Supporting Information

Bacterial Magnetosomes – Nature's powerful contribution to MPI tracer research

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Characterization methods

TEM images and SAED patterns were obtained using a Zeiss EM 912 Ω operating at an acceleration voltage of 120 kV. Dynamic light scattering was performed using a Submicron Particle Sizer Model 370 (Nicomp Particle Sizing Systems; USA). The magnetic particle spectra were measured applying a drive field with an amplitude of 10 mT and a frequency of $f_0 = 25.25$ kHz with a commercial MPS system (Bruker BioSpin GmbH, Germany). Static M(H)-curves were measured with a commercial MPMS XL system (Quantum Design Inc., USA). MRX measurements were performed using a homebuilt device. MRX is a measurement technique which probes the dynamics of the magnetic moments of the MNP in the sample are aligned by an external magnetic field of about 2 kA/m. After switching off the field within 420 µs, the decay of the magnetisation is measured by a low-Tc SQUID system at T = 295 K for 0.5 s.⁴

Sample	$\sigma_{ m c} nm$	$rac{\sigma_{ m h}}{ m nm}$	$\sigma_{ m m}$ nm
LMU1	5.4	51.4	0.181 ± 0.002
LMU2	8.1	45.5	0.200 ± 0.007
LMU3	6.6	40.3	0.326 ± 0.02

Table SI1. Standard deviation σ_c , σ_h and σ_m of crystal diameter, intensity-weighted hydrodynamic diameter and magnetic diameter (from M(H) data), respectively.



Figure SI1. Distributions of the effective magnetic diameters of all magnetosome samples and Resovist[®] calculated from their static magnetization curves normalized to the volume fraction of magnetite using the Moment Superposition Model assuming a bimodal size distribution.

Table SI2. $d_{spacing}$ values obtained from the SAED pattern of magnetosomes on TEM grid (corresponding to figure 1d in the manuscript). The calculated $d_{spacing}$ values (ref) for Fe₃O₄ are reported for comparison.

hkl	d (ref) ^[a]	d (measured)		
111	4.85	4.84		
220	2.97	2.97		
311	2.53	2.53		
400	2.10	2.09		
422	1.71	1.70		
511	1.61	1.61		
440	1.48	1.47		
[a] ref. ICDD-PDF4+ 04-001-7822				

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