Supplementary Materials

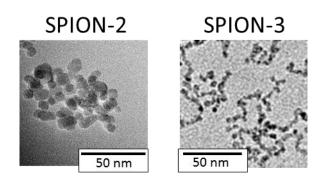
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Manuscript ID: molecules-520234

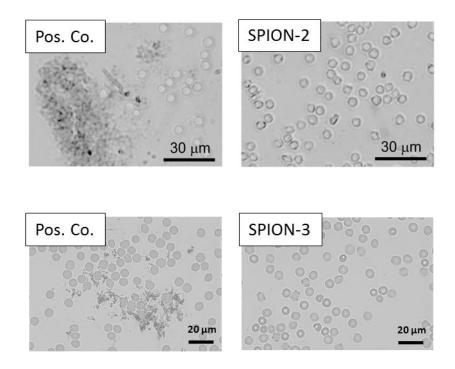
Supplementary Table 1. Physicochemical characterisation of the tested SPIONs.

	SPION-2	SPION-3
Core size [nm]	8.9 ± 2.2	4.3 ± 0.9
Hydrodynamic diameter [nm]	71	74
ζ-potential [mV]	-18.1	-3.8
Coating	LA/BSA	crosslinked dextran T-40

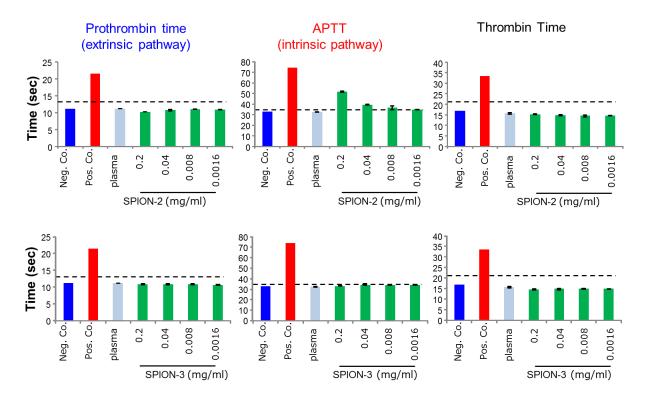
The hydrodynamic size (Z-average size) and ζ-potential of the nanoparticles were determined with a Zetasizer Nano ZS (Malvern). BSA, bovine serum albumin; LA, lauric acid; SPIONs, superparamagnetic iron oxide nanoparticles.



Supplementary Figure 1. Transmission electron microscopy (TEM) images of the tested SPIONs [1,2].



Supplementary Figure 2. Blood stability of the tested SPIONs. Blood stability was investigated in EDTA-anticoagulated sheep (SPION-2 [3]) or rabbit (SPION-3 [2]) whole blood. Blood sample was mixed with SPIONs to an iron concentration of 1 mg/mL. Lauric acid-coated SPIONs served as positive control. Microscopic images were taken after 45 min incubation.



Supplementary Figure 3. Effects of SPIONs on plasma coagulation. Platelet poor human plasma pooled from 3 donors was treated with SPIONs for 30 minutes. Afterwards, the respective coagulation activation reagent was added to each sample (Neoplastine for prothrombin time, CaCl₂ for activated partial thromboplastin time (APTT), or thrombin for thrombin time) and their respective coagulation time was measured. Means ± SD of replicate samples are shown. Upper panel: SPION-2 (data courtesy of Dr. Christina Janko, SEON, University Hospital Erlangen); Lower panel: SPION-3 [2].

References

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