Supplementary file

Combining photothermal therapy and immunotherapy against melanoma by polydopaminecoated Al₂O₃ nanoparticles

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Figure S1. Characterization of pD-Al₂O₃ nanoparticles. (A) Average sizes and zeta potentials of pD-Al₂O₃ nanoparticles at various concentrations. (B) Photograph of pD-Al₂O₃ nanoparticles synthesized with the indicated concentrations of dopamine. (C) Average sizes, PDI and zeta potentials of pD-Al₂O₃ nanoparticles at 1,000 μ g/mL before or after irradiation at 1.18 W/cm² for 300 s.

A					
		Al ₂ O ₃ nanoparticles		pD-Al ₂ O ₃ nanoparticles	
	Chemical element	Weight percent %	Atomic percent %	Weight percent %	Atomic percent %
	AI	52.44	37.92	23.87	13.68
	0	37.47	45.70	36.09	34.87
	С	10.09	16.38	39.53	50.88
	Ν	-	-	0.52	0.57



Figure S2. Interactions between Al₂O₃ nanoparticles and polydopamine. (A) We analyzed the weight percent and atomic percent (%) of four chemical elements (Al, O, C, N) in our Al₂O₃ and pD-Al₂O₃ nanoparticles by EDX mapping. Spectrogram for Al, O, C and N in Al₂O₃ (B) and pD-Al₂O₃ (C) nanoparticles. Element mapping for Al (red), O (green), C (yellow) and N (blue) in Al₂O₃ (D) and pD-Al₂O₃ (E) nanoparticles.



Figure S3. The ratio of IgG1/IgG2a *in vivo*. We calculated the IgG1/IgG2a ratio as an indicator of Th1/Th2 immune responses.



Figure S4. Comparison of anti-tumor effects and immune responses induced by $pD-Al_2O_3$ or $pD-Fe_3O_4$ nanoparticles *in vivo*. Animals were exposed to the combination therapy of PTT followed by immunotherapy in the presence of CpG. (A) Tumor growth (8 mice per group). (B) Survival of mice (8 mice per group). (C) Proliferation rates of lymphocytes and splenocytes from mice treated as indicated. (D) Levels of IL-4 in the supernatants of cultures of lymphocytes and splenocytes cultures, based on ELISA.



Figure S5. Toxicity of pD-Al₂O₃ nanoparticles *in vivo*. Mouse body weight was measured after the indicated treatments (8 mice per group).