Preparation and antioxidant properties of selenium nanoparticles-loaded chitosan microspheres

Kaikai Bai¹,²
Bihong Hong¹,²
Jianlin He¹,²
Zhuan Hong¹,²
Ran Tan¹,²

¹Third Institute of Oceanography, ²Engineering Research Center of Marine Biological Resource, Comprehensive Utilization, State Oceanic Administration, Xiamen, People's Republic of China

Abstract: Selenium nanoparticles (SeNPs), as a special form of selenium (Se) supplement, have attracted worldwide attention due to their favorable properties and unique bioactivities. Herein, an eco-friendly and economic way to prepare stable SeNPs is introduced. SeNPs were synthesized in aqueous chitosan (CTS) and then embedded into CTS microspheres by spray-drying, forming selenium nanoparticles-loaded chitosan microspheres (SeNPs-M). The physicochemical properties including morphology, elemental state, size distribution and surface potential were investigated. Institute of Cancer Research mice were used as model animal to evaluate the bioactivities of SeNPs-M. Trigonal-phase SeNPs of ~35 nm were synthesized, and SeNPs-M physically embedding those SeNPs were successfully prepared. Amazingly, acute toxicity test indicated that SeNPs-M were much safer than selenite in terms of Se dose, with a LD₅₀ of around 18-fold of that of selenite. In addition, SeNPs-M possessed powerful antioxidant activities, as evidenced by a dramatic increase of both Se retention and the levels of glutathione peroxidase, superoxide dismutase and catalase. The design of SeNPs-M can offer a new way for further development of SeNPs with a higher efficacy and better biosafety. Thus, SeNPs-M may be a potential candidate for further evaluation as an Se supplement with antioxidant properties and be used against Se deficiency in animals and human beings.

Keywords: selenium, nano, microsphere, chitosan, antioxidant

Introduction

As a dietary nutrient, Se is an indispensable trace element required for most of the living organisms including animals and human beings.¹ Se is present in at least 25 human selenoproteins and enzymes as selenocysteine, which plays an essential role in preventing various diseases such as cardiovascular disease, hypercholesterolemia and certain cancers.¹,² To meet the daily requirement of Se, extra Se supplementation is necessary, especially for those having Se deficiency.¹,² Some Se compounds in both inorganic and organic forms, such as selenite, selenomethionine and Se-methylselenocysteine, have been used for decades to avoid Se deficiency in animals and human beings.² However, these Se supplements, especially the inorganic ones, are usually toxic when taken above their nutritional dosage.²

In the recent years, SeNP, a kind of elemental Se particle at a nano-size scale with a bright red color, has aroused worldwide attention due to its unique properties and excellent biological activities.²⁻⁵ It not only scavenges the free radicals in vitro⁶ but also improves growth, serum oxidant status and Se concentration in vivo.³⁻⁵ Amazingly, compared with Se compounds such as selenite,⁵ selenomethionine,⁴ Se-yeast⁵ and Se-methylselenocysteine,⁷ SeNPs exhibit much lower acute toxicity while increasing...
the activities of selenoenzymes.\textsuperscript{2–5} Besides, SeNPs are able to inhibit the growth of microorganisms.\textsuperscript{8} They also exhibit antitumor activities both in vitro\textsuperscript{9,10} and in vivo,\textsuperscript{11} by inducing mitochondria-mediated apoptosis\textsuperscript{9} and stimulating immune reaction against cancer cells.\textsuperscript{11} Thus, SeNPs are considered as a prospective Se formulation for nutritional supplement use, chemoprevention and chemical therapy against cancer.

However, the application of SeNPs is limited by the following facts: (1) economic and environment-friendly ways to synthesize SeNPs are needed\textsuperscript{10,12–14} and (2) bare SeNPs usually enlarge, aggregate and finally transform into a gray/black analog that is thermodynamically stable, but biologically inert.\textsuperscript{15,16} Actually, even an increase in the size of SeNPs can lead to significant reduction in the activities of these nanoparticles such as scavenging multiple radical species in vitro\textsuperscript{5} and enhancing Se accumulation and GSH S-transferase activity in vivo.\textsuperscript{17} It has also been proved that SeNPs are most effective as a chemopreventive agent at a smaller particle size.\textsuperscript{17} Thus, it seems that only SeNPs with a smaller size can guarantee high bioactivities.

Over the past decade, much effort had been devoted to synthesize and stabilize SeNPs with the use of polysaccharides,\textsuperscript{15,18–20} monosaccharides,\textsuperscript{21} proteins,\textsuperscript{22–24} amino acids,\textsuperscript{25} polyphenols,\textsuperscript{26–28} melatonin,\textsuperscript{29} ATP,\textsuperscript{30} complex plant extract\textsuperscript{10,31} and even microorganism culture.\textsuperscript{8} Among them, polysaccharide is regarded as an appropriate template for fabricating SeNPs when considering energy efficiency and eco-friendliness, according to Zhang et al.\textsuperscript{15} However, the SeNPs prepared using polysaccharides were found to be unstable, due to enlargement of size and decrease of zeta potential during storage in aqueous environment.\textsuperscript{15,19} Moreover, SeNPs preserved using aqueous polysaccharide often encounter and form aggregates. Hence, an improved method of synthesizing SeNPs is needed.

CTS is the only positively charged natural polysaccharide, having excellent biodegradable and biocompatible characteristics.\textsuperscript{32} It has been extensively examined in the pharmaceutical industry for its potential in the development of medicine delivery systems.\textsuperscript{32} Herein, a novel form of SeNPs, named SeNPs-M, was designed and manufactured by two steps: (1) SeNPs were synthesized in the presence of CTS, and (2) the resultant CTS-SeNPs were embedded into CTS microspheres by spray-drying method to form SeNPs-M. Each SeNP was physically separated and preserved into solid SeNPs-M. Besides, the physical and chemical characteristics of SeNPs-M were investigated for a deeper understanding of the bioactivities of SeNPs-M to evaluate the application of SeNPs in this from.

### Materials and methods

#### Reagents and animals

CTS (90.32% deacetylated, average molecular weight of 37kDa) of food grade was purchased from Aoxin Pharmaceutical Co. Ltd (Taizhou, People’s Republic of China). The assay kits for measuring TBARS, GSH, CAT, SOD, GSH-Px and protein were provided by Jiancheng Bio-engineering Institute (Nanjing, People’s Republic of China). Acetic acid, sodium selenite, Vc and other reagents of analytical reagent-grade quality were obtained from Sigma-Aldrich (Shanghai) Trading Co. Ltd (Shanghai, People’s Republic of China).

ICR mice of specific-pathogen-free grade, 4–5 weeks old, 18–22 g bw, were supplied by Xiamen University Animals Laboratory Center (Xiamen, People’s Republic of China) with the license no SCXK 2013-0001. The procedures used in animal experiments were approved by the Animal Ethics Committee of Xiamen University, and were also compliant with the Provisions and General Recommendation of Chinese Experimental Animals Administration Legislation.

#### Synthesis of SeNPs and SeNPs-M

SeNPs were synthesized by reducing selenite as described by Zhang et al\textsuperscript{17} with small modification. In brief, 0.5 g of CTS and 1.6 g of Vc were dissolved in 100 mL of 1% (w/w) acetic acid. Then, 10 mL of selenite aqueous solution containing 0.4 g of sodium selenite was slowly added to theCTS/Vc solution and vigorously stirred (600–800 rpm) to obtain a CTS-SeNPs colloid (shown in Scheme 1A). The SeNPs could be stable at 25°C for at least 4 weeks, without any visible precipitation. Bare SeNPs were gained by replacing CTS with deionized water during the preparation of CTS-SeNPs.

Consequently, the CTS-SeNPs colloid was dialyzed against 1% (w/w) acetic acid for 6 hours to remove the excess Vc and other by-products. After that, the colloid was well mixed with another CTS solution, with the final concentrations of Se and CTS being 0.09% (w/w) and 2.5% (w/w), respectively. The spray-drying process was applied to evaporate the moisture of the new mixture (shown in Scheme 1A).\textsuperscript{33} A laboratory spray dryer (SY-6000; Shiyuan Biological Equipment Engineering Co. Ltd., Shanghai, People’s Republic of China) with a standard 0.7 mm nozzle was used under the working condition: inlet temperature 160°C, pumping flow rate 500 mL/hour, spray air pressure 0.35 kg/cm² and drying air flow discharge 600 m³/hour. Thus, SeNPs-M were generated with the expected structure shown in Scheme 1B. The Blank-M without any SeNP was prepared by replacing selenite with deionized water. The Se content of each sample was determined by ICP-MS assay according to Dufailly et al.\textsuperscript{34}
Morphology observation and EDS

TEM (JEM-2100; JEOL, Tokyo, Japan) was applied to study the morphological characteristics of SeNPs. Briefly, diluted SeNPs solution was dropped onto copper grids and dried in clean air. TEM observation was carried out at an accelerating voltage of 200 kV. Additionally, the details of microspheres were observed by SEM (S-4800; Hitachi, Tokyo, Japan). The microspheres were coated with platinum in a vacuum state, and the photographs were taken at an accelerating voltage of 2–10 kV. In particular, samples were frozen at −30°C and then cut into frozen sections by a Microtome Cryostat (CM1850; Leica, Wetzlar, Germany), each 6 μm thick. The frozen sections were observed by SEM. Meanwhile, EDS was applied during TEM and SEM.

FTIR

Each sample was dried and then ground into a homogeneous powder with potassium bromide to record the infrared spectra on a Nicolet Nexus 470 spectrometer (Thermo Fisher Scientific, Waltham, MA, USA). The spectra were acquired at 400–4,000 cm⁻¹ wavenumbers with a 4 cm⁻¹ resolution.

XPS

XPS measurement was conducted by a high-resolution photoelectron spectrograph (Escalab 250Xi; Thermo Fisher Scientific) which was equipped with a monochromatic Al Kα X-ray source featuring a highly focused beam size of 500×500 μm. The energy resolution was 0.05 eV. A dual-beam charge neutralization system composed of a low-energy electron flood gun (~1 eV) and an Argon ion gun (≤10 eV) was used.

XRD

The XRD patterns were recorded with an X-ray diffractometer (X’Pert Pro MPD system; Philips, Amsterdam, the Netherlands), using a Cu Kα source (λ=0.15406 nm). Besides, crystallographic identification was accomplished by comparing the experimental XRD patterns with those of Joint Committee on Powder Diffraction Standards (JCPDS) database.¹³

Measurement of particle size and surface charge

Hydrodynamic sizes and zeta potentials of nanoparticles were measured using a Zetasizer Nano ZS particle analyzer (VEM3600; Malvern Instruments, Malvern, UK) with a 173°
scattering angle according to Kong et al. The measurements based on DLS theory were performed under the following conditions: particle refractive index 1.590, particle absorption coefficient 0.01, dispersant (water) refractive index 1.33 and temperature 25°C. In addition, another particulate size analyzer (LS-POP(6); Zhuhai OMIC Instruments Co. Ltd., Zhuhai, People’s Republic of China) was utilized to determine the size distribution of microspheres, with a particle reflective index of 1.70 and fluid reflective index of 1.33.

Animal experiments
ICR mice were used to evaluate the bioactivities of samples in two experiments. In each experiment, mice were housed in plastic cages (only one gender each) in a room with controlled temperature (25°C ±2°C) and humidity (50±10%) and a 12-hour light/dark cycle. After the oral administration of candidate samples, mice were allowed free access to low-Se diet (<0.1 mg Se/Kg diet) and water.

For the first experiment, the acute lethal properties of sodium selenite, Blank-M and SeNPs-M were measured. After adaption for 3 days, 110 ICR mice were randomly divided into 11 groups with 10 mice per group. Among them, one group was orally administrated saline as control, while the other groups were intragastrically administered sodium selenite or SeNPs-M as shown in Table 1. Cumulative mortality within 14 days after the treatment was recorded to calculate median lethal dose (or LD₅₀) by Bliss method. Besides, another group of 20 ICR mice was given Blank-M at an accumulated dose of 15 g/kg bw within 24 hours to determine the MTD of Blank-M.

In the second experiment, the bioactivities of SeNPs-M were evaluated. Forty-eight male mice were randomly divided into six groups with eight mice in each group. The mice were orally administered saline, sodium selenite, Blank-M and SeNPs-M, respectively, at the doses shown in Table 2 with the frequency of once per 2 days. After seven consecutive Se treatment cycles, the mice were fasted overnight and then were sacrificed to obtain their blood and livers. Se in the blood or liver was determined by ICP-MS assay. Moreover, liver samples were homogenized in ice-cold saline and centrifuged at 12,000×g at 4°C for 15 minutes. The resulting supernatants and the plasma samples were used to determine the levels of TBARS, GSH, CAT, SOD, GSH-Px and total protein, following the instructions of commercial kits.

Statistical analysis
In all the experiments, data were presented as mean ± SD. The difference between two groups was analyzed by Student’s t-test, while the difference between three or more groups was analyzed by one-way analysis of variance (ANOVA) with Tukey’s post hoc test. All statistical analysis was performed using GraphPad Prism (version 8.0 for Windows). A P-value of <0.05 was considered statistically significant.

Results and discussion
Morphology and formation of CTS-SeNPs
Many methods were applied to prepare elemental Se. Thus, Se obtained were of various morphologies, such as amorphous, sphere, wire, rod and tube. Herein, aqueous Se(IV) was chemically reduced by Vc to synthesize SeNPs in the presence of CTS. The initial colloid nucleated and then

<table>
<thead>
<tr>
<th>Sodium selenite</th>
<th>SeNPs-M</th>
</tr>
</thead>
<tbody>
<tr>
<td>Se dose (mg/kg bw)</td>
<td>Mouse mortality (%)</td>
</tr>
<tr>
<td>11.0</td>
<td>100</td>
</tr>
<tr>
<td>5.9</td>
<td>70</td>
</tr>
<tr>
<td>3.2</td>
<td>50</td>
</tr>
<tr>
<td>1.7</td>
<td>20</td>
</tr>
<tr>
<td>0.9</td>
<td>0</td>
</tr>
</tbody>
</table>

Notes: *mg/kg bw means mg Se per kilogram of body weight. **LD₅₀ refers to median lethal dose. *The number in brackets represents LD₅₀ of 95% confidence interval.

Abbreviations: ICR, Institute of Cancer Research; Se, selenium; SeNPs-M, selenium nanoparticles-loaded chitosan microspheres; bw, body weight.

<table>
<thead>
<tr>
<th>Group</th>
<th>Plasma Se (µg/mL)</th>
<th>Hepatic Se (µg/g liver)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>0.57±0.27*</td>
<td>1.36±0.09*</td>
</tr>
<tr>
<td>Blank-M</td>
<td>0.49±0.23*</td>
<td>1.29±0.15*</td>
</tr>
<tr>
<td>SeNPs-M (0.5)</td>
<td>0.71±0.23*</td>
<td>2.18±0.39*</td>
</tr>
<tr>
<td>SeNPs-M (2)</td>
<td>0.61±0.16*</td>
<td>4.12±1.45*</td>
</tr>
<tr>
<td>SeNPs-M (8)</td>
<td>1.46±0.47*</td>
<td>8.35±1.08*</td>
</tr>
<tr>
<td>Sodium selenite (2)</td>
<td>1.12±0.26*</td>
<td>5.49±1.27*</td>
</tr>
</tbody>
</table>

Notes: Data were presented as mean ± SD. *The number in brackets refers to the dose of selenite or SeNPs-M, based on Se dose (mg Se/kg body weight). **Means within a row with different letters differ significantly (P<0.05).

Abbreviations: ICR, Institute of Cancer Research; Blank-M, blank chitosan microspheres; SeNPs-M, selenium nanoparticles-loaded chitosan microspheres; SD, standard deviation.
assembled into SeNPs (Scheme 1A). In this process, CTS adhered to Se atoms and controlled the size of their aggregates. Finally, monodispersed and homogeneous spherical CTS-SeNPs were obtained (Figure 1A). The colloid’s color was dependent on the concentration of SeNPs (Figure 1A inset). Besides, the size distribution analysis conducted using TEM and DLS calculated the average diameters of CTS-SeNPs to be 36 and 95 nm, respectively (Figure 1B). It suggested that there is an invisible layer covering SeNPs since DLS can only measure hydrodynamic size.\(^{31}\)

Elemental formation and microstructure of CTS-SeNPs were studied. Clear Se peaks (1.37, 11.22 and 12.49 keV) were found in the EDS spectra, confirming the elemental nature of SeNPs (Figure 1C). Besides, the structure of SeNPs synthesized in 0.5 mg/mL CTS solution was analyzed by high-resolution transmission electron microscopy. As shown in Figure 1D inset, similar crystalline structures with a lattice fringe of 0.35 nm were observed as compared with the two fringes of 0.149 and 0.31 nm obtained in the presence of 0.08 mg/mL CTS and 0.1% (w/w) CTS, respectively.\(^{15,37}\)

Figure 1 (Continued)
Moreover, the selective area electron diffraction patterns demonstrated the orderliness and symmetry of CTS-SeNPs (Figure 1D). It seemed that CTS was helpful to form crystall Se with a nano size. Its concentration seemed to have played an important role in forming various crystalline structures.

The impacts of CTS on the shape and size of SeNPs were also studied. Fresh CTS-SeNPs were monodispersed spherical particles with a high uniformity (Figure 1E), whereas bare SeNPs aggregated to bulks in defect of CTS (Figure 1E inset). A higher concentration of CTS resulted in smaller SeNPs (Figure 1F), similar to the fact that higher concentration of BSA generally leads to SeNPs of a smaller size.\(^6,17\) Obviously, the morphology of SeNPs could be easily mediated by CTS. A smaller size of CTS-SeNPs, however, might be preferred instead of the bigger ones, since better thermostability,\(^38\) more powerful antioxidant activities,\(^6\) and more potent chemopreventive abilities\(^17\) could be achieved.

**Stability of CTS-SeNPs**

Stability is an important factor for nanomaterials.\(^{15,16}\) SeNPs decorated by sialic acid\(^{21}\) or gum arabic\(^{19}\) could be stable for at least 30 days. CTS-SeNPs reported by Yu et al\(^{17}\) remained stable for 60 days when stored in 0.08 mg/mL CTS solutions, but the size dramatically increased from 120 to 400 nm and aggregation began to occur. In this study, bare SeNPs (without CTS) quickly agglomerated in few days, while CTS-SeNPs were stable for more than 28 days in 1\% (w/w) CTS solution at 25°C (Figure 2A). However, CTS-SeNPs turned to black bulks after 42 days of storage (Figure 2A), accompanied by the enlargement of CTS-SeNPs (Figure 2B). Similar instability of SeNPs within gum arabic solution was reported by Kong et al.\(^{19}\) Furthermore, the TEM images of CTS-SeNPs were studied. Enlarged size of original CTS-SeNPs was very common at the 28th day (Figure 2C), while many aggregates with the size of several micrometers were generated after 42 days of storage (Figure 2D). The aggregates comprising numerous single SeNPs were actually dark bulks that appeared during the long-term preservation. However, the narrow size distribution, highly faceted morphology and uniform edge-to-edge distances between particles induced by heating 40 nm SeNPs\(^{38}\) were not found among these enlarged SeNPs shown in Figure 2C and D. Perhaps, sharp reconstruction of original SeNPs could not be triggered under such a mild storage condition (25°C). However, size enlargement and agglomeration of SeNPs seemed to be unavoidable in an aqueous environment.

The zeta potentials of SeNPs were compared for further comprehension of the stability. As presented in Figure 2E, bare SeNPs were slightly negative (−4.3 mV) in the reducing water environment, but the surface decoration by CTS elevated...
the zeta potential of SeNPs to be +38.6 mV, probably due to the positively charged $\sim\text{NH}_3^+$ groups from CTS.$^{15,25,39}$ It was why the fresh CTS-SeNPs were able to remain very stable. However, the zeta potential decreased from 40.8 mV at the initial day to 18.4 mV at the 28th day (Figure 2F), indicating that CTS-SeNPs become more unstable during storage. Similar results were reported by Kong et al.$^{19}$ and Zhang et al.$^{15}$ Considering the relationship between nanoparticles’ stability and their zeta potential, it was difficult to preserve CTS-SeNPs in an aqueous solution for a longer time.

Figure 2 The effects of CTS upon SeNPs. CTS-SeNPs and bare SeNPs were synthesized in the presence and absence of CTS, respectively. (A) Photographs of SeNPs in the presence or absence of CTS during a 42-day storage. Day 0: immediately after sample synthesis day. (B) Size distribution of CTS-SeNPs after 0, 14, 28 and 42 days, respectively. The data of 42nd day were obtained using LS-PD60 particle analyzer, while other data within the panel were recorded by VeM3600 particle analyzer. (C) TEM image of CTS-SeNPs after a 28-day storage. (D) TEM image of CTS-SeNPs after a 42-day preservation period. (E) Zeta potentials of new bare SeNPs and fresh CTS-SeNPs recorded by VEM3600 particle analyzer. (F) Zeta potentials of CTS-SeNPs during a 42-day preservation period (n=3). **$P<0.01$ versus day 0, and ***$P<0.01$ versus day 14.

Abbreviations: Se, selenium; CTS, chitosan; SeNPs, selenium nanoparticles; TEM, transmission electron microscopy.
Characterization of SeNPs-M

Bare SeNPs and CTS-SeNPs were unavailable for commercial application in oral administration system since they were not very stable. To address this problem, SeNPs were maintained in solid state. As illustrated in Scheme 1A, CTS-SeNPs were spray-dried, and SeNPs were finally embedded into solid SeNPs-M. White Blank-M (without any Se) and red SeNPs-M were obtained, both of which were free-flowing powders. The Se content of SeNPs-M was determined by ICP-MS, and it could be adjusted by modifying the material ratio. An SeNPs-M sample (29 g Se/kg) was selected to conduct the rest of the study.

The morphology of SeNPs-M was studied at first. SeNPs-M were a collection of regular spheres with a smooth surface, while Blank-M were more wrinkled (Figure 3A vs Figure 3A inset). It suggested that SeNPs play an unknown role in forming SeNPs-M. Besides, the average size of SeNPs-M was calculated as 6.24 μm in water and 4.21 μm in ethanol (Figure 3B). Apparently, SeNPs-M might swell in water. Moreover, EDS analysis was performed on both the surfaces and the cross sections of SeNPs-M to study the Se distribution. As shown in Figure 3C, apparent Se information can hardly be found on the surfaces; however, clear Se signals were observed on the transections (Figure 3C inset). Thus, there is no doubt that Se was distributed inside SeNPs-M.

Additionally, the spectral features of SeNPs-M were investigated. The FTIR spectra of SeNPs-M were similar to that of Blank-M, without any new chemical structure (Figure 3D).

Figure 3 (Continued)
However, significant difference was observed in the XPS patterns of Se 3d and 3p orbitals (Figure 3E and F). Typical Se 3d peaks of Se (0) and Se (IV) were detected at 55.3 and 59.5 eV, respectively, confirming that the valence state of Se in SeNPs-M was zero, though it needed an Argon ion etching lasting 60 seconds before Se XPS detection to expose the area under the surfaces of the microspheres. Interestingly, the characteristic peaks of $-\text{NH}_2$ and $-\text{NH}_3$ displayed clear variation between SeNPs-M and Blank-M (Figure 3G). This was consistent with some previous studies reporting that two chemical states of nitrogen could be observed by N 1s XPS.

Finally, XRD was conducted to analyze the amorphous/crystalline nature of SeNPs. Typical XRD peaks of crystal Se were found in the XRD patterns of SeNPs-M, while Blank-M were amorphous (Figure 3H). These peaks were
in accordance with the trigonal phase of Se having lattice
constants \(a=4.366 \text{ Å}, b=4.366 \text{ Å} \) and \(c=4.954 \text{ Å}\), which corre-
tioned to the reported value (JCPDS File No 06-0362). Similar
XRD patterns had been found by Sharma et al\(^{31}\) when
using dried \textit{Vitis vinifera} (raisin) to prepare SeNPs, whereas
different XRD patterns of SeNPs mediated by 3 or 200 kDa
CTS had been reported by Zhang et al.\(^{15}\) This suggested that
manufacturing condition may greatly affect the crystalline
feature of SeNPs.

Based on the characteristics of SeNPs-M, nano-crystal
SeNPs were successfully embedded into solid CTS micro-
spheres by spray-drying, generating globular SeNPs-M,
which might guarantee the physical isolation of each SeNP
(shown in Scheme 1B).

**Acute toxicity**

In the acute lethal test, SeNPs-M and sodium selenite at
increasing doses (1.78-fold and 1.86-fold, respectively)
were administered by single intragastric administration, and
mortality were recorded over 14 days. The results presented
in Table 1 illustrated that selenite at a dose of 11.0 mg Se/kg
bw killed all the mice. However, SeNPs-M did not cause any
mortality at a dose of 14.6 mg Se/kg bw. The \(LD_{50}\) of selenite
and SeNPs-M was 3.4 and 62.3 mg Se/kg bw, respectively,
indicating that the acute toxicity of SeNPs-M was only 1/18
of that of selenite based on Se dose. Considering the safety
of Blank-M with an MTD exceeding 15 g/kg bw (data not
shown), it was not difficult to deduce that it was SeNPs within
the SeNPs-M that were responsible for the acute toxicity.
This was consistent with some previous studies performed on
mice and rat.\(^{40-42}\) Remarkably, SeNPs in the form of SeNPs-M
were much safer than selenite.

CTS might be superior to BSA\(^{40,41}\) in terms of the safety
of SeNPs (18-fold vs sevenfold as compared to selenite). In
addition to the different species of experiment animals, the
ability to release SeNPs might account for the difference
between these two forms of SeNPs. As a hydrosoluble pro-
tein, BSA could be easily dissolved and enzymolysized in the
digestive tract of animals;\(^{43}\) thus, the SeNPs might be released
quickly, inducing the acute toxicity. However, the release of
SeNPs might be retarded by the absorption characteristics
of CTS towards many materials\(^{32}\) and the deficiency of the
enzymes to deal with CTS in some species of animals and
human beings.\(^{32}\)

**Growth inhibition**

The bw of mice was recorded and compared during the
treatment with saline (as control), sodium selenite, Blank-M
and SeNPs-M for 14 consecutive days. It was reported that
selenite at the dose of 6 mg Se/kg bw resulted in complete
growth suppression.\(^3\) As shown in Figure 4A, however,
mice generally kept growing during the administration of

![Figure 4](https://www.dovepress.com/)

**Figure 4** The body weights of male ICR mice during a 14-day treatment period of different selenium source. Se-deficient mice were orally administered saline (Control), sodium selenite (Selenite, 2 mg Se/kg bw), blank microspheres (Blank-M) and SeNPs-M once per 2 days for 14 consecutive days. (A) The change of body weight during SeNPs-M treatment (n=8). (B) The impacts of different Se sources on the last body weights (n=8). The number in brackets refers to the dose of selenite or SeNPs-M, based on Se dose (mg Se/kg bw). *\(P<0.05\) versus control. \(^\text{a-c}\) Means within a panel with different letters differ significantly (\(P<0.05\)).

**Abbreviations:** bw, body weight; Blank-M, blank chitosan microspheres; ICR, Institute of Cancer Research; SeNPs-M, selenium nanoparticles-loaded chitosan microspheres.
SeNPs-M, though this Se form at a high dose of 8 mg Se/kg bw might suppress the growth of mice (P<0.05 vs control). After the last treatment, it was found that sodium selenite (2 mg Se/kg bw) significantly inhibited the growth of mice (Figure 4B). It induced more growth inhibition than SeNPs-M at the same Se dose. No statistical significance (P>0.05) could be found between SeNPs (2 mg Se/kg bw) and control. Considering the safety of Blank-M shown in Figure 4B, it was clear that (1) SeNPs in SeNPs-M were responsible for the growth inhibition of mice and (2) that SeNPs-M (≤2 mg Se/kg bw) were safe while selenite at the same dose was toxic.

Se retention
In order to investigate the Se distribution within the body of mice, the Se contents of the blood and liver were determined. As shown in Table 2, it was evident that (1) SeNPs in SeNPs-M contributed to the Se retention, (2) Se retention occurred in a dose-dependent manner, and (3) Se retention of SeNPs-M was weaker than that of sodium selenite. The last result was different from some reports5,44 reporting the greater Se retention of nano-elemental Se (called SeNPs in the present study) over selenite in broiler chickens and male goats, respectively. It implied that the Se retention in animals is complex and is affected by a variety of animal species. Possibly weaker accumulation of Se could partly explain the low toxicity of SeNPs-M compared with selenite according to Wang et al.4 Oral administration of SeNPs-M could be an effective way to supply Se.

Biomarkers
The levels of GSH, TBARS (MDA equivalent), GSH-Px, SOD and CAT were measured to evaluate the antioxidant activities of SeNPs-M. As shown in Table 3, sodium selenite (2 mg Se/kg bw) increased TBARS content and decreased GSH content vs control (P<0.05). Compared with selenite, however, SeNPs-M at the same Se dose not just reduced TBARS levels but elevated GSH levels. This was consistent with some previous studies,3-5 which had confirmed the advantages of SeNPs over selenite. In particular, SeNPs-M significantly enhanced plasma or hepatic GSH-Px activity in a dose-dependent manner, resulting in some twofold increase of GSH-Px activities in Se-deficient mice as compared with control. Based on the data in Table 3, SeNPs-M and selenite had a similar efficacy in increasing GSH-Px activity in the blood and liver of mice. Similar results had been observed by Zhang et al.15 and by Shi et al.4 using mice and goats as the model animal to study nano-elemental Se (SeNPs) controlled by BSA, respectively. Probably, similar GSH-Px activity could be achieved if the amount of Se was equal, no matter whether the SeNPs form was decorated by CTS or BSA. Nevertheless, GSH-Px activity could be saturated at nutritional levels, and further supplementation with supranutritional levels of Se did not further increase its activity.17 It could explain why increasing the supplementation of SeNPs-M did not necessarily result in further increase of GSH-Px (Table 3). With respect to other enzymes, selenite (2 mg Se/kg bw) had a little effect on SOD and CAT, but SeNPs-M were helpful in increasing the activities of SOD and CAT even at a low dose of 0.5 mg Se/kg bw. More enzymatic activities could be achieved by SeNPs-M vs selenite. In short, SeNPs-M outweighed selenite in terms of antioxidant activities.

Conclusion
In this study, we introduced a simple way to synthesize and preserve SeNPs. Crystal SeNP spheres were manufactured in the presence of CTS and then were embedded into CTS microspheres by using spray-drying method, giving birth to SeNPs-M. With an LD50 of around 18-fold of that of selenite, SeNPs-M were much safer but had an equal efficacy in increasing GSH-Px activity in vivo when compared with

Table 3 Effects of different Se sources upon antioxidant activities within ICR mice (n=8)

<table>
<thead>
<tr>
<th>Group</th>
<th>Blood</th>
<th>Liver</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>GSH (mg/L)</td>
<td>TBARS (nmol/L)</td>
</tr>
<tr>
<td>Control</td>
<td>11.20±1.58a</td>
<td>0.49±0.08a</td>
</tr>
<tr>
<td>Blank-M</td>
<td>12.95±2.57a</td>
<td>0.46±0.07a</td>
</tr>
<tr>
<td>SeNPs (0.5)a</td>
<td>11.85±3.10b</td>
<td>0.43±0.12c</td>
</tr>
<tr>
<td>SeNPs (2)a</td>
<td>17.99±4.82a</td>
<td>0.44±0.08a</td>
</tr>
<tr>
<td>SeNPs (8)a</td>
<td>13.99±3.26b</td>
<td>0.51±0.14a</td>
</tr>
<tr>
<td>Sodium selenite (2)a</td>
<td>8.30±2.77c</td>
<td>0.94±0.27a</td>
</tr>
</tbody>
</table>

Notes: Data were presented as mean ± SD. aThe number in brackets refers to the dose of selenite or seNPs-M, based on Se dose (mg Se/kg body weight). bMeans within a row with different letters differ significantly (P<0.05).

Abbreviations: ICR, Institute of Cancer Research; GSH, glutathione; TBARS, thiobarbituric acid-reactive substances; GSH-Px, glutathione peroxidase; SOD, superoxide dismutase; CAT, catalase; Blank-M, blank chitosan microspheres; SeNPs-M, selenium nanoparticles-loaded chitosan microspheres.
selenium. At the same time, levels of intracorporal Se, GSH, SOD and CAT were enhanced by SeNPs-M, while lower level of TBARS (MDA equivalent) was achieved. In summary, this CTS microspheres design of SeNPs opens a new path for oral delivery of Se with a higher efficacy and better biosafety. SeNPs-M can be a candidate Se source worthy of further development for nutrient supplement.

**Abbreviations**
Blank-M, blank chitosan microspheres; BSA, bovine serum albumin; bw, body weight; CAT, catalase; CTS, chitosan; DLS, dynamic light scattering; EDS, energy-dispersive X-ray spectroscopy; FTIR, Fourier transform infrared spectroscopy; GSH, glutathione; GSH-Px, glutathione peroxidase; ICR, Institute of Cancer Research; ICP-MS, inductively coupled plasma mass spectrometry; MDA, malondialdehyde; MTD, maximum tolerated dose; SEM, scanning electron microscopy; Se, selenium; SeNPs, selenium nanoparticles; SeNPs-M, selenium nanoparticles-loaded chitosan microspheres; SOD, superoxide dismutase; TBARS, thiobarbituric acid-reactive substances; TEM, transmission electron microscopy; Vc, ascorbic acid; XPS, X-ray photoelectron spectroscopy; XRD, X-ray diffraction.

**Acknowledgments**
This work was supported by grants from Scientific and Technological Projects of Fujian Province, People’s Republic of China (No 2016N0018), the Scientific Research Foundation of Third Institute of Oceanography, State Oceanic Administration, People’s Republic of China (No 20140101), the Science & Technology Major Projects of Fujian province (No 2014NZ0001) and the Co-sponsored Project by Xiamen Oceanic Research and Development Institute. The authors also thank the support from Southern Ocean Research Center Project of Xiamen and the Co-sponsored Project by Xiamen Oceanic Research and Development Institute. The authors thank the support from Southern Ocean Research Center Project of Xiamen Oceanic Research and Development Institute. The authors also thank the support from Southern Ocean Research Center Project of Xiamen Oceanic Research and Development Institute. The authors also thank the support from Southern Ocean Research Center Project of Xiamen Oceanic Research and Development Institute.

**Disclosure**
The authors report no conflicts of interest in this work.

**References**