Improving the solubility and in vitro cytotoxicity (anticancer activity) of ferulic acid by loading it into cyclodextrin nanosponges

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Purpose: Ferulic acid (FA) is a poorly water-soluble natural antioxidant with anticancer activity. This poor solubility limits the application of FA in the food and pharmaceutical industry. Cyclodextrin nanosponges (CD-NSs) are a novel group of cross-linked CD derivatives which can be used to enhance the solubility of low-soluble bioactive compounds.

Methods: In this study, FA was encapsulated into the NSs in the proportion of 1:4 (FA:NS). Diphenyl carbonate was used as a cross-linker in different proportions with β-CD. Characterization of obtained NSs was performed using scanning electron microscopy, X-ray diffraction (XRD), differential scanning calorimetry (DSC), and Fourier transform infrared (FTIR) analysis.

Results: Our results revealed that the solubility of encapsulated FA was increased up to fifteenfold compared with pure FA in the proportion of 1:4 (CD:cross-linker). The results of FTIR, XRD, and DSC confirmed the interaction of FA with NSs. The cytotoxicity of encapsulated FA against MCF7 and 4T1 breast cancer cell lines was investigated using different concentrations of FA in 24, 48, and 72 hrs. The cytotoxicity assay indicated that FA treatment reduced viability and enhanced apoptosis of cancer cells. IC50 value of encapsulated FA (250 ppm) was decreased by threefold when compared with pure FA (750 ppm).

Conclusion: In general, CD-NS was found to be a suitable delivery system for poorly soluble bioactives such as FA.

Keywords: nanoencapsulation, nanosponges, cyclodextrins, ferulic acid, cytotoxicity

Introduction

4-hydroxy,3-methoxy cinnamic acid known as ferulic acid (FA) is a monophenolic compound which is found plentifully in plant cell walls, such as grains, vegetables, and fruits.1 Also, it can be found in the beverages such as beer and coffee.2 FA is present as a free form or linked to hemicellulosic polysaccharides (in grain) or hydroxy acids (in fruits and vegetables).3 It is described to have antioxidant, antimicrobial, antiangiogenic, anticancer, antidiabetic, wound healing, cholesterol-lowering, and anti-inflammatory activities.2,4-7 In Japan, FA has been approved as a sunscreen due to its photoprotective properties.8 The antioxidant activity of FA is related to its phenolic nucleus and unsaturated side chain which contributes to the formation of resonance compounds and acts as an antiproliferative agent.5

There are many evidences about the antitumor activity of FA against cervical,9 breast,6 colon,10 prostate,11 and osteosarcoma.12 FA performs its anticancer activity by interference in cell cycle and apoptotic pathways11 or inhibition of metastasis in cancer
Breast cancer is the first reason of cancer mortality in women. The major strategy for the recovery and treatment of cancer is radiotherapy but because of its undesirable efficacy on normal tissues and also other limitations, such as chemoresistance, immune escape, metastasis, and recurrence, researchers are trying to find new drugs from natural plant-based sources.\textsuperscript{5,6}

Natural antioxidants such as FA have indicated cytotoxic traces in various cancers either alone or combined with other bioactive compounds.\textsuperscript{13} Despite of the many beneficial effects of FA, its application has been limited similar to other hydrophobic phenolic compounds due to low water solubility, low bioavailability, and phytochemical instability.\textsuperscript{14,15} Therefore, some efforts have been made to improve the solubility and bioavailability of FA using different nanocarriers such as biopolymeric nanoparticles\textsuperscript{6,7} or electrospun nanofibers.\textsuperscript{16–18} Panwar et al (2016) encapsulated FA into chitosan nanoparticles and showed that the solubility of FA was increased by 28% and 25% in water and acetic acid (1%, v/v), respectively.\textsuperscript{5} Other researchers have also encapsulated poorly soluble compounds, such as biopolymeric nanoparticles,\textsuperscript{7} chitosan-polyacrylate nanofibers,\textsuperscript{18} or poly (d,l-lactide-co-glycolide)/polyethylene oxide nanofibers,\textsuperscript{16} but the solubility of FA was not investigated.

Cyclodextrins (CDs) are oligosaccharides with a cyclic structure that have a hydrophilic outer surface and a hydrophobic central part.\textsuperscript{19,20} Therefore, they can be used to encapsulate both hydrophobic and hydrophilic bioactive compounds.\textsuperscript{21–23} Cyclodextrin-based nanosponges (CD-NSs) are nanoporous structures which are obtained through hyper-cross-linking of the CDs using different cross-linkers. Recently, CD-NSs are specially used as innovative nanodelivery vehicles to enhance the water solubility of poorly soluble compounds,\textsuperscript{24,25} to improve their bioavailability,\textsuperscript{26} and to prolong the release of encapsulated compounds.\textsuperscript{27} CD-NSs can form colloidal nanosuspensions in aqueous solutions and enhance the release of bioactive compound.\textsuperscript{28} CD-NSs have a higher capability than the natural CDs to improve the water solubility of hydrophobic bioactive compounds.\textsuperscript{28,29} Some of the advantages of CD-NSs compared to other solubilization technologies are that CD-NSs are solid particles and can easily be used in formulations, their production is easy and affordable and also have a higher loading capacity than other nanocarriers.

The aim of this study was to encapsulate FA into the CD-NSs for the first time and then, to characterize the obtained NSs in terms of encapsulation efficiency, structure, and physicochemical properties. Also, the toxicity of encapsulated FA against MCF7 and 4T1 cancer cell lines was investigated.

### Materials and methods

B-CD was provided by SD-fine corporation, India. FA and diphenyl carbonate were obtained from Sigma Aldrich (Sigma, USA). DMEM, FBS, PBS, EDTA, and trypsin were purchased from Biodia corporation, Iran. Penicillin-streptomycin and 3 (4,5-dimethylthiazol-2-yl)-2,5- diphenyltetrazolium bromide (MTT) were procured from Sigma-Aldrich (Sigma, USA). MCF7 (human breast cancer) and 4T1 (mouse breast cancer) cell lines were purchased from Pasteur Institute of Iran. All other chemicals and reagents were of analytical grade.

#### Preparation of β-CD-NSs

Three types of β-CD-NSs were produced according to the different molar ratios of β-CD to diphenyl carbonate as cross-linker (1:2, 1:4, and 1:8).\textsuperscript{28} Briefly, determined amounts of anhydrous β-CD were reacted with calculated amounts of melted diphenyl carbonate at 90°C for at least 5 hrs. Once the reaction was completed, the obtained compound was cooled at room temperature and washed by extra amount of double distilled water to eliminate unreacted β-CD. After this, the obtained product was Soxhlet extracted with ethanol up to 4 hrs to remove unreacted or impurities of diphenyl carbonate. After purification, the nanospores were dried under vacuum. This reaction also was done in with sonication for 10 mins.

#### Preparation of FA-loaded NSs

Obtained NSs from the previous stage were dissolved into distilled water and then calculated amounts of FA were added into the solution in a w/w ratio of 4:1 (NS:FA). Then, this mixture was stirred for 24 hrs in dark conditions at room temperature. The uncomplexed FA was separated by centrifuging the suspension at 2,000 rpm for 10 mins; the supernatant which was containing the complexed (encapsulated) FA was finally freeze dried (Dena Vacuum, Iran) at ~40°C for 24 hrs.

### Determination of encapsulation efficiency and loading efficacy

Specified amounts of FA-NSs as well as fixed amounts of NSs were dissolved in 10 mL ethanol and then sonicated for 10 mins at ambient temperature, individually. Next, the
solutions were filtered through a 0.45 μm filter. The amount of FA was obtained by reading the solution absorbance using a UV spectrophotometer at 319 nm. The dispersion of NS was used as the blank sample. Encapsulation efficiency and loading efficacy of FA within NSs were determined according to the following equations:

$$EE = \frac{\text{Weight of loaded FA in NS}}{\text{weight of initial used FA for loading}} \times 100$$ (1)

$$\text{Loading efficacy} = \frac{\text{Weight of loaded FA in NS}}{\text{total weight of FA - NS}} \times 100$$ (2)

**Analysis of solubility**

An excess amount of FA-NS as well as a fixed amount of NS were dispersed in 10 mL distilled water and agitated by a stirrer for 24 hrs at 100 rpm and room temperature, separately. Further, the suspensions were centrifuged at 10,000 rpm for 10 mins. The absorbance of the supernatant was read at 319 nm after filtration through a 0.45 μm filter. The dispersion of NS was used as the blank sample.

**Instrumental analysis of FA-loaded nanosponges**

The morphology of FA-NSs was investigated by scanning electron microscopy (SEM). The samples were fixed onto the glass slides and prepared by sputtering with a thin film of gold before imaging (HITACHI S-4160, Japan). Particle size was specified using dynamic light scattering by a Zetasizer ZEN 3600 (Malvern, UK). The apparatus worked at a fixed temperature of 25°C with a scattering angle of 90°. Suitable dilution of the samples was prepared with ethyl acetate before measurement.

Fourier transform infrared (FTIR) spectroscopy analysis of FA, NS, FA-NSs, and diphenyl carbonate were done with a Tensor 27 spectrometer (Bruker, Germany) using the ATR pellet method. The frequency range and the resolution of the instrument were 4000–600 cm⁻¹ and 4 cm⁻¹, respectively. The X-ray diffraction (XRD) patterns of FA, NS, and FA-NSs were determined using a Philips X’Pert diffractometer and Cu Kα radiation (λ=1.54 Å). The operating range was 2θ =10–60° and step size was 0.05°. The XRD spectra were analyzed using X’Pert and Origin software. Differential scanning calorimetry (DSC) (STA 503, Germany) was applied for analyzing the thermal properties of FA, NS, and FA-NSs. For the DSC analysis, the heating rate was 20°C/min and the samples heated from 25°C to 350°C under nitrogen gas.

**In vitro release of FA**

FA release from NSs was studied using a dialysis membrane (molecular weight cutoff between 12,000 and 14,000 Da, Himedia Lab, India) in phosphate buffer (pH=6.8) at 37°C and 100 rpm in a shaking thermostatic incubator. To study the FA release, 15 mg of FA-NS was dispersed in 2 mL phosphate buffer and then poured into the dialysis tube. Then, the dialysis tube was suspended into the 30 mL phosphate buffer. At specified time intervals, 5 mL of the solution was removed for sampling and substituted with the same volume of the release medium to keep a fixed volume. The absorbance of the samples was recorded by UV spectrophotometer at 319 nm after filtration through a 0.45 μm filter.

The cumulative amount (%) of released FA from NSs was obtained using the following equation:

$$FA \text{ release} \% = \frac{\text{The amount of released FA}}{\text{Total amount of FA in the NS}} \times 100$$ (3)

**Cell cytotoxicity of FA-loaded nanosponges**

The cell cytotoxicity of FA-NSs, blank NS, and free FA were specified using MTT assay in MCF7 (human breast cancer) and 4T1 (mouse breast cancer) cell lines. Briefly, cells were grown in 90% DMEM supplemented with 10% FBS, sodium bicarbonate, L-glutamine, non-essential amino acid, sodium pyruvate, 100 mg/mL streptomycin, and 100 U/mL penicillin incubated at the temperature of 37°C and under 5% CO₂ atmosphere. Further, the cells were gathered using 0.25 w/v% trypsin-EDTA solution and seeded in a 96-well culture plate at a density of 5×10³ cell/well and incubated for 24 hrs to confirm the attachment of the cells. Then, 100 μL of different concentration (5, 10, 20, 40, 80, 125, 250, 500, 750, and 1,000 μM) from FA-NS, free FA, and blank NS in the medium culture were added into the wells and incubated for 24, 48, and 72 hrs for evaluating a time and dose-depending manner. Next, cell viability was
estimated by MTT assay at 570 nm. Untreated cells were taken as control. The cell viability was calculated as:

\[
\text{Cell viability} = \left( \frac{OD_{570\text{nm}} \text{treated cells}}{OD_{570\text{nm}} \text{control}} \right) \times 100
\]

Finally, half maximal inhibitory concentration (IC50) of FA-NSs and free FA was obtained.

Statistical analysis

All the experiments were performed in triplicates and data were expressed as mean value± standard deviation. Statistical analysis was done using the SPSS program (version 20). The significance of difference \((P<0.05)\) was analyzed using one-way ANOVA.

Results and discussion

Solubility and encapsulation efficiency of FA loaded into nanosponges

Three different proportions of CD:cross-linker (1:2, 1:4, 1:8) were used for the preparation of nanosponges. The solubility, loading capacity, and encapsulation efficiency of FA were evaluated to find the suitable proportion of CD: cross-linker. As can be seen in Figure 1, the solubility of FA increased after encapsulation into the NSs. Also, FA solubility was enhanced by increasing the proportion of CD:cross-linker from 1:2 to 1:4. This higher solubility may be due to entrapment of FA as an inclusion complex within CDs and in the porous structure of NSs. In the proportion of 1:4, the higher cross-linker resulted in a more porous structure compared to lower proportions (1:2). By increasing the proportion up to 1:8, the solubility was decreased which could be due to the hyper cross-linking of CDs preventing the interaction of FA with inclusion cavity. Also, it can be due to saturation solubility of FA in NSs. The proportion of 1:4 (CD: cross-linker) produced NSs with the highest solubility of FA; the solubility increased up to 15 folds compared with free FA.

The loading capacity and encapsulation efficiency of FA are presented in Table 1. By increasing the proportion of CD:cross-linker from 1:2 to 1:4, loading capacity and encapsulation efficiency were remarkably enhanced which may be due to the more porous structures resulted from the higher cross-linking; while by increasing the proportion to 1:8, encapsulation efficiency and loading capacity did not change significantly \((P>0.05)\) possibly because of the hyper cross-linking of CDs.

According to the higher encapsulation efficiency, loading capacity and also solubility results, the proportion of 1:4 (CD:cross-linker) was selected to produce the NSs for encapsulation of FA and the characterization stages at the following steps.

Characterization results of FA loaded into nanosponges

SEM and particle size

SEM images of FA-NSs are presented in Figure 2. As can be seen, the obtained NSs had a porous structure. The particle size analysis indicated that 78.4% of the samples with sonication treatment and 88.2% the samples without sonication had a particle size around 369±27 and 443±44 nm, respectively. Therefore, the sonicated samples which had a lower particle size were selected for further analysis.

Results of FTIR analysis

The FTIR spectra of the samples are presented in Figure 3. Characteristic peaks of FA were detected at 3,437 cm\(^{-1}\) (-OH stretching), 1,692 cm\(^{-1}\) (C=O stretching vibration), 1,619,

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<tr>
<th>Proportion of CD:cross-linker</th>
<th>Encapsulation efficiency of FA (%)</th>
<th>Loading capacity of FA (%)</th>
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<tr>
<td>1:2</td>
<td>33.33</td>
<td>16.85</td>
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<tr>
<td>1:4</td>
<td>45.75</td>
<td>25.7</td>
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<td>1:8</td>
<td>44.5</td>
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Table 1 Encapsulation efficiency and loading capacity of ferulic acid (FA) at different proportions of cyclodextrin (CD): cross-linker.

Figure 1 Solubility of ferulic acid (FA) at the different proportions of cyclodextrin: cross-linker. Abbreviation: NS, nanosponge.
1,514 cm\(^{-1}\) (C=C aromatic ring), and 1,272, 1,205 cm\(^{-1}\) (C-H bending modes). These results were similar to the FTIR peaks presented earlier for FA.\(^{15}\) In the spectra of CD and diphenyl carbonate, the characteristic peaks of each component are differentiable. The NS spectrum is highly similar to CD spectrum and their difference is only in the
intensity of peaks. In the spectrum of FA-NS, the characteristic peaks of FA with a little shifting (1,683, 1,626, 1,595, 1,518, 1,269, and 1,206 cm$^{-1}$) are visible which indicate FA has been properly incorporated into the NSs.

XRD results

Figure 4 indicates the XRD pattern of FA, NS, and FA-NSs. The sharp characteristic peaks of FA revealed its crystalline structure (Figure 4B). The XRD pattern of FA-NS (Figure 4C)
has different crystalline structures compared with its original constituents, namely, FA and NS (Figure 4A) which confirms the complex formation. Furthermore, the reduction in the intensity and difference in diffraction peaks and interplanar spacing indicates successful encapsulation of FA into the nanoporous structures of NS.

DSC results
Thermal properties of FA, NS, and FA-NSs were determined using DSC and are depicted in Figure 5. FA showed an endothermic point at 174.2°C related to the melting point of FA followed by an exothermic effect corresponding to the thermal decomposition at higher temperatures (Figure 5B). Thermal analysis of NS and FA-NSs indicated that their thermograms are very similar and showed an endothermic point at 297.9°C due to the CD dehydration and decomposition. The components of NS did not show separate endothermic points which reveals the formation of NSs (Figure 5A). FA-NSs did not show the endothermic point of FA that again confirms FA has been completely included into the NSs and also there is a tight interaction between CD and FA (Figure 5C).²⁶,⁸

Results of in vitro release of FA
The in vitro release profile of FA from NSs is presented in Figure 6. As can be seen, the NSs have been resulted in a controlled and slow release of FA. The initial burst release in the first hour (around 15%) indicated surface loading of FA and then followed by a slow release up to 70% after 24 hrs. This slow release may be due to the inclusion complexation of FA into the NSs and also cross-linking structure of NSs; this confirms the optimum complexation between FA and NSs. Other researchers have also reported a slow release of encapsulated compounds from NSs.⁸,³²

In vitro cytotoxicity results of FA loaded into nanosponges
The in vitro cytotoxicity of FA, FA-NSs, and blank NS was investigated against MCF7 (human breast cancer) and 4T1 (mouse breast cancer) cell lines and cell viability was specified using MTT assay (Figure 7). For this purpose, the mentioned cancer cell lines were treated with different concentrations of FA-NSs at different times (24, 48, 72 hrs). The concentrations of 5, 10, 20, 40, and 80 µM of FA-NS did not have a significant effect (P>0.05) on the viability of cells; therefore, we used higher concentrations. Our results revealed that the toxicity of FA-NSs was concentration-dependent and by increasing the concentration of FA-NS up to 500 µM, the cell viability decreased; the higher concentrations did not have a significant influence (P>0.05) on cell viability. Other researchers have also indicated that the higher concentration of phenolic compounds such as FA may be unprofitable and even result in oxidative damage of ordinary cells because these compounds can act as pro-oxidants in high concentrations.²⁹,³⁵ Also, the toxicity of FA-NSs was time dependence meaning that by increasing the time from 24 to 72 hrs, the cell viability was decreased. As can be seen in Figure 7, 250 µM of FA-NS in both MCF7 and 4T1 cell lines caused around 50% inhibition of cell proliferation (IC₅₀). For free FA, the concentration of 750 µM was determined as IC₅₀. Panwar et al (2016) indicated that ME-180 cervical cancer cells after treatment with 40 µM of FA-loaded chitosan nanoparticles showed a 50% inhibition in their growth. This low concentration was due to the synergistic effect of FA and chitosan nanoparticles. Our results indicated that encapsulation of FA into the NSs enhanced its toxicity and decreased the inhibitory concentration. Encapsulation of FA into the NSs leads to a higher solubility and stability of FA and may have contributed to higher anticancer properties of FA. The higher solubility of FA-NSs may be resulted in higher cell permeability and also higher toxicity.³⁴ In vitro cytotoxicity of blank NS revealed that it was nontoxic.

Figures 8 and 9 show the impact of different formulations on cell morphology after 24 hrs. The morphology of cancer cells treated with NS was very similar to
control and displayed a normal morphology (Figure 8A and C and Figure 9A and C). Our results revealed that the morphology of cancer cell lines which were treated with 250 µM of free FA (Figures 8B and 9B) did not show a significant change compared with the control cells; while for the cancer cell lines treated with 250 µM of FA-NS (Figures 8D and 9D), they had significant structural changes compared with other formulations and also induced 50% cell death. Panwar et al (2015) reported that the cytotoxicity of encapsulated FA into the chitosan nanoparticles was higher than the free FA against cervical cancer cell lines and resulted in cell apoptosis along with cytoplasmic remnants and damaged wrinkled cells. Other researchers have also indicated that FA caused cell apoptosis, enhanced lipid peroxidation, affected on the mitochondrial activity, disrupted the cell organelle such as mitochondria and resulted in the cell death.

**Conclusion**

In this study, we successfully encapsulated FA into the NSs which resulted in a significant improvement of the FA solubility up to 15 folds compared with the free form of FA. The physicochemical and structural characterization
using FTIR, XRD, and DSC revealed that FA was properly encapsulated into the NS structures. MTT assay showed an enhanced antiproliferation activity of FA-NS as compared to free FA against MCF7 and 4T1 breast cancer cell lines. In vitro release data indicated a slow controlled release of FA from NSs. Our results confirmed that CD-NS is a suitable nano delivery system for FA which could increase its cytotoxicity, solubility, and anticancer potential.

Figure 8 MCF7 cell lines treated with different formulations: control cells (A); ferulic acid, 250 µM (B); nanosponges, 250 µM (C); ferulic acid-loaded nanosponges, 250 µM (D). Microscopic magnification ×10 K, after 24 hrs of incubation.

Figure 9 4T1 cell lines treated with different formulations: control cells (A); ferulic acid, 250 µM (B); nanosponges, 250 µM (C); ferulic acid-loaded nanosponges, 250 µM (D). Microscopic magnification ×10 K, after 24 hrs of incubation.
Disclosure

The authors report no conflicts of interest in this work.

References


