poor intracellular uptake efficiency. In our previous report, we prepared CM-SF NPs by using SEDS for the first time, and their solubility was remarkably increased over the original CM. Here, we further studied the controlled drug-release properties of SF nanoplatforms for smart colon cancer therapy.

In this study, small (<100 nm) spherical CM-SF NPs were prepared by SEDS. The chemical structure of CM-SF NPs was confirmed by FTIR results; it is believed that the incorporated CM most likely interacted with the hydrophobic residues in the silk nanoplatforms. Moreover, the 3,495 cm⁻¹ (O–H) peak in CM shifted to 3,290 cm⁻¹, suggesting that CM linked to the SF nanoplatforms, maybe via hydrogen bonds. Importantly, the solubility of CM-SF NPs was greatly improved, which was mainly due to size reduction, supersaturation of CM, water solubility of the SF nanoplatform, better dispersibility and wettability of CM, and hydrogen bonds between CM and the SF nanoplatform.27,28

The intracellular uptake result of CM-SF NPs is in agreement with previous studies.26,29 In those studies, the introduction of SF improved the retention of loaded drugs in cancer cells, thus leading to a higher anticancer efficacy. In our opinion, there are two reasons for the better cellular uptake
efficiency of CM-SF NPs: 1) the endocytosis pathway, which is believed to be energy-costly for NP uptake (Figure 3B), and 2) the SF nanoplatforms are able to facilitate intracellular uptake.

In our opinion, the combination of improved intracellular uptake efficiency and enhanced permeability and retention effect of CM-SF NPs contributes to the higher toxic effect on colon cancer cells. Moreover, it is important to ascertain whether their anticancer effects would be retained or diminished during a therapeutic period. The cell viability of the treated cells (HCT116) for 6 days (at the IC$_{50}$ of samples) was calculated (Figure S3). The results showed that CM-SF NPs and CM-DMSO had similar cell viability on days 2 and 4. However, a significant difference in cell viability was obtained on day 6, CM-SF NPs displaying an improved anticancer effect (>98%) compared to CM-DMSO (~66%) and 5-Fu (~63%), due to the sustained-release property of CM-SF NPs. Moreover, SF NPs did not display significant toxic effects. Therefore, CM-SF NPs possess both dose- and time-dependent anticancer effects, mainly due to sustained release. Our findings are similar to those of previous studies on colon cancer therapy by mucoadhesive CM-containing chitosan NPs.²⁷,³⁰

Meanwhile, CM-SF NPs showed reduced toxicity on normal human colon mucosal epithelial cells at an effective concentration (~10 µg/mL) in the treatment of colon cancer cells. This phenomenon could be explained by the slow and sustained drug-release potentials of SF nanoplatforms, thereby reducing their cytotoxicity. Our observation is in agreement with an earlier report on colon cancer therapy with CM-poly(lactic-co-glycolic acid) (PLGA) conjugates.²⁸,³¹ In conclusion, sc-CO$_2$-developed SF nanoplatforms showed a smart colon cancer therapy potential (Figure 4C).

The cell-cycle results indicated that the inhibition effect on colon cancer cells by CM-SF NPs was partly owing to cell-cycle arrest in the G$_0$/G$_1$ and G$_2$/M phases. Our findings are in line with previous studies on doxorubicin and CM encapsulated in liposomes for tumor therapy²⁸,³² and CM-PLGA NPs for breast cancer treatment.³⁰,³³ Our cell apoptosis results were similar to the previous CM nanoformulation study.³⁴ In the present study, we hypothesized that the better apoptosis induction ability of CM-SF NPs was owing to their higher cellular uptake efficiency, resulting in greater accumulation of released CM within cells in conjunction with continuous delivery that arrested a greater percentage of apoptotic cells. This result was also consistent with the MTS assays (Figure 4A). It has been reported that inhibition of the key antiapoptotic proteins Mcl1 and BclxL and induction of PARP cleavage are possible mechanisms of apoptosis induction by CM.³⁵

Conclusion

In conclusion, CM-SF NPs with controllable particle size (<100 nm) were prepared by SEDS in this study, with results showing that they showed time-dependent intracellular uptake ability due to SF nanoplatforms facilitating endocytosis, which led to improved inhibition effects on colon cancer cells. The mechanism can be explained by cell-cycle arrest in the G$_0$/G$_1$ and G$_2$/M phases in association with inducing apoptotic cells. Interestingly, the anticancer effect of CM-SF NPs was improved while the side effect was reduced at concentrations of ~10 µg/mL, which may have been caused by the combination of the slow-release property of the SF nanoplatform and intrinsic properties of CM. In conclusion, NC-loaded SF nanoplatforms prepared by SEDS showed promising colon cancer therapy potential.

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Disclosure

The authors report no conflicts of interest in this work.

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Supplementary materials

Formation of curcumin–silk fibroin nanoparticles

The solution-enhanced dispersion by supercritical CO₂ (SEDS) apparatus was composed of the following parts (Figure S1): a precipitation vessel for particle formation, a CO₂ container and delivery part, and a solution transport and spray part. CO₂ was cooled to 0°C and then pumped into a precipitation vessel, which was placed in a heated gas bath.

In the SEDS process, a CO₂ flow was maintained to keep the system pressure stable when temperature and pressure reached the desired values. Then, an enhanced mixing effect between the curcumin (CM)–silk fibroin (SF) blend solution and supercritical CO₂ resulted when they met within a specially designed coaxial nozzle. Subsequently, the mixed flow was sprayed into the precipitation vessel quickly. Afterward, a fresh CO₂ flow was maintained to completely clear the HFIP residue. Then, the precipitation vessel was depressurized slowly. Finally, dry CM-SF nanoparticles were obtained and vacuum-dried, then stored at 4°C for further characterization.

Surface morphology, particle size, and size distribution

An aluminum sample holder with a thin self-adherent carbon film was used to load the samples, and then they were coated with a thin layer of gold. The surface morphology of the samples was visualized by using field-emission scanning electron microscopy (SEM, JSM-6490; JEOL, Tokyo, Japan). The equivalent diameters of 500 particles in the obtained SEM photographs were measured, and particle size and size distribution were analyzed by Nano Measurer software.

Physicochemical characterization

KBr powders were used to mix the obtained samples (ratio ~1:100), and then the mixtures were transformed to a thin tablet. Fourier-transform infrared spectra of the obtained samples were collected and analyzed with a 1720 spectrophotometer (PerkinElmer, Waltham, MA, USA) in transmission mode with the wave number ranging 400–4,000 cm⁻¹. For solubility test results, see Figure S2.

Cell culture and proliferation

Both NCM460 and HCT116 cells were incubated in RPMI 1640 medium (Thermo Fisher Scientific, Waltham, MA, USA).
USA) combined with 1% penicillin–streptomycin (100 U/mL penicillin and 100 μg/mL streptomycin; Thermo Fisher Scientific) and 10% fetal bovine serum (Thermo Fisher Scientific). Cells were cultured in an incubator with a humidified atmosphere of 37°C±0.2°C and 5% CO₂.

Figure S3 Cell proliferation of HCT116 after treatment with CM-SF NPs, CM-DMSO, 5-Fu, SF NPs, and control.

Note: Results are shown as means ± SD, n=6.

Abbreviations: CM, curcumin; SF, silk fibroin; NPs, nanoparticles; DMSO, dimethyl sulfoxide; 5-Fu, 5-fluorouracil.