## **Supporting Information**

## Bioevaluation of magnetic mesoporous silica rods: cytotoxicity, cell uptake and biodistribution in zebrafish and rodents.

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Fig. S1. TEM images of magnetic mesoporous silica rods. Long rods –  $Fe_2O_3@LR$  (a, b) and short rods –  $Fe_2O_3@SR$  (c, d).



Fig. S2. Magnetic hysteresis loops measured at 10 K. a)  $Fe_2O_3@SR$ , b) pellet of ZFL cells incubated with  $Fe_2O_3@SR$  (100  $\mu$ g/ml).



Table S1. Uptake of MSRs by ZFL cells studied by SQUID magnetometry. Mean values of the remanent magnetic
moment at 10 K were measured for pellets of ZFL cells treated with different concentrations of Fe <sub>2</sub> O <sub>3</sub> @SR and
$Fe_2O_3(a)SR-NH_2$ (n=4).

MSR sample	[MSR] (µg/ml)	m <sub>r</sub> (10 <sup>.6</sup> emu)	m <sub>r</sub> /cell (10 <sup>-11</sup> emu/cell)	MSR uptake (pg/cell)
Fe₂O₃@SR	20	$2.2 \pm 0.3$	1.8 ± 0.3	5 ± 1
	50	24 ± 4	19.± 3	58 ± 9
	100	42 ± 12	34 ± 9	102 ± 28
Fe <sub>2</sub> O <sub>3</sub> @SR- NH <sub>2</sub>	20	6 ± 6	5 ± 5	16 ± 17
	50	13 ± 4	11 ± 3	35 ± 11
	100	39 ± 10	31 ± 8	105 ± 26

**Fig. S3.** Zebrafish lifecycle. a) Zebrafish. b) Zebrafish eggs. c) Zebrafish eggs hatching. d) Zebrafish larvae. Scale bars: 1 cm. Zebrafish larvae survival after 96 h of exposure to fluorescamine-functionalized MSRs.



**Fig. S4.** *In vivo* fluorescence images of mice injected with Fe<sub>2</sub>O<sub>3</sub>@SR-Cy5 at (a) 30 min, (b) 60 min, c) *in vivo* fluorescence values measured in the abdomen at 30 min and 60 min post-injection.



**Fig. S5.**  $T_2$  relaxation time maps of the abdominal area of (a, b, c) animal 1 and (d, e, f) animal 2 recorded at various times: t = 0 (pre-injection), t = 20-25 min and t = 60 min post-injection of Fe<sub>2</sub>O<sub>3</sub>@SR.  $T_2$  values in the liver of the two studied rats, before and 60 min after MSR injection (g). Respiration and body temperature of the two studied rats were monitored after intravenous injection of Fe<sub>2</sub>O<sub>3</sub>@SR (h).

