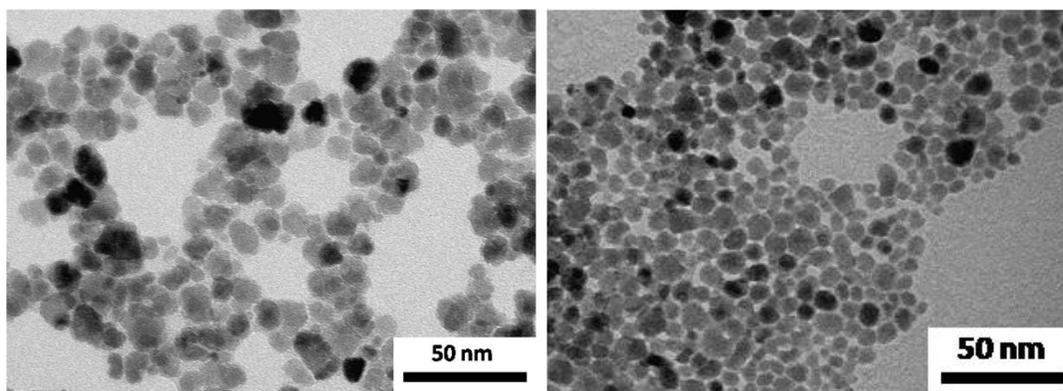
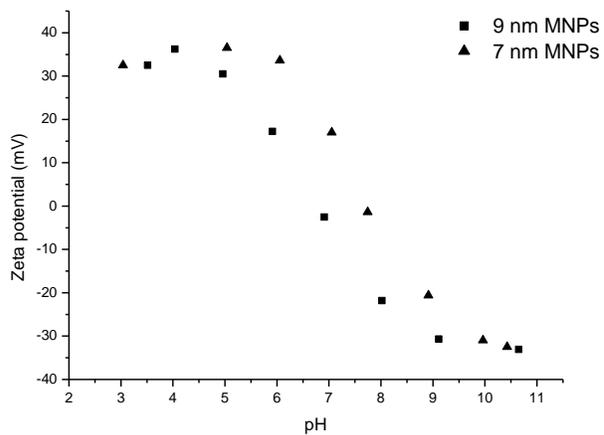


# Supporting Information to Surface Decoration of Catanionic Vesicles by Superparamagnetic Iron Oxide Nanoparticles: a Model System for Triggered Release with Moderate Temperature Conditions

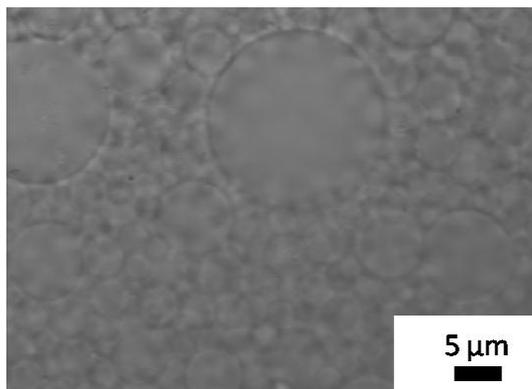
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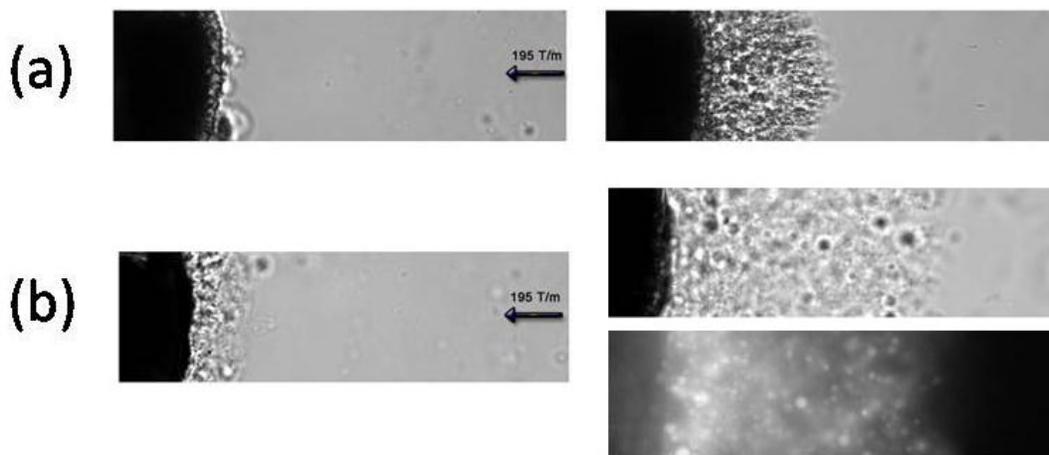
**Fig. S11.** TEM micrographs of 9 nm superparamagnetic nanoparticles (left) and 7 nm superparamagnetic nanoparticles (right).



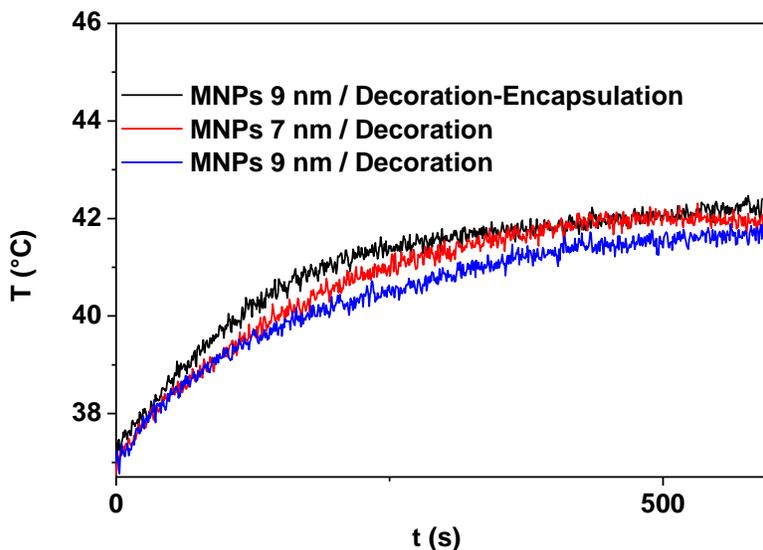
**Fig. S12.** Determination of the isoelectric point of  $\gamma$ - $\text{Fe}_2\text{O}_3$  9 nm and 7 nm superparamagnetic nanoparticles.



**Fig. S13.** Microscopy picture in transmission mode of catanionic vesicles with their surface covered with MNPs.



**Fig. S14.** Magnetophoresis experiments under a 195 T/m magnetic field gradient on (a) decorated vesicles encapsulating 9 nm maghemite nanoparticles (optical microscopy) and (b) decorated vesicles encapsulating Rhodamine 6G (optical microscopy and fluorescence microscopy)



**Fig. SI5.** Evolution of the temperature of each sample as a function of time. For the experiment, the iron concentration was adjusted to 25mM. Temperatures were recorded every 0.7 s. Measurements were performed applying a 520 kHz oscillating magnetic field with an amplitude of 28 kA.m<sup>-1</sup>.

#### Appendix SI1. Calculation of the concentration of MNPs for the surface saturation of vesicles

The concentration of surfactants  $C_S$  is given by

$$C_S = \left( \frac{m_{MA}}{M_{MA}} + \frac{m_{CTA}}{M_{CTA}} \right) \times \frac{1}{V} \quad \text{Equation (1)}$$

where  $m_{MA}$  and  $m_{CTA}$  are the mass of myristic acid and CTACl introduced, respectively.

From this concentration, the total surface area of vesicles per volume unit of sample is given by:

$$S_{vesicles} = C_S \times N_A \times a_0 \quad \text{Equation (2)}$$

where  $N_A$  is the Avogadro number and  $a_0$  the surface of one surfactant in the bilayer (20 Å<sup>2</sup> [14])

The total surface area of MNPs per volume unit of sample is given by:

$$S_{MNP_s} = C_{Fe} \times N_A \times \frac{a_{MNP}}{n_{Fe}} \quad \text{Equation (3)}$$

with  $C_{Fe}$  the concentration of iron,  $a_{MNP}$  the surface of a single nanoparticle and  $n_{Fe}$  the amount of iron in one nanoparticle.

$n_{Fe}$  is given by:

$$n_{Fe} = 2 \times N_A \times \frac{d \times v_{MNP}}{M_{Fe_2O_3}} \quad \text{Equation (4)}$$

where  $d$  is the density of maghemite ( $5240 \text{ kg.m}^{-3}$ ) and  $v_{MNP}$  the volume of one nanoparticle. Substitution in Equation 3 leads to:

$$S_{MNP_s} = \frac{1}{2} \times C_{Fe} \times \frac{M_{Fe_2O_3}}{d_{Fe_2O_3}} \times \frac{a_{MNP}}{v_{MNP}} \quad \text{Equation (5)}$$

The condition to completely cover all the vesicles with nanoparticles is that the total surface of MNPs ( $S_{MNP_s}$ ) is superior to the total surface of vesicles ( $S_{vesicles}$ ). Using equations 2 and 5, this leads to:

$$C_{Fe} > C_S \times 2 \times N_A \times \frac{a_0}{a_{MNP}} \times \frac{d_{Fe_2O_3} \times v_{MNP}}{M_{Fe_2O_3}} \quad \text{Equation (6)}$$

**Table SI 1. SAR values and  $\Delta T$  of the samples.**

	[Fe] mM	$\Delta T$	SAR (W/g)
MNPs 9 nm decoration / encapsulation	25	5.6	116
MNPs 7 nm decoration	23	4.8	174
MNPs 9 nm decoration	27	4.8	154

The specific absorption rate (SAR) was calculated from the initial slope of the curve  $\frac{dT}{dt}$ , using Equation (7):

$$SAR = C_p \times \frac{m_{sample}}{m_{Fe}} \times \frac{dT}{dt} \quad \text{Equation (7)}$$

where  $C_p = 4.18 \text{ J.g}^{-1}.\text{K}^{-1}$  is the heat capacity of water,  $m_{sample}$  is the mass of the sample and  $m_{Fe}$  is the mass of iron in the samples. For all samples, the SAR values found were in the range 115-180 W/g.