Reduction of Nanoparticle Avidity Enhances the Selectivity of Vascular Targeting and PET Detection of Pulmonary Inflammation

Blaine J. Zern^{1,2}, Ann-Marie Chacko^{2,3}, Jin Liu⁶, Colin F. Greineder^{1,2,4}, Eric R. Blankemeyer³, Ravi Radhakrishnan⁵, Vladimir Muzykantov^{1,2}*

¹Department of Pharmacology, ²Center for Targeted Therapeutics and Translational Nanomedicine, Institute for Translational Medicine and Therapeutics, ³Department of Radiology, ⁴Department of Emergency Medicine, Perelman School of Medicine and ⁵Department of Bioengineering University of Pennsylvania, Philadelphia, PA 19104, ⁶School of Mechanical and Materials Engineering, Washington State University, Pullman, WA 99164

*Address correspondence to: muzykant@mail.med.upenn.edu

Supporting Information:



Fig. S1. Antibody coverage analysis of NPs. [¹²⁵I]IgG adsorbed onto NPs was calculated as the number of IgG molecules adsorbed per NP as a function of the number of IgG molecules added to NPs. Full coating (~200 Ab/NP, highlighted in red) denotes theoretical saturating coverage.

TABLE ST. CHARACLERIZATION OF AIGAMINF	TABLE S1.	Characterization	of aICAM-NP ^a
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	D _{avg} (nm)	Polydispersity (PD)
IgG NP	187 ± 8	0.17
5 Ab/NP	183 ± 6	0.11
50 Ab/NP	186 ± 7	0.12
100 Ab/NP	192 ± 7	0.18
200 Ab/NP	198 ± 9	0.19

^aAverage hydrodynamic radius, D_{avg} and PD for anti-ICAM-1/NP were determined by DLS

50 Ab/NP



Fig. S2. Coronal sections of real-time *in vivo* CT and PET images acquired after administration of ICAM-1-targeted [¹²⁴I]-NP (50 Ab/NP) in naïve and LPS-challenged mice. Each animal is a 1 h summed PET image overlaid with an anatomical CT scan for a total of n=4 for each group.



Fig. S3. Coronal sections of real-time *in vivo* CT and PET images acquired after administration of ICAM-1-targeted [¹²⁴I]-NP (200 Ab/NP) in naïve and LPS-challenged mice. Each animal is a 1 h summed PET image overlaid with an anatomical CT scan for a total of n=4 for each group.

IgG/NP-LPS



Fig. S4. Coronal sections of real-time *in vivo* CT and PET images acquired after administration of IgG-control [¹²⁴I]-NP (200 Ab/NP) in LPS-challenged mice. Each animal is a 1 h summed PET image overlaid with an anatomical CT scan for a total of n=4 for each group.