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Scherrer equation for spherically-shaped particles:

(S1)

 $D(nm) = \frac{0.89 \cdot \lambda}{\Delta 2\theta \cdot \cos{(\theta)}}$ 

Where  $\lambda$  - X-ray source wavelength. nm

 $\Delta 2\theta$  - peak width at half its height. °

## Table S1

Parameters of the Mössbauer spectra for the samples.

Temperature . K		77.7±0.3						296±3				D	ST E O	Compoun		
Sample	№	*δ	ε (Δ=2ε)	Γ <sub>exp</sub>	H <sub>eff</sub>	S	α	*δ	ε (Δ=2ε)	Γ <sub>exp</sub>	Heff	S	α	1	δ in Fe <sub>3</sub> O <sub>4</sub>	d d
		mm/s		kOe	%			mm/s		kOe	%	nm	1			
MOF**	1	0.52±0.0 1	(0.73±0.06	0.54±0.0 1		80±20		0.41±0.0 1	(0.7±0.1)	0.62±0.0 1		80±30				
	2	0.53±0.0 1	(0.44±0.02)	0.30±0.0 7		20±20		0.41±0.0 1	(0.45±0.05 )	0.4±0.2		20±30				
Fe <sub>3</sub> O <sub>4</sub> - MOF	1	0.51±0.0 1	0.07±0.01	0.23±0.0 4	529.4±0. 7	28±8	8	0.27±0.0 1	0.01±0.01	0.47±0.0 1	485.3±0. 3	63±7				
	2	0.52±0.0 2	-0.11±0.03	0.39±0.0 2	525.5±0. 6	34±6	7.3±0. 1	0.47±0.0 2	-0.04±0.01	0.41±0.0 4	484.4±0. 6	24±7	3.66±0.0 3	2.01±0.0 1	0.317±0.00 8	Fe <sub>2.683</sub> O <sub>4</sub>
	3	0.30±0.0 1	0.03±0.01	0.38±0.0 2	522.0±0. 8	37±5		0.28±0.0 2	-0.03±0.02	0.76±0.0 8	172±2	5.9±0. 3				
	4	0.40±0.0 4	(0.65±0.01	0.57±0.0 1		1.3±0. 1		0.33±0.0 1	(0.65±0.01 )	0.56±0.0 1		6.8±0. 1				
Fe <sub>3</sub> O <sub>4</sub> - AA- MOF	1	0.61±0.0 3	0.01±0.01	0.32±0.0 6	518±2	15±5	2.0±0. 5	0±0. 0.33±0.0	3±0.0 0 02 0 02	0.8±0.1	470±10	52±1	0.5±0.2	3.5±0.4	0.290±0.00 8	Fe <sub>2.710</sub> O <sub>4</sub>
	2	0.37±0.0 2	-0.01±0.01	0.39±0.0 4	513±1	31±6		5 4	4							
	3	0.49±0.0 1	(0.91±0.06	0.57±0.0 2		42±7		0.37±0.0 1	(1.03±0.08	0.53±0.0 4		25±7				
	4	0.51±0.0 1	(0.55±0.03	0.35±0.0 6		11±7		0.39±0.0 1	(0.62±0.04	0.43±0.0 3		23±7				

 $^*\delta$  — the isomeric shift.  $\Delta$  — the quadrupole splitting.  $\Gamma_{exp}$  — the linewidth.  $H_{eff}$  — the hyperfine magnetic field. S — the relative area of the subspectrum.  $\alpha$  – the quotient of particle anisotropy energy to thermal energy. D - magnetic domain diameter.

\*\* The hyperfine parameters for this sample were determined from the experimental spectrum obtained in a narrow speed range. The spectra in Figure MS1 (a.d) are for illustration purposes only.

<b>N</b>			1
Sampl	MOF	Fe <sub>3</sub> O <sub>4</sub> -MOF	Fe <sub>3</sub> O <sub>4</sub> -AA-
Element			MOF
Iron (Fe)	5.8	18.7	6.4
Oxygen (O)	33.5	76.8	44.2
Carbon (C)	60.7	4.5	49.4

Table S2. Concentrations of elements in the samples. at.%.

Table S3. parameters of spectra deconvolution of MOF.  $Fe_3O_4$ -MOF and  $Fe_3O_4$ -AA-MOF samples.

Sampla	Bond	E aV	FWUM aV	Bond
Sample		$E_b$ . C v		portion. at.%
	$Fe^{2+}(2p_{3/2})$	710.0	2.88	22.25
	$Fe^{2+}(2p_{1/2})$	723.6	2.88	11.13
	$Fe^{3+}(2p_{3/2})$	712.1	4.45	25.10
MOF	$Fe^{3+}(2p_{1/2})$	726.3	4.45	9.94
	$Fe^{2+}$ sat (2p <sub>3/2</sub> )	715.5	4.56	13.51
	$Fe^{2+}$ sat (2p <sub>1/2</sub> )	729.0	4.56	4.13
	C-C	284.6	1.94	71.91

	С-ОН	286.2	1.94	2.46
	С=О	288.5	1.94	22.72
	pi-pi* satellite	291.2	1.94	2.90
	Fe-OOH	530.0	2.19	14.85
	Fe-O	531.2	2.19	21.93
	Fe-OH ads.	531.8	2.19	63.22
	Fe <sup>2+</sup> (2p <sub>3/2</sub> )	709.9	3.22	19.54
	$Fe^{2+}(2p_{1/2})$	723.5	3.21	9.77
	$Fe^{3+}(2p_{3/2})$	711.9	4.15	18.48
	$Fe^{3+}(2p_{1/2})$	725.8	4.15	9.96
	$Fe^{2+}$ sat (2p <sub>3/2</sub> )	715.4	6.25	13.48
	Fe <sup>2+</sup> sat $(2p_{1/2})$	729.0	6.25	5.65
Fe <sub>3</sub> O <sub>4</sub> -MOF	Fe <sup>3+</sup> sat $(2p_{3/2})$	719.9	6.32	16.73
	Fe <sup>3+</sup> sat $(2p_{1/2})$	732.3	6.32	6.39
	C-C	284.6	1.94	71.91
	С-ОН	286.2	1.94	2.46
	C=O	288.5	1.94	22.72
	O-C=O	293.1	2.82	5.84
	Fe-OOH	530.3	2.77	100.00
	$Fe^{2+}(2p_{3/2})$	710.6	3.22	32.98
	$Fe^{2+}(2p_{1/2})$	724.2	3.22	13.07
	$Fe^{3+}(2p_{3/2})$	712.9	4.01	16.52
	$Fe^{3+}(2p_{1/2})$	726.2	4.01	7.78
	$Fe^{2+}$ sat (2p <sub>3/2</sub> )	716.2	8.05	12.69
	$Fe^{2+}$ sat (2p <sub>1/2</sub> )	730.2	4.08	1.8
	$Fe^{3+}$ sat (2p <sub>3/2</sub> )	720.9	8.05	12.1
Fe <sub>3</sub> O <sub>4</sub> -AA-MOF	Fe <sup>3+</sup> sat $(2p_{1/2})$	733.2	4.08	3.07
	C-C	284.6	1.93	65.4
	С-ОН	286.6	1.93	7.47
	C=O	288.6	1.93	22.98
	pi-pi* satellite	291.1	2.97	4.15
	Fe-OOH	530.4	2.13	41.49
	Fe-OH ads.	531.9	2.13	50.97
	-OH (ads. $H_2O$ )	533.5	2.13	7.54



Figure S1. The pH of 3% NaCl solution exposed to nanocomposite of  $Fe_3O_4$ -MOF nanoparticles functionalized by ascorbic acid,  $Fe_3O_4$ -AA-MOF (1), and ascorbic acid (2). The pH of 3% NaCl solution (control) was 5. 0.



**Figure S2.** Bioluminescence intensity.  $I^{rel}$  (green curves), and ROS content.  $ROS^{rel}$  (red curves), in bacterial suspension vs. concentration of MOF (A) in the absence of 1.4-benzoquinone, (B) in the presence of 1.4-benzoquinone ( $E_{C50}^{Bq}=10^{-7}M$ ). ROS content in the control bacterial suspensions was **3**·10<sup>-7</sup>M (A) and **4**·10<sup>-7</sup>M (B)