

Biological effects induced by Gadolinium nanoparticles on Lymphocyte A20 cell line

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Abstract

Gadolinium nanoparticles (GdNPs) are potential agents for MRI of lymph nodes. The aim of this study was to evaluate the *in vitro* effects of 1 μ M, 2.5 μ M and 5 μ M of GdDOTACCS-TPP/HA and GdDOTPCCS-TPP/HA NPs on A20 lymphocyte cells exposed for 6 and 24 hours. The total cellular biomass (SRB), lactate dehydrogenase activity (LDH) and oxidative stress parameters, such as reactive oxygen species generation (ROS), reduced glutathione (GSH), malondialdehyde (MDA) and advanced oxidation protein products (AOPP) were analyzed by spectrophotometric and fluorimetric methods. After cells exposure to 1 μ M, 2.5 μ M and 5 μ M of GdDOTPCCS-TPP/HA NPs their viability decreased in a time- and dose-dependent manner, whereas for GdDOTACCS-TPP/HA no significant changes were noticed. Both NPs formulations in doses of 1 μ M, 2.5 μ M, 5 μ M did not affect the plasma membrane at each time point tested. The levels of ROS, MDA and AOPP increased proportionally with the concentration and exposure time. GSH concentration decreased significantly for all doses of both NPs tested. Taken together our data suggest that, GdDOTPCCS-TPP/HA and GdDOTACCS-TPP/HA NPs induced oxidative stress in A20 lymphocyte cells which was counteracted by the cells antioxidant defense system to a certain extend.

Introduction

In the last years, the development of gadolinium-based nanoparticles (GdNPs) as contrast agents (CAs) for Magnetic Resonance Imaging (MRI) has grown exponentially, due to their interesting magnetic properties (1). Those MRI probes, that associated gadolinium chelates (GdCAs) and nano-sized objects (by covalent grafting or by encapsulation) act as catalysts to enhance the relaxation rates of tissue water proton molecules and then, to improve the image contrast in MRI. The advantage to combine GdCAs with nanoparticles is not only to enhance MRI sensitivity by improvement of image contrast (2, 3) but also to diminish the injected Gd doses. Therefore, possible side effects (4, 5) reported for some GdCAs of Nephrogenic Systemic Fibrosis (NSF) in patients with renal failure (4) can be avoided, for the benefit of the patients. Recently, confinement of GdCAs within a hydrophilic polymeric nanostructures has allowed to pass a milestone in MRI probes efficiency (6, 7). The corresponding objects that combine an important concentration of GdCAs within high-water content nanostructure help to improve the efficacy of commercially available GdCAs by a factor of 10 to 24. These highly hydrophilic NPs also called hydrogels are obtained by ionic gelation (8-12) between two biocompatible (13) polysaccharides chitosan (CS) and hyaluronic acid (HA) in the presence of sodium tripolyphosphate (TPP) as a cross-linker (6, 7). The biocompatibility of the polymers does not presuppose that the corresponding hydrogels are biocompatible and the biocompatibility of the hydrogels incorporating the GdCA of DOTAREM, GdDOTA (DOTA = 1,4,7,10-tetraazacyclododecane-1,4,7,10-tetraacetic acid) and GdDOTP (DOTP = 1,4,7,10-tetraazacyclododecane-N,N',N'',N'''-tetrakis(methylenephosphonic acid) is still poorly explored. Biodegradation, biodistribution and toxicity of chitosan were reported by T Kean et al. in a recent review (14). In this context, before considering their applica-

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tions in clinical area, studies regarding nanoparticles toxicity are necessary (15).

This study aims to investigate the potential toxicity of two well-known, low risk contrast agents towards NSF i.e. GdDOTA and GdDOTP, encapsulated in a nanohydrogels constituted of CS and HA (GdDOTACCS-TPP/HA and GdDOTPCCS-TPP/HA NPs, respectively). The *in vitro* effects of these new formulations have been evaluated on A20 murine model. The cytotoxicity and biochemical changes induced by the exposure of A20 murine lymphocyte B cells to increasing concentrations of GdDOTACCS-TPP/HA and GdDOTPCCS-TPP/HA NPs were analyzed. For that, the total cellular biomass (SRB), lactate dehydrogenase activity (LDH) and oxidative stress parameters, such as reactive oxygen species generation (ROS), reduced glutathione (GSH), malondialdehyde (MDA) and advanced oxidation protein products (AOPP) were investigated in a time dependent manner.

Materials and Methods

Reagents

Sterile water was purchased from Laboratoire Aguettant, Lyon, France and sodium tripolyphosphate from Acros Organics. DOTA and DOTP were purchased from Macrocyclics (Dallas, USA). Chitosan (KiOmedine CsU-A), sodium hyaluronate were provided by Sigma (France) and bovine serum albumin by Sigma (New Zealand). Fetal bovine serum, RPMI-1640 medium and phosphate-buffered saline were purchased from Gibco and 2',7'-dichlorodihydrofluorescein diacetate from Invitrogen by Thermo Fisher Scientific Inc. (Waltham, MA, USA). HCl (37%), HNO₃ (69%), acetic acid glacial, potassium iodide, glucose, ethylenediaminetetraacetic acid, 5-sulfosalicylic acid, antibiotic-antimycotic solution stabilized with 10.000 units penicillin, 10 mg streptomycin and 25 µg amphotericin B per mL, *In Vitro* Toxicology Assay Kit Sulforhodamine B based, *In Vitro* Toxicology Assay Kit Lactic Dehydrogenase based, Reduced glutathione, 5,5-dithio-bis-(2-nitrobenzoic acid), thio-barbituric acid, 1,1,3,3-tetramethoxypropane and chloramine T were purchased from Sigma (St. Louis, MO, USA). Bradford reagent was provided by Bio-Rad.

Synthesis and characterization of nanoparticles

Stock solutions of CS (KiOmedine CsU-A) were prepared by dissolution of the CS powder (2.5 mg.mL⁻¹) in a 10% (m/v) citric acid aqueous solution and stirred overnight.

Nanoparticles were obtained by an ionotropic gelation process (6, 7). The polyanionic phase i.e. HA (0.8 mg.mL⁻¹) and TPP (1.2 mg.mL⁻¹) in water (4.5 mL), were added dropwise to the CS solution (9 mL) under sonication (750W, amplitude 32%) to obtain stable nanosuspensions. The gadolinium complex (GdDOTA or GdDOTP) was previously dissolved in the polyanion solution. At the end of the addition, magnetic stirring was maintained for 10 min. Unloaded nanoparticles were obtained in the same way, omitting the gadolinium complex. Nanosuspensions were then freeze-dried, using glucose 15%

(m/v) as a cryoprotectant. NP average hydrodynamic diameters (D_H) and polydispersity indexes (PdI) were determined by Dynamic Light Scattering (DLS) (Malvern Zetasizer Nano-ZS, Malvern Instruments, Worcestershire, UK). Each nanosuspension was analyzed in triplicate at 20 °C at a scattering angle of 173°, for each sample, after 1/20 dilution in water. Pure water was used as a reference dispersing medium. ζ-(zeta) potential data were collected through electrophoretic light scattering at 20 °C, 150 V, in triplicate for each sample, after 1/20 dilution in water. The instrument was calibrated with a Malvern - 68 mV standard before each analysis cycle. Gadolinium nanoparticle loading was determined on nanoparticle suspensions by ICP-OES. The nanoparticle suspension was incubated overnight in a 1:3 (v:v) mixture of HCl (37%) and HNO₃ (69%). Volumetric dilutions were carried out to achieve an appropriate Gd concentration within the working range of the method. Samples were analysed using Thermo Scientific iCAP 6300 series Duo ICP spectrometer. Gd emission intensity was correlated to Gd concentration by means of a calibration curve which was previously established from Gd(NO₃)₃ ICP-OES standard. Solutions used for the calibration were obtained by dilution of increasing amounts of Gd(NO₃)₃ standard with unloaded nanoparticles previously incubated under acidic conditions, as described above.

Cell culture and treatment

A20 is a BALB/c B cell lymphoma line derived from a reticulum cell sarcoma found in an BALB/cAnN mouse. The cells were cultured in RPMI-1640 medium (Roswell Park Memorial Institute medium) supplemented with 1% antibiotic (penicillin, streptomycin and amphotericin B) and 10% fetal bovine serum maintained at 37°C in a humidified air atmosphere with 5% CO₂. They were seeded at a density of 2 × 10⁵ cells/mL onto 75 cm² culture flasks and the experiments were carried out using cells between passages 3 and 5. The cells were exposed to 1 µM, 2.5 µM, 5 µM of GdDOTACCS-TPP/HA and GdDOTPCCS-TPP/HA NPs for 6h and 24h. CS-TPP/HA NPs were used as controls.

Cellular lysate preparation and determination of protein concentration

A20 cells, collected from culture flasks were centrifuged at 1500 rpm for 5 min at 4°C. After removing the culture medium, the cell pellet was washed with phosphate-buffered saline (PBS) two times and centrifuged again in the same conditions. Cell pellets were resuspended in 0.3 mL of PBS and then, homogenized by sonication using Hielscher Ultrasonic processor UP50H on ice three times, for 30 seconds each. The cellular lysate was centrifuged at 5000 rpm for 10 min at 4°C. The supernatant was decanted, aliquoted and stored at -80°C and used for further biochemical determinations. The total protein concentration was measured according to the method described by Bradford (16) using bovine serum albumin (BSA) as a standard. The absorbance was spectrophotometrically measured at a wavelength of 595 nm using the Appliskan Thermo Scientific Multireader and the results were expressed as mg. mL⁻¹.

Sulforhodamine assay (SRB)

The viability of A20 cells was assessed using the *in vitro* Toxicology Assay Kit, Sulforhodamine B based (Sigma-Aldrich, St. Louis, MO, USA) and following manufacturer's instructions. This method is based on the measurement of total biomass by staining cellular proteins with the Sulforhodamine B dye. After 6 and 24h of exposure to 1 μ M, 2.5 μ M, 5 μ M Gd-DOTACCS-TPP/HA and GdDOTPCCS-TPP/HA NPs, cells were fixed with 50% trichloroacetic acid (TCA) solution (1/4 volume of growth medium) for 1 hour at 4°C. Then, cells were washed with distilled water several times and the plates were air dried. Cells were then stained for 20 minutes at room temperature in the presence of a 0.4% sulforhodamine B (Acid Red 52) solution. The dye was then removed and the cells washed with a solution of acetic acid 1% (v:v). In order to solubilize the sulforhodamine B dye incorporated into the cells a 10mM Tris base solution was used. The absorbance was spectrophotometrically measured at a wavelength of 550 nm using the Appliskan Thermo Scientific Multireader.

Lactate dehydrogenase release (LDH)

Lactate dehydrogenase (LDH) activity was evaluated using a commercial kit, *in vitro* Toxicology Assay Kit, Lactic Dehydrogenase based (Sigma-Aldrich, St. Louis, MO, USA) according to the manufacturer's instructions. The cell membrane integrity was estimated as a function of the quantity of LDH released into the cell culture medium. The method is based on the reduction of nicotinamide adenine dinucleotide (NAD⁺) in the presence of lactate and LDH, followed by the conversion of a tetrazolium dye into a colored formazan compound which can be spectrophotometrically quantified at 450 nm. A total of 2×10^5 A20 cells per mL were seeded in 24-well cell culture plates. After 24h of incubation, the cells were exposed to concentrations of 1 μ M, 2.5 μ M, 5 μ M of GdDOTACCS-TPP/HA and GdDOTPCCS-TPP/HA NPs for 6 and 24 h. A volume of 50 μ L of each sample was placed on a 96-well microtiter plate and then 100 μ L of lactate dehydrogenase assay mixture were added. After 30 min at room temperature, the reaction was stopped by adding 1/10 volume of HCl 1M to each well and the absorbance was read spectrophotometrically at a wavelength of 450 nm using the Tecan GENios Multireader. The LDH activity was expressed as % of controls (cells treated with unloaded CS-TPP/HA NPs).

Lipid peroxidation measurement (MDA)

The level of malondialdehyde (MDA), a marker for oxidative degradation of lipids was assessed using thiobarbituric acid (TBA) as a reagent according to Dinischiottu et al. (17). A volume of 200 μ L of cell lysate with a total protein concentration of 3 mg. mL⁻¹ was mixed with 700 μ L of HCl 0.1 M and incubated at room temperature for 20 min. Afterwards, 900 μ L of 0.025 M thiobarbituric acid were added and the mixture was maintained for 65 min at 37°C. Finally, 400 μ L of PBS were added. The fluorescence of MDA-TBA adduct was quantified fluorometrically ($\lambda_{exc/em}$ = 520/549 nm) using a Jasco FP-750

spectrofluorometer (Tokyo, Japan). MDA concentration was calculated using a 1,1,3,3-tetramethoxy propane calibration curve with concentrations in the range of 0.5-5 μ M. The results were expressed as nmoles of MDA per mg of protein.

Reduced glutathione assay (GSH)

The GSH concentration was determined using the Glutathione Assay Kit (Sigma-Aldrich, St. Louis, MO, USA) following manufacturer's instructions. Briefly, cellular lysates were deproteinized with an equal volume of 5% 5-sulfosalicylic acid (SSA) and centrifuged at 10000 rpm for 10 minutes at 4°C. A volume of 10 μ L of each supernatant was placed on a 96-well microtiter plate. Then 150 μ L of working mixture containing 8 mL of 100 mM potassium phosphate buffer, pH 7.0, with 1 mM ethylenediaminetetraacetic acid (EDTA) - assay Buffer and 228 μ L of 1.5 mg.mL⁻¹ 5,5'-dithiobis-(2-nitro benzoic acid) (DTNB) were added. After 5 minutes of incubation at room temperature, the absorbance of 5-thio-2-nitrobenzoic acid (TNB) was measured spectrophotometrically at a wavelength of 412 nm using the Tecan GENios Multireader. In order to determine the amount of glutathione in the samples, a calibration curve of GSH in the range of 0.5-0.03 nmoles was used. The results were expressed as nmoles of GSH per mg of protein.

Determination of intracellular reactive oxygen species (ROS)

Detection of intracellular ROS in A20 cells was assessed using a chemically reduced, acetylated form of fluorescein, 2',7'-dichlorodihydrofluorescein diacetate (H₂DCFDA) (18). This nonfluorescent compound is converted to a green-fluorescent form upon cleavage of the acetate groups by cellular esterases and oxidation occurs within the cell. A20 cells were seeded in six-well plates at a density of 2×10^5 cells mL⁻¹.

After exposure to 1 μ M and 5 μ M of GdDOTACCS-TPP/HA and GdDOTPCCS-TPP/HA NPs for 6h and 24h, the cells were incubated with 10 μ M of H₂DCFDA in the dark for 30 minutes at 37°C. Then, cells were collected by centrifugation at 250 g for 4 minutes, the pellets were washed twice and resuspended in 2 mL of PBS solution. The fluorescence of dichlorofluorescein (DCF) was quantified fluorometrically ($\lambda_{exc/em}$ = 488/515 nm) using a Jasco FP-750 spectrofluorometer (Tokyo, Japan). The results were calculated by dividing the fluorescence intensity to the number of viable cells and expressed as % of controls (cells treated with unloaded CS-TPP/HA NPs).

Advanced oxidation protein products assay (AOPP)

The level of AOPP, a marker of oxidative stress was analyzed using the method described by Witko-Sarsat et al. (19). 200 μ L of cell lysate was mixed with 10 μ L of 1.16 M potassium iodide and vortexed for 5 minutes at room temperature. A volume of 20 μ L of glacial acetic acid was added and then the mixture was vortexed again for 30 seconds. The sample's optical density was read at a wavelength of 340 nm using a Tecan GENios Multireader. Spectrophotometric determination of AOPP was calculated based on a chloramine T calibration curve with con-

centrations up to 100 μM . The results were expressed as nmoles of AOPP per mg of protein.

Statistical analysis

The results were expressed as mean value of triplicate experiments \pm standard deviation (SD) and compared by Student's *t*-test using standard statistical packages. Data were represented as graphics relative to control which was considered 100%. Statistical significance was accepted for a *p* value less than 0.05.

Results

Sulforhodamine B assay (SRB)

A20 murine B lymphocytes viability in the presence of different concentrations (1, 2.5 and 5 μM) of GdDOTA \subset CS-TPP/HA and GdDOTP \subset CS-TPP/HA NPs for 6 and 24h was presented in Fig. 1.

When A20 cells were treated with GdDOTA \subset CS-TPP/HA NPs, no significant NPs impact was encountered on cell viability at any doses tested up to 24h. In the case of cell exposure to GdDOTP \subset CS-TPP/HA NPs, a time- and dose-dependent decrease of cell viability was observed. Thus, after 24h, the cell viability decreased by 18% at the lowest concentrations and 28% at the highest.

LDH release

The cytotoxic potential of Gd \subset CS-TPP/HA NPs on A20 cells was evaluated by testing the cell membrane integrity via LDH leakage in the culture medium.

Fig. 2 showed that both nanoparticle formulations in doses of 1 μM , 2.5 μM , 5 μM did not affect the plasma membrane at each time points, since no significant amount of LDH was released to cell culture medium, as compared to control (CS-TPP/HA NPs).

Determination of intracellular ROS

ROS level proved that exposure of A20 cells to both types of nanoparticles generated a rise of ROS production in a time dependent manner for the higher concentration (Fig. 3).

A significant increase in ROS levels by 16% and 25% was noticed in A20 cells exposed to 5 μM GdDOTP \subset CS-TPP/HA after 6h and 24h, respectively, as compared to control (CS-TPP/HA NPs) whereas for GdDOTA \subset CS-TPP/HA treatment the increase was insignificant.

Lipid peroxidation (MDA), reduced glutathione (GSH) and advanced oxidation protein products (AOPP) in A20 cells

A time- and dose-dependent increase in the MDA level in A20 cells was registered compared to control (Table 1).

Significant MDA level increases of 15% and 17% were observed after 6h of exposure to 5 μM GdDOTA \subset CS-TPP/HA and GdDOTP \subset CS-TPP/HA respectively, and by 16% and 21% after 24h.

The concentrations of GSH, a non-enzymatic antioxidant decreased in a significant statistically manner, compared to

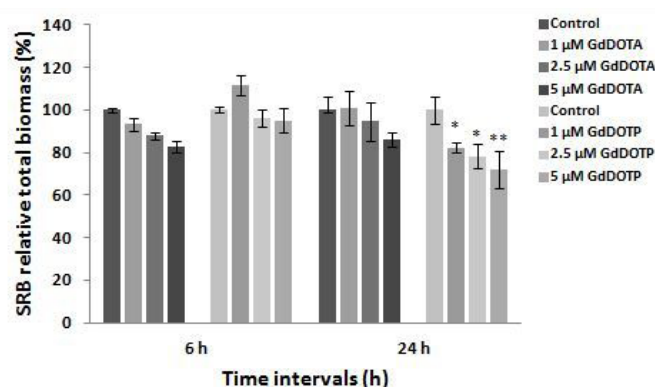


Figure 1. Viability cell (SRB assay) of A20 cells after exposure to 1 μM , 2.5 μM , 5 μM GdDOTA \subset CS-TPP/HA and GdDOTP \subset CS-TPP/HA NPs for 6h and 24 h. Unloaded CS-TPP/HA NPs were used as control. Results are calculated as means \pm SD (*n* = 3) and expressed as % from controls. **p* < 0.05; ***p* < 0.01

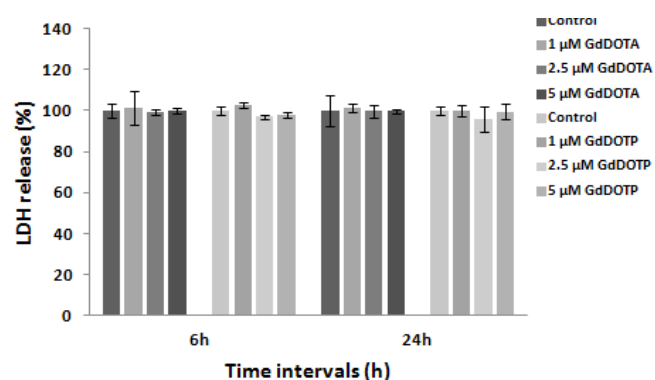


Figure 2. LDH release from A20 cells after exposure to 1 μM , 2.5 μM , 5 μM GdDOTA \subset CS-TPP/HA and GdDOTP \subset CS-TPP/HA NPs for 6h and 24 h. Unloaded CS-TPP/HA NPs were used as control. Results are calculated as means \pm SD (*n* = 3) and expressed as % from controls.

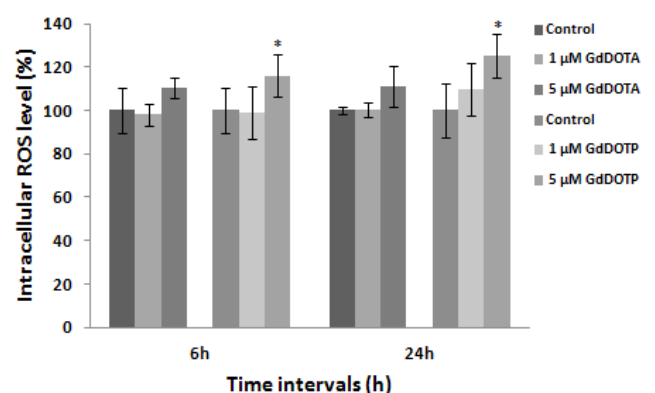


Figure 3. Reactive oxygen species (ROS) production in A20 cells, after exposure to 1 μM and 5 μM NPs GdDOTA \subset CS-TPP/HA and GdDOTP \subset CS-TPP/HA for 6h and 24 h. Unloaded CS-TPP/HA NPs were used as control. Results are calculated as means \pm SD (*n* = 3) and expressed as % from controls.

Table 1. Relative values of lipid peroxidation (MDA), reduced glutathione (GSH) and advanced oxidation protein products (AOPP) in A20 cells, after exposure to 1 μ M and 5 μ M GdDOTA \subset CS-TPP/HA and GdDOTP \subset CS-TPP/HA NPs for 6h and 24 h. Unloaded CS-TPP/HA NPs were used as control.

Time (h)	Nanoparticles	MDA (nmoles/mg)			GSH (nmoles/mg)			AOPP (nmoles/mg)		
		Control	1 μ M	5 μ M	Control	1 μ M	5 μ M	Control	1 μ M	5 μ M
6 h	GdDOTA	100 \pm 8	102 \pm 2	115 \pm 8	100 \pm 4	96 \pm 8	93 \pm 7	100 \pm 5	101 \pm 5	106 \pm 2
				*						
6 h	GdDOTP	100 \pm 10	102 \pm 4	117 \pm 6	100 \pm 5	95 \pm 4	90 \pm 3	100 \pm 6	102 \pm 9	119 \pm 4
				*			*			**
24 h	GdDOTA	100 \pm 5	110 \pm 5	116 \pm 10	100 \pm 9	95 \pm 8	89 \pm 7	100 \pm 9	108 \pm 3	111 \pm 11
				*			*			
24 h	GdDOTP	100 \pm 6	113 \pm 9	121 \pm 3	100 \pm 6	89 \pm 2	82 \pm 6	100 \pm 7	109 \pm 9	148 \pm 12
				**			**			**

Note: Results are calculated as means \pm SD ($n = 3$) and expressed as % from controls; * $p < 0.05$; ** $p < 0.01$

controls (Table 1). Thus, GSH levels were reduced after 6h of exposure for both NP doses tested. The decrease was amplified till 5% and 11% after 24h for 1 μ M and 11% and 18% for 5 μ M GdDOTA \subset CS-TPP/HA and GdDOTP \subset CS-TPP/HA, respectively.

Finally, AOPP concentrations were not significantly changed at any GdDOTA \subset CS-TPP/HA doses tested up to 24h of exposure, whereas significant increases up to 19% and 48% were recorded after 6h and 24h of cell exposure to the higher concentration of GdDOTP \subset CS-TPP/HA NPs, compared to control.

Discussion

Despite the gaining popularity of Gd(III) contrast agents in the medical field, their potential toxicity and long-term adverse side effects represent an important concern (20-24). In order to improve these aspects, new nanocarriers which are characterized by a good biodegradability, water-solubility, controlled drug release and low toxicity are now receiving a growing interest for Gd-based contrast agents vectorization (25-29).

Being designed for *in vivo* applications, especially for lymph node imaging, the evaluation of the potential toxicity generated by GdDOTA and GdDOTP nanocarriers (GdDOTA \subset CS-TPP/HA and GdDOTP \subset CS-TPP/HA) is essential for their future applications.

The first step towards understanding the way in which nanoparticles interact with cells often involves *in vitro* cell culture studies. Murine models of B-cell lymphomas are considered to be promising tools for development of cancer studies (30, 31) and can determine a massive invasion of liver, spleen and lymph nodes (32, 33) Previously it was demonstrated that some nanoparticles induce oxidative stress on lymphocytes and lymph nodes which could lead to their damage (34). For these reasons A20 cell line was selected as a model for *in vitro* experiments.

Several studies have revealed that Gd-based contrast agents

induced apoptosis and necrosis in different cell types (renal tubular cells, human foreskin fibroblasts (HFFs), NIH/3T3 fibroblasts) in a time (35) -and dose (36, 37) -dependent manner. By contrast, another research group has demonstrated the safety of a new designed multifunctional MRI contrast agents against HepG2 cells at the test concentrations, after 24h of incubation (38) Lack of toxic effects induced by different doses of gadolinium loaded chitosan and chitosan-hyaluronic acid nanoparticles in three cell lines (B16, HepG2, and A549 cells) at the same time point, were also reported (39, 40). Likewise, hydrogels incorporating GdDOTA and GdDOTP did not exhibit any toxicity towards C6 glioma or primary fibroblasts cells (6, 7).

Sulforhodamine B viability tests suggested that GdDOTP \subset CS-TPP/HA NPs were more toxic for A-20 cells compared to GdDOTA \subset CS-TPP/HA ones. These results were not confirmed by LDH activity in the culture medium probably due to a lesser sensitivity of the enzymatic method.

Feng X. et al. have reported a remarkable increase in LDH leakage, after exposure of rat cortical neuron cells to GdCl₃ (0.2, 2, 20 and 100 μ M) even at low micromolar doses (41). Almost in the same time, Xia Q. et al. (42) revealed that GdCl₃ (0.2, 2, and 20 μ M) induced in neurons a decrease of cell viability and an increase of LDH released onto the culture medium in a time- and concentration-dependent manner (43). Our data are not in accordance with these two studies, due to the fact that in our experiments Gd³⁺ was firstly complexed by DOTA or DOTP and secondly embedded in the nanogels.

The mechanisms by which Gd-based nanoparticles enters the cells are not entirely understood. Some studies demonstrated that, these nanoparticles are generally transported to cells by endocytic pathway depending on their size, shape and nature (44). Thus, mechanisms such as macropinocytosis (45), clathrin-mediated endocytosis (46), receptor-mediated endocytosis (47) were proposed. Dodane and Vilivalam reported that, chitosan and chitosan nanoparticles due to their positive charge,

enter the cells via cell membrane depolarization, leading to an increased Ca^{2+} influx (48).

Gd-based contrast agents have been intensively investigated with main focus on nephrotoxicity and nephrogenic systemic fibrosis (NSF) in patients with severe renal impairment (49–53). Our experiment showed that, both types of nanoparticles at a concentration of 5 μM induced moderate levels of reactive oxygen species in A-20 cells.

Lipid peroxidation is a process generated by the action of reactive oxygen species on polyunsaturated fatty acids. Malondialdehyde (MDA) the end product of lipid peroxidation is considered a mutagenic secondary product (54). In the present work, significant increases of MDA level, by 16% and 21% after 24h of exposure to 5 μM GdDOTACCS-TPP/HA and GdDOTPCCS-TPP/HA, respectively show that, the antioxidant system of A20 cells counteracted an extensive damage of lipids. After exposure of human bladder tumor cells 5637 to chitosan, it was suggested that chitosan could induce apoptosis via caspase-3 activation, without the generation of MDA (55). This suggests that the peroxidation cascade occurs differently in a cell type dependent manner.

GSH is the principal non-protein thiol in cells and a ROS scavenger molecule, being involved in maintaining intra- and extracellular redox environments (56, 57). GSH depletion, as a consequence of the oxidative stress was observed in peripheral blood mononuclear cells (PBMC) exposed to low-dose Gd-Tex (58).

In our experiment, the decrease of GSH level registered in cells exposed to GdDOTACCS-TPP/HA and GdDOTPCCS-TPP/HA- was correlated with the increase of MDA concentration.

The disturbance of the natural balance between pro-oxidant species and antioxidant systems in favor of the former, causes proteins oxidation. AOPP are formed via myeloperoxidase- H_2O_2 -halide system (59) and their role in activation of human neutrophil and monocyte oxidative metabolism was previously shown (60).

In our work, significant increases of AOPP, by 19% and 48%, were recorded after exposure to the higher concentration of GdDOTPCCS-TPP/HA NPs for 6h and 24h, respectively related to control, which are in a close correlation with GSH and MDA results.

Conclusions

Our data suggest that GdDOTPCCS-TPP/HA NPs and GdDOTACCS-TPP/HA NPs induced a moderate oxidative stress in A 20 cells which resulted in a slight decrease of GSH level and increase of MDA and AOPP concentrations. These effects were more pronounced in the case of GdDOTPCCS-TPP/HA NPs exposure. According to these, it appears that the cells antioxidant defense system counteracted to a certain extend the induced oxidative stress. Taking into account all of these, it appears that these NPs could be used in injectable formulations for MRI diagnostic.

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Conflict of interest statement

The authors declare no conflict of interest.

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