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Supporting Information

Inert Gas Condensation made bimetallic FeCu nanoparticles - plasmonic response and magnetic ordering

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Figure S1 Schematic view of a) the vacuum deposition system based on IGC technique containing the NPs source (Mantis Deposition Ltd), b) two type of deposited systems: NPs without (top) and with 3 nm thick film of Au (bottom). c) Distribution of the electrical current carried by Cu, Fe and bimetallic FeCu NPs collected at the exit grid of the quadruple mass spectrometer as a function of the diameter of the selected mass of NPs. The frequency of the quadruple mass spectrometer at which the electrical current reaches a maximum (given by dashed vertical line) was chosen as a desire NPs diameter.

Sample	Working	Input	Argon	Insertion	Time	
	preassure	current	flow	length	[min]	
	[mTorr]	[mA]	[sccm]	[mm]	լոոոյ	
Fe	4.7	70	70	10	40	
Fe50Cu50	5.1	100	100	0	40	
Fe25Cu75	5.0	100	100	0	40	
Cu	60.0	110	100	30	40	



Figure S2 HR-TEM image of the iron, cooper and bimetallic a) Fe50Cu50 b) Fe25Cu75, c) Fe NPs with the proposed lattice planes of the detected structural phases together with the electron diffraction pattern.

Table S1 Deposition parameters for studied NPs.



Figure S3 SEM image with marked 10 nanoparticles for which EDS analysis was performed for a) Fe25Cu75 and b) Fe50Cu50 NPs.

Table S2 The atomic% content of Cu and Fe in the studied bimetallic samples of Fe25Cu75, Fe50Cu50 NPs based on the EDS method for 10 selected nanoparticles.

	Fe25Cu75		Fe50Cu50				
No	Cu (atomic %)	Fe (atomic %)	No	Cu (atomic %)	Fe (atomic %)		
1	74.8	25.2	1	55.6	44.4		
2	68.4	31.6	2	50.4	49.6		
3	76.3	23.7	3	46.1	53.9		
4	80.5	19.5	4	49.2	50.8		
5	76.8	23.2	5	56.7	43.3		
6	73.4	26.6	6	58.7	41.3		
7	76.7	23.3	7	45.7	54.3		
8	76.6	23.4	8	41.3	58.7		
9	77.9	22.1	9	56.4	43.6		
10	77.7	22.3	10	56.3	43.7		
Average	75.9±3.2	24.1±3.2	Average	51.6±5.9	48.4±5.9		



Figure S4 PFY XAS at Fe and Cu L absorption edges. Chemical and structural characteristics of FeCu NPs can be deduced by comparison with spectra of reference samples, namely iron(III) oxide and metallic film as well as copper(II) oxide and CuO.

Table S3. Wavenumbers and proposed band assignment for the Raman, FT-IR, SERS and SEIRA spectra of Phe adsorbed onto Cu, Fe25Cu75, and Fe50Cu50 NPs.

Assignment	Wavenumber	[cm ⁻¹]						
	Phe		Cu		Fe25Cu75		Fe50Cu50	
	Raman	FT-IR	SERS	SEIRA	SERS	SEIRA	SERS	SEIRA
v(NH)	3169							
v(NH)		3125		3120				
v_2 , $v(CH)_{ring}$	3065	3060	3061	3061	3059		3061	
v(CH)	3040	3028	3034	3030	3039	3030		3030
$v_{as}(CH)$	2967		2962				2967	
v _{as} (CH)	2927	2918	2925		2938	2918	2923	2922
v _s (CH)	2858		2867		2870		2860	
ρ _{bas} (NH)		1634						
$v_{8a} v(CC)_{ring i. p.}$	1600	1609	1604	1605	1604	1603	1601	1604
$v_{as}(COO); v_{8b} v(CC)_{ring}$	1583	1589	1583	1585	1583	1579	1585	1580
$\nu_{19a} \nu(CC)_{ring}; \rho_{bs}(NH)$		1503		1497		1518		1518
$v_{19b} v(CC)_{ring}, \rho_s(CH_2)$	1445	1448	1439	1455	1451	1456	1438	1457
v _s (COO)	1408	1408		1402		1398		1399
v ₃	1340	1366		1365		1345		
δ(CO)	1306	1325	1318	1322	1324	1314	1312	1313
v _{7a}	1212	1210	1209	1208	1206		1212	1207

v _{9a}	1182	1197	1177	1195			1182	1162
δ (CH)		1143						
v(CN)		1074		1073		1073		1074
v _{18a, i. p.}	1035	1028	1029		1028	1031	1032	1031
v ₁₂	1002		1001		1000		1002	
v(COO)	912							
δ _{o.o.p.(} C-H) _{ring}	850							
pr(CH ₂)	832		822		810		824	
δ(ring)			773		773			
ν ₁₁ , δ _{0.0.p.} (C-H) _{ring}	745		745		746		744	
ρ_r (COO)	682		672		667		668	
v _{6b}	619		619		619		620	
Abbreviations:	v - stretching, δ - deformation, ρb - bending, ρr - rocking, ρs - scissoring, i.p in-plane, o.o.p out-of-							
	plane, s – symmetric, and as – asymmetric vibrations;							