Supporting information for

## Manganese-impregnated mesoporous silica nanoparticles for signal enhancement in MRI cell labelling studies

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Figure S1. TEM images of Mn-M48SNs after (a) thermal reduction treatment and after suspension in  $H_2O$  for (b) 24 h or (c) 96 h and (d) corresponding particle size distributions.



**Figure S2.** TEM image of Mn-M48SNs after thermal reduction treatment (corresponding to the product in Figure S1.a).



**Figure S3.** XRD patterns of Mn-M48SNs after (a) thermal reduction treatment and after suspension in  $H_2O$  for (b) 3 h, (c) 24 h, or (d) 96 h.



**Figure S4.**  $H_2$ -TPR profiles of Mn-M48SNs after (a) thermal reduction treatment and after suspension in  $H_2O$  for (b) 3 h, (c) 24 h, or (d) 96 h.



Figure S5. Magnetometric profiles of Mn-M48SNs at different temperatures.



Figure S6. Colloidal stability assay for Mn-M48SN particles suspended in DMEM.



**Figure S7.**  $T_1$  and  $T_2$  relaxation times (60 MHz, 37 °C) of Mn-M48SNs suspended for prolonged times in (a) H<sub>2</sub>O and (b) acetate buffer (20 mM, pH 5). Before each measurement, suspensions were centrifuged at 5000 g and materials were suspended in fresh media.



**Figure S8.** NMRD profiles of Mn-M48SNs suspended in acetate buffer (20 mM, pH 5) performed at 37°C.



Figure S9. P388 cell proliferation assay, after 4 hours of incubation with 7 % (v/v) Mn-M48SNs.



**Figure S10.**  $T_1$ -w. MR images of P388 cells incubated 4 h with a 7 % (v/v) stock solution of M48SNs obtained with different echo delay times and repetition times. No difference of signal was found between cells incubated with 7 % (v/v) MSNs, water and gelatin.