

Targeting experimental orthotopic glioblastoma with chitosan-based superparamagnetic iron oxide nanoparticles (CS-DX-SPIONs)

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Background: Glioblastoma is the most devastating primary brain tumor of the central nervous system in adults. Magnetic nanocarriers may help not only for a targeted delivery of chemotherapeutic agents into the tumor site but also provide contrast enhancing properties for diagnostics using magnetic resonance imaging (MRI).

Methods: Synthesized hybrid chitosan-dextran superparamagnetic nanoparticles (CS-DX-SPIONs) were characterized using transmission electron microscopy (TEM) and relaxometry studies. Nonlinear magnetic response measurements were employed for confirming the superparamagnetic state of particles. Following in vitro analysis of nanoparticles cellular uptake tumor targeting was assessed in the model of the orthotopic glioma in rodents.

Results: CS-DX-SPIONs nanoparticles showed a uniform diameter of 55 nm under TEM and superparamagnetic characteristics as determined by T_1 (spin-lattice relaxation time) and T_2 (spin-spin relaxation time) proton relaxation times. Application of the chitosan increased the charge from +8.9 to +19.3 mV of the dextran-based SPIONs. The nonlinear magnetic response at second harmonic of CS-DX-SPIONs following the slow change of stationary magnetic fields with very low hysteresis evidenced superparamagnetic state of particles at ambient temperatures. Confocal microscopy and flow cytometry studies showed an enhanced internalization of the chitosan-based nanoparticles in U87, C6 glioma and HeLa cells as compared to dextran-coated particles. Cytotoxicity assay demonstrated acceptable toxicity profile of the synthesized nanoparticles up to a concentration of 10 µg/ml. Intravenously administered CS-DX-SPIONs in orthotopic C6 gliomas in rats accumulated in the tumor site as shown by high-resolution MRI (11.0 T). Retention of nanoparticles resulted in a significant contrast enhancement of the tumor image that was accompanied with a dramatic drop in T_2 values ($P < 0.001$). Subsequent histological studies proved the accumulation of the nanoparticles inside glioblastoma cells.

Conclusion: Hybrid chitosan-dextran magnetic particles demonstrated high MR contrast enhancing properties for the delineation of the brain tumor. Due to a significant retention of the particles in the tumor an application of the CS-DX-SPIONs could not only improve the tumor imaging but also could allow a targeted delivery of chemotherapeutic agents.

Keywords: brain tumor, glioblastoma, tumor targeting, chitosan, superparamagnetic iron oxide nanoparticles, magnetic resonance imaging

Introduction

Despite multimodal treatment approach for glioblastoma multiforme that currently includes surgical resection of the tumor with radio- and pharmacotherapy with temozolomide (TMZ), the survival of patients does not exceed 15 months.^{1,2} Introduction of the tumor-treating fields (TTFields) as locoregional treatment only slightly improved

the median overall survival in subgroups of patients, up to 20.5 months.³ These data indicate the necessity for novel diagnostic and therapeutic modalities in neuro-oncology. Presumably, theranostic approaches could be based on the chitosan nanoparticles that show a high biocompatibility at relatively low costs. Previously, several studies reported the application of chitosan particles as nanocarriers of anti-glioma agents.^{4–10} Thus, Liu et al⁴ showed that chitosan surface-modified poly(lactide-co-glycolide)–chitosan nanoparticles (PLGA/CS NPs) loaded with carmustine (bis-chloroethylnitrosourea, BCNU) and its sensitizer (O⁶-benzylguanine, BG) had a strong antitumor effect in the F98 glioma-bearing mice model. In another study by Lollo et al,¹¹ the authors could demonstrate similar effects with paclitaxel (PTX)-loaded particles in GL261 glioma-bearing mice. Subsequent functionalization of the chitosan nanoparticle surface with bioligands further enhanced the ability of the particles to cross the blood–brain barrier (BBB) and their brain tumor accumulation level. Up-to-date, various ligands were proposed for decoration of the surface, including iRGD (tumor targeting peptide [CRGDKGPDC]) with high affinity to α v integrins,⁴ transferrin receptor targeting agents,⁵ and lactoferrin.¹²

Alternative strategies based on chitosan particles could be their application, not only for the delivery of antitumor agents, but also for the detection of glioma. For the imaging of the glioblastoma in the current work, dextran-coated superparamagnetic iron oxide nanoparticles (DX-SPIONs) that were further coupled with chitosan were produced. The coupling of chitosan to preliminary activated dextran magnetic nanoparticles (MNPs) can increase the surface charge of the latter and thus improve the tumor uptake of SPIONs. Hence, as shown by Barrow et al,¹³ an increase in charge of the 40 kDa dextran-based iron oxide nanoparticles (IONPs) from -1.5 to $+18.2$ mV provided a sixfold increase in uptake in mouse mesenchymal stem cells (MSCs) with respect to neutral dextran-coated particles. Furthermore, the hybrid chitosan–dextran-coated nanoparticles exert stronger membrane interactions as compared to monodextran particles.¹⁴ Due to their unique magnetic and optical characteristics, such as high paramagnetism, low magnetic coercivity, and high Curie temperature, SPIONs show favorable characteristics compared to other types of magnetic nanocarriers.¹⁵ Magnetic properties of the SPIONs were also employed for the development of the contrast agents for the magnetic resonance imaging (MRI) of brain tumors.^{16,17}

In our study, the tumor targeting ability and the MRI contrast-enhancing properties of the synthesized hybrid dextran–chitosan SPIONs (CS-DX-SPIONs) were assessed in an orthotopic C6 Wistar rat model. In this study, we report

that chitosan-based MNPs could be successfully employed for the MRI detection of brain tumors.

Materials and methods

Synthesis of the DX-SPIONs

The superparamagnetic (SPM) nanoparticles were obtained from iron salt solutions by coprecipitation at 80°C in alkaline media, as described previously.¹⁸ Briefly, FeSO₄ and FeCl₃ (Fe²⁺/Fe³⁺ ratio of 1:2) salts were dissolved together with CsCl in water.¹⁹ The magnetite precipitation was induced in an inert atmosphere by titration with NH₄OH solution and collected using a permanent magnet. The nanoparticle suspension was treated by ultrasound at 22 kHz for 15 minutes. To prevent the sedimentation of the nanoparticles, 10 kDa molecular weight (MW) dextran (Sigma-Aldrich Co., St Louis, MO, USA) was added into dispersion in the process of ultrasound application. The obtained SPION stock solution was washed and separated into fractions by centrifugation and microfiltration using 0.2 μ m pore size membranes (EMD Millipore, Billerica, MA, USA). The concentration of Fe in the suspension was assessed by ultraviolet (UV) absorption (at $\lambda=480$ nm) of the thiocyanate–Fe(+3) complex.

Preparation of the hybrid chitosan nanocarriers (CS-DX-SPIONs)

About 94% deacetylated chemical-grade chitosan (MW 30 kDa) was prepared from crab shells in formate solution at pH 6–7.4. DX-SPIONs were transferred into glass and aminated with epichlorohydrin. Chitosan was dissolved in 100 mL of water under magnetic stirring at pH 7.0. Hybrid nanoparticles were prepared by ionotropic gelation by mutual interaction of chitosan and dextran with negatively charged ion tripolyphosphate (TPP) (Vekton, St. Petersburg, Russia). An aqueous solution of TPP was added dropwise to the chitosan solution with suspended SPIONs. The hybrid nanoparticles were formed instantly after addition of TPP. The tertiary composition in reaction medium was as follows: polyvinylpyrrolidone (MW 3.6 MD; BASF), polyvinylpyrrolidone (PVP) (3% g/mL), TPP (0.04% g/mL), and chitosan (0.25% g/mL).

Physicochemical characterization of CS-DX-SPIONs

The particle size of SPIONs and size distribution were analyzed by transmission electron microscopy (TEM) using the JEOL-2000 microscope (JEOL, Tokyo, Japan) and dynamic light scattering (DLS) with the help of Malvern Instruments (Malvern, UK). Electrophoretic properties and hydrodynamic size were measured on Zetasizer Nano (Malvern Instruments). The relaxation times T_1 , T_2 , and T_2^*

and ^1H nuclear magnetic resonance (NMR) spectra were measured with the NMR spectrometer CXP-300 (Bruker Optik GmbH, Ettlingen, Germany) at the magnetic field of 7.1 T (corresponding to 300 MHz for ^1H). The NMR spectra were registered with a one-pulse sequence by applying Fourier transform of free induction decay. The T_2^* values were estimated from line width at half height. To assess the T_1 and T_2 relaxation times, the inversion recovery and Carr–Purcell–Meiboom–Gill (CPMG) pulse sequences were employed. Proton relaxation times in buffer solution were recorded as a function of magnetic conjugate concentration. The R_1 , R_2 , and R_2^* (relaxivity) coefficients of relaxation efficiency were defined from the slopes of concentration plots. The ^{31}P NMR spectra were obtained on a Bruker CXP-300 at 121 MHz using a 5 mm phosphorus probe and at 162 MHz on a WM 400 spectrometer using a 10 mm broadband probe. Phosphorus chemical shifts are referred relative to H_3PO_4 solution at pH 2. The ^{31}P spin–lattice relaxation rate constants were obtained from 10 to 12 relaxation intervals using the inversion recovery pulse sequence at 120 MHz without proton decoupling.

Magnetic characterization of CS-DX-SPIONs

To characterize the magnetic state of the CS-DX-SPIONs (single-/multi-domain) and to estimate their average magnetic moment, we employed our home-built setup, registering the nonlinear longitudinal response (NLR) of MNPs to a weak *ac* magnetic field $h(t) = h \sin(\omega t)$ parallel to steady field H (the frequency $f = \omega/2\pi = 15.7$ MHz and $h = 13.8$ Oe). The setup included two coaxial Helmholtz coils (providing the linear scanning of steady field H with frequency, F_{sc} , varied in the range $8\text{--}10^{-2}$ Hz) combined with the electromagnet (for the creation of a constant field H_0 if it is necessary) and a transmitter–receiver high frequency coil of a two-mode resonant system. The resonant system, on the one hand, creates *ac* field on a sample under study and, selects from the NLR the second harmonic M_2 of the magnetization (NLR- M_2) generated by a sample.²⁰ The M_2 signal components, ie, $\text{Re}M_2(H)$ and $\text{Im}M_2(H)$, are recorded simultaneously as functions of H . The frequency of H -scan, F_{sc} , can be changed within $8\text{--}0.02$ Hz, which allows the testing of the state (single-/multi-domain) of MNPs. Indeed, in the multi-domain state, the magnetization reversal is determined by movement of domain boundaries, which is fast, and H -hysteresis will not depend on F_{sc} . However, in single-domain MNPs, the field hysteresis appears in the blocking regime below blocking temperature, T_b . The latter can be found from condition $\tau = \tau_{mes}$ (where τ is the time of the magnetic relaxation of MNPs ensemble and τ_{mes} is the time of measurements), and in our

experiments, it depends on F_{sc} . Indeed, in the case of the uniaxial anisotropy applicable to our MNPs, $1/\tau = f_0 \exp(-E_b/kT)$ (where anisotropy energy $E_b = KV$, k is the Boltzmann constant, K is the anisotropy constant, V is an average volume of a particle, and the frequency f_0 is of the order of 10^9 s $^{-1}$) and $T_b = KV/[k \cdot \ln(f_0/F_{sc})]$.²¹ The dependence of H -hysteresis on F_{sc} evidences the single-domain state of MNPs in our study. Usually, NLR- M_2 measurements are carried out under the condition $M_2 \propto h^2$, which allows the application of the perturbation theory for analysis of the results. The main contribution to $\text{Re}M_2(H)$ arises due to the nonlinear behavior of $M(H)$, so in the regime of the MNP ensemble close to the SPM one (a weak H -hysteresis/an hysteretic of $M_2(H)$ response), the $\text{Re}M_2(H)$ dependence can be semiquantitatively described by $\partial^2 L(H)/\partial H^2 + k_{pM}H$, where $L(H)$ is the Langevin function and the term $k_{pM}H$ takes into account possible small linear contributions from low-molecular paramagnetic fraction.²² The $\text{Re}M_2(H)$ reaches extreme values near the inflection point of $M(H)$, which is dependent on the average moment of MNPs that allows to estimate the latter. This description is not quantitative since it does not take into account the effect of the magnetic field on relaxation processes. This could be done by utilizing the Gilbert–Landau–Lifshitz formalism, but this is out of the scope of our study.²³

Assessment of cellular interactions of CS-DX-SPIONs

Cells

All employed cell lines (rat C6 glioma, human U87 glioma, and human cervix carcinoma HeLa cells) were obtained from the Russian Cell Culture Collection of the Institute of Cytology of the Russian Academy of Sciences (St Petersburg, Russia). C6 and HeLa cells were cultured in a CO_2 incubator (at 37°C , 6% CO_2) in DMEM culture medium supplemented with 2 mM L-glutamine, 10% fetal bovine serum (FBS), and antibiotics (ie, streptomycin [100 units/mL] and penicillin G [100 $\mu\text{g}/\text{mL}$]). Human U87 glioma cells were harvested in complete RPMI-1640 culture medium (Sigma-Aldrich Co.) supplemented with antibiotics and 10% FBS.

Cytotoxicity assay

For the analysis of cytotoxicity of CS-DX-SPIONs, the commercial non-radioactive cytotoxicity assay kit, CytoTox 96[®] (Promega Corporation, Fitchburg, WI, USA), was employed in accordance with the manufacturer's protocol. Briefly, tumor cell lines (ie, C6 glioma, U87, and HeLa cells) were cocultured (for 1, 3, 12, and 24 hours) with SPIONs or hybrid CS-DX-SPIONs taken at different Fe concentrations (of 1, 5, 10, 50, and 150 $\mu\text{g}/\text{mL}$). The obtained supernatants from

the samples were analyzed for the concentration of lactate dehydrogenase (LDH).

Confocal microscopy

For the assessment of cellular interactions of synthesized nanoparticles with tumor cells (ie, C6 glioma, HeLa cells, and U87 cells [1×10^6 cells/mL]), the latter were permitted to settle on poly-L-lysine-coated glass slides. SPIONs or CS-DX-SPIONs (Fe concentration: $150 \mu\text{g/mL}$) were coincubated with the tumor cells for 1, 3, 12, and 24 hours in a CO_2 incubator (37°C , 6% CO_2). Prior to cell imaging, the nuclei of the cells were stained with 4,6-diamidine-2-phenylindole (DAPI) solution (Vector Laboratories, Burlingame, CA, USA). A series of the fluorescence *z*-images were recorded using a Leica TCS SP5 confocal microscope (Leica Microsystems, Heidelberg, Germany).

Flow cytometry analysis

For assessment of the nanoparticle internalization by tumor cells, a flow cytometer equipped with a coherent red solid state diode (635 nm), uniphase argon laser (488 nm), side scatter detector (SSD), and forward scatter detector (FSD), Cytomics FC500 (Beckman Coulter, Indianapolis, IN, USA) was used. The setups of the cytometer measured forward scattering (FSC) and side scattering (SSC) linearly and logarithmically. After an incubation period of 24 hours of the tumor cells (C6, U87, and HeLa) with DX-SPIONs, CS-DX-SPIONs (Fe concentration of $150 \mu\text{g/mL}$), or phosphate buffered saline (PBS), the cells were analyzed by flow cytometry. The scatter parameters derived from the gated scatter region were used for counting cells within a large SSC (log) versus FSC (linear) histogram.

Gel-phantom model

For the assessment of the magnetic properties of the CS-DX-SPIONs following cellular uptake, the C6 glioma cells were coincubated with DX-SPIONs (Fe concentration: $150 \mu\text{g/mL}$), CS-DX-SPIONs (Fe concentration: $150 \mu\text{g/mL}$), or PBS for 24 hours. Following coincubation and washing with PBS, the cells were embedded into 1% agarose gel. For the assessment of the intracellular nanoparticle localization, several anatomical high-resolution turbo spin echo (SE) T_2 -weighted coronal images (20 slices in total) were obtained using the following parameters: repetition time (TR)/echo time (TE) 4,200/36 ms, flip angle (FA) 180° , slice thickness of 1.0 mm, interslice distance 1.2 mm, field of vision (FOV) 3.0×3.0 cm, and matrix 256×256 . Subsequently, gradient echo (GE) scans (FLASH) (TR/TE 350/5.4 ms, FA 40° , slice thickness 1.0 mm, 3.0×3.0 cm, and matrix 256×256) and SE T_1 -weighted images (TR/TE 1,500/7.5 ms, FA 180° , slice thickness 1.0 mm, FOV 3.0×3.0 cm, and matrix 256×256)

were obtained in the coronal plane. Accumulation of the DX-SPIONs or CS-DX-SPIONs in the samples was assessed using multi-scan multi-echo (MSME) sequences.

Orthotopic C6 glioma model

Male Wistar rats with a body weight of 280–300 g were obtained from the “Rappolovo” RAMN animal nursery (St Petersburg, Russia). For all animal experiments, the approval has been provided by the local ethical committee of First Pavlov Saint Petersburg State Medical University (St Petersburg, Russia) in accordance with institutional guidelines for the welfare of animals. Before mounting in a stereotactic frame (David Kopf Instruments, Tujunga, CA, USA), the rats were anesthetized with 0.2 mL 2% Rometar (xylazine hydrochloride; “Bioveta”, Ivanovice na Hané, Czech Republic) and 10 mg tiletamine hydrochloride and zolazepam (“Virbac santé Animale”, Carros, France) intraperitoneally. C6 rat glioma cells (1×10^6 cells in 10 μL of PBS) were stereotactically infused into the nucl. caudatus dexter (right caudate nucleus) using a Hamilton microsyringe.

Magnetic resonance assessment of CS-DX-SPIONs' tumor-targeting potential

On the 15th day following the intracranial injection of the C6 tumor cells for the assessment of the accumulation of DX-SPIONs or CS-DX-SPIONs in the tumor site, the animals were randomly divided into three groups as follows (six rats in each group): 1) control group – intravenous (i.v.) injection of PBS via the tail vein; 2) i.v. administration of DX-SPIONs (Fe concentration of 2.5 mg/kg); and 3) i.v. administration of the CS-DX-SPIONs (Fe concentration of 2.5 mg/kg). Twenty-four hours after i.v. injection, intratumoral retention of MNPs was assessed using a high-field MRI scanner (11.0 T) (Avance III 400 WB; Bruker Optik GmbH) in the following conventional sequences: 1) T_1 -weighted fast-field echoes (TR/TE [TR] 1,500/7.5 ms, FOV 3.0×3.0 cm, matrix 256×256 , FA 180° number of signals averaged [NSA]); 2) T_2 -weighted turbo SE (FA 180° , TR 4,200 ms, and TE 36 ms, FOV 3.0×3.0 cm, matrix 256×256); 3) FLASH scans (GE) (FA 40° , TR 350 ms, and TE 5.4 ms); and 4) MSME regimen. All images were obtained with a slice thickness of 1.0 mm in the coronal plane. Assessment of the MRI scans was carried out using the Paravision 3.1 software package (Bruker BioSpin GmbH, Rheinstetten, Germany) and Analyze software (AnalyzeDirect, Inc., Overland Park, KS, USA).

Histology

For histological analysis, the rats were sacrificed using an intraperitoneal injection of pentobarbital (150–200 mg/kg)

with subsequent perfusion with PBS (100 mL) and 4% paraformaldehyde (PFA). For the analysis of the intratumoral distribution of nanoparticles, the obtained serial cryosections (5–7 μm) were additionally stained with DAPI and assessed by using a Leica TCS SP5 confocal microscope in a reflecting laser scanning regimen (laser 488 nm [Ar/Kr]).

Statistical analysis

For the analysis of the statistical differences, one-way analysis was carried out by the Kruskal–Wallis test. Student's *t*-test was performed to compare the obtained animal data using Statistica Version 9.2 program. Differences were considered to be statistically significant when $P < 0.05$.

Results

Synthesized CS-DX-SPIONs exhibit high-contrast-enhancing properties

The sequence of the steps in the synthesis of hybrid SPIONs is represented in Figure 1. Following the addition

of dextran to the prepared SPIONs for stabilization of the particles, the average size ranged between 30 and 40 nm. According to TEM measurements, DX-SPIONs are presented as round-shaped particles. For further coating of the particles with chitosan, the surface of DX-SPIONs was activated by NH_2 groups through addition of epichlorohydrin. Chemical activation did not change the stability of nanoparticle dispersion (as assessed by DLS). Size distribution measured by the DLS method was 55 nm. The hybrid dextran–chitosan coat of the magnetite core was obtained in the cross-linking procedure with the help of TPP shown in Figure 1 as Step III. Negatively charged ions of TPP induce ionotropic gelation of chitosan and dextran. To study the molecular mechanism of interaction between TPP and chitosan, ^{31}P NMR measurements were conducted with gradual increase in TPP concentration in the reaction solution. According to the ^{31}P NMR spectra, three α -, β -, and γ - PO_4 groups produce two well-resolved lines at different chemical shifts (Figure 2). The low-shift

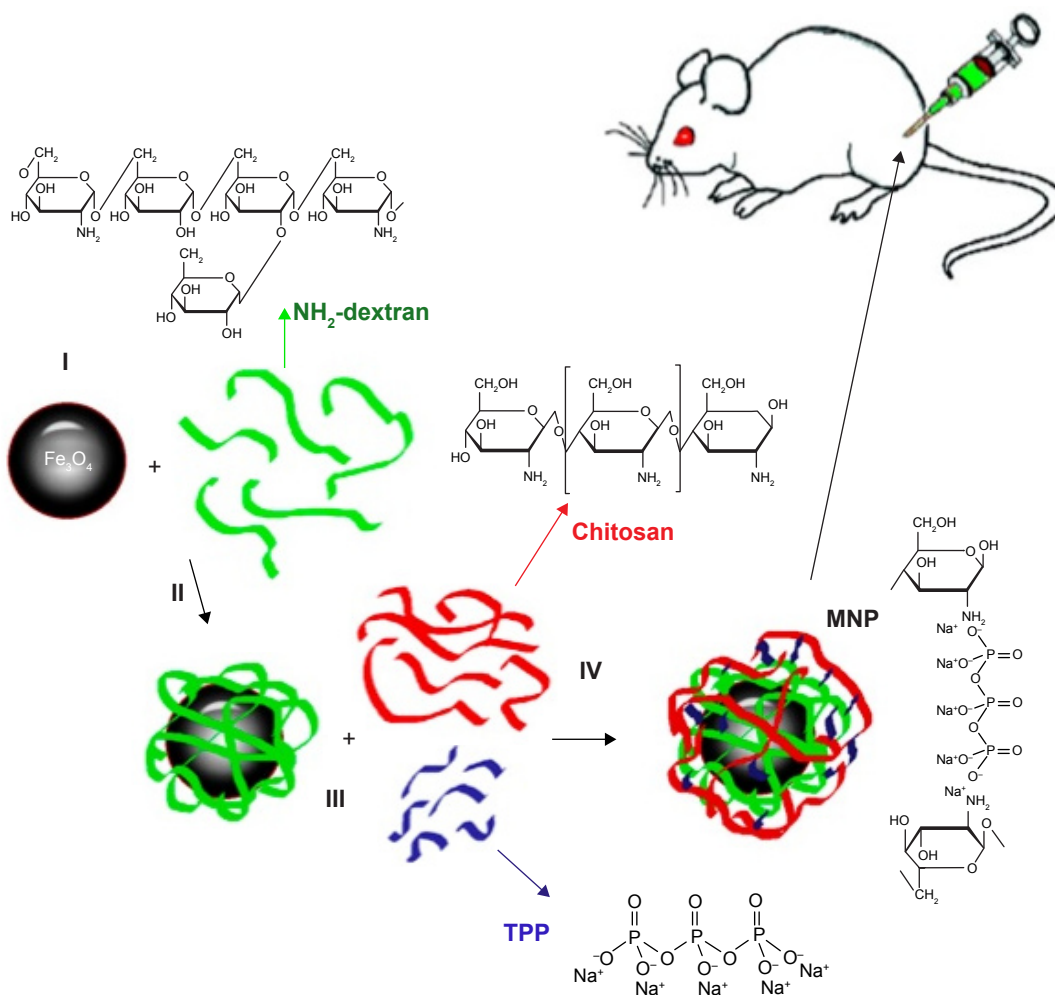


Figure 1 Schematic representation of the synthesis of chitosan-based superparamagnetic iron oxide nanoparticles (CS-DX-SPIONs).

Abbreviations: CS-DX-SPIONs, chitosan–dextran superparamagnetic iron oxide nanoparticles; MNP, magnetic nanoparticle; TPP, triphosphosphate.

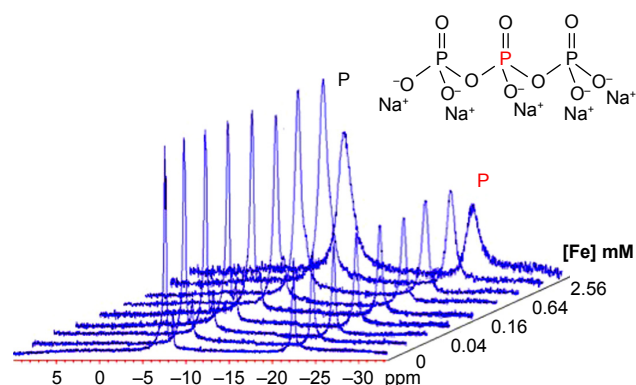


Figure 2 ^{31}P Phosphorus NMR spectroscopy of the TPP.
Abbreviations: NMR, nuclear magnetic resonance; TPP, tripolyphosphate.

line (ppm) belongs to α and γ phosphate groups. In the process of titration, initial addition of TPP did not lead to the corresponding increase in intensity resonance. According to the theoretical analysis, the ^{31}P relaxation rates of phosphate groups in strong magnetic field (7.1 T) due to dipole–dipole (DD) and chemical shift anisotropy (CSA) mechanisms are well described by Bloembergen et al.^{24,25} In line with this theory, ion–ion interaction of PO_4 groups with protonated NH_3^+ groups reduces molecular motion of TPP and leads to strong decrease in magnetic relaxation times T_1 and T_2 and thus loss of spectral resolution. At low TPP concentrations, the resonance lines are invisible. At saturation threshold of all free NH_3^+ groups, excess of TPP agent gives rise to narrow PO_4 lines in spectra with intensity proportional to TPP concentration. The observed concentration dependencies are in good agreement with the hypothesis that one molecule of TPP binds 2–3 positively charged NH_3 groups. Coverage of magnetic nuclei by the dextran–chitosan coat results in an increase in the hydrodynamic diameter of nanoparticles and initiation of magnetic moment in the whole nanostructure. With respect to the small size of the nanoparticles (40 nm) one might assume that the magnetic particle consists of a single domain nanocrystal. In this case, the nanoparticle would behave as an SPM unit exposing classical hysteresis behavior in accordance to the Langevin law.²⁶ Otherwise, the particles behave as a ferromagnetic material. To elucidate the magnetic nature of hybrid CS-DX-SPIONs, the novel NLR- M_2 approach was applied to assess the particles for hysteresis criteria. Both components, $\text{Re}M_2$ and $\text{Im}M_2$ of nonlinear response, were measured versus steady field H at $T=294$ K at different rates of field scan, F_{sc} . The obtained H -dependencies display the characteristic signals, exhibiting extremes at a

weak magnetic field H in $H>0$ range of H -scan, opposite signs in $\text{Re}M_2(H)$ and $\text{Im}M_2(H)$ components, and the weak field hysteresis, which decreases with a reduction in F_{sc} . These features correspond to M_2 response of ensemble of MNPs in the single domain state (Figure 3).²⁷ A weak field hysteresis is predictive for a close to SPM regime, which is in accordance with a single domain structure. M_2 responses yielded by the Fe_3O_4 nuclei and magnetic parameters of the latter in the first approximation are not affected by the formation of the hybrid dextran–chitosan shell. The response of an ensemble of NPs in SPM regime is characterized by the absence of field hysteresis. It is obvious that in this case the magnetization M will be directed along the field, parallel to the z axis. As for complex magnetic materials such as manganites, cobaltites, and composite substances on co-compounds, the shape of hysteresis curves depends on F_{sc} and temperature, and SPM regime occurs above blocking temperature T_B .^{28,29} The hybrid CS-DX-SPIONs have high magnetic moment to accelerate magnetic relaxation of aqueous protons in suspension due to large perturbation of the magnetic field in the neighborhood of MNPs. Inhomogeneous magnetic fields in the probe can be calculated as the sum of dipole fields induced by magnetite cores. The inhomogeneous

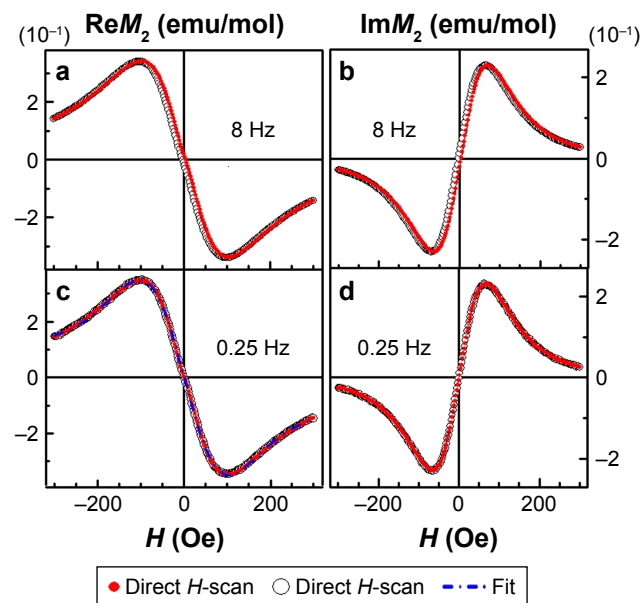


Figure 3 Second harmonic measurements of CS-DX-SPIONs in water at different frequencies of H -scan, $F_{sc}=8$ (panels (a) and (b)) and 0.25 Hz (panels (c) and (d)), $T=294$ K.

Notes: Solid and open circles display direct and reverse H -scans, respectively. Dashed-dot curve in panel (c) presents fit $\text{Re}M_2(H)$ dependence obtained at $F_{sc}=0.25$ Hz by $\partial^2 L(H)/\partial H^2 + k_{PM}H$.

Abbreviation: CS-DX-SPIONs, chitosan–dextran superparamagnetic iron oxide nanoparticles.

character of the field shortens the magnetic relaxation times of H_2O .

Assessment of cellular interactions of CS-DX-SPIONs

Uptake of the synthesized DX-SPIONs and hybrid CS-DX-SPIONs was analyzed in tumor cells (C6, U87, and HeLa) following coincubation of the latter with DX-SPIONs or CS-DX-SPIONs at various concentrations of Fe^{3+} (ie, 1, 10, 50, and 150 $\mu g/mL$) and incubation periods (ie, 1, 3, 6, 12, and 24 hours). Following an overnight coincubation, the confocal microscopic views show that DX-SPIONs were incorporated into the cytosol of the tumor cells (Figure 4). Magnetic conjugates on the reflective laser scanning are presented as

green dots throughout the cytosol of the tumor cells but are not detected in the nucleus of the cells (Figure 4A). In the case of hybrid CS-DX-SPIONs, an increase in the internalization of nanoparticles as compared to DX-SPIONs was detected (Figure 4A). The uptake of the CS-DX-SPIONs was further evaluated as related to their contrast-enhancing properties when coincubated with C6 glioma cells. Accumulation of the hybrid SPIONs in the cells led to the hyperintense T_2 -weighted MRI images and subsequent decrease in T_2 values (Figure 4B). This indicates that CS-DX-SPIONs preserve their magnetic contrast-enhancing properties after internalization into cells. Thus, T_2 values following internalization of the CS-DX-SPIONs constituted 6.0 ± 1.34 ms which was significantly lower in comparison to the DX-SPIONs and

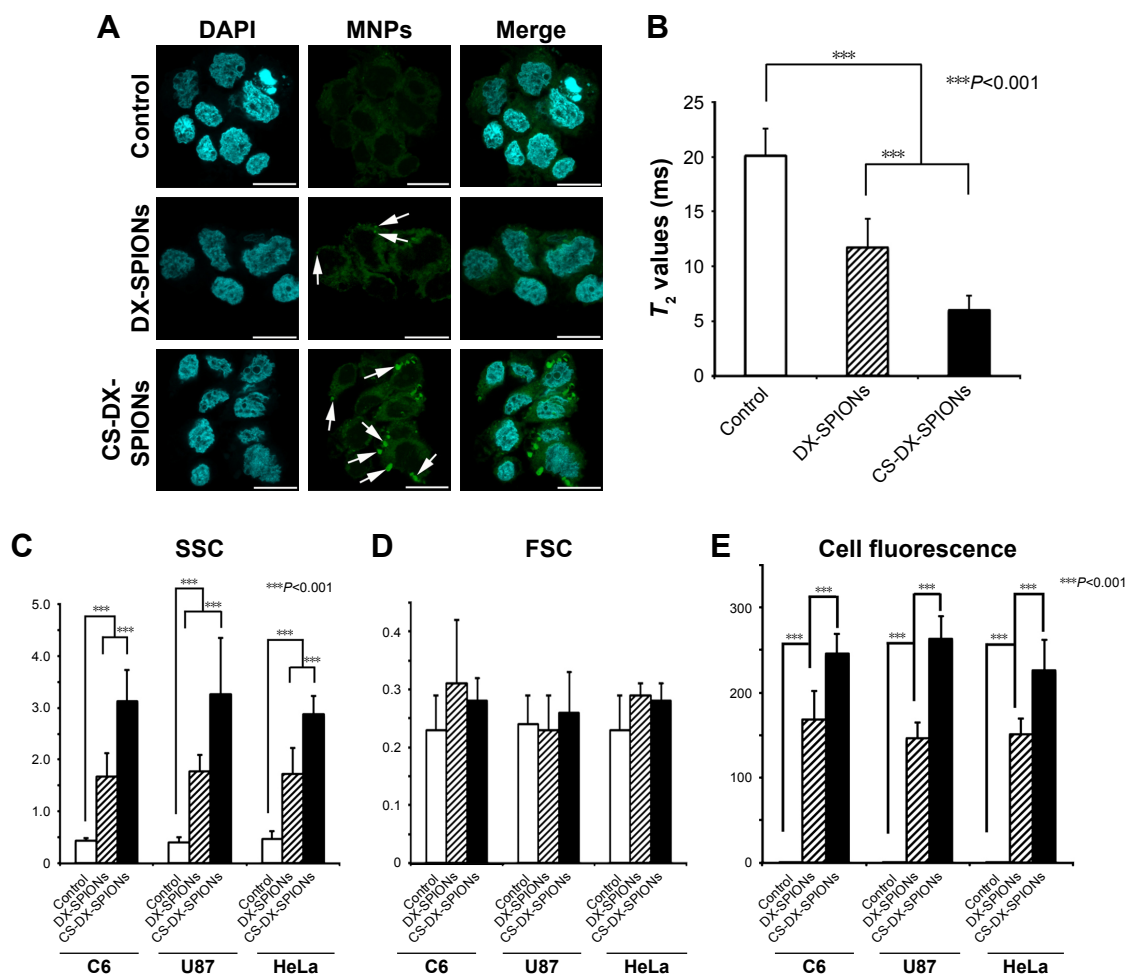


Figure 4 Interaction of CS-DX-SPIONs with cells.

Notes: (A) Confocal microscopy images of the rat C6 glioma cells coincubated for 24 hours with PBS (control), DX-SPIONs, or CS-DX-SPIONs. Nanoparticles were detected using reflecting laser scanning at 488 nm (presented as green dots). Nuclei were stained with DAPI (blue). Scale bar, 7 μm . (B) T_2 values (ms) of the C6 cells following coincubation with PBS (control), DX-SPIONs, or CS-DX-SPIONs for 24 hours in 0.1% agarose gel. Data are presented as mean \pm SD. (C) SSC, (D) FSC, and (E) cell fluorescence for three cell lines (C6, U87, and HeLa) following coincubation with PBS (control), DX-SPIONs, or CS-DX-SPIONs for 24 hours. The mean and SD were calculated for the three experiments. T_2 is the spin-spin relaxation time.

Abbreviations: CS-DX-SPIONs, chitosan–dextran superparamagnetic iron oxide nanoparticles; DAPI, 4,6-diamidine-2-phenylindole; DX-SPIONs, dextran-coated superparamagnetic iron oxide nanoparticles; FSC, forward scattering; PBS, phosphate buffered saline; SSC, side scattering.

control (PBS), 11.72 ± 2.65 and 20.1 ± 2.45 ms, respectively ($P < 0.001$) (Figure 4B). To further assess the internalization of nanoparticles, flow cytometry was applied. These data clearly showed an enhanced accumulation of CS-DX-SPIONs in the C6, U87, and HeLa cells as compared to the dextran-coated particles after 24 h (Figure 4C and E). The retention of the particles significantly increased the granularity (SSC) of the tumor cells, but not their size (FSC) (Figure 4D). After internalization, the cytotoxicity of the nanoparticles, by the detection of LDH in the samples, was analyzed. A dose and time-dependent toxicity of the CS-DX-SPIONs after co-incubation with tumor cells was observed (Figure 5). Twenty-four hours after co-incubation with CS-DX-SPIONs (at a Fe concentration of $150 \mu\text{g/mL}$), the cytotoxicity of C6, U87,

and HeLa cells was $30.1 \pm 6.4\%$, $36.2 \pm 7.9\%$, and $27.4 \pm 6.2\%$, respectively.

Targeting of the brain tumor by hybrid CS-DX-SPIONs

A total of 15 days after orthotopic injection of C6 glioblastoma cells, the animals were randomly divided into three groups (six animals each): 1) control group injected with PBS; 2) i.v. administration of the DX-SPIONs; and 3) i.v. injection of CS-DX-SPIONs. Twenty-four hours after the injection of the nanoparticles, brain tumors were imaged on the high-field MRI scanner. In control animals, the tumor was presented as hypointense on the T_1 -weighted images and hyperintense on T_2 images (Figure 6). In the

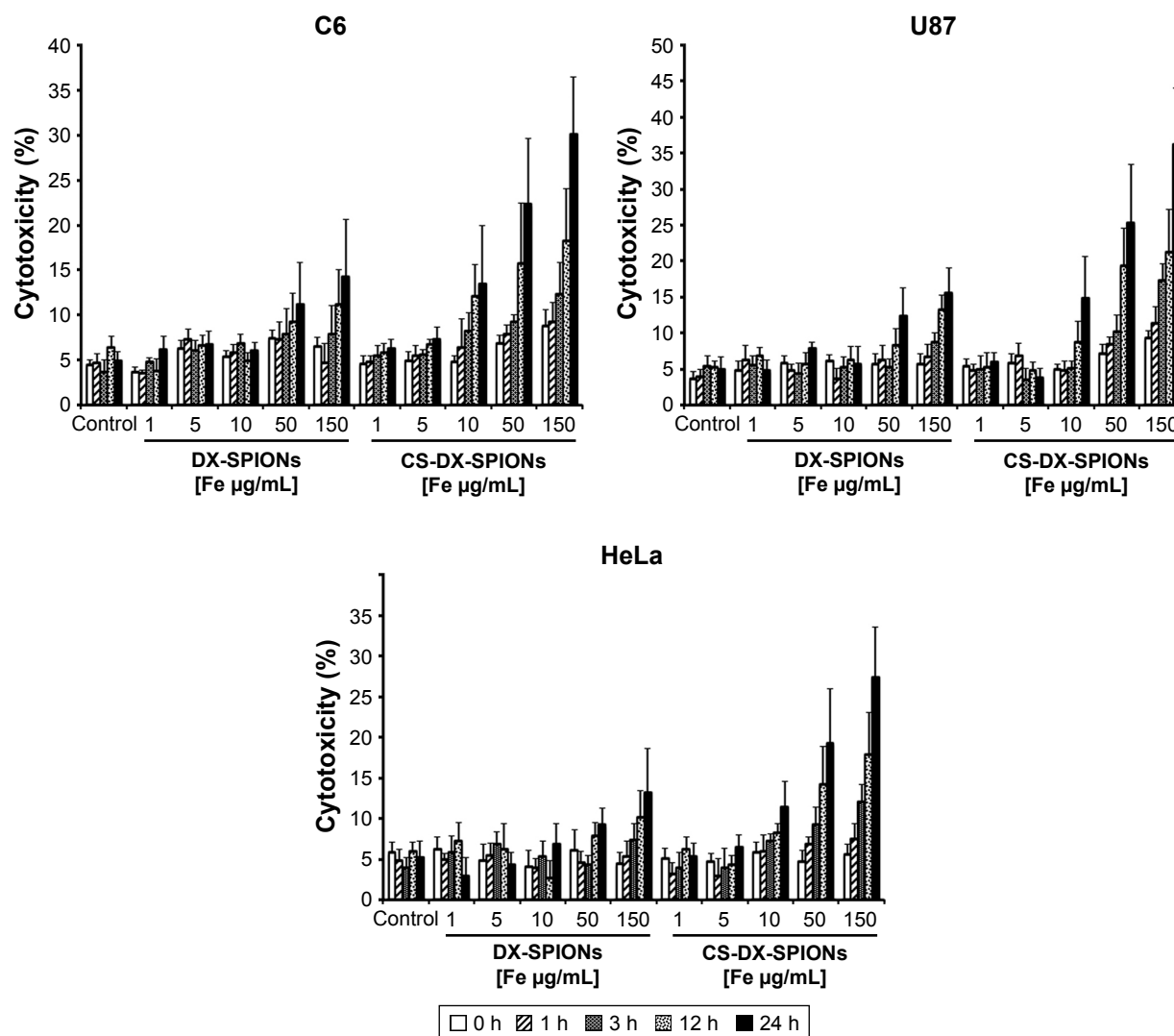


Figure 5 Cytotoxicity assay of the tumor cells (C6, U87, and HeLa) co-incubated with PBS (control), DX-SPIONs, or CS-DX-SPIONs.

Note: The mean and SD was calculated for the three experiments.

Abbreviations: CS-DX-SPIONs, chitosan–dextran superparamagnetic iron oxide nanoparticles; DX-SPIONs, dextran-coated superparamagnetic iron oxide nanoparticles; h, hour; PBS, phosphate buffered saline.

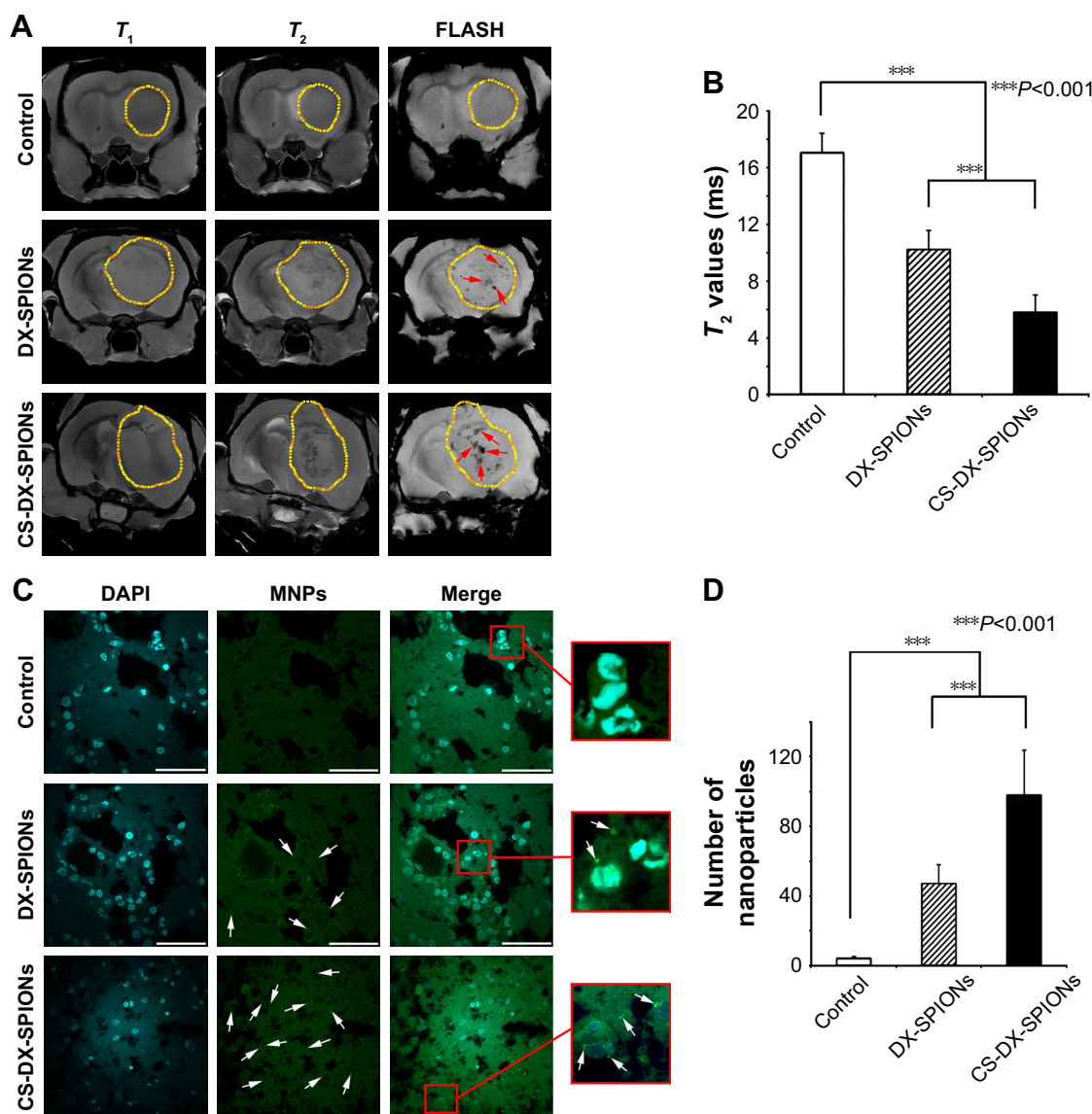


Figure 6 Targeting of the C6 glioma with CS-DX-SPIONs.

Notes: (A) Magnetic resonance images of the brain tumors of the control (PBS), animals treated with DX-SPIONs and CS-DX-SPIONs. T_1 -weighted and T_2 -weighted and gradient echo (FLASH) images of the glioblastoma were obtained. Accumulation of the nanoparticles is shown by solid red arrows. (B) T_2 values (ms) of the brain tumors in the control (PBS), animals treated with DX-SPIONs and CS-DX-SPIONs. Data are presented as mean \pm SD. (C) Immunofluorescent images of the glioma. Nuclei stained with DAPI (blue). Nanoparticles detected using reflective laser scanning at 488 nm (green). Scale bar, 40 μ m. (D) Quantification of the nanoparticles in the glioma tissue. Data are presented as mean \pm SD. T_1 – spin-lattice relaxation time and T_2 – spin-spin relaxation time.

Abbreviations: CS-DX-SPIONs, chitosan–dextran superparamagnetic iron oxide nanoparticles; DAPI, 4,6-diamidine-2-phenylindole; DX-SPIONs, dextran-coated superparamagnetic iron oxide nanoparticles; MNPs, magnetic nanoparticles; PBS, phosphate buffered saline.

DX-SPIONs-injected animal group, glioma was characterized by a hypointense signal in the T_2 -weighted imaging. Twenty-four hours after i.v. infusion of CS-DX-SPIONs, a significant decrease in the T_2 relaxivity has been observed in comparison to control and SPION-injected animals (17.1 ± 1.4 , 10.2 ± 1.4 , and 5.8 ± 1.2 ms, respectively; $P < 0.001$). Reduction of the transverse relaxation time T_2 is accompanied by the growth of image contrast due to the darkening of accumulation sites of MNPs. Subsequent histological assessment of brain tumor cryosections confirmed the accumulation of the

MNPs inside glioma cells (Figure 6C and D). The distribution pattern of DX-SPIONs or hybrid CS-DX-SPIONs retention in the tumor did not differ.

Discussion

Nanotechnology can provide a novel theranostic approach in managing glioblastoma multiforme.^{30,31} In our studies, SPIONs with a high MRI contrast enhancement capacity have been used.³² For an improved biocompatibility, nanoparticles were coated with chitosan that increased the charge

of the dextran-coated nanoparticles to +19.2 mV. Chitosan, a polysaccharide with a positive charge at neutral or acidic pH, has advantageous biocompatibility as a carrier material,³³ a strong permeation enhancing capability for hydrophilic compounds,³⁴ and bioadhesive properties.³⁵ An increase in charge of the nanoparticles also resulted in an enhanced uptake of CS-DX-SPIONs by tumor cells (Figure 4). The same effect was previously reported for positively charged pullulan-coated NPs that were incorporated by bone marrow-derived rat MSCs³⁶ or mouse MSCs.¹³ On the other hand, the decrease in the surface charge abrogated the internalization of the nanoparticles.³⁶

Although we observed an enhanced intracellular uptake of the particles, we subsequently determined an increased cytotoxicity of CS-DX-SPIONs (Figure 5). This could be explained by the production of reactive oxygen species (ROS) and subsequent DNA damage.^{37,38} Increased amounts of ROS might also negatively affect mitochondrial DNA (mDNA).³⁹

Synthesized CS-DX-SPIONs also act as a negative MRI contrast agent. The measured proton relaxation rates of water in magnetic suspensions of CS-DX-SPIONs clearly indicated a high contrast property of the particles (Figure 3). The registered relaxation rates in the hybrid CS-DX-SPIONs resembled the main relaxation features of previously reported dextran-coated MNPs and commercial NP formulations.^{16,17} The composite shell (between chitosan and dextran) cross-linked by TPP does not hamper the diffusion of protons into dephasing volume around MNPs thus preserving the MRI contrast-enhancing properties of the particles. Following coincubation with tumor cells, CS-DX-SPIONs accumulated in cytosol, which resulted in a decrease in T_2 values when the samples were analyzed using MRI scanning (Figure 4B). This indicates that the obtained NPs preserve their magnetic properties after internalization into cells. In the current study, for the analysis of accumulation of nanoparticles in the glioma, standard MRI sequences (ie, T_2 -weighted scans, GE [FLASH], and multi-scan multi-echo [MSME]) were applied. Recently, other MRI methods for visualization of the MNPs were proposed including investigation of transverse relaxation rates R_2 and R_2^* ,⁴⁰ quantitative susceptibility mapping (QSM),⁴¹ and susceptibility-weighted imaging (SWI).⁴²⁻⁴⁴ In the study by Blasiak et al, the highest contrast-to-noise ratio for the detection of the iron deposition in an U87MG glioma model was achieved when the SWI was employed in comparison to any other pulse sequences (ie, SE and GE).⁴² The magnitude of magnetic relaxation rate of hybrid CS-DX-SPIONs corresponded to the contrast enhancement of the agent in MRI scans of brain tumors in rats (Figure 6A).

Following intravenous administration, CS-DX-SPIONs accumulated in glioma cells, which significantly enhanced the contrasts of the tumor. Retention of the particles resulted in a significant decrease in T_2 values compared to DX-SPIONs.

Accumulation of the particles inside glioma cells could be explained by an enhanced permeation and retention (EPR) effect.⁴⁵ Thus, particles <100 nm in diameter extravasate preferentially from the chaotic tumor vasculature, due to the increased permeability of vessels compared to normal tissues.⁴⁶ Presumably, further modifications of the surface of CS-DX-SPIONs with targeting bioligands (eg, antibodies, peptides, proteins, etc) could further increase the accumulation of particles inside glioma tissue. Previously, several studies demonstrated that coating with tumor-targeting molecules including anti-Hsp70 antibody,¹⁶ peptide-targeting interleukin-13 receptor $\alpha 2$ (IL-13R $\alpha 2$),⁴⁷ anti-VEGF antibodies,⁴⁸ antibodies toward EGFR deletion mutant (EGFRvIII),⁴⁹ anti-insulin-like growth factor binding protein 7 (anti-IGFBP7) single domain antibody,⁵⁰ and many others could show an enhanced particle retention inside tumor cells.⁵¹⁻⁵⁴

Conclusion

Hybrid CS-DX-SPIONs synthesized by ionotropic gelation at the SPM state have a high MRI contrast-enhancing activity and tumor-targeting potential. In vitro experiments clearly demonstrated that the increase in the surface charge of the nanoparticles due to the chitosan coating subsequently enhanced the intracellular uptake of particles and thus increased their cytotoxic activity. Administration of the magnetic nanocarriers via clinically relevant routes resulted in a significant contrast enhancement of glioma. Diagnostic properties of the synthesized hybrid nanoparticles could be further explored for the development of targeted therapies.

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Disclosure

The authors report no conflicts of interest in this work.

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