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Supplementary Information

Biocompatible ligands modulate nanozyme activity of CeO₂ nanoparticles

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Outline

S1 – Experimental dependence of the change in the parameter S on the concentration of SOD.

S2 – Fluorescence spectra of $CeO_2(a)\gamma$ -IgG NPs.

S3 – Experimental dependencies of the change in the parameter S on the concentration of CeO₂ sols.

S4 – Experimental dependence of the change in the parameter S on the concentration bare γ -IgG. S5 – Chemiluminograms for CeO₂@ γ -IgG NPs in a system with alkylperoxyl radicals and luminol.



Figure S1. (a) Chemiluminograms of xanthine (20 μ M) oxidation with xanthine oxidase (4.4 mU/mL) in the presence of lucigenin (20 μ M) upon the addition of various amounts of superoxide dismutase (SOD); (b) the corresponding light sums plotted against SOD concentration.



Figure S2. Fluorescence spectra ($\lambda_{ex} = 280 \text{ nm}$) of γ -IgG (6.5 μ M) before and after loading with CeO₂ nanoparticles.



Figure S3. Dependencies of light sums (*S*) on the concentration of CeO₂ modified with ammonium citrate (**a**), maltodextrin (**b**), dextran (**c**), phosphatidylcholine (**d**), γ -IgG (**e**).



Figure S4. Chemiluminescence light sum measured upon the addition of xanthine oxidase to the phosphate buffer solution containing xanthine, lucigenin and bare γ -IgG with various concentrations.



Figure S5. Chemiluminograms of bare CeO₂ sol (90 μ M), CeO₂ sol modified with γ -IgG (90 μ M) and an individual γ -IgG solution (0.3 μ M) after addition to a system containing luminol and alkyl peroxyl radicals formed as a result of AAPH thermolysis.