Supporting information

Targeted Delivery of Soluble Guanylate Cyclase (sGC) Activator Cinaciguat to Renal Mesangial Cells via Virus-Mimetic Nanoparticles Potentiates Anti-Fibrotic Effects by cGMP-Mediated Suppression of the TGF-β Pathway

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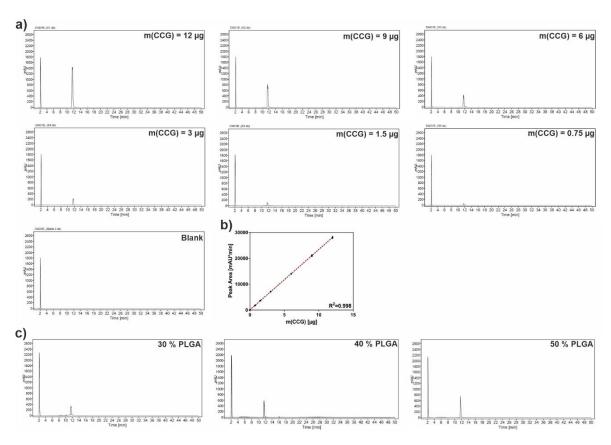


Figure S1. HPLC chromatograms. **a)** Representative HPLC chromatograms for the calibration of free Cinaciguat. **b)** HPLC calibration. Results represent mean \pm SD (n = 3). **c)** Representative chromatograms for NP species carrying 30-50% PLGA and an initial CCG addition of 10 μ g per 4 mg NP batch.

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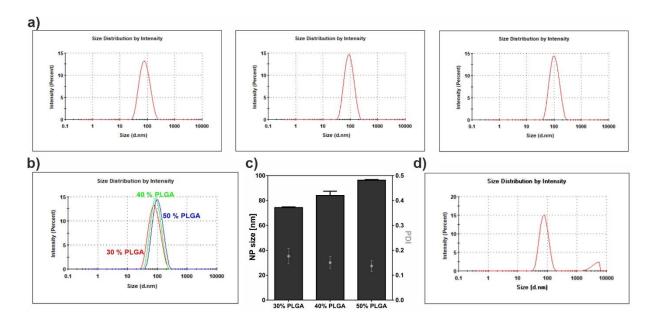
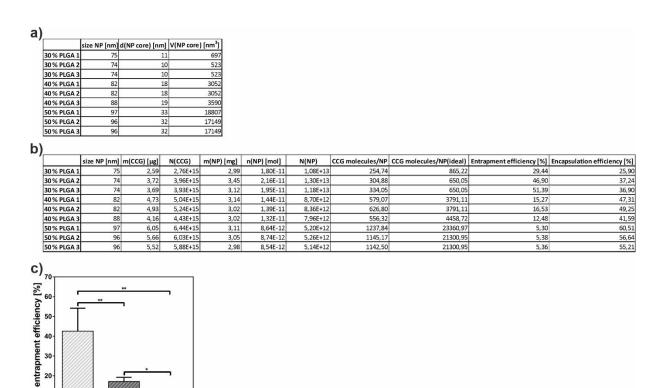
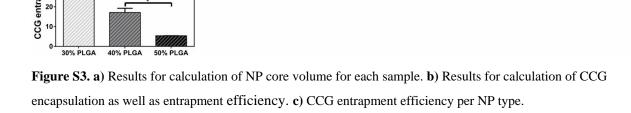


Figure S2. a) DLS size distribution of NP species containing 30/40/50% PLGA (from left to right). b) Merged DLS graphs. c) Size and polydispersity index (PDI) data for different NP species. d) Representative DLS graph of 40% PLGA NPs and an addition of 17.5 µg per 4 mg NP batch, showing a strong tendency for larger aggregates (see right peak).





Results represent mean \pm SD (n = 3). *P<0.05 **P<0.01.

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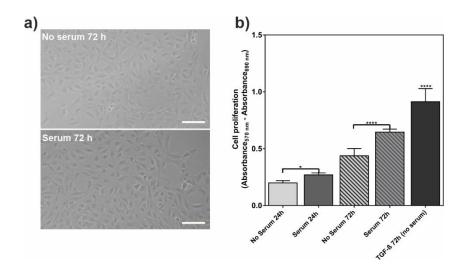


Figure S4. a) Brightfield images of rMCs cultured for 72 h during the cell proliferation assay either without (top) or with (bottom) addition of 10 % FBS. (Scale bar 100 μm). **b**) MTT assay results of rMC proliferation after 24 h and 72 h with and without addition of 10 % FBS. (Note: The results for samples "No serum 72 h" and "TGF-β 72 h (no serum)" represent the blank-normalized samples "Blank cells" and "TGF-β" in Figure 7a of the main manuscript.) Results represent mean \pm SD (n = 3). *P<0.05 ****P<0.0001.