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High performance magneto-fluorescent nanoparticles assembled from terbium and gadolinium 1,3-diketones

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Polyelectrolyte-coated nanoparticles consisting of terbium and gadolinium complexes with calix[4] arene tetra-diketone ligand were first synthesized. The antenna effect of the ligand on Tb(III) green luminescence and the presence of water molecules in the coordination sphere of Gd(III) bring strong luminescent and magnetic performance to the core-shell nanoparticles. The size and the core-shell morphology of the colloids were studied using transmission electron microscopy and dynamic light scattering. The correlation between photophysical and magnetic properties of the nanoparticles and their core composition was highlighted. The core composition was optimized for the longitudinal relaxivity to be greater than that of the commercial magnetic resonance imaging (MRI) contrast agents together with high level of Tb(III)-centered luminescence. The tuning of both magnetic and luminescent output of nanoparticles is obtained via the simple variation of lanthanide chelates concentrations in the initial synthetic solution. The exposure of the pheochromocytoma 12 (PC 12) tumor cells and periphery human blood lymphocytes to nanoparticles results in negligible effect on cell viability, decreased platelet aggregation and bright coloring, indicating the nanoparticles as promising candidates for dual magneto-fluorescent bioimaging.

Noninvasive diagnostics provides *in situ* insight to the structural and functional features of the investigated systems and organs. The use of magnetic and optical techniques is emerging for these purposes. Improved sensing in tissues, which in turn can differentiate normal tissue from diseased one, can be achieved by using magnetic contrast agents (CA)^{1–4} and optical emissive probes^{5–9}. The f-elements (rare earth elements) are outstanding candidates for this, since their chelates combine unique magnetic and luminescent characteristics^{3,10,11}.

It is well known that trivalent lanthanide chelates are very good alternatives to organic luminescent stains or quantum dots in view of their extraordinary properties. Their emission bands span both the visible and near infrared (NIR) ranges and can be easily discriminated from the organic background by both emission wavelength and photoluminescence lifetime. They possess enhanced photostability, larger Stokes shift and longer excited state lifetime over conventional organic fluorophores^{11,12}.

Typically, all lanthanide luminescent labels contain an organic chromophore, which serves as antenna or sensitizer to absorb the excitation light and to transfer this energy to the emission levels of lanthanide ions. This antenna acts as a shield for the lanthanide ion from the solvent quenching effects as well as a reactive group for coupling the chelate complex to biotargets^{12,13}.

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Having seven unpaired electrons at the valence shell, gadolinium in the form of chelates provides high positive contrast between healthy and tumor tissues¹⁴. Despite the proven clinical efficacy of Gd-DTPA, Gd-DOTA and other gadolinium-based molecular MRI agents, they suffer from dechelation and transmetallation processes taking place in the presence of chelating anions and endogenous cations, correspondingly^{15,16}. At the same time, luminescent function of lanthanide-containing optical labels is quenched in presence of organic background in the biological sample due to antennae-ligands replacement^{11,17,18}.

Nanoparticulate approach has attracted increasing interest in recent years^{13,19–22}. Novel MRI CA based on gadolinium-containing nanoparticles show enhanced relaxation parameters compared to commercial specimens^{13,23–26}. The slowed down rotation of magnetic ions is the main reason for better relaxivity of the nanoparticulate CA versus mononuclear gadolinium complexes. Enhanced permeation, retention, lowered toxicity of nanoparticulate CA contribute to the enhanced positive imaging, as well^{27–29}.

Large number of attempts to combine the benefits of simultaneous magnetic and luminescent properties of lanthanides(III) were undertaken on oxide^{30–33}, fluoride^{34–37} and phosphate^{38–40} matrices, targeting the development of multimodal imaging materials. However, the development of novel nanoparticulate platforms with possibility to conjoin efficient magnetic and optical performance of lanthanides is still a challenge.

The precipitation of lanthanide complexes from water miscible organic solvents to polyelectrolyte-containing water solutions under effective stirring is an example of easy and effective synthetic pathway to embed large amount of lanthanide functional complexes to the nanoparticle. The synthesis of europium-doped luminescent polyelectrolyte colloids was first described by Mustafina *et al.*⁴¹ and later applied to various terbium⁴², and gadolinium⁴³ complexes.

Herein we report for the first time the synthesis and characterization of polystyrenesulfonate-coated nanoparticles consisting of terbium and gadolinium complexes with calix[4]arene tetra-diketone ligand. The new nanoparticles are rationally designed to simultaneously exhibit the optical properties of Tb and the magnetic behavior of Gd ions, towards bifunctional system with strong luminescence and high relaxivity. Colloidal, photophysical and magnetic studies were performed on nanoparticles with varying core composition, which is controlled by changing the Tb:Gd atomic ratio. We compared the relaxation parameters of the produced nanoparticles with commercial MRI agents. Particular attention was paid to the simplicity and universality of the strategy for the synthesis of bifunctional species. Tuning both magnetic and luminescent output of nanoparticles is possible via simple variation of composition of initial synthetic solution of lanthanides chelates. This synthetic strategy opens the door to new biomedical applications of lanthanide complexes.

Results and Discussion

Nanoparticle morphology. We synthesized polyelectrolyte-coated nanoparticles consisting of terbium(III) and gadolinium(III) complexes with calix[4]arene tetra-diketone ligand **1** (Fig. 1) according to the procedure described in the experimental section. According to our previous reports, the complex formation of **1** with both Gd(III)⁴³ and Tb(III)⁴⁴ occurs in 1:1 stoichiometry in alkaline DMF solutions. Our previous report⁴³ on the synthesis of PSS-stabilized Gd-**1** colloids demonstrated the ability of keeping the full amount of Gd-**1** complex within NP shell, thanks to the poor solubility of the complex, experimentally confirmed by luminescence measurements of the supernatant solutions in the synthesis of PSS-stabilized Tb-**1** colloids. This tendency provides great synthetic advantage, since the Tb-to-Gd ratio in the colloids can be easily driven by their ratio in the organic (DMF) solution. Transmission electron microscopy (TEM) and dynamic light scattering (DLS) techniques were applied to examine the morphology of the fabricated colloids. The aliquot of as-prepared colloidal solution was diluted ten times by deionized water to adjust the concentration of Ln(III) ions in the colloidal form to 0.075 mM to perform the DLS study.

The DLS measurements reveal hydrodynamic diameter being 100–170 nm in the obtained colloids (Table 1). These values are similar to those previously reported for other polyelectrolyte-coated lanthanide chelates^{17,42}. Their size distribution is narrow even without any prefiltration or fractionizing, which is clearly seen from the polydispersity (PDI) indices. Moreover, the sizes are stable during the period of one week at least.

The molar ratio of Ln (Tb or Gd) in the core of Ln-**1** based colloids was calculated via equation (1):

$$\chi_{\text{Gd}} = C_{\text{Gd}} / (C_{\text{Gd}} + C_{\text{Tb}}) \quad (1)$$

where C_{Gd} and C_{Tb} are the concentrations of the lanthanide chelates.

DLS and electrokinetic potential measurements confirm hard core-soft shell morphology of the colloids (Table 1), where the high negative charge of PSS-based exterior layer is the reason for their high colloidal stability. The DLS data reveal the size of hydrated PSS-coated colloids, while their functionality results from the hard cores inside the polyelectrolyte coating^{17,43,45}. The previously published reports^{17,43} highlight hard-soft morphology of the colloids. The TEM image of the dried colloids presented in Fig. 1g confirms the hard-soft nature of the colloids, where soft polyelectrolyte shell is evident from grey color spheres, while denser lanthanide-based cores can be clearly defined from the darker shade of grey color. The nanoparticle diameters are in the range 27–37 nm and the polyelectrolyte shell thickness is 2–3 nm. Sizes revealed from the TEM images of the dried aqueous colloids (Fig. 1g) are lower than those estimated by DLS measurements, because of their aggregation and swelling of polyelectrolyte coating in aqueous solutions. Moreover, the hard templates within the soft PSS coating are not uniform, consisting of smaller (2–10 nm sized) cores, as previously reported for PSS-stabilized Gd-**1** colloids⁴³. The TEM images of PSS-stabilized Tb-**1** colloids confirm similar morphology (Fig. 1h). Figure 1i schematically illustrates the core-shell morphology of PSS-coated Gd-**1**/Tb-**1** nanoparticle.

The size distribution of PSS-stabilized Gd-**1**/Tb-**1** colloids at various Gd content from the TEM data (Fig. S1 in SI) revealed no detectable effect of the Gd content on their size. The uniformity of both Gd- and Tb-based colloids is fully consistent with the isomorphism of Tb-**1** and Gd-**1** complexes. It is worth noting that the previously

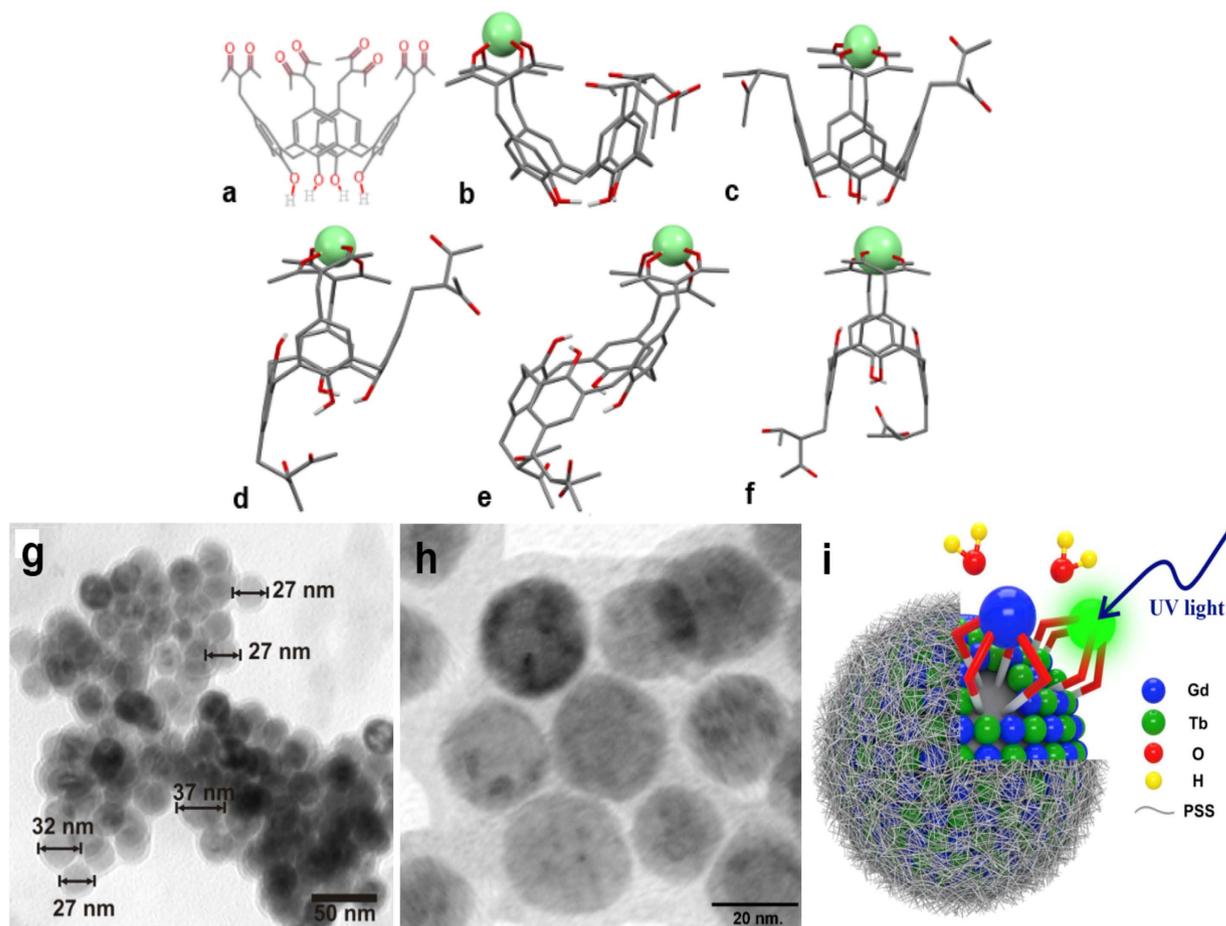


Figure 1. Structure of 5,11,17,23-tetrakis[(acetylaceton-3-yl)methyl]-25,26,27,28-tetrahydroxy-calix[4]arene **1** (a) and possible coordination modes of Ln-1 in the alkylated DMF solution (b–f). For more details see ref. 44. TEM image of dried PSS-coated Gd-1 colloids (g), higher resolution TEM image of the same colloids (h), and schematic illustration of PSS-coated Gd-1/Tb-1 nanoparticle and interactions on its core surface with water molecules and UV light (i).

χ_{Gd}	Diameter (nm)	PDI	ζ (mV)
0	113.6 ± 1.6	0.168	−29.9 ± 0.1
0.2	106.4 ± 2.0	0.153	−32.6 ± 0.2
0.4	144.4 ± 3.4	0.354	−31.9 ± 0.2
0.6	166.4 ± 2.8	0.238	−33.1 ± 0.2
0.8	123.5 ± 2.2	0.168	−28.3 ± 0.1
1	101.8 ± 1.7	0.180	−29.5 ± 0.1

Table 1. The hydrodynamic diameter, electrokinetic potential (ζ) and polydispersity indices (PDI) of PSS-covered polyelectrolyte nanoparticles as a function of Gd content ($\chi_{Gd} = 1, 0.8, 0.6, 0.4, 0.2, 0$; $\chi_{Tb} = 1 - \chi_{Gd}$).

reported amorphous nature of the hard templates of PSS-stabilized Gd-1 colloids results from metastability of the hard cores, probably due to reprecipitation of the complex from organic to aqueous solutions⁴³. Similar XRD measurements (for detailed description of the procedure see SI) were performed here for PSS-stabilized Tb-1 colloids. Both diffraction pattern and two-dimensional diffraction picture (Fig. 2) confirm the amorphous nature of the hard cores of the core-shell colloids. Moreover, XRD measurements of the ligand **1** (5,11,17,23-tetrakis[(acetylaceton-3-yl)methyl]-25,26,27,28-tetrahydroxy-calix[4]arene) also indicate its amorphous nature (Fig. S2 in SI).

Luminescence properties. It is known that the intraconfigurational 4f–4f transitions are formally forbidden, and thus possess very low molar absorptivities, limiting direct excitation of lanthanide-centered emission. Complexation with a sensitizing chromophore (referred as the antenna) allows overcoming this restriction,

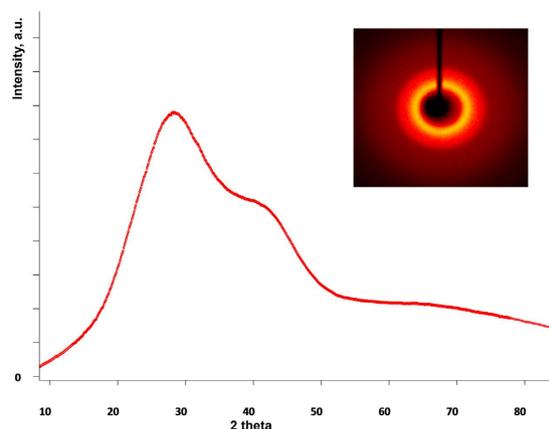


Figure 2. X-ray diffraction pattern of the PSS-stabilized Tb-1 colloids. Two-dimensional X-ray diffraction picture (inset).

χ_{Tb}	χ_{Gd}	I (a.u.)	τ_1 (ms)	τ_2 (ms)	r_1	r_2
1	0	5609000	0.271 ± 0.005	0.072 ± 0.002	—	—
0.8	0.2	4567000	0.277 ± 0.004	0.066 ± 0.002	13,46	17,17
0.6	0.4	4918000	0.284 ± 0.004	0.067 ± 0.002	10,67	12,72
0.4	0.6	5204000	0.296 ± 0.006	0.072 ± 0.003	7,62	9,02
0.2	0.8	2585000	0.291 ± 0.005	0.066 ± 0.002	8,15	9,36
0	1	3000	—	—	6,83	7,86

Table 2. Luminescence intensities (I), excited state lifetimes (τ), as well as r_1 and r_2 of PSS-stabilized Ln-1 (Ln = Tb, Gd) nanoparticles at various χ_{Tb} and χ_{Gd} .

providing indirect excitation of Ln(III) emission levels via energy transfer from singlet and triplet levels of organic ligand. 1,3-Diketones have been reported as a very promising antenna-ligands for the Ln(III)-centered emission^{46–48}. The luminescent properties of Tb-1 have recently been discussed⁴⁴.

The Ln-1 aqueous colloids with various χ_{Tb} exhibit emission under excitation at 320 nm with the spectrum patterns $^5D_4 \rightarrow ^7F_6$ (494 nm), $^5D_4 \rightarrow ^7F_5$ (545 nm), $^5D_4 \rightarrow ^7F_4$ (587 nm) peculiar for Tb(III)-centered luminescence (Fig. S3 in SI). The intensity of the main emission at 546 nm coming from the $^5D_4 \rightarrow ^7F_5$ transition is applied for the quantitative evaluation of the change of luminescence intensity under the variation of χ_{Tb} . The gradual increase of luminescence intensity with increase of number of Tb emission centers in the core of nanoparticles is expected. This trend is observed when χ_{Tb} increases from 0.2 to 0.6, while it is less pronounced when χ_{Tb} increases from 0.6 to 1.0 (Table 2). The time resolved luminescence measurements were performed in PSS-stabilized Tb-1/Gd-1 colloids at various χ_{Tb} . The decay curves of Tb(III)-centered luminescence (Figs S4–S8 in SI) are well fitted by a biexponential decay. Monoexponential decay results in poorer fitting parameters. The obtained excited state lifetime values shown in Table 2 point to the presence of Tb(III)-containing species with longer ($\tau_1 = 0.271$ – 0.296 ms) and shorter ($\tau_2 = 0.066$ – 0.072 ms) values. The origin of the double τ values most probably arises from chelated and dechelated forms of Tb(III) species within PSS-stabilized Tb-1/Gd-1 nanoparticles. Taking into account the size of PSS-stabilized Tb-1/Gd-1 nanoparticles (Fig. 1) a dechelation of Tb(III) complexes at the interface is the most probable reason for the origin of the dechelated Tb(III) forms. The higher τ_1 values tend to remain unchanged with the χ_{Tb} increase from 0.2 to 0.6, while gradually decrease with the increase in χ_{Tb} from 0.6 to 1.0, while the smaller τ_2 values remain almost constant (Table 2). It is also worth noting that τ_1 and luminescence intensity present opposite trends as a function of χ_{Tb} . The decreased intensity as a function of increasing χ_{Tb} can be explained by the concentration-induced quenching of Tb(III)-centered luminescence^{49,50}. The concentration-induced quenching occurs when the distance between emission centers is small enough for efficient cross relaxation ($^5D_3 \rightarrow ^7F_6$) \rightarrow ($^5D_4 \rightarrow ^7F_0$) between two neighboring Tb(III) ions⁵¹.

The above-mentioned amorphous nature of PSS-stabilized Ln-1 colloids (Fig. 2) confirms the dense packing of lanthanide centers within hard cores of the colloids. The obtained results reveal the optimal composition ($\chi_{Tb} = 0.4$) of PSS-stabilized Tb-1/Gd-1 nanoparticles for the best combination of steady state intensity and excited state lifetime values.

Magnetic properties. Longitudinal and transverse relaxation times T_1 and T_2 of water molecule protons in the presence of studied paramagnetic Gd(III)-containing colloids were registered at 20 MHz magnetic field frequency (Tables S1–S6 in SI). The $1/T_1$ and $1/T_2$ plots versus Gd(III) concentration of PSS-stabilized Tb-1/Gd-1 aqueous colloids at various χ_{Gd} are presented in Fig. 3a,b and Table 2. The corresponding longitudinal and transverse relaxivities (r_1 and r_2) of PSS-stabilized Ln-1 nanoparticles were calculated as a tangent of plots' incline angle at $1/T_{1(2)}$ vs C_{Gd} coordinates and presented in Fig. 3c as a function of χ_{Gd} .

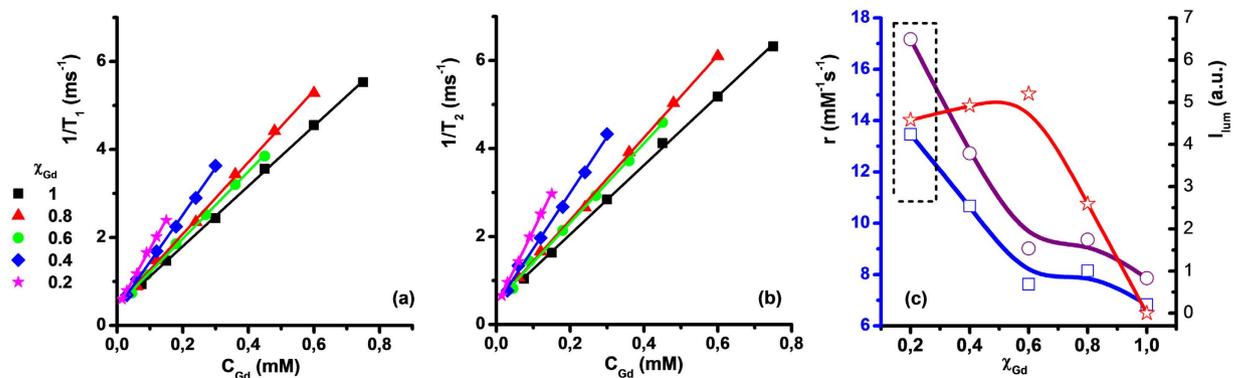


Figure 3. $1/T_1$ (a) and $1/T_2$ (b) of PSS-stabilized Ln-1 nanoparticles with various χ_{Gd} : 1.0 (1), 0.8 (2), 0.6 (3), 0.4 (4), 0.2 (5) versus Gd(III) concentration. Straight lines are linear fitting of the experimental data. Relaxivity values r_1 (1-blue) and r_2 (2-purple), luminescence intensities (3-red) of PSS-stabilized Ln-1 colloids versus χ_{Gd} (c). Dashed rectangle reflects the optimal χ_{Gd} in the colloids core exhibiting the best magnetic and luminescent parameters.

It is worth noting that r_1 and r_2 values keep almost constant, when χ_{Gd} decreases from 1.0 to 0.6. The further decrease in χ_{Gd} from 0.6 to 0.2 results in the increase of r_1 and r_2 (Fig. 3c). The explanation of this tendency should be preceded by discussing on main mechanisms contributing to relaxation of water protons at the colloid/water interface. Albeit no theoretical framework is known for accurate interpretation of relaxivities in Gd(III)-based colloids, several mechanisms are proposed as predominantly contributing to enhancing water protons relaxivity in Gd(III)-based aqueous colloids. Slow rotation resulting from inclusion of mononuclear Gd(III) complexes into nanoparticles should be claimed as the main reason for high relaxivity in Gd(III)-based aqueous colloids. This factor is guided by size of the hard cores inside the polyelectrolyte coating. Taking into account that exchange between inner-sphere and bulk water molecules is another key factor affecting relaxivity of Gd(III)-based colloids, size-dependent ratio of Gd(III)-centers localized close to the surface to those localized inside the hard cores is also worth noting. The size of the hard cores inside the PSS-coating varies from 2 to 10 nm for PSS-stabilized Gd-1 colloids⁴³. According to TEM results, the size of the colloids is not greatly affected by the change in χ_{Gd} , although no exact assignment of the smaller versus larger nanoparticles within the polyelectrolyte coating to Tb-1 or Gd-1 complex, as well as their mixture, can be done. The increase in r_1 and r_2 for χ_{Gd} from 0.6 to 0.2 points to predominant localization of Gd(III) centers close to the colloid/water interface at low χ_{Gd} values. The reasons for this behavior are not clear, and further investigations are ongoing.

The luminescence intensities (I) of PSS-stabilized Ln-1 colloids were plotted in Fig. 3c together with the r_1 and r_2 values versus χ_{Gd} in order to show the correlation between the property and the composition of the colloids. Dashed rectangle shows the optimal χ_{Gd} in the colloids core exhibiting the best magnetic and luminescent parameters.

Taking into account that both molecular and nanoparticulate (PSS-stabilized colloids) forms of Tb-1 suffer from different radiationless decay mechanisms, it is interesting to evaluate the ratio of luminescence intensity measured in aqueous PSS-stabilized colloids of Tb-1 (I_{NP}) to that in DMF solution of Tb-1 (I_{MOL}) at the same complex concentration (0.75 mM). The measured spectra (Fig. S9 in SI) point to the stronger luminescence intensity of the colloids versus the complex in solution. The ratio I_{NP}/I_{MOL} is equal to 1.7 ± 0.2 , which argues in favor of application of PSS-stabilized Tb-1 colloids versus Tb-1 complexes as contrast agents in bioimaging.

Effect of the colloids on cell viability and platelets aggregation. The toxic effect of PSS-stabilized Tb-1 and Gd-1 colloids on human blood lymphocytes and pheochromocytoma (PC12) cells were determined by means of MTT assay (based on tetrazolium dye MTT 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) and trypan blue viability test, respectively. The obtained results indicate no toxic effect of the colloids on human blood lymphocytes at 32 $\mu\text{g}/\text{mL}$ of PSS-coated Gd-1 colloids and 66 $\mu\text{g}/\text{mL}$ of PSS-coated Tb-1 colloids (Fig. 4a). Both 50 $\mu\text{g}/\text{mL}$ and 500 $\mu\text{g}/\text{mL}$ of Tb-1 colloids show no negative effect on PC12 cell viability. The number of viable cells was calculated as 99.4% and 97.7%, respectively.

Effect of the colloids on platelet aggregation is of great importance for their biomedical applicability due to the risk of formation of blood clots. The effect of the colloids on platelet aggregation was determined in the 200 μl mixture of 50 μl of platelets mass in Hanks balanced salt solution (HBSS), and 150 μl of PBS (0.1 M, pH 7.2) in the presence of PSS-stabilized Ln-1 colloids. The values in Fig. 4b indicate the decrease in the rates of platelet aggregation after the admixture with the colloids. The obtained results indicate low thrombogenic potential of PSS-stabilized Tb-1 colloids, which along with their low cytotoxicity effect point to potential applicability of the colloids in bioimaging.

Interaction of colloids with cells using fluorescent microscopy. A cellular uptake behavior of luminescent nanoparticles is affected by several factors, including size, exterior charge and aggregation of nanoparticles in physiological media^{52,53}. Taking into account high negative electrokinetic potential values of PSS-stabilized colloids (Table 1), very poor, if any, cellular uptake behavior of the nanoparticles is anticipated due to weak

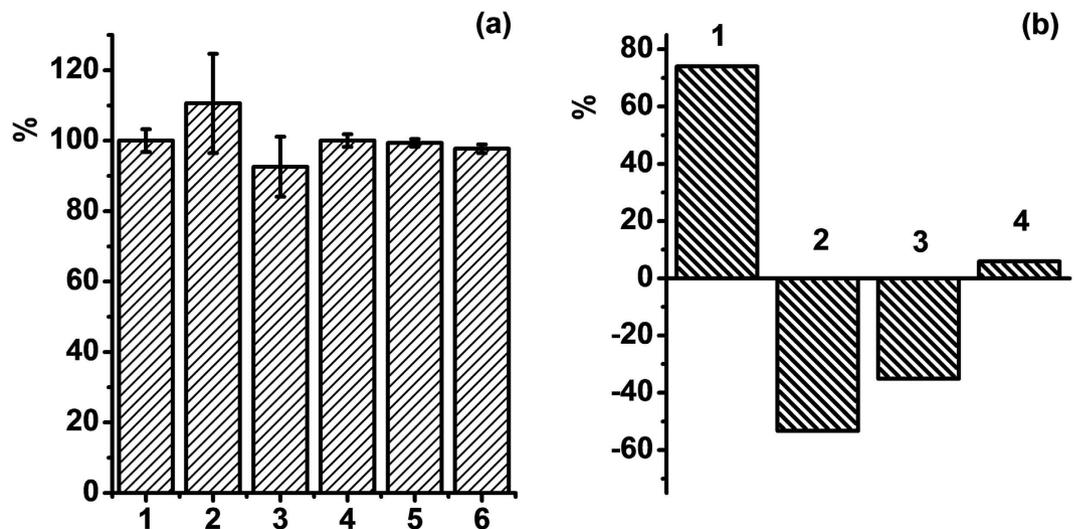


Figure 4. (a) Cell viability of periphery human blood lymphocytes without (1) and in the presence of 32 $\mu\text{g}/\text{mL}$ of PSS-coated Gd-1 (2), and 66 $\mu\text{g}/\text{mL}$ of Tb-1 (3) nanoparticles; PC 12 tumor cells (4) in the presence of 50 $\mu\text{g}/\text{mL}$ of PSS-coated Tb-1 colloids (5) and 500 $\mu\text{g}/\text{mL}$ of PSS-coated Tb-1 colloids (6). (b) Platelet aggregation in the presence of 10 μM adrenaline (1), 86 $\mu\text{g}/\text{mL}$ PSS-coated Gd-1 (3), and 193 $\mu\text{g}/\text{mL}$ PSS-coated Tb-1 (4) nanoparticles and without any additives (2) as a per cent of absorptivity changes at 540 nm after 5 min.

interaction of negatively charged nanoparticles with cellular membrane. Nevertheless, interactions of the nanoparticles with inorganic and biological background in physiological media can affect the localization of nanoparticles on cell membrane. Thus, DLS measurements were performed for the aqueous colloids in PBS buffer before and after the protein admixture, since protein corona is well documented factor, affecting both aggregation and cellular uptake behavior of nanoparticles^{54,55}. The size increase from ~ 100 to ~ 500 nm (Table 1 and S7 in SI) is observed from DLS measurements of PSS-stabilized Tb-1 colloids in water and in PBS buffer (0.1 M). This tendency indicates that the exterior charge neutralization resulted from the binding with counterions. Aqueous buffer solutions of bovine serum albumin (BSA), which was applied as a model protein, are characterized by 10–11 nm aggregates (Table S7 in SI). The admixture of BSA ($1 \text{ g}\cdot\text{L}^{-1}$) to PSS-stabilized colloids in the buffer solutions results in the size decrease to ~ 400 nm, although this trend can be explained by the contribution of the smaller aggregates of BSA.

Figure 5 illustrates bright field and confocal fluorescent images of rat PC12 cells after their exposure for 2 hours at 37 $^{\circ}\text{C}$ to PSS-stabilized Tb-1 colloids. The images indicate predominant localization of the emissive nanoparticles at the cellular membrane, although some cellular uptake cannot be excluded. It is worth noting that cellular uptake behavior can be enhanced by the recharging of the exterior layer of PSS-stabilized Tb-1 colloids, which will be done in the nearest future, although the represented results highlight a potential of PSS-stabilized Tb-1 colloids as a basis for imaging applications.

Conclusions

Core-shell morphology nanoparticles with terbium and gadolinium complexes with calix[4]arene tetra-diketone ligand (1) as hard core and polystyrenesulfonate coating as soft shell as promising basis for magneto-luminescent imaging are introduced for the first time. Water dispersed polyelectrolyte-stabilized Ln-1 nanoparticles exhibit high colloidal stability. Their size is about 30 nm in the dried state, increasing to about 100 nm in aqueous solutions due to hydration effect. Luminescent and magnetic relaxation properties of the colloids are greatly dependent on Gd-1:Tb-1 ratio. The latter can be easily tuned for best bifunctional magneto-luminescent performance by simple mixing of Tb(III) and Gd(III) complexes in the initial DMF solution at the desired ratio. Both of Ln-1 colloids are improved by appropriate variation of Gd-1:Tb-1 ratio. The optimized composition (Gd-1:Tb-1 = 0.2/0.8) of the colloids for optimal Tb(III)-centered luminescence and Gd(III)-based relaxivity results in better performance than the commercial Gd(III)-contrast agents. The low cytotoxicity and thrombogenic potential of PSS-stabilized Tb-1 colloids open the door for their application in biomarking. Confocal microscopy imaging reveals predominant localization of the emissive nanoparticles at the cellular membrane, which is in agreement with the negative exterior charge of PSS-stabilized Tb-1 colloids.

Experimental section

Reagents and materials. Gadolinium nitrate $\text{Gd}(\text{NO}_3)_3\cdot 6\text{H}_2\text{O}$ (99.9%) (Alfa Aesar), terbium nitrate hydrate $\text{Tb}(\text{NO}_3)_3\cdot 5\text{H}_2\text{O}$ (Alfa Aesar) triethylamine (TEA) (Acros Organics), poly(sodium 4-styrenesulfonate) (PSS) ($MW_{\text{average}} = 70000$) (Acros Organics), sodium chloride (Sigma-Aldrich), were used as commercially received without further purification. N,N-Dimethylformamide (DMF) (Acros Organics) was twice distilled over P_2O_5 .

Synthesis of 5,11,17,23-tetrakis[(acetylaceton-3-yl)methyl]-25,26,27,28-tetrahydroxy-calix[4]arene was reported previously⁴³.

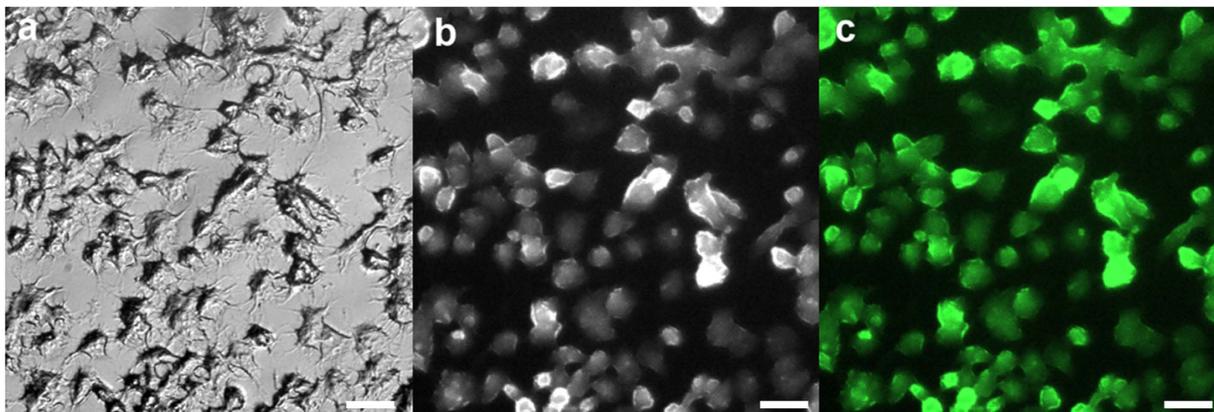


Figure 5. Images of PC 12 tumor cells: optical (a); confocal in the presence of PSS-coated Tb-1 nanoparticles (b); colored rework of b using GFP emission filter (510–550 nm) (c), $\lambda_{\text{ex}} = 360\text{--}370\text{ nm}$. Scale bar indicates 20 μm .

Synthesis of colloids. The colloids were synthesized via precipitation of water non-soluble lanthanide(III) complexes from DMF solution to polyelectrolyte containing aqueous solution. Four equivalents of triethylammonia additives ($C = 18\text{ mM}$) to DMF solution of 4.5 mM calix[4]arene 1 were followed by lanthanide(III) ($C = 4.5\text{ mM}$) addition to promote the formation of 1:1 complex between lanthanide(III) cations and anions of 1. The aliquote of 1 ml DMF dissolved lanthanide(III) complex was added to the 5 ml of polystyrenesulfonate sulfonate aqueous solution ($1\text{ g}\cdot\text{L}^{-1}$) in the presence of NaCl ($C = 0.5\text{ M}$) at the effective stirring (2200 rpm) dropwisely using the syringe pump. The solution becomes turbid since the dispersion of polyelectrolyte stabilized nanoprecipitate forms. The obtained colloids were washed from the DMF and excess amounts of PSS and NaCl via triple centrifugation (11000 rpm, 10 min.)/decantation and redispersion (deionized water addition and ultrasonication for 30 min.) procedures.

Six syntheses were performed from DMF solutions with different Gd/Tb molar ratios ($\chi_{\text{Gd}} = 0, 0.2, 0.4, 0.6, 0.8$ and 1) within the core of polyelectrolyte coated colloids. Two equimolar solutions of Gd-1 and Tb-1 in DMF were prepared ($C = 4.5\text{ mM}$) for this purpose. The Gd/Tb core composition was varied via volume of aliquots of Gd-1 and Tb-1 in initial DMF solution. The colloids were ultrasonicated for 30 min before measurements. All measurements have been performed in triplicates.

Methods of characterization of colloids. DLS measurements were performed using Malvern Mastersize 2000 particle analyzer operating with a He-Ne laser (633 nm) and emitting vertically polarized light as a light source. The transmission electron microscopy (TEM) images have been obtained with Hitachi HT7700, Japan. Samples have been sonicated in water for 30 min and then dispersed on 200 mesh copper grids with continuous formvar support films. The images have been acquired at an accelerating voltage of 100 kV.

The steady-state luminescence and time-resolved spectra have been recorded on a spectrofluorometer FL3-221-NIR (Jobin Yvon). Excitation of samples has been performed at 320 nm, and emission detected at 545 nm with 1 nm slit width for both excitation and emission.

Bruker “Minispec mq20” NMR analyzer was employed to measure T_1 and T_2 of water molecule protons in studied solutions at 20 MHz magnetic field. Inversion-recovery and Carr-Purcell-Meiboom-Gill (CPMG) pulse sequences⁵⁶ were used for longitudinal T_1 and transverse T_2 relaxation times accordingly with 20 points data collected for fitting⁵⁷.

Powder X-ray diffraction (PXRD) measurements were performed on a Bruker D8 Advance diffractometer equipped with Vario attachment and Vantec linear PSD, using Cu radiation (40 kV, 40 mA) monochromated by the curved Johansson monochromator ($\lambda\text{ Cu K}_{\alpha 1} 1.5406\text{ \AA}$). Room-temperature data were collected in the reflection mode with a flat-plate sample. The samples were loaded on a standard zero diffraction silicon plate, which was kept spinning (15 rpm) throughout the data collection. Patterns were recorded in the 2θ range between 3° and 60° , in 0.008° steps, with a step time of 0.1–4.0 s. Several diffraction patterns in various experimental modes were collected and summed for the sample. Processing of the obtained data performed using EVA⁵⁸ software packages. Additional experiments were carried out on a single-crystal X-ray diffractometer Bruker Smart Apex II CCD using Mo radiation (MoK_{α} , graphite monochromator, $\lambda 0.71073\text{ \AA}$) at the 23°C with the liquid samples in a standard glass 1 mm capillary. CCD-detector of the diffractometer was kept on the fixed values of 2θ angles (0° and 30°), recording time was varied from 30 to 400 seconds. Standard powder diffractograms were obtained by integration of several two-dimension scattering patterns with the use of software packages APEX2⁵⁹.

Cell viability test. Cell viability of human blood lymphocytes towards nanoparticles was estimated using MTT proliferative test (Promega, USA). Yellow MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium bromide) turns into purple formazan under the effect of mitochondrial NADN reductase of living cells. After the lysis of the cells formazan dissolves in the DMSO. The activity of the mitochondrial reductase (and cell viability respectively) has been defined as the function of the formazan absorbance (500–600 nm).

Human blood lymphocytes (7500 cells per well) were cultivated in $100\mu\text{l}$ of DMEM in the presence of nanoparticles under the standard conditions in a 96-well cultural plate during 3.5 hours. Cells were washed using PSB

(phosphate-buffered saline (PBS; 10 mM/L; pH 7.2) and the mixture of 80 μ l of DMEM and 20 μ l of nanoparticles was added. Then the 20 μ l aliquot of 5 mg/ml MTT was added to each well and incubated in CO₂-incubator under the 37 °C for 3.5 hours. The medium was removed then and 150 μ l of DMSO were added. The absorbance of the resulted solutions was measured at 590 nm after 10 minutes using plate spectrophotometer Stat Fax 2100 (Awareness Technology, USA).

The effect of NPs on viability of PC12 cells was assessed by the trypan blue exclusion test. All NPs samples were sonicated for 30 min prior the experiment. Solutions of NPs were prepared by diluting the stock solution of NPs in cell growth media and adjusting to pH 7.4. Cells were seeded in flasks and, after confluence was reached, they were exposed for 2 h with 0 (control), 50 μ g/mL and 500 μ g/mL of Tb-1 colloids. Conditions of exposure were 5% CO₂, 37 °C, 100% humidity. Thereafter, cells were trypsinized, by standard trypsinization, and resuspended in equal volumes of culture medium and trypan. Viable (unstained) and nonviable (blue-stained) cells were counted using a haemocytometer, and cell viability was calculated as (N viable cells/N total cells) \times 100%.

Platelets aggregation. The citrated human blood was centrifuged at 120-g for 15 min at room temperature to obtain platelets mass containing 25-10⁷ of platelets per ml. Platelets concentration was determined with automatic hematology analyzer Abacus (Diatron, Austria) according to the manufacturer method.

The platelets aggregation was detected in a 200 μ l mixture of 50 μ l of platelets mass in HBSS (125-10⁵ of platelets), 50 μ l of 0.188 mM/L water dispersion of nanoparticles and 100 μ l of PBS (0.1 M, pH 7.2). Platelet aggregation was recorded after 5 min.

Positive control 200 μ l mixture contained 50 μ l of platelets mass in HBSS, adrenaline (10 μ M, Sigma) and 100 μ l of PBS (0.1 M, pH 7.2).

The spontaneous platelets aggregation was examined in a 200 μ l mixture of 50 μ l of platelets mass in HBSS, and 150 μ l of PBS (0.1 M, pH 7.2). The absorbance of the resulted solutions at 540 nm was measured using spectrophotometer Shimadzu (Japan) at 37 °C. The extent of aggregation was calculated quantitatively as a percentage of the maximal change in optical density.

The authors confirm that all methods were carried out in accordance with relevant guidelines and regulations. A Kazan Federal university committee approved all experimental protocols. Informed consent was obtained from all volunteers. There are no animal experiments included in the project.

References

- Kandasamy, G. & Maity, D. Recent advances in superparamagnetic iron oxide nanoparticles (SPIONs) for *in vitro* and *in vivo* cancer nanotheranostics. *Int. J. Pharm.* **496**, 191–218 (2015).
- Jung, J. *et al.* Europium-doped gadolinium sulfide nanoparticles as a dual-mode imaging agent for T1-weighted MR and photoluminescence imaging. *Biomaterials* **33**, 5865–5874 (2012).
- Heffern, M. C., Matosziuk, L. M. & Meade, T. J. Lanthanide Probes for Bioresponsive Imaging. *Chem Rev* **114**, 4496–4539 (2014).
- Shokrollahi, H. Contrast agents for MRI. *Mater Sci Eng, C* **33**, 4485–4497 (2013).
- Lim, S.-Y., Hong, K.-H., Kim, D. I., Kwon, H. & Kim, H.-J. Tunable Heptamethine-Azo Dye Conjugate as an NIR Fluorescent Probe for the Selective Detection of Mitochondrial Glutathione over Cysteine and Homocysteine. *J Am Chem Soc* **136**, 7018–7025 (2014).
- Song, B. *et al.* Time-resolved lanthanide luminescence for lab-on-a-chip detection of biomarkers on cancerous tissues. *Analyst* **134**, 1991–1993 (2009).
- Naczynski, D. J., Tan, M. C., Riman, R. E. & Moghe, P. V. Rare earth nanoprobe for functional biomolecular imaging and theranostics. *J Mater Chem B* **2**, 2958–2973 (2014).
- Naccache, R. *et al.* High Relaxivities and Strong Vascular Signal Enhancement for NaGdF₄ Nanoparticles Designed for Dual MR/Optical Imaging. *Advanced Healthcare Materials* **2**, 1478–1488 (2013).
- Laurent, S., Vander, E. L. & N., M. R. Lanthanide complexes for magnetic resonance and optical molecular imaging. *Q J Nucl Med Mol Imaging* **53**, 586–603 (2009).
- Eliseeva, S. V. & Bunzli, J.-C. G. Rare earths: jewels for functional materials of the future. *New J Chem* **35**, 1165–1176 (2011).
- Bünzli, J.-C. G. Lanthanide Luminescence for Biomedical Analyses and Imaging. *Chem Rev* **110**, 2729–2755 (2010).
- Amoroso, A. J. & Pope, S. J. A. Using lanthanide ions in molecular bioimaging. *Chem Soc Rev* **44**, 4723–4742 (2015).
- Wang, X. *et al.* Recent developments in lanthanide-based luminescent probes. *Coord Chem Rev* **273–274**, 201–212 (2014).
- Caravan, P., Ellison, J. J., McMurry, T. J. & Lauffer, R. B. Gadolinium(III) Chelates as MRI Contrast Agents: Structure, Dynamics, and Applications. *Chem Rev* **99**, 2293–2352 (1999).
- Caroline, R., Sarah, C., Marie-Christine, G., Jean-Marc, I. & Marc, P. The role of phosphate on Omniscan[®] dechelation: an *in vitro* relaxivity study at pH 7. *BioMetals* **24**, 759–768 (2011).
- Perazella, M. A. Nephrogenic Systemic Fibrosis, Kidney Disease, and Gadolinium: Is There a Link? *Clinical Journal of the American Society of Nephrology* **2**, 200–202 (2007).
- Shamsutdinova, N. *et al.* Interfacial interactions of hard polyelectrolyte-stabilized luminescent colloids with substrates. *Colloids Surf, A* **482**, 231–240 (2015).
- Davydov, N. *et al.* Determination of fluoroquinolone antibiotics through the fluorescent response of Eu(III) based nanoparticles fabricated by layer-by-layer technique. *Anal Chim Acta* **784**, 65–71 (2013).
- Na, H. B., Song, I. C. & Hyeon, T. Inorganic Nanoparticles for MRI Contrast Agents. *Adv Mater* **21**, 2133–2148 (2009).
- Yang, M. *et al.* Characterization of Fe₃O₄/SiO₂/Gd₂O₃(CO₃)₂ core/shell/shell nanoparticles as T1 and T2 dual mode MRI contrast agent. *Talanta* **131**, 661–665 (2015).
- Gao, Z. *et al.* Small is Smarter: Nano MRI Contrast Agents – Advantages and Recent Achievements. *Small* **12**, 556–576 (2016).
- Pihlasalo, S., Kirjavainen, J., Hänninen, P. & Härmä, H. High Sensitivity Luminescence Nanoparticle Assay for the Detection of Protein Aggregation. *Anal Chem* **83**, 1163–1166 (2011).
- Peng, E., Wang, F. & Xue, J. M. Nanostructured magnetic nanocomposites as MRI contrast agents. *J Mater Chem B* **3**, 2241–2276 (2015).
- Lee, S. H., Kim, B. H., Na, H. B. & Hyeon, T. Paramagnetic inorganic nanoparticles as T1MRI contrast agents. *Wiley Interdisciplinary Reviews: Nanomedicine and Nanobiotechnology* **6**, 196–209 (2014).
- Caravan, P. Strategies for increasing the sensitivity of gadolinium based MRI contrast agents. *Chem Soc Rev* **35**, 512–523 (2006).
- Ahrén, M. *et al.* Synthesis and Characterization of PEGylated Gd₂O₃ Nanoparticles for MRI Contrast Enhancement. *Langmuir* **26**, 5753–5762 (2010).
- Zhou, S., Wu, Z., Chen, X., Jia, L. & Zhu, W. PEGylated Polyethylenimine as Enhanced T1 Contrast Agent for Efficient Magnetic Resonance Imaging. *ACS Applied Materials & Interfaces* **6**, 11459–11469 (2014).

28. Arias, J. L. Advanced methodologies to formulate nanotheragnostic agents for combined drug delivery and imaging. *Expert opinion on drug delivery* **8**, 1589–1608 (2011).
29. Klasson, A. *et al.* Positive MRI contrast enhancement in THP-1 cells with Gd₂O₃ nanoparticles. *Contrast Media & Molecular Imaging* **3**, 106–111 (2008).
30. Hu, Z. *et al.* Highly Water-Dispersible Surface-Modified Gd₂O₃ Nanoparticles for Potential Dual-Modal Bioimaging. *Chemistry – A European Journal* **19**, 12658–12667 (2013).
31. Pectoral, R. M. *et al.* Synthesis and Characterization of Tb³⁺-Doped Gd₂O₃ Nanocrystals: A Bifunctional Material with Combined Fluorescent Labeling and MRI Contrast Agent Properties. *J Phys Chem C* **113**, 6913–6920 (2009).
32. Shi, Z., Neoh, K. G., Kang, E. T., Shuter, B. & Wang, S.-C. Bifunctional Eu³⁺-doped Gd₂O₃ nanoparticles as a luminescent and T1 contrast agent for stem cell labeling. *Contrast Media & Molecular Imaging* **5**, 105–111 (2010).
33. Chen, F., Chen, M., Yang, C., Liu, J., Luo, N., Yang, G., Chen, D. & Li, L. Terbium-doped gadolinium oxide nanoparticles prepared by laser ablation in liquid for use as a fluorescence and magnetic resonance imaging dual-modal contrast agent. *Phys. Chem. Chem. Phys.* **17**, 1189–1196 (2015).
34. Qiao, X.-F. *et al.* Triple-functional core-shell structured upconversion luminescent nanoparticles covalently grafted with photosensitizer for luminescent, magnetic resonance imaging and photodynamic therapy *in vitro*. *Nanoscale* **4**, 4611–4623 (2012).
35. Grzyb, T. *et al.* Synthesis, characterization, and cytotoxicity in human erythrocytes of multifunctional, magnetic, and luminescent nanocrystalline rare earth fluorides. *J Nanopart Res* **17**, 1–18 (2015).
36. Passuello, T. *et al.* PEG-capped, lanthanide doped GdF₃ nanoparticles: luminescent and T2 contrast agents for optical and MRI multimodal imaging. *Nanoscale* **4**, 7682–7689 (2012).
37. Ju, Q. *et al.* Amine-Functionalized Lanthanide-Doped KGdF₄ Nanocrystals as Potential Optical/Magnetic Multimodal Bioprobes. *J Am Chem Soc* **134**, 1323–1330 (2012).
38. Chen, F., Huang, P., Zhu, Y.-J., Wu, J. & Cui, D.-X. Multifunctional Eu³⁺/Gd³⁺ dual-doped calcium phosphate vesicle-like nanospheres for sustained drug release and imaging. *Biomaterials* **33**, 6447–6455 (2012).
39. Zhang, L. *et al.* Multifunctional GdPO₄:Eu³⁺ Hollow Spheres: Synthesis and Magnetic and Luminescent Properties. *Inorg Chem* **50**, 10608–10613 (2011).
40. Ren, W. *et al.* Lanthanide ion-doped GdPO₄ nanorods with dual-modal bio-optical and magnetic resonance imaging properties. *Nanoscale* **4**, 3754–3760 (2012).
41. Mustafina, A. *et al.* Synthesis and photophysical properties of colloids fabricated by the layer-by-layer polyelectrolyte assembly onto Eu(III) complex as a core. *Colloids Surf, B* **88**, 490–496 (2011).
42. Shamsutdinova, N. A. *et al.* A facile synthetic route to convert Tb(III) complexes of novel tetra-1,3-diketone calix[4]resorcinarene into hydrophilic luminescent colloids. *New J Chem* **38**, 4130–4140 (2014).
43. Shamsutdinova, N. *et al.* Polyelectrolyte-stabilized nanotemplates based on Gd(III) complexes with macrocyclic tetra-1,3-diketones as a positive MR contrast agents. *Chemistry Select* **1**, 1377–1383 (2016).
44. Zairov, R. *et al.* Structure impact in antenna effect of novel upper rim substituted tetra-1,3-diketone calix[4]arenes on Tb(III) green and Yb(III) NIR-luminescence. *Tetrahedron* **72**, 2447–2455 (2016).
45. Zairov, R. *et al.* Impact of polyelectrolyte coating in fluorescent response of Eu(III)-containing nanoparticles on small chelating anions including nucleotides. *Surf Coat Technol* **271**, 242–246 (2015).
46. Binnemans, K. In *Handbook on the Physics and Chemistry of Rare Earths* Vol. 35 (eds Jean-Claude, G., Bünzli Karl, A. Gschneidner & K. Pecharsky Vitalij) 107–272 (Elsevier, 2005).
47. Mironov, L. Y., Sveshnikova, E. B. & Ermolaev, V. L. Energy transfer from the singlet levels of diketones and dyes to lanthanide ions in nanoparticles consisting of their diketone complexes. *Opt Spectrosc* **116**, 933–940 (2014).
48. Smith, L. F., Blight, B. A., Park, H.-J. & Wang, S. Sensitizing Tb(III) and Eu(III) Emission with Triarylboron Functionalized 1,3-Diketone Ligands. *Inorg Chem* **53**, 8036–8044 (2014).
49. Zhao, C.-J. *et al.* White light emission from Eu³⁺/Tb³⁺/Tm³⁺ triply-doped aluminoborate glass excited by UV light. *J Non-Cryst Solids* **358**, 604–608 (2012).
50. Su, Q. *et al.* The Effect of Surface Coating on Energy Migration-Mediated Upconversion. *J Am Chem Soc* **134**, 20849–20857 (2012).
51. Zhang, Y. *et al.* Tunable luminescence and energy transfer from Gd³⁺ to Tb³⁺ ions in silicate oxyfluoride scintillating glasses via varying Tb³⁺ concentration. *J Non-Cryst Solids* **423–424**, 30–34 (2015).
52. Shang, L., Nienhaus, K. & Nienhaus, G. U. Engineered nanoparticles interacting with cells: size matters. *J Nanobiotechnology* **12**, 1–11 (2014).
53. Graf, C. *et al.* Surface Functionalization of Silica Nanoparticles Supports Colloidal Stability in Physiological Media and Facilitates Internalization in Cells. *Langmuir* **28**, 7598–7613 (2012).
54. Lynch, I. *et al.* The nanoparticle–protein complex as a biological entity; a complex fluids and surface science challenge for the 21st century. *Adv Colloid Interface Sci* **134–135**, 167–174 (2007).
55. Zairov, R. R. *et al.* Nanoparticles on the basis of complexes of gadolinium(III) and europium(III) for bioimaging *Russ. Chem. Bull.* **55**, 1325–1331 (2016).
56. Meiboom, S. & Gill, D. Modified Spin-Echo Method for Measuring Nuclear Relaxation Times. *Rev Sci Instrum* **29**, 688–691 (1958).
57. Henoumont, C., Laurent, S. & Vander Elst, L. How to perform accurate and reliable measurements of longitudinal and transverse relaxation times of MRI contrast media in aqueous solutions. *Contrast Media & Molecular Imaging* **4**, 312–321 (2009).
58. DIFFRAC Plus Evaluation package EVA v. 11 (Bruker AXS, Karlsruhe, Germany, 2005).
59. APEX2 v. 2.1 SAINTPlus. Data Reduction and Correction Program (Ver. 7.31A) (Bruker AXS Inc., Madison, Wisconsin, USA, 2006).

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Author Contributions

Polystyrenesulfonate-coated nanoparticles consisting of terbium and gadolinium complexes with calix[4] arene tetra-diketone ligand were synthesized and characterized using luminescent spectroscopy, DLS and magnetic relaxation measurements by R.R. Zairov, N.A. Shamsutdinova and A.R. Mustafina. I.R. Nizameev and

A. Vomiero were responsible for TEM studies. A.B. Moreira and G.R. Safina - cell viability of nanoparticles towards PC12 cells, confocal microscopy imaging. S.N. Sudakova, S.N. Podyachev - calix[4]arene tetra-diketone ligand synthesis. A.N. Fattakhova – cell toxicity of nanoparticles towards human blood lymphocytes and platelets aggregation studies. A.T. Gubaidullin performed XRD experiments. I. Lundstrom and all other authors reviewed the manuscript.

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