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Triptolide and Celastrol Loaded Silk Fibroin Nanoparticles Show Synergistic Effect against Human Pancreatic Cancer Cells

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Abstract

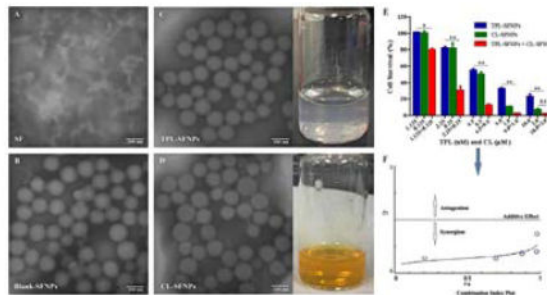
Pancreatic cancer is a lethal disease with a dreadful 5-year survival rate of only 5%. In spite of several treatment options, the prognosis still remains extremely poor. Therefore, novel therapy strategies with combinations of drugs are urgently required to combat this fatal disease. Triptolide (TPL) and celastrol (CL), two main compounds in traditional Chinese medicine isolated from Thunder God Vine, have a broad range of bioactivities including anticancer activity. Silk fibroin (SF), a naturally occurring protein with several unique properties, is an ideal carrier material. In this study, we prepared TPL and CL loaded silk fibroin nanoparticles (TPL-SFNPs and CL-SFNPs) by a modified desolvation method and evaluated their synergistic effects against human pancreatic cancer cells. Both SFNPs were characterized for particle size and zeta potential. The entrapment efficiency, drug loading, and drug release profiles were evaluated by HPLC. The cytotoxicity and synergistic effect of SFNPs were investigated in MIA PaCa-2 and PANC-1 human pancreatic cells. The results showed that the particle sizes of TPL-SFNPs and CL-SFNPs were 166.4 ± 4.6 nm and 170.4 ± 2.3 nm, with a mean zeta potential -27.2 ± 2.0 mV and -25.5 ± 2.57 mV, respectively. TPL-SFNPs and CL-SFNPs have a drug loading of 57.0 ± 4.7 $\mu\text{g}/\text{mg}$ and 63.5 ± 3.8 $\mu\text{g}/\text{mg}$ along with an encapsulation efficiency of $81.8 \pm 2.8\%$ and $87.0 \pm 5.1\%$, respectively. Drug release studies revealed that a rapid release of the drugs from SFNPs was observed at pH 4.5 (lysosomal pH) and a delayed release was observed at pH 7.4 (plasma pH). TPL-SFNPs (IC_{50} 3.80 and 4.75 nM) and CL-SFNPs (IC_{50} 0.38 and 0.64 μM) were 2–3 fold more potent against MIA PaCa-2 and PANC-1 cells than free TPL (IC_{50} 11.25 and 11.58 nM) and CL (IC_{50} 0.84 and 1.23 μM). Furthermore, co-treatment with TPL-SFNPs and CL-SFNPs increased the growth inhibition of the same cells significantly in comparison with TPL-SFNPs or CL-SFNPs alone. Almost all combination index (CI) values, calculated using the Compusyn software, were < 1 , suggesting that the growth inhibition effect of TPL-SFNPs in combination with CL-SFNPs was synergistic rather than additive, further suggesting that this novel combination may offer a potential treatment for pancreatic cancer.

Graphical Abstract

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Novel combination of triptolide and celastrol loaded silk fibroin nanoparticles show synergistic anti-pancreatic cancer effect *in vitro*



Keywords

Pancreatic cancer; triptolide; celastrol; silk fibroin nanoparticles; synergistic effect

1. Introduction

Pancreatic cancer is one of the most aggressive human malignancies with a 5-year survival rate of only 5%, which has a rather low morbidity of 2% among all cancers diagnosed but 6% of cancer deaths worldwide.^{1, 2} Pancreatic cancer is mainly treated with chemotherapeutic drugs, which may result in side effects and potential drug resistance that can further lead to the unfortunate demise of the patient.^{3–5} The current standard drug for chemotherapy to treat pancreatic cancer is gemcitabine; however, its efficacy is far from satisfactory. To date, there is no other agent that has promoted a clinically meaningful prolongation of overall survival. Studies have indicated that a single inhibition of signaling pathway or single chemotherapy is not sufficient to effectively kill pancreatic cancer cells, while the combination of two or more signaling pathways and chemotherapy can reduce the number of pancreatic cancer cells to almost undetectable levels.^{3, 6–8} Thus, novel therapy strategies to treat pancreatic cancer are still to be proposed, and there is a critical need for the development of novel therapeutic agents to improve the treatment of this disease.

Triptolide (TPL, Fig. S1A in the Supporting Information) and celastrol (CL, Fig. S1B in the Supporting Information), two principal bioactive compounds derived from traditional Chinese herb Thunder God Vine, have attracted immense attention due to their unique chemical structures and broad range of bioactivities, particularly their anticancer activities.^{9–11} TPL has been shown to effectively induce cell cycle arrest and apoptosis in various types of cancers including pancreatic cancer, breast cancer, melanoma and lung cancer.^{12–15} It could effectively decrease the viability of pancreatic cancer cells *in vitro* and reduce the growth and metastases of tumors *in vivo*.^{9, 12, 16} Celastrol can suppress the invasion of pancreatic cancer cells by downregulating the expression of CXCR4 chemokine receptor.¹⁷ However, owing to their poor water solubility and severe toxicity, TPL and CL cannot be used systemically in the clinic. Therefore, there is an urgent need to develop alternative TPL and CL formulations for clinical use.

Combination therapy is the routine strategy of cancer chemotherapy with significant advantages including lower treatment failure rate and slower development of drug resistance. TPL can sensitize cancer cell lines to several chemotherapeutic drugs, such as cisplatin, adriamycin, temozolomide, and TRAIL in both *in vivo* and *in vitro* settings.^{13–16} CL also shows synergistic anticancer effects in combination with several chemotherapeutic agents on suppressing the proliferation of various cancers including pancreatic cancer, melanoma, lung cancer and breast cancer.^{17–20} Combination of low doses of TPL and CL can induce significant pancreatic cancer cell death compared to treatment with either drug alone.²¹ Therefore, TPL and CL combination is a viable strategy to enable effective treatment of pancreatic cancer. However, there are still some critical issues that remain to be solved during combination treatment. There is a lack of defined optimal dose and schedule of administration. Administration of multiple drugs in free solution at normal therapeutic doses often results in increased normal tissue toxicity.²²

Nanocarrier-based drug delivery systems, nanoparticles in particular, have generated great excitement in the field of drug delivery since they provide advantages over the administration of free drugs, including targeting ability to enhance accumulation at tumor sites, overcoming resistance by intracellular drug delivery, and realizing controlled and sustained release to enhance drug bioavailability.^{23–26} Well-designed nanoparticles can also alter the pharmacokinetics and toxicity profiles of the free drug combination, enable drug accumulation in the tumor and release the drugs at a synchronized rate, thereby maintaining synergistic drug ratios to achieve enhanced anticancer effects. Silk fibroin (SF) protein, a natural product regenerated from silkworm cocoons, possesses several unique properties including biocompatibility, tunable biodegradation, stabilizing effects, water-based processing and diverse material formats, making it a potential material for incorporation into a variety of drug delivery vehicles capable of delivering a range of therapeutic agents.^{27–29} SF-based nanoparticles (SFNPs) have been shown to successfully deliver anticancer drugs, including small molecules and biomolecules.^{29–32}

In this study, we developed and characterized TPL-loaded silk fibroin nanoparticles (TPL-SFNPs) and CL-loaded silk fibroin nanoparticles (CL-SFNPs) separately, which not only over-came the pitfall of hydrophobicity, but also facilitated passive accumulation of TPL and CL in cancerous tissues based on the enhanced permeability and retention (EPR) effect as well as optimal dose and schedule of administration. Formulation and characterization of TPL-SFNPs and CL-SFNPs followed by optimization of the process of drug loaded SFNPs in terms of particle size, encapsulation efficiency and drug loading were evaluated. Shape and dispersity were evaluated using TEM. The time dependent cumulative drug release of CL in different pH media from SF nanoparticles using HPLC was performed. Evaluation of the biocompatibility of the silk fibroin nanoparticles (SFNPs) using hemolytic assay was conducted using mice blood *in vitro*. The cellular uptake behavior of SF nanoparticles was investigated in MIA PaCA-2 and PANC-1 cells with Rhodamine-B-Isothiocyanate (RITC) as the fluorescent probe confirmed by confocal microscopy and flow cytometer. The superiority of the combined regimen of CL-SFNPs mixed with TPL-SFNPs was demonstrated by anti-proliferation and cell apoptosis detection in human pancreatic cancer cell lines, MIA PaCA-2 and PANC-1.

2. Materials and methods

2.1 Materials

TPL and CL were purchased from Chengdu Biopurify Phytochemicals Ltd. (Chengdu, China). Cocoons were kindly supplied by Tongxiang mulberry silk base of Zhejiang Province (Tongxiang, China). Dialysis membrane with a cut off of 7,000 Da (MWCO) was obtained from Viskase Companies, Inc. (Chicago, USA). Hoechst 33342 and 3-[4, 5-dimethylthiazol-2-yl]-2, 5-diphenyltetrazolium bromide (MTT) were obtained from Sigma-Aldrich (St. Louis, MO, USA). RITC and Annexin V-FITC apoptosis detection kits were supplied by Sigma Chemicals (St. Louis, MO, USA). HPLC grade acetonitrile and methanol were obtained from BDH Chemicals (Gibbstown, NJ, USA). Other chemicals used in this study were all analytical pure grade and used as received. Ethanol, acetone and other chemicals used in this study were acquired from VWR international (Darmstadt, Germany) unless specified otherwise.

Dulbecco's modified Eagle's medium (DMEM) and fetal bovine serum (FBS) were purchased from Gibco BRL (Carlsbad, CA, USA). Penicillin–streptomycin, 0.25% trypsin-EDTA and non-essential amino acids were obtained from Invitrogen Co. (Carlsbad, CA, USA). Human pancreatic cancer cells MIA PaCa-2 and Panc-1 cells were generous gifts from Dr. Prabhu's lab, originally obtained from American Type Culture Collection (Rockville, MD, USA).

2.2 Regenerated silk fibroin preparation, purification and molecular weight

The regenerated SF protein solution was prepared as previously reported with minor modification³³. Briefly, the purchased *B. mori* cocoons were cut into small pieces and degummed twice in 0.02 M Na₂CO₃ for 30 min each and thoroughly rinsed with deionized water. After air-drying, the degummed SF was dissolved in a ternary system, CaCl₂-CH₃CH₂OH-H₂O (1:2:8, mole ratio), under constant stirring at 75 °C for 2 h, 4 h and 8 h. Furthermore, the resulting SF solution was centrifuged at 4000 rpm for 5 min. The supernatant was collected carefully and dialyzed in Viskase dialysis membrane (MWCO 7 000 Da) against deionized water for 3 days to remove salts and ethanol. After centrifugation at 14000 rpm for 15 min, the concentration of prepared silk fibroin solution was determined using NanoDrop™ 2000/2000c spectrophotometers. The molecular weight range of SF extracted was investigated by SDS-PAGE with 8% separation gel and 5% condensing gel, followed by staining with Coomassie brilliant blue R-250.³⁴ The silk fibroin solution was stored at 4 °C for further use.

2.3 Silk fibroin protein nanoparticles

2.3.1 Fabrication of silk fibroin protein nanoparticles—TPL-SFNPs and CL-SFNPs were formulated using a modified desolvation technique.^{35, 36} Typically, TPL and CL were dissolved in a mixture of acetone:ethanol (3:2, v/v) individually at a certain concentration, and the prepared TPL or CL solution was added into SF solution dropwise under gentle constant stirring. The resulting suspension was incubated in a refrigerator at -20 °C for 16 h and defrosted quickly in warm water (40 °C). The suspension was subsequently centrifuged at 14000 rpm for 15 min and washed three times with deionized water to collect the

nanoparticles. Finally, the purified nanoparticles were suspended in deionized water using an ultrasound processor at 10% amplitude for 2 min with an ultrasonic cell disruptor to disperse the clustered silk spheres, followed by lyophilization with 5% (w/v) trehalose as the cryoprotectant. The prepared freeze-dried nanoparticle formulations are stored at 2–8 °C.

The Rhodamine-B-Isothiocyanate (RTIC)-loaded SFNPs (RITC-SFNPs) were prepared by the same method, and blank SFNPs (Blank-SFNPs) were prepared according to the above mentioned method but omitting the TPL and CL.

2.3.2 Experimental design for formulation optimization—Taguchi's L9 orthogonal array experimental design was used to optimize the formulation parameters of TPL-SFNPs and CL-SFNPs. A 3-factor, 3-level design was employed for studying the interaction and quadratic effects of the formulation variables. The initial concentrations of SF, TPL and CL and volume ratio of organic/SF solution for formulation optimization were selected based on preliminary experiments (data not shown). The three factors and their levels selected for formulation optimization are shown in Table S1. A Design-Expert® (Version 10.1, Stat-Ease Inc., USA) software was used for analyzing the results.

2.4 Nanoparticle characterization

2.4.1 Particle size, zeta potential and morphology—Freeze-dried SFNPs (Blank-SFNPs, TPL-SFNPs, CL-SFNPs) were dispersed in deionized water (pH 7.0). Average size and zeta potential of SFNPs were measured by a dynamic light-scattering detector (Nanobrook Omni, Brookhaven Instrument Corp, USA). All measurements were performed at room temperature in triplicate. The morphological examination of SF and SFNPs was conducted via transmission electron microscopy (TEM, JEOL JEM-1230, Japan).

2.4.2 Infrared spectra IR absorption and β -sheet content—The FTIR spectra of Blank-SFNPs, drug-loaded SFNPs, as well as free drug were obtained via a Fourier transform infrared spectrophotometer (FTIR, Varian, USA). Lyophilized, regenerated SF was also examined. For each measurement, the spectra were generated from 32 scans with a resolution of 4 cm^{-1} . The β -sheet content of SF in SFNPs or regenerated SF was obtained by deconvolution of amide I band using PeakFit 4.12 software.^{36, 37}

2.4.3 Drug loading capacity and encapsulation efficiency—The encapsulation efficiency and drug loading capacity of TPL-SFNPs and CL-SFNPs were analyzed by Agilent 1050 HPLC (Agilent Technologies, Palo Alto, CA, USA). Analyses were performed at 25 °C using a C18 column (250 mm \times 4.6 mm, 5 μm , Agilent Technologies, USA). Methanol:water (58:42, v/v) and methanol:water (90:10, v/v) were used as mobile phases for TPL and CL, respectively, at a flow rate of 1 mL/min. The detection wavelengths were 218 nm and 430 nm, respectively.

Encapsulation efficiency (EE) and drug loading (DL) of nanoparticles were calculated according to equations (1) and (2):

$$DL (\%) = \frac{\text{amount of the drug in NPs}}{\text{amount of the NPs}} \times 100 \quad (1)$$

$$EE (\%) = \frac{\text{amount of the drug in NPs}}{\text{amount of the feeding drug}} \times 100 \quad (2)$$

2.4.4 *In vitro* drug release—The cumulative release kinetics of TPL and CL from SFNPs were determined in phosphate buffered saline (PBS), at pH 7.4 and pH 5, respectively. Equal amount of SFNPs was suspended in PBS and separated in capped glass bottles, followed by an incubator at 37 °C with a shaking speed of 120 strokes/min. At predetermined time intervals (1, 4, 8, 24, 48, 72, 120 and 168 h), three glass bottles of each formulation were withdrawn and drug release was monitored by separating nanoparticles and release media via centrifugation (14000 rpm, 15 min) repeated three times. The amounts of residual TPL or CL in the nanoparticles were determined by HPLC using the same procedure as described above. The profile that shows the cumulative drug release as a function of time was plotted.

2.5 *In vitro* hemolysis assay

The hemolysis assay was performed on SFNPs *in vitro*. In brief, 0.2 mL of 3.8% sodium citrate was mixed with 4 mL of fresh mouse blood, followed by gentle centrifugation at 3,000 rpm for 10 min; the sediment was collected and washed 3 times by suspending it in PBS (pH 7.4) at 37 °C to get the final 5% (v/v) RCBs suspension. Furthermore, 10 mg/mL of suspended NPs (Blank-SFNPs, TPL-SFNPs and CL-SFNPs) in PBS were incubated with RBCs for 2 h at room temperature. Moreover, 0.1% Triton X-100 and PBS in RBCs solution were used as positive and negative control, respectively. Subsequently, samples were centrifuged for 5 min, and 100 μL supernatants were collected carefully and transferred to a 96-well plate to enable the quantification of hemoglobin by a micro plate reader (μQuant™, BioTek® Instruments, Inc.). The percentage of hemolysis was calculated according to the following equation:

$$\text{Hemolysis (\%)} = \frac{\text{optical density- negative control optical density}}{\text{positive control optical density- negative control optical density}} \times 100$$

2.6 Cell culture

The pancreatic cancer cells PACA-2, PANC-1, HEK 293 and HGF-1 were cultivated in monolayers to 70–80% confluence in DMEM or EMEM medium supplemented with 10% fetal bovine serum at 37 °C in a humidified environment of 5% CO₂. The medium was replenished every other day, and the cells were sub cultured after reaching confluence.

2.7 *In vitro* cellular uptake

2.7.1 Intracellular localization imaging by confocal microscopy—To investigate cellular uptake of SFNPs, 5×10⁴ cells/well MIA PaCa-2 and PANC-1 cells were seeded onto

sterilized microscope cover slips placed in 6-well plates and incubated overnight in DMEM medium supplemented with 10% fetal bovine serum at 37°C with 5% CO₂. Seeded cells were then treated with free RTIC and RTIC-SFNPs (equaled to 0.5 μM in medium) and incubated for 5 min, 30 min and 60 min (untreated cells were used as control). Subsequently, the cells were washed three times with PBS, followed by fixation with 4% p-formaldehyde for 15 min and washed three times again with PBS. Hoechst 33342 was used to stain the nuclei of the cells; furthermore, the cells were washed three times with PBS and imaged using Leica TCS SP2 confocal microscopy, and the images were analyzed by Leica confocal software.

2.7.2 Cellular uptake determined by flow cytometry—Flow cytometry analysis was performed to quantify the cellular uptake of SFNPs in MIA PACA-2 and PANC-1. Briefly, 5 × 10⁴ cells/well were seeded in 12-well plates and incubated for 24 h. The culture medium was then replaced by free RTIC and RTIC-SFNPs (equaled to 0.5 μM in medium), incubated for 5 min, 30 min and 60 min and washed twice with warm (37 °C) Dulbecco's Phosphate-Buffered Saline (D-PBS). Thereafter, cells were detached using trypsin and washed with ice-cold D-PBS. Finally, cells were re-suspended in 500 μL of ice-cold D-PBS and kept on ice in dark for 10 min. The samples were analyzed by BD Accuri™ C6 flow cytometer (BD Biosciences, San Jose, CA, USA).

2.8 Cell cytotoxicity studies by MTS assay

To evaluate the cytotoxicity of Blank-SFNPs *in vitro*, 5×10³ cells/well of MIA PaCa-2 and PANC-1 cells, 6×10³ cells/well of HEK 293 and HGF-1 cells were seeded in 96-well plates and incubated for 24 h before being treated with Blank-SFNPs. To evaluate the cell viability, MIA PaCa-2 and PANC-1 cells were treated with various concentrations of TPL, CL, TPL-SFNPs and CL-SFNPs.

After incubation for 72 h, the growth medium was removed, followed by addition of 100 μl serum-free culture medium containing 20% MTS and 1% of phenazine methosulfate (PMS) and further incubated for 1 h. The absorbance of each well was measured by micro plate reader at 490 nm. Viability of untreated cells was set at 100%, and absorbance of wells with medium and without cells was set as zero. IC₅₀ values were analyzed by Prism software (San Diego, CA). Each experiment was performed at least three times.

2.9 Clonogenic assay

Clonogenic assay was used to determine the growth inhibition of pancreatic cancer cells by TPL-SFNPs and CL-SFNPs. Furthermore, 1×10³ MIA PaCa-2 and PANC-1 cells/well were seeded in 6-well plates. On the third day, cells were treated with different concentrations of CL and CL-SFNPs (C_{CL}: 0.05–0.45 μM) and TPL and TPL-SFNPs (C_{TPL}: 0.3–3.0 nM) and incubated for 4 days. The media was then replaced with fresh media and maintained for another 7 days. Cells were fixed with ice-cold mixture of methanol and acetic acid (3:1, v/v) and incubated for 15 min. Furthermore, colonies were stained with 0.5% crystal violet. Colonies with more than 50 cells were counted manually under a microscope and the data were analyzed using Graph Pad Prism software. Untreated cells were considered as controls.

2.10 Determination of combination index value

To evaluate the combination effect of TPL and CL and TPL-SFNPs and CL-SFNPs, pre-incubated MIA PaCa-2 and PANC-1 cells were co-treated with various concentrations of drug and drug-loaded SFNPs for 72 h. The experiment was designed as constant combination ratio based on the suggestion of CompuSyn software and the IC_{50} values obtained in section 2.8 (Table S2). In addition, the cell viability was determined by the same MTS assay method described in section 2.8 and analyzed by CompuSyn software, which follows the median effect principle to identify the combination index (CI) value. The equation to calculate the combination index is $CI = (D)_1/(Dx)_1 + (D)_2/(Dx)_2$, where $(Dx)_1$ and $(Dx)_2$ indicate the individual dose of either free drugs or drug loaded SFNPs required to inhibit a given level of cell growth, and $(D)_1$ and $(D)_2$ are the doses of free drugs and drug loaded SFNPs necessary to produce the equal effect in combination, respectively.^{38–40} Combination index (CI) value <1 , $=1$ and >1 indicate synergism, additive effect and antagonism, respectively.

2.11 Apoptosis assay

Apoptosis assay was performed using the Annexin V-FITC Apoptosis Detection Kit according to the manufacturer's protocol supplied by Sigma Chemicals (St. Louis, MO, USA). 5×10^4 MIA PaCa-2 and PANC-1 cells were pre-incubated in 24 well plates and incubated for 24 h, and treated with TPL (4.5 nM), CL (0.5 μ M), TPL+CL, and TPL-SFNPs, CL-SFNPs, TPL-SFNPs + CL-SFNPs loading the same amount of TPL and CL for 48 h. Afterwards, the cells were washed twice with ice-cold PBS, trypsinized, centrifuged and then re-suspended in cold binding buffer. A total of 500 μ l of this cell suspension was then subjected to the addition of 5 μ l of Annexin V FITC and 10 μ l of propidium iodide solution, followed by incubation on ice in the dark for 10 min. After incubation, the cell samples were analyzed by flow cytometer immediately. The tests were carried out in triplicate.

2.12 Statistical analysis

GraphPad Prism software (La Jolla, CA) was used for statistical analysis and plotting. The data were presented as mean \pm standard deviation. Statistical significance was determined using the one-way ANOVA or two-way ANOVA, followed by Tukey's test, where p-value of 0.05 was considered statistically significant.

3. Results

3.1 Regenerated silk fibroin

SF is a biomacromolecule consisting of 5509 amino acids and composed of a heavy chain (391 kDa) and a light chain (26 kDa). It has been reported that SF molecules undergo hydrolysis during degumming procedure.⁴¹ Similarly, in our study, SDS-PAGE showed that the regenerated SF was a mixture of polypeptides with a wide molecular distribution (Fig. 2S).

3.2 Experimental design for formulation optimization

The formulation parameters of TPL-SFNPs and CL-SFNPs were optimized using Taguchi's L9 orthogonal array experimental design (three-factor, three-level). The responses selected for experimental design are shown in Table S3. A numerical desirability function was used to select the optimized formulation from the experimental design.⁴² The significance of the experimental model used in formulation optimization of TPL-SFNPs and CL-SFNPs was validated by comparing the predicted values with experimental values (Table S4). The results from the optimization study indicate that the particle size, EE and DL of TPL-SFNPs and CL-SFNPs are comparable to the predicted values demonstrating the accuracy and validity of the model used in the experimental design.

3.3 Nanoparticle characterization

3.3.1 Particle size, zeta potential and morphology—As shown in the TEM images in Fig. 1, the SFNPs are spherical in shape and monodisperse. The observed particle sizes of SFNPs were around 100 nm, smaller than those measured by DLS, which indicated that the particle sizes of TPL-SFNPs and CL-SFNPs were 166.4 ± 4.6 nm (PDI<0.3) and 170.4 ± 2.3 nm (PDI<0.3), respectively (Table 1). This was mainly due to the dehydration process that led to the shrinkage of the particles in the preparation of TEM samples. TPL-SFNPs and CL-SFNPs have a mean zeta potential of -27.2 ± 2.0 mV and -25.5 ± 2.57 , respectively (Table 1).

3.3.2 IR absorption and β -sheet content—FTIR spectra of TPL, CL, TPL-SFNPs, CL-SFNPs, SF and SFNPs, are shown in Fig. 2. TPL-SFNPs and CL-SFNPs exhibit characteristic absorption peaks at 1735 cm^{-1} and 1500 cm^{-1} of TPL and CL which confirms the successful encapsulation of drug into SFNPs.^{36, 43, 44} It is found that the β -sheet content in SF is about 16.8%, but it increases to 35.3% in the SFNPs. Moreover, the β -sheet content in TPL-SFNPs and CL-SFNPs increases to 57.7% and 77.6% in the presence of TPL and CL (Table 1, Fig. S3). This means that TPL and CL addition is also favorable for the conformation transition of SF from random coil/helix to β -sheet probably because of the hydrophobic interactions between TPL, CL and SF macromolecules. Furthermore, the increased β -sheet formation induced by TPL or CL may benefit their encapsulation in the SFNPs leading to slow drug release in later applications.⁴⁰

3.3.3 Drug loading capacity and encapsulation efficiency—Drug loading capacity (DL) and encapsulation efficiency (EE) play a critical role in drug delivery. To obtain high drug loading and meet the therapeutic needs, we used the optimized formulations to prepare SFNPs. The prepared TPL-SFNPs and CL-SFNPs have a satisfactory DL of 57.0 ± 4.7 $\mu\text{g}/\text{mg}$ and 63.5 ± 3.8 $\mu\text{g}/\text{mg}$, along with an EE of $81.8 \pm 2.8\%$ and $87.0 \pm 5.1\%$, respectively (Table 1).

3.4 *In vitro* drug release profile

Cumulative drug release from TPL-SFNPs and CL-SFNPs were studied at two different pH conditions; at pH 7.4, which is comparable to the extracellular physiological pH of normal cells and pH 5, the lysosomal pH of cancer cells. As shown in Fig. 3A & 3B, both drugs were gradually released from the nanoparticles at pH 7.4 indicating that encapsulation of the

drug in SFNPs may significantly prolong the release in plasma. However, at pH 5, the release was significantly faster throughout the early time points (1 to 24 h) and the 7-day release period. The cumulative TPL and CL release was approximately 53% and 55% within 24 h and a total 95% and 80% in 7 days at pH 5 which showed significantly faster release pattern in comparison to 82% and 65% in pH 7.4 in 7 days, indicating that SFNPs could serve as a lysosomotropic delivery platform for drugs.

3.5 Safety evaluation of Blank-SFNPs and drug loaded SFNPs

Hemolytic test was performed on Blank-SFNPs and drug loaded SFNPs. As shown in Fig. 4A, compared to positive control Triton X-100 that showed the highest toxicity, breaking 100% of RBCs present in solution, almost no hemolysis was observed when treated with PBS as negative control. Similar to PBS, Blank-SFNPs and drug loaded SFNPs at the high concentrations of 10 mg/mL did not cause any hemolysis on mouse erythrocytes.

To study the cytotoxicity, Blank-SFNPs at a concentration of 0.125 to 1 mg/mL were used to detect aggregation and sedimentation that could potentially result in cell toxicity. The study results showed no significant inhibition of pancreatic cancer cells, MIA PaCa-2, PANC-1, normal cells, HEK293 and HGF-1 (Fig. 4B), indicating that Blank-SFNPs alone do not contribute to any quantifiable toxicity in the studied concentration. The study demonstrates high safety and biocompatibility of the Blank-SFNPs *in vitro*.

3.6 *In vitro* cellular uptake

3.6.1 Intracellular localization of nanoparticles—To study the uptake, PANC-1 and MIA PaCa-2 cancer cells were exposed to RITC-SFNPs for 5, 30 and 60 min. RITC-SFNPs are efficiently taken up by both cell lines in a time dependent manner and are predominately distributed in the cytoplasm or perinuclear region following 60 min of incubation (Fig 5A & 5B). The RITC-SFNPs internalized more rapidly and significantly compared to free RITC at 30 and 60 min, indicating that drug encapsulated in SFNPs could be efficiently taken up by cancer cells enabling consistent intracellular release of the loaded drug. The nanoparticle internalization process into the cells can be considered as a binding (adsorbing) process, wherein the formed nanoparticles are internalized by endocytosis.^{41, 43} Once the nanoparticles were endocytosed, they were found preliminarily in the cytoplasm around the nuclear membrane. We found that a large fraction of the incubated nanoparticles was internalized by the cells and remained stable during vesicular transport despite their negative surface. We also observed that both cell lines remain viable during the study, supporting our previous results that silk fibroin nanoparticle alone did not result in any measurable toxicity.⁴⁴

3.6.2 Cellular uptake determined by flow cytometry (FCM)—The cellular uptake of RITC-SFNPs is quantitatively evaluated and depicted in Fig. 6A & 6B. The results show that RITC-SFNPs exhibited significantly high fluorescence intensity compared to the free RITC following incubation with PANC-1 cells for 5, 30 and 60 min, and about two-fold increase of fluorescence intensity was observed with RITC-SFNPs compared to free RITC (Fig. 6B). On the contrary, RITC-SFNPs incubated with MIA PaCa-2 cells also indicated significant

difference in fluorescence intensity compared to free RITC treatment cells (Fig. 6A). However, the intensity was not as prominent as seen in PANC-1 cell line.

3.7 Cytotoxicity of drug-loaded SFNPs

To study the efficacy of CL-SFNPs and TPL-SFNPs on cancer cells, the inhibitory concentration (IC_{50}) values of the free drug and SFNPs were compared by MTS assay. The cell viability studies of CL-SFNPs and TPL-SFNPs are presented in Fig. 7A–7D, respectively. Both free drug and SFNPs exhibited dose dependent decrease in the proliferation of pancreatic cancer cells. However, it was observed that the SFNPs exhibited lower IC_{50} values in comparison to the free drug for both TPC and CL drugs, demonstrating higher efficacy with SFNPs. In case of MIA PaCa-2 cells, free TPL exhibited IC_{50} value of 11.58 μ M. However, for TPL-SFNPs, the IC_{50} value was significantly reduced to 3.8 μ M, showing approximately three-fold reduction compared to the free drug (Fig. 7A). Similarly, for PANC-1 cells, the IC_{50} values obtained for free TPL and TPL-SFNPs were 11.25 μ M and 4.57 μ M, respectively (Fig. 7B). Similar trend was observed with CL-SFNPs. The viability study for free CL with MIA PaCa-2 cells, exhibited IC_{50} value of 1.32 μ M and for CL-SFNPs, the IC_{50} value was reduced to 0.65 μ M, showing approximately two-fold reduction compared to the free form of the drug (Fig. 7C). Similarly, for PANC-1 cells, the IC_{50} values obtained for free CL and CL-SFNPs were 0.83 μ M and 0.38 μ M, respectively (Fig. 7D). Therefore, our results demonstrate that the drugs, when encapsulated in SFNPs, exhibited cytotoxicity at lower concentrations compared to the free drug.

3.8 Inhibition of colony formation

Clonogenic assay or colony formation assay is an *in vitro* cell survival assay based on the ability of a single cell to grow into a colony. Clonogenic assay was performed to determine the anti-cancer activity of the free drug, SFNPs of TPL and CL through cell growth inhibition. In this study, the untreated MIA PaCa-2 and PANC-1 cell lines produced large colonies while the treated cells with free drug and SFNPs showed a dose dependent inhibition of colony formation. The SFNP formulations of TPL and CL showed higher inhibition of colony formation at 0.15 and 0.45 μ M concentrations, compared to free TPL and CL (Fig. 8A, 8C & 8B, 8D), respectively at equivalent doses. At 0.15 μ M concentration, TPL-SFNPs and CL-SFNPs showed significant reduction in colony formation in MIA PaCa-2 and PANC-1 cancer cells compared to free TPL and CL. Similarly, TPL-SFNPs and CL-SFNPs are also more effective at the 0.45 μ M concentrations compared to free TPL and CL, indicating superior anticancer activity after incorporation of TPL and CL into SFNPs. We also observed that PANC-1 cells are more sensitive to CL-SFNP. At a dose of 0.45 μ M, colony formation is inhibited by 90% compared to 70% in MIA PaCa-2 cell line. However, both cell lines are sensitive to TPL-SFNPs, and about 95% colony formation is inhibited at 3.0 nM dose (Fig. 8B & 8D). In summary, the rapid internalization and sustained release of CL and TPL from nanoparticles may substantially improve anticancer effects of celastrol and triptolide at the same dose.

3.9 Synergistic effect of TPL-SFNPs and CL-SFNPs

MTS assay was performed to study the combination effects of TPL and CL or TPL-SFNPs and CL-SFNPs on MIA PaCa-2 and PANC-1 cell lines. After determining the dose response

SFNPs, individually, showed increased apoptosis compared to free CL and TPL at equivalent doses. The total apoptosis was 24.0% and 25.2%, respectively when MIA PaCA-2 cells were treated with TPL-SFNPs and CL-SFNPs individually. However, when used in combination at the same concentration, the total apoptosis of MIA PaCA-2 cells was significantly higher (58.6%) (Fig. 11A & 11B). The same trend was found in PANC-1 cells when treated with the combination (Fig. 11C & 11D). Overall this study confirmed that nanoparticle combination of TPL and CL was extremely effective in enhancing apoptosis of cancer cells and demonstrating the synergistic effect of TPL-SFNPs and CL-SFNPs combination.

4. Discussion

Multidisciplinary treatment strategies are commonly used to combat pancreatic cancer. However, these remedies are often inadequate in contending the disease due to drug resistance and safety issues associated with normal healthy tissues. Diagnosis using combination drug therapy has been widely employed in clinical settings to treat the lethal diseases and attain synergistic therapeutic effect, to minimize drug toxicity and to overcome drug resistance. Triptolide and celastrol are reported to inhibit the growth of cancer cells synergistically when treated in combination both *in vitro* and *in vivo*.²¹ However, their use is limited because of their highly hydrophobic nature and non-selective toxicity to healthy cells. Therefore, to address these issues, triptolide and celastrol loaded silk fibroin nanoparticles were developed and evaluated for their effect on pancreatic cancer cell lines. Silk fibroin, a natural biopolymer extracted from cocoons, with proven clinical safety record, mechanical properties and biocompatibility has been used as carrier to attain targeted drug delivery of triptolide and celastrol. A modified desolvation method was used to prepare silk fibroin nanoparticles.

SFNPs were prepared using a modified process with a mixture of acetone and ethanol produced the desired particle size. The selected solvent mixture might have facilitated dehydration of silk fibroin, leading to closer chain packaging of GLY-X or nanofabrication, thereby allowing mild folding of SF polymer to generate nanoparticles of less than 200 nm in contrast to those prepared using other reported methodologies.⁴⁵

The formulations were prepared systematically using a statistical experimental design and the resulting formulation variables such as size, entrapment efficiency and drug loading were systematically analyzed using Taguchi's orthogonal array design to identify the parameters that have optimum influence on the formulation (Fig. S4 & S5).^{42, 45-47} The data analyzed using Design Expert® software demonstrated the reproducibility of the optimized formulation. According to previous reports, negatively charged nanoparticles have longer circulation lifetimes and less toxicity towards the cell.^{48, 49} The zeta potentials of TPL-SFNPs and CL-SFNPs were -27.2 ± 2.0 mV and -25.5 ± 2.57 mV, respectively, which might result in less aggregation and long circulation of SF nanoparticles in the blood. FTIR spectra showed that TPL-SFNPs and CL-SFNPs exhibit characteristic absorption peaks at 1735 cm^{-1} and 1500 cm^{-1} of TPL and CL, respectively, indicating successful encapsulation of the drug into SFNPs. TEM images suggest that the formulated SFNPs could be spherical and monodisperse.

The release study results indicate that silk fibroin nanoparticle can release the drug in a slow and sustained manner at physiological pH and releases rapidly at lower pH (Fig. 5). At low pH, silk loses its overall acidic surface properties and negative net charge leading to rapid drug release. The pH dependent drug release may also be attributed to the efficient packing as well as loading of drug inside the SFNPs. In the previous report, the extent of release at different pH conditions was predominantly governed by the electrostatic interaction between doxorubicin and SF.⁵⁰ Since doxorubicin is a weak base and positively charged, it is adsorbed to silk in part by strong electrostatic bonding. However, unlike doxorubicin, TPL is neutral and CL possesses a weak negative charge and therefore, may be weakly adsorbed or bonded to the SFNPs (in part). Due to this difference in electrostatic interaction, TPL and CL showed higher release (50 %) at pH 5.0 compared to doxorubicin (20 %) at pH 4.5 in 24 hr. Similar studies by Wang et al. also showed that positively-charged molecules exhibit a more prolonged or sustained *in vitro* release of the drugs from the SFNPs compared to negatively charged molecules due to strong electrostatic interactions.⁵¹ The increase of TPL and CL release from SFNPs might also be attributed to the balance of the negative charges inside and outside the silk aggregate. Xiao et al. reported that metal ions usually interact with functional groups containing negative charges outside the aggregates while hydrogen ions could neutralize the negative charges both outside and inside the aggregates simultaneously due to their differential capacity of entering into the aggregates. When the negative charge outside silk aggregates is shielded, the repulsive force of the negative charge inside the aggregates could result in the destruction of the aggregates.⁵² Since the balance of the negative charge was shielded at acidic pH, the aggregates may be destructed and result in increased release of the drug. We will further investigate the mechanism of drug release and *in vivo* efficacy of the TPL-SFNPs and CL-SFNPs in a follow up study.

Delivering the anticancer drug inside the tumor cells while avoiding the p-gp mediated efflux of free drug to overcome the resistance and achieving the therapeutic concentration are important considerations for any nanoparticle formulation.⁵³ Cellular uptake of nanoparticles is influenced by size, shape, material, surface charge and hydrophobicity.⁴⁹ Cellular uptake assay by using RITC loaded SFNPs reiterated the potential uptake of SF nanoparticles. Confocal laser scanning microscope results indicated the superiority of the SF nanoparticles uptake inside the cells over free RITC (Fig. 5), which may be attributed to the rapid uptake of nanoparticles by endocytosis mechanism. Time dependent uptake of the nanoparticles was also observed in MIA PaCa-2 and PANC-1 cell lines. Superiority and time dependent uptake of SFNPs using flow cytometry analysis (Fig. 6) robustly support that SF nanoparticle approach could be an efficient way to deliver drugs to cancer cells.

The cytotoxic assay performed on PANC-1, MIA PaCa-2, HEK 293 and HFG-1 cell lines employing MTS assay exhibited safety and biocompatibility of placebo or Blank-SFNPs. At all concentrations, cell viability was > 95% after 72 h of treatment, clearly demonstrating the safe and non-toxic nature of Blank-SFNPs. Furthermore, hemolysis test was performed using fresh mouse blood for empty nanoparticles and drug loaded nanoparticles to confirm the biocompatibility of newly formed nanoparticles.^{54, 55} Hemolysis assay which gives an indication of the interactions between SFNPs and RBCs showed no significant hemolysis for the formulations indicating the use of safe, biocompatible, and biodegradable silk fibroin nanoparticles.

In both MIA PaCa-2 and PANC-1 cells, 72 h cytotoxicity assays revealed the IC₅₀ of CL, TPL, CL-SFNPs and TPL-SFNPs (Fig. 7). Notably, the delivery of the drugs using SF-based CL and TPL formulations showed lower IC₅₀ values as compared to free drugs IC₅₀ indicating that nanoparticle formulations were more potent in inhibiting the cancer cell growth. This can be attributed to rapid uptake of nanoparticles inside the cells followed by releasing their high payload in cytosol.^{56, 57} The colony formation assay performed with the same cell line indicated superiority of CL-SFNPs and TPL-SFNPs over the free drug (Fig. 8).

In this study, we estimated the combination index (CI) value for both free drug combination and drug-loaded SFNPs combination using CompuSyn software® to evaluate the synergistic effect and the results demonstrated that TPL-SFNPs and CL-SFNPs have significant synergistic effect at low dose compared to free drug in both pancreatic cancer cell lines; all CI values were below 0.7 (Fig. 9 & 10). As shown in Fig. 9A2 & 9B2, Fig. 10A2 & 10B2, the calculated combination index values of TPL-SFNPs and CL-SFNPs (CI: 0.369–0.630) are much smaller than the combination index values of free drug TPL and CL (CI: 1.600–0.680) indicating much greater synergistic effect of SFNPs' compared to that of free drugs. This synergistic effect may be attributed to the increase in the drug concentration inside the cell. The inhibition of cell growth at low dose drug combination may translate to a decrease in the toxicity *in vivo* condition. Apoptosis study with equivalent dose of drug and drug-loaded nanoparticles was conducted to confirm the potent synergistic effects of this nanoparticle combination. The results demonstrated that even at low dose combination, nanoparticle treatment is more potent in terms of inducing apoptosis and cell death.

5. Conclusion

In summary, TPL-SFNPs and CL-SFNPs with desired particle size and drug loading were successfully prepared to overcome the poor water solubility and high toxicity of TPL or CL as well as to facilitate the simultaneous delivery to cancer cells. The drug-loaded SFNPs have sustained a pH-sensitive drug release profile along with biocompatibility characteristics. Moreover, when compared with free TPL and CL, TPL-SFNPs and CL-SFNPs showed enhanced antitumor activity, higher inhibition of colony formation and could induce more apoptosis of cancer cells, which might be due to the sustained release of TPL and CL from SFNPs as well as the improved stability and cellular uptake of drugs encapsulated inside the nanoparticles. Further reduction in cell viability and significant increase of apoptosis when treated with a combination of TPL-SFNPs and CL-SFNPs provides strong proof of better anticancer efficacy of combination nanoparticle therapy than either TPL-SFNPs or CL-SFNPs individually. More importantly, the combination index value below 0.7, calculated by CompuSyn software, indicates that in certain ratio or drug concentration the combination therapy effect is strongly synergistic rather than additive. The results indicate that combination therapy of TPL and CL encapsulated in SFNP_s could be a promising strategy for the treatment of pancreatic cancer.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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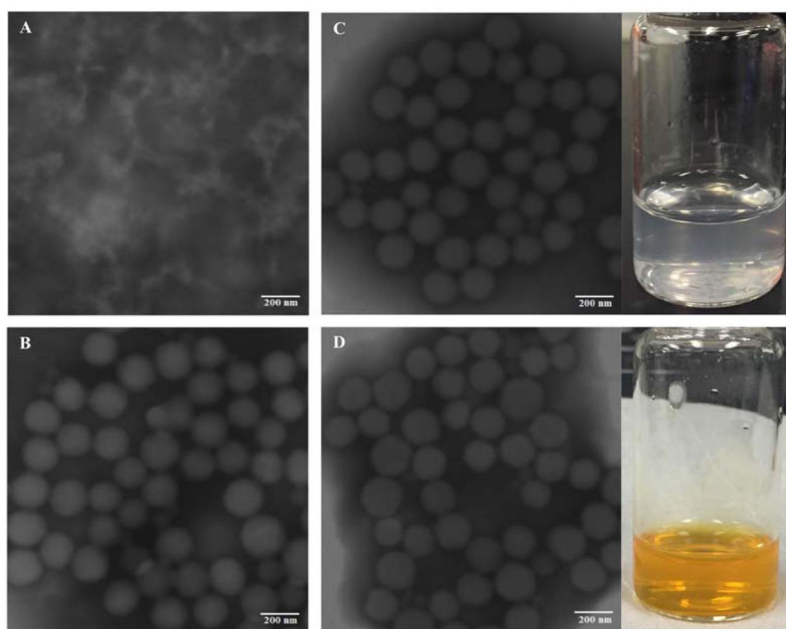


Fig. 1. Characterization of SF and SFNPs

A. TEM image of SF. B. TEM image of Blank-SFNPs. C. TEM image and photograph of TPL-SFNPs formulation. D. TEM image and photograph of CL-SFNPs formulation. Scale bar 200 nm.

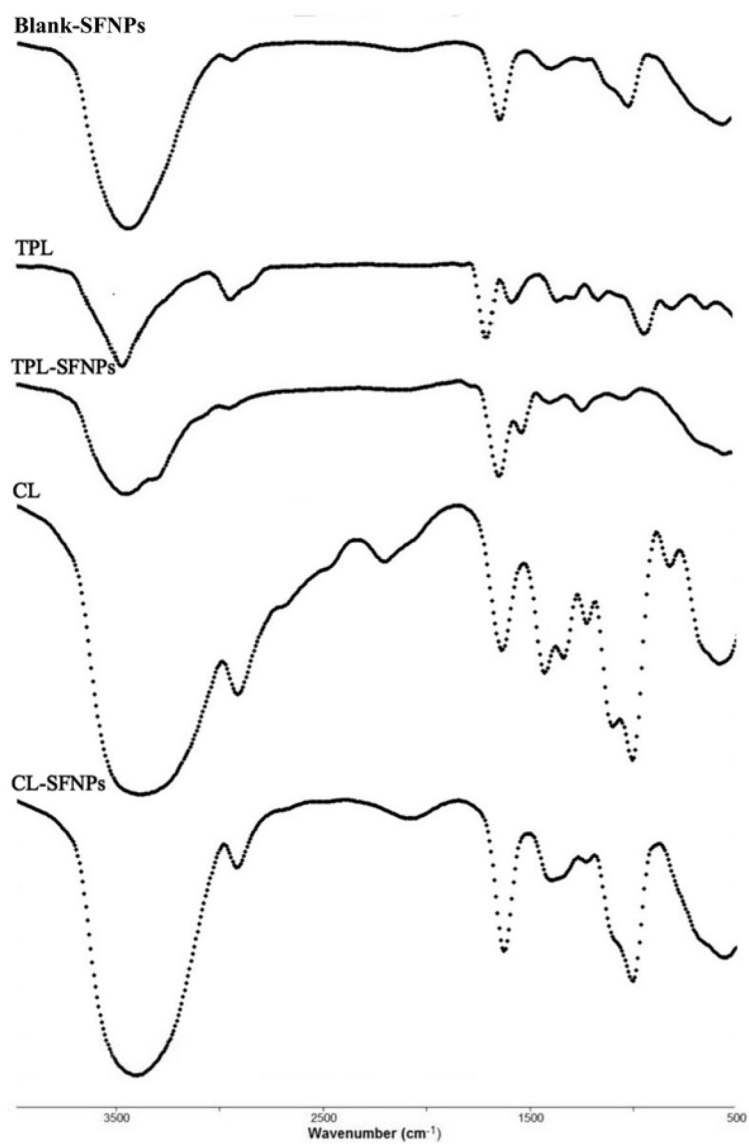


Fig. 2. FTIR spectra of Blank-SFNPs, TPL, TPL-SFNPs, CL and CL-SFNPs

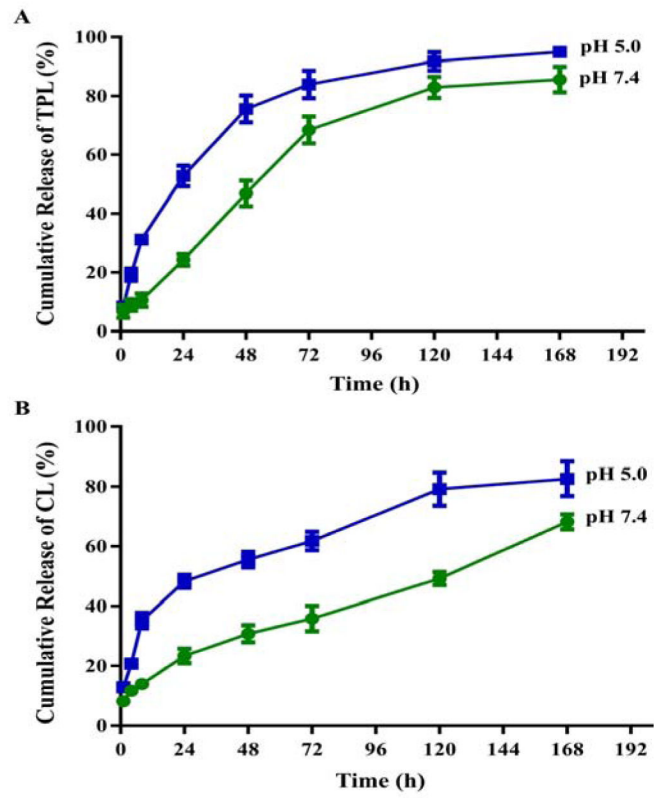


Fig. 3. The cumulative release profiles of TPL from TPL-SFNPs (A) and CL from CL-SFNPs (B) in PBS of pH=7.4 and pH=5.0 at 37 °C.

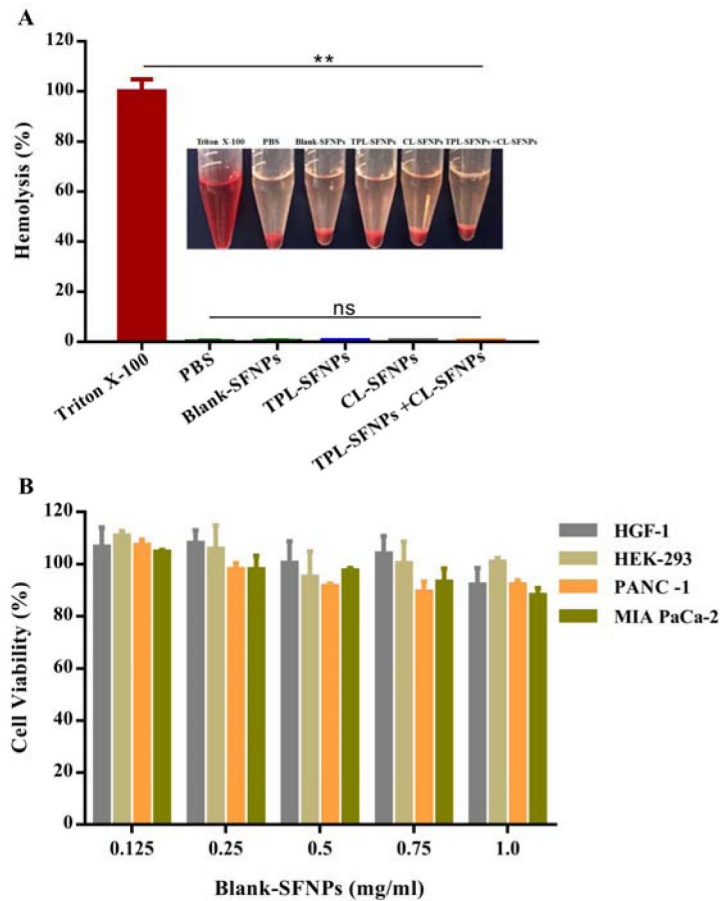


Fig. 4. Safety evaluation of Blank-SFNPs, TPL-SFNPs and CL-SFNPs
 A. Biocompatibility of TPL-SFNPs, CL-SFNPs and their combination in fresh mouse blood. Triton X-100 and PBS were used as positive control and negative control, respectively. B. Cytotoxicity of Blank-SFNPs in MIA PaCa-2, PANC-1, HEK-293 and HGF-1 cell line using MTS assay. (Mean ± SD, n = 5; ns: no significance, *: P<0.05, **: P<0.01)

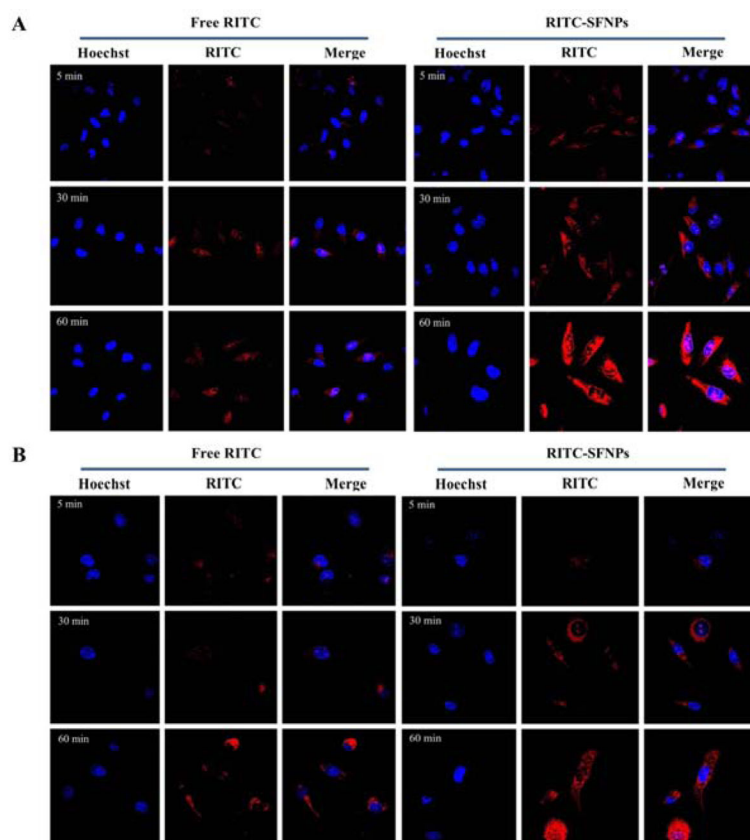


Fig. 5. Cellular uptake analysis of RITC-SFNPs using confocal laser scanning microscopy, free RITC was used as control
A. MIA PaCa-2, with an incubation time of 5 min, 30 min and 60 min. B. PANC-1, with an incubation time of 5 min, 30 min and 60 min. Original magnification: $\times 40$.

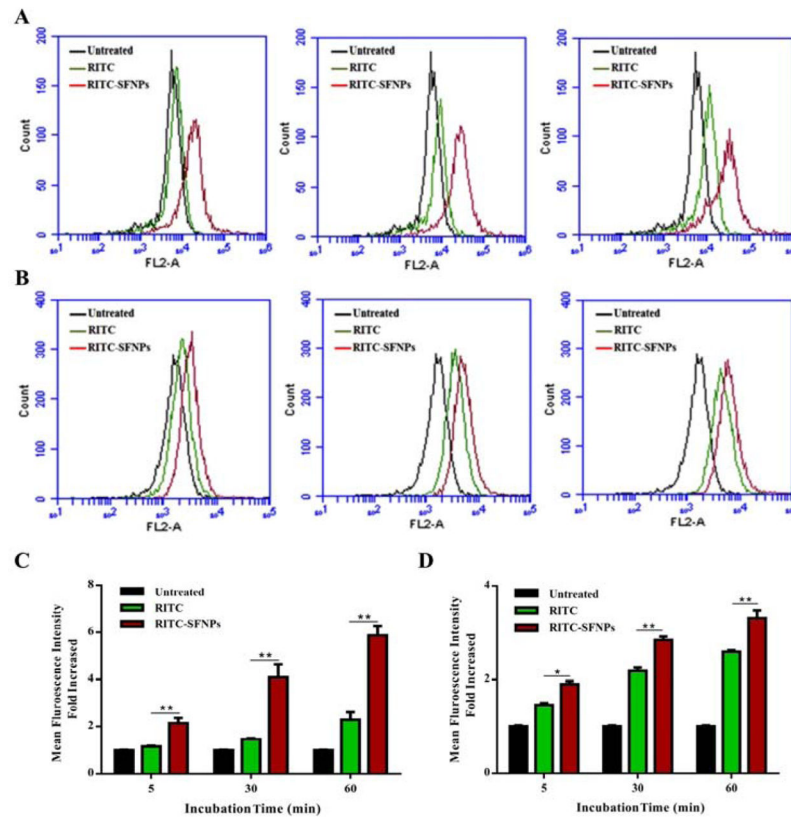


Fig. 6. Quantification of cellular uptake of SFNPs. Uptake analysis of RITC-SFNPs using flow cytometry. Untreated cells and the cells treated with free RITC were used as negative control and positive controls respectively

A. MIA PaCa-2, with an incubation time of 5 min, 30 min, and 60 min. B. PANC-1, with an incubation time of 5 min, 30 min and 60 min. (Mean \pm SD, n = 5; *: P<0.05, **: P<0.01)

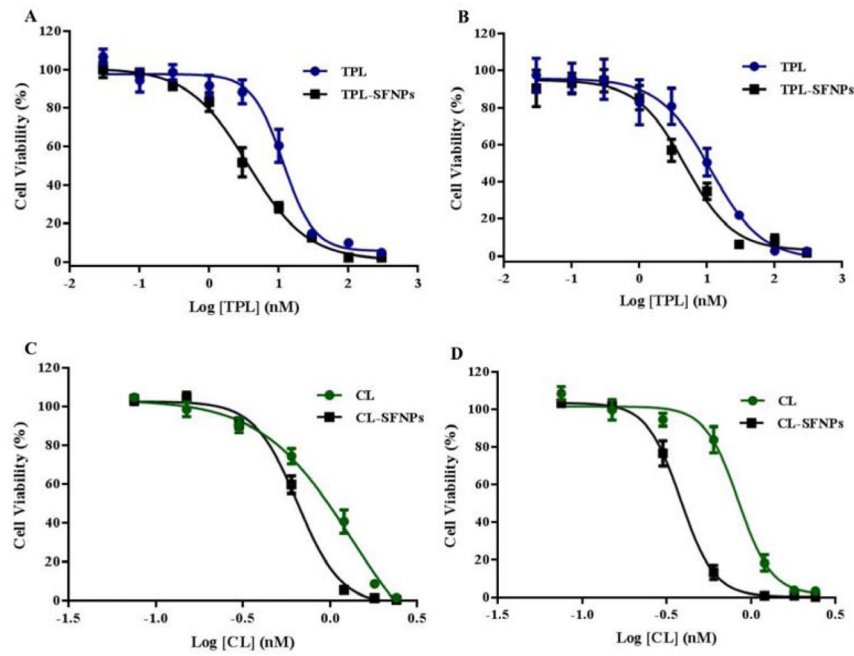


Fig. 7. Dose dependent inhibition of cell viability and IC_{50} determination in MIA PaCa-2 and PANC-1 cell line treated with TPL, TPL-SFNPs, CL & CL-SFNPs using MTS assay
 A. IC_{50} profile of TPL and TPL-SFNPs in MIA PaCa-2 cell line. B. IC_{50} profile of TPL and TPL-SFNPs in PANC-1 cell line. C. IC_{50} profile of CL and CL-SFNPs in MIA PaCa-2 cell line. D. IC_{50} profile of CL and CL-SFNPs in PANC-1 cell line.

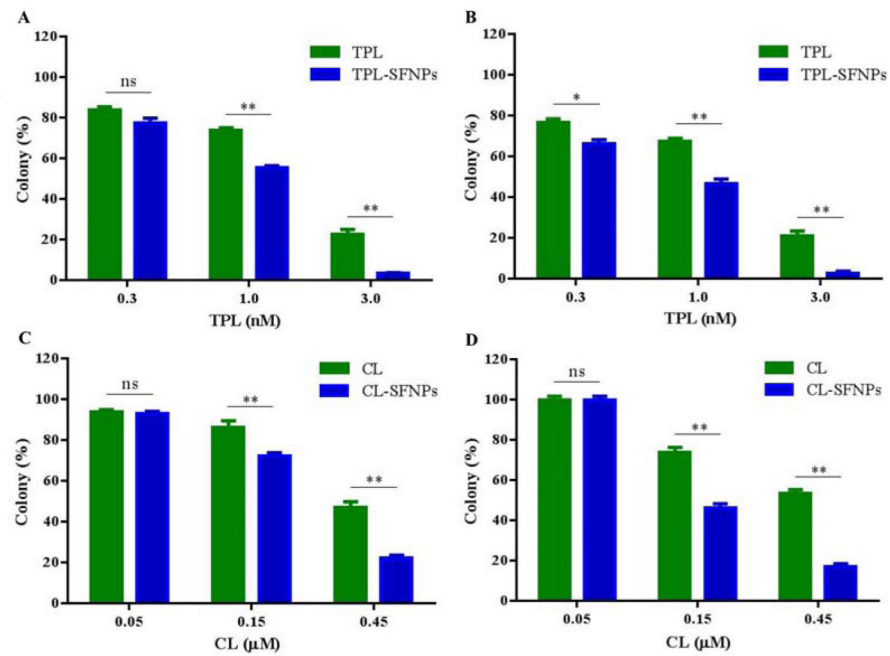


Fig. 8. Dose dependent colony formation inhibition of MIA PaCa-2 and PANC-1 cell line treated with different drugs

A. Colony formation inhibition of TPL and TPL-SFNPs in MIA PaCa-2 cell line. B. Colony formation inhibition of TPL and TPL-SFNPs in PANC-1 cell line. C. Colony formation inhibition of CL and CL-SFNPs in MIA PaCa-2 cell line. D. Colony formation inhibition of CL and CL-SFNPs in PANC-1 cell line. (Mean \pm SD, n = 3; ns: no significance, *: P < 0.05, **: P < 0.01)

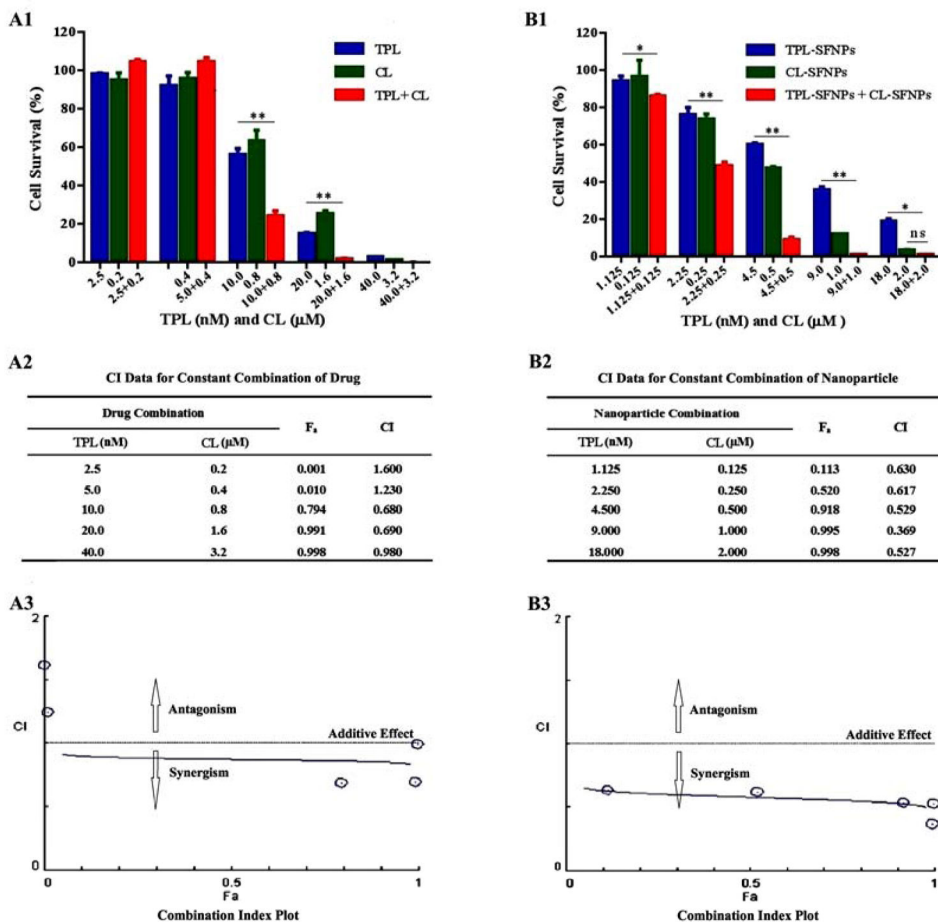


Fig. 9. Synergistic effect of TPL and CL, TPL-SFNPs and CL-SFNPs in MIA PaCa-2 cell line (A1) Cell viability determination and (A2) combination index (CI) value identified using Compusyn software and (A3) combination index plot of CL, TPL and in combination. (B1). Cell viability determination and (B2) combination index (CI) value identified using Compusyn software and (B3) combination index plot of TPL-SFNPs, CL-SFNPs and in combination.

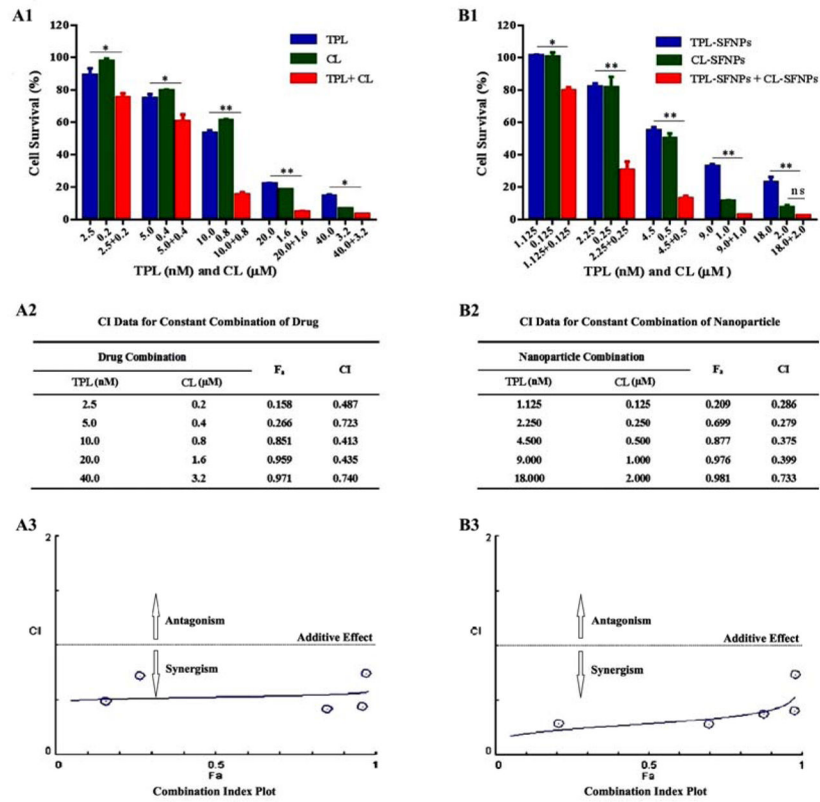


Fig. 10. Synergistic effect of TPL and CL, TPL-SFNPs and CL-SFNPs in PANC-1 cell line (A1) Cell viability determination and (A2) combination index (CI) value identified using Compusyn software and (A3) combination index plot of CL, TPL and in combination. (B1) Cell viability determination and (B2) combination index (CI) value identified using Compusyn software and (B3) combination index plot of TPL-SFNPs, CL-SFNPs and in combination.

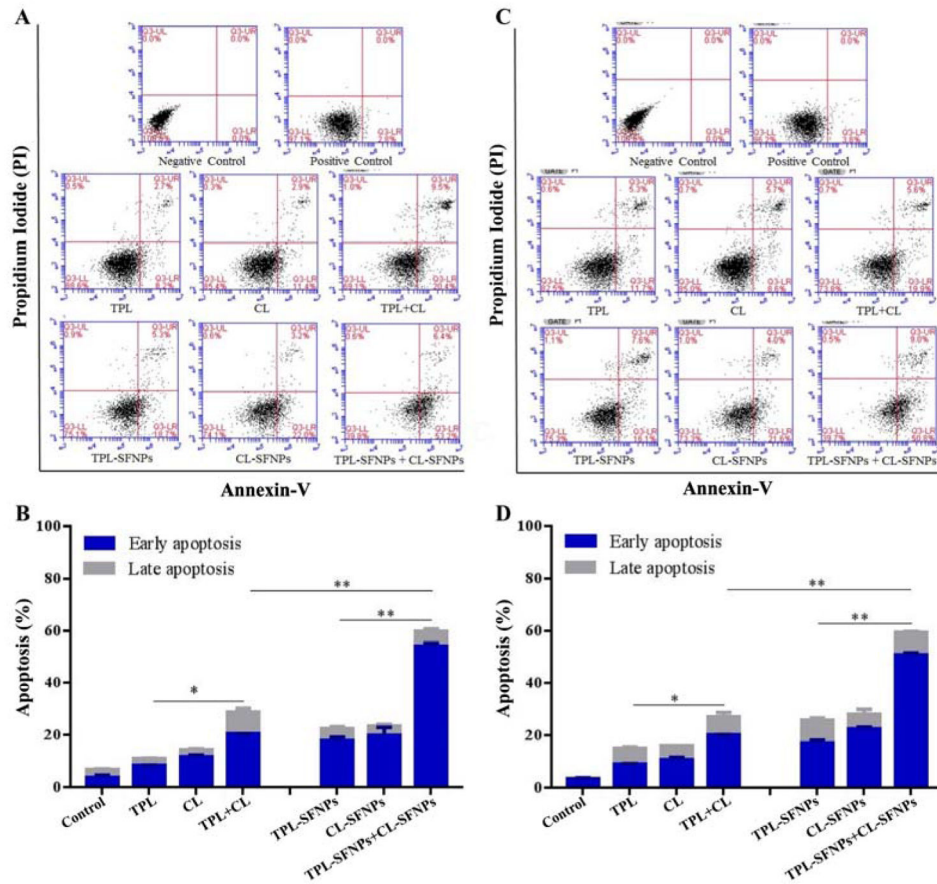


Fig. 11. Apoptotic effects of CL-SFNPs and TPL-SFNPs in MIA PaCa-2 and PANC-1 cell line
 A–B. Determination of apoptosis in MIA PaCa-2 cell line after treatment with equivalent dose of CL, TPL individually and in combination and CL-SFNPs and TPL-SFNPs individually and in combination using flow cytometer. C–D. Determination of apoptosis in PANC-1 cell line after treatment with equivalent doses of CL, TPL individually and in combination and CL-SFNPs and TPL-SFNPs individually and in combination using flow cytometer. (Mean \pm SD, n = 3; *: P<0.05, **: P<0.01)

Particle size, zeta potential, encapsulation efficiency, drug loading and β -sheet content of SF, SFNPs, TPL-SFNPs, CL-SFNPs (n=3)

Table 1

Sample	Particle size (nm \pm s.d)	Zeta Potential (mV \pm s.d)	EE (% \pm s.d)	DL (μ g/mg \pm s.d)	β -sheet content
SF	-	-	-	-	16.8
SFNPs	201.0 \pm 5.1	-21.1 \pm 3.3	-	-	35.3
TPL-SFNPs	166.4 \pm 4.6	-27.2 \pm 2.0	81.8 \pm 2.8	57.7 \pm 4.7	57.7
CL-SFNPs	170.4 \pm 2.3	-25.5 \pm 2.6	87.0 \pm 1.1	63.5 \pm 3.8	77.6

Novel combination of triptolide and celastrol loaded silk fibroin nanoparticles show synergistic anti-pancreatic cancer effect *in vitro*