

Supporting Information

Asymmetrically substituted triazenes as poor electron donor ligands in the precursor chemistry of iron(II) for iron-based metallic and intermetallic nanocrystals

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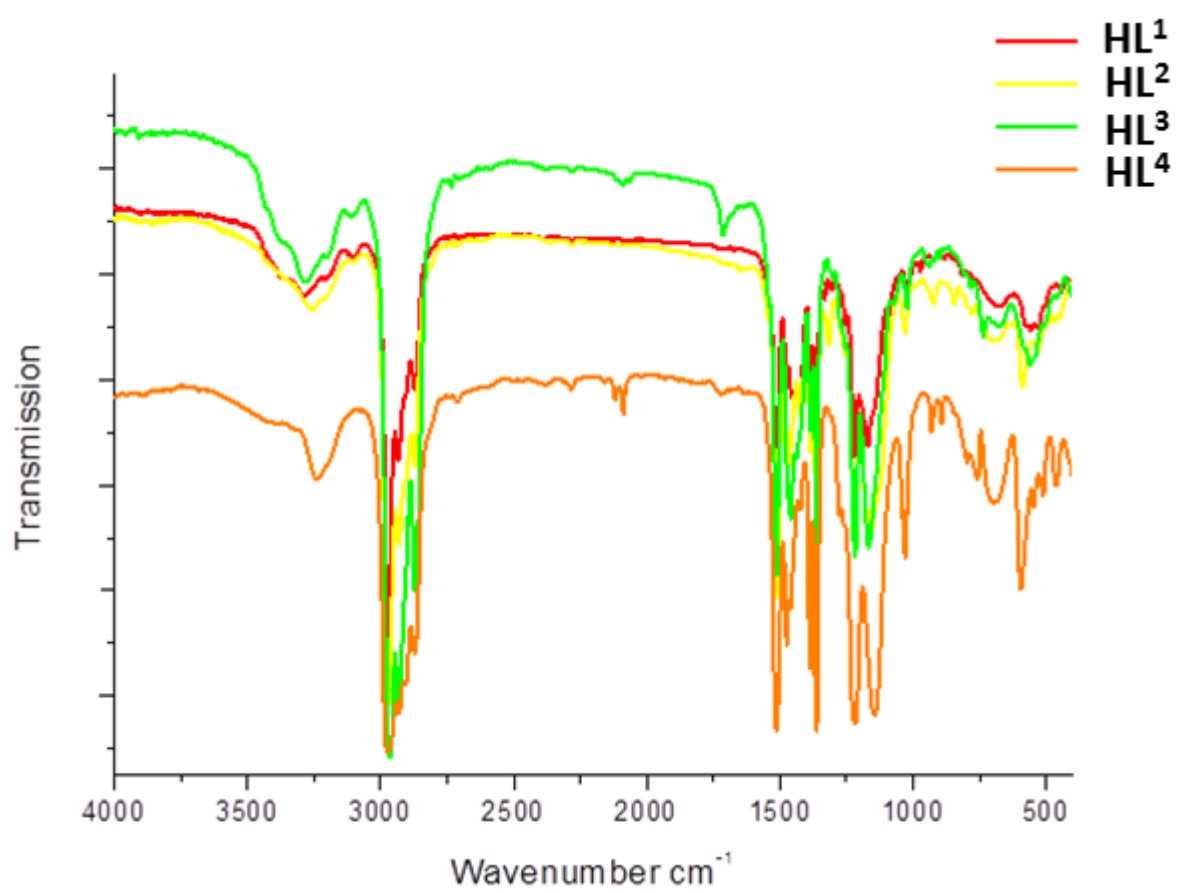


Figure S1: FT-IR spectra of the new ligands **HL**¹-**HL**⁴.

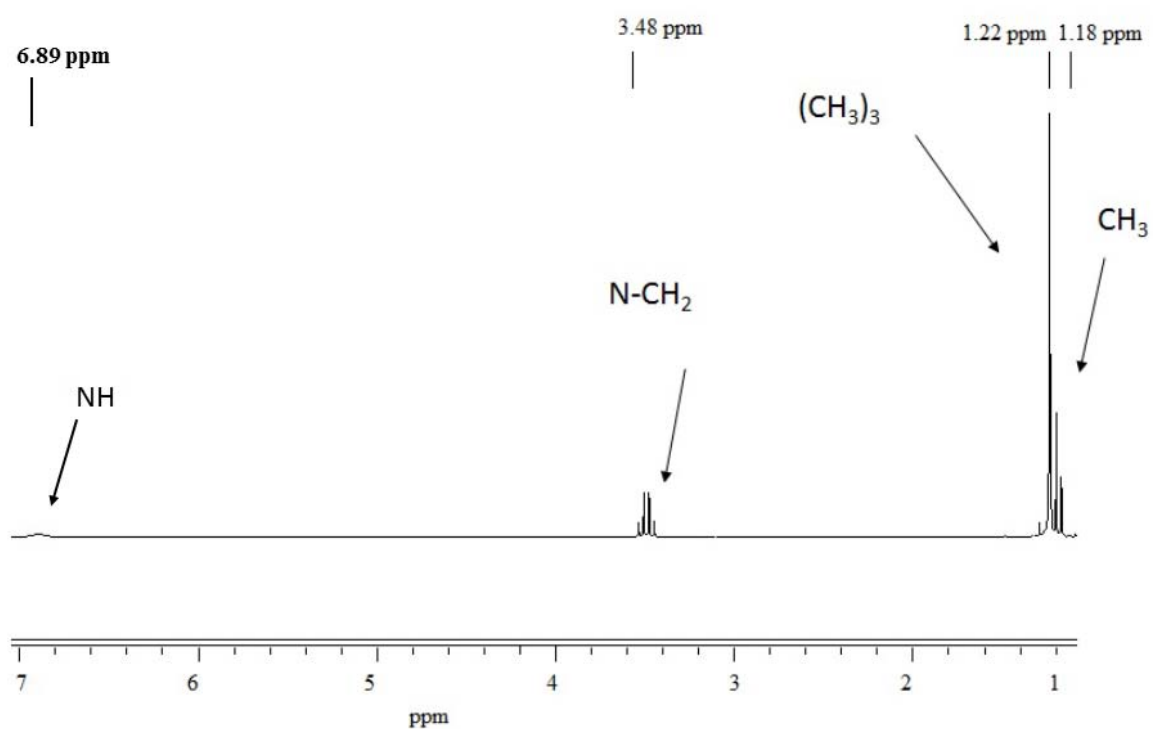


Figure S2: ¹H NMR spectrum of the ligand **HL**¹.

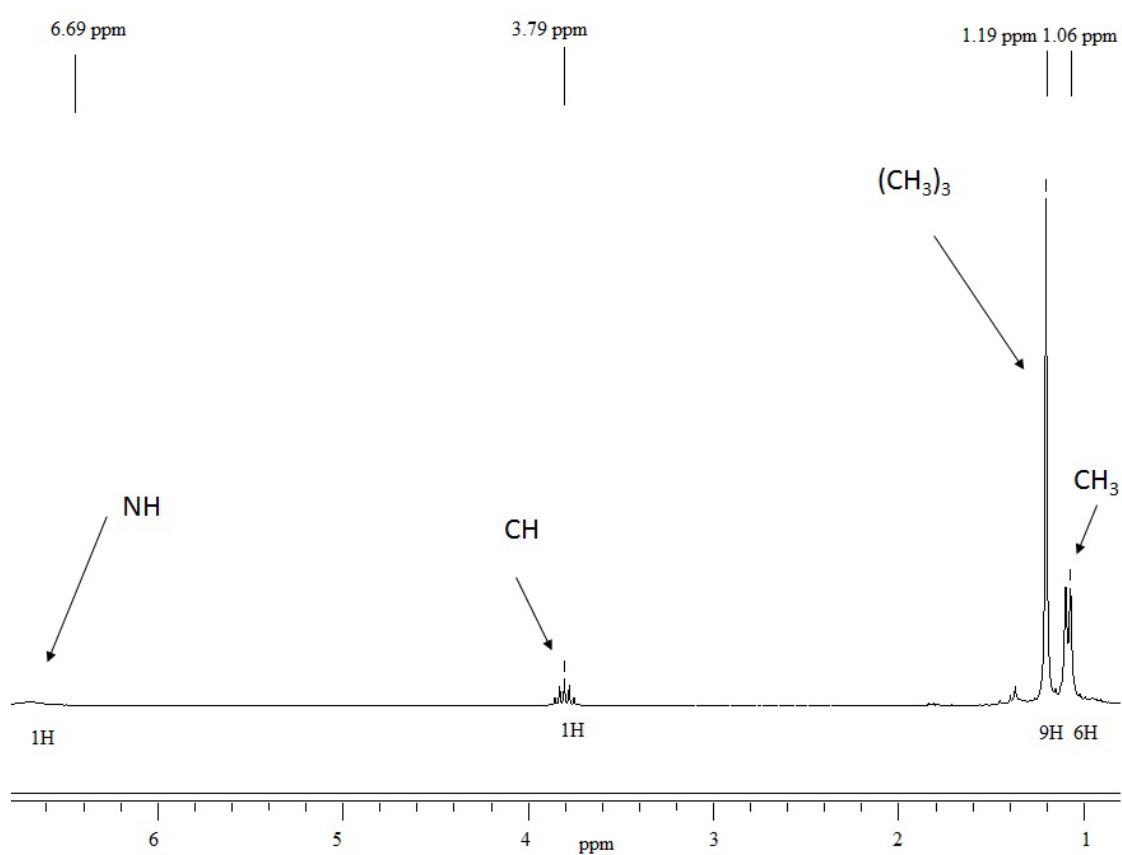


Figure S3: ¹H NMR spectrum of the ligand **HL**².

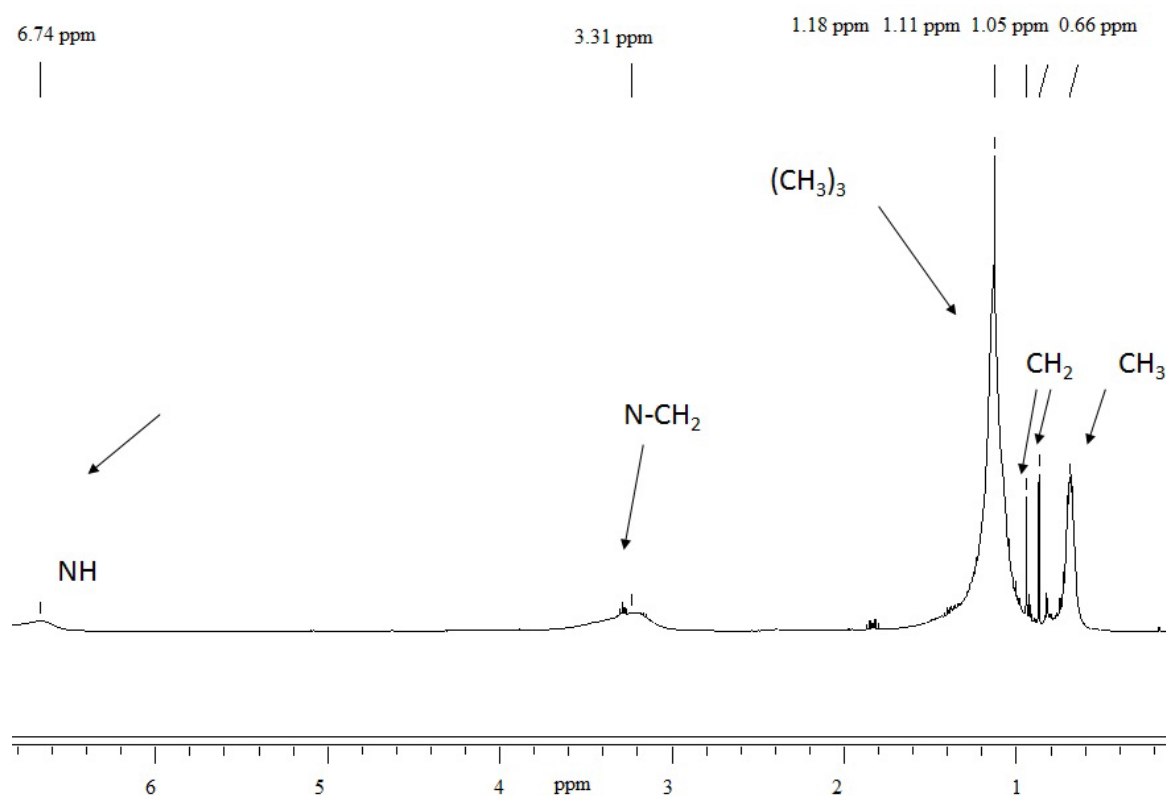


Figure S4: ¹H NMR spectrum of the ligand **HL**³.

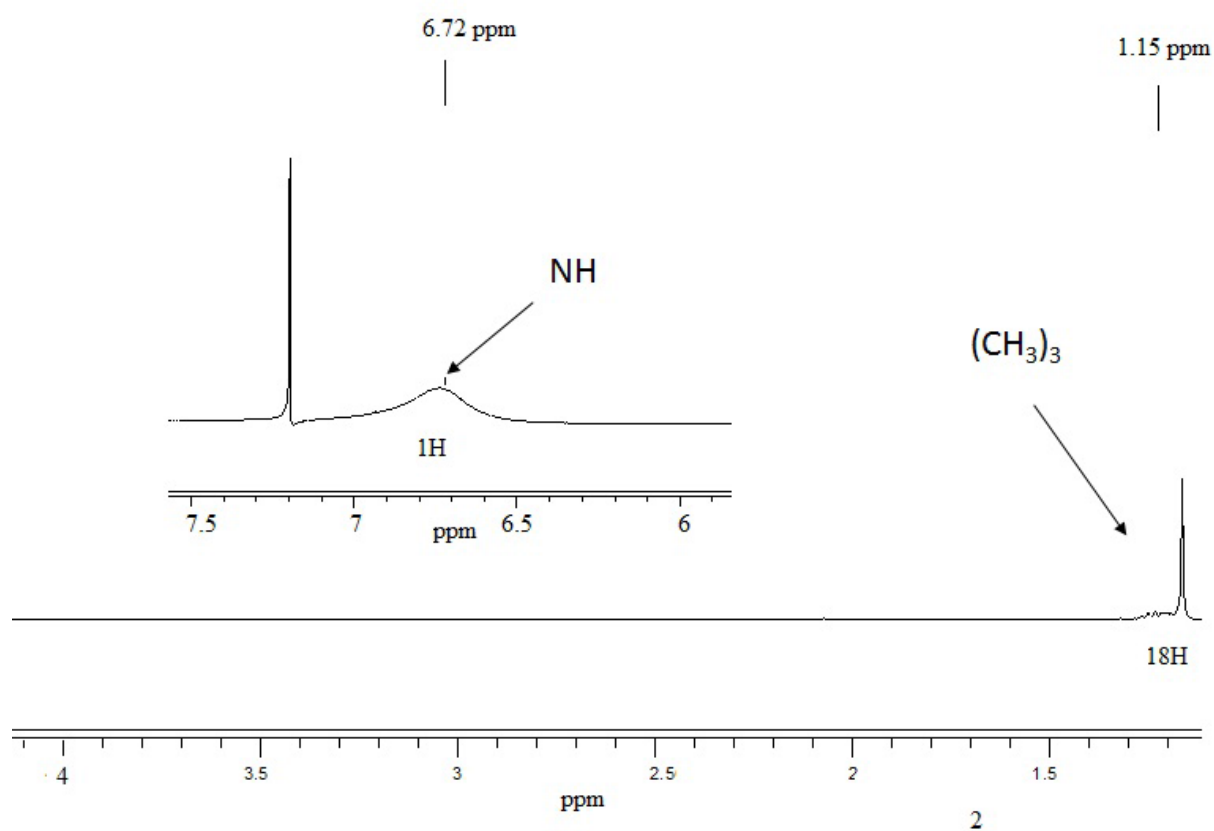


Figure S5: ¹H NMR spectrum of the ligand **HL**⁴.

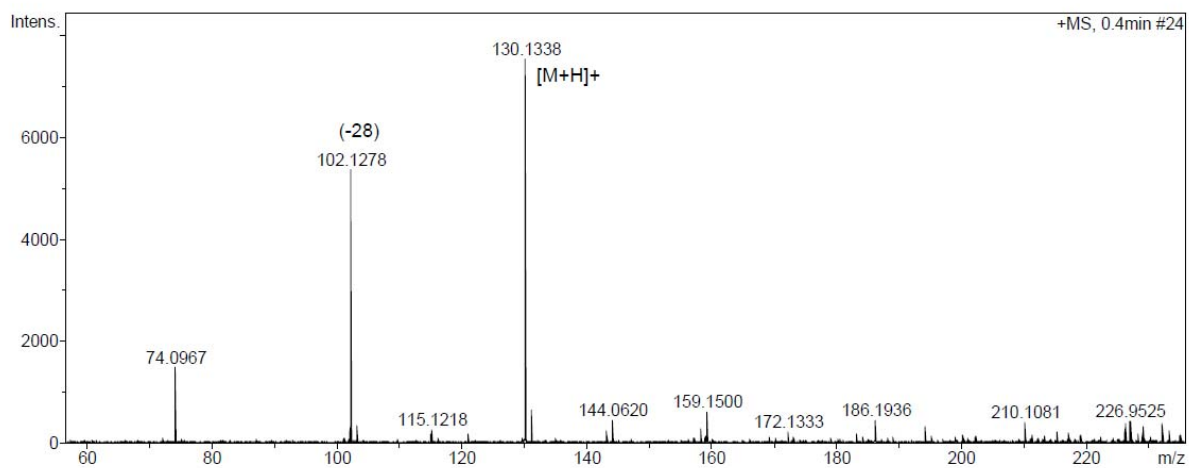


Figure S6: Mass Spectrum of **HL¹**.

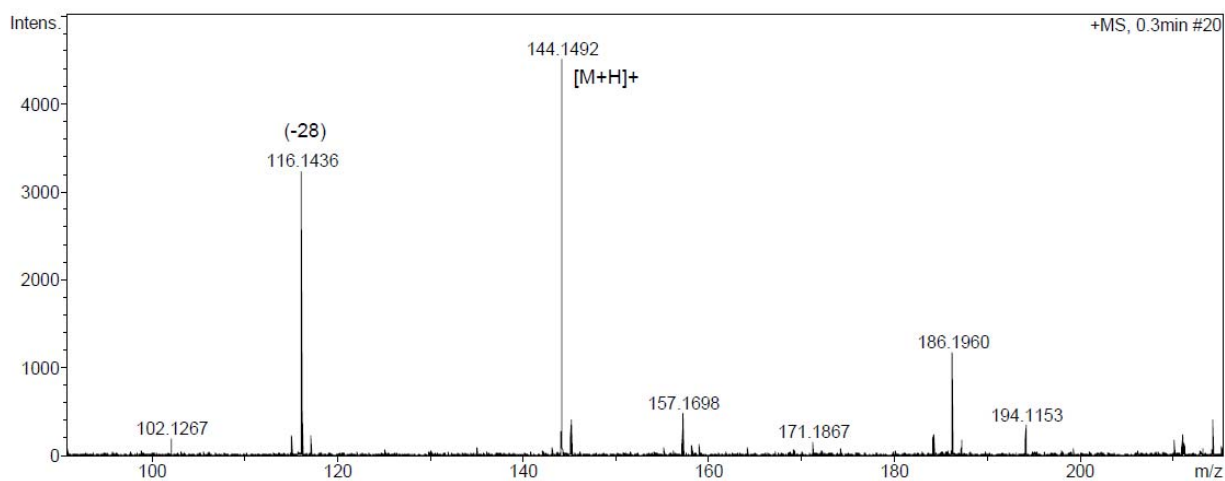


Figure S7: Mass Spectrum of **HL²**.

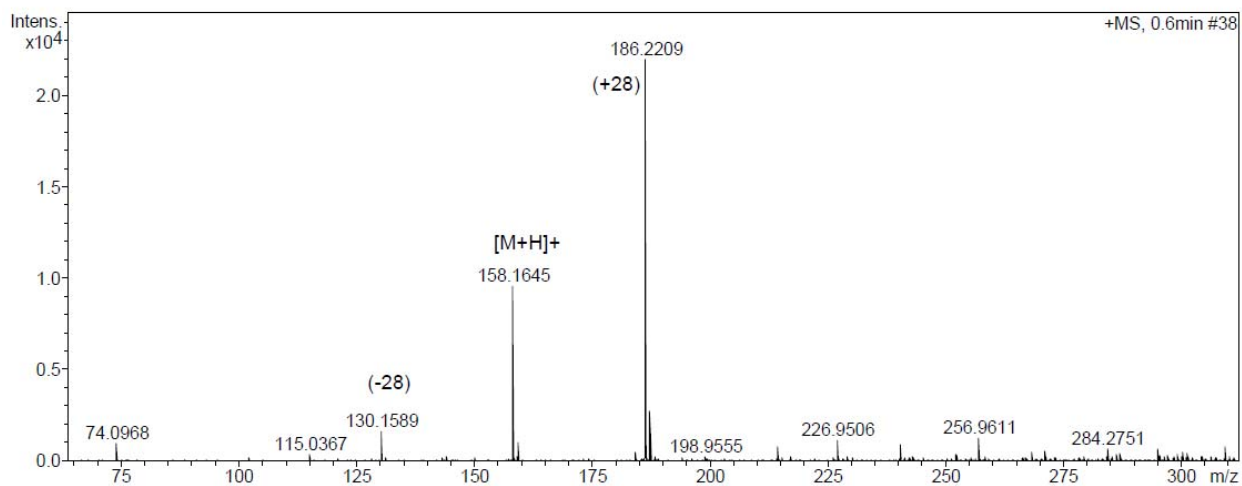


Figure S8: Mass Spectrum of **HL³**.

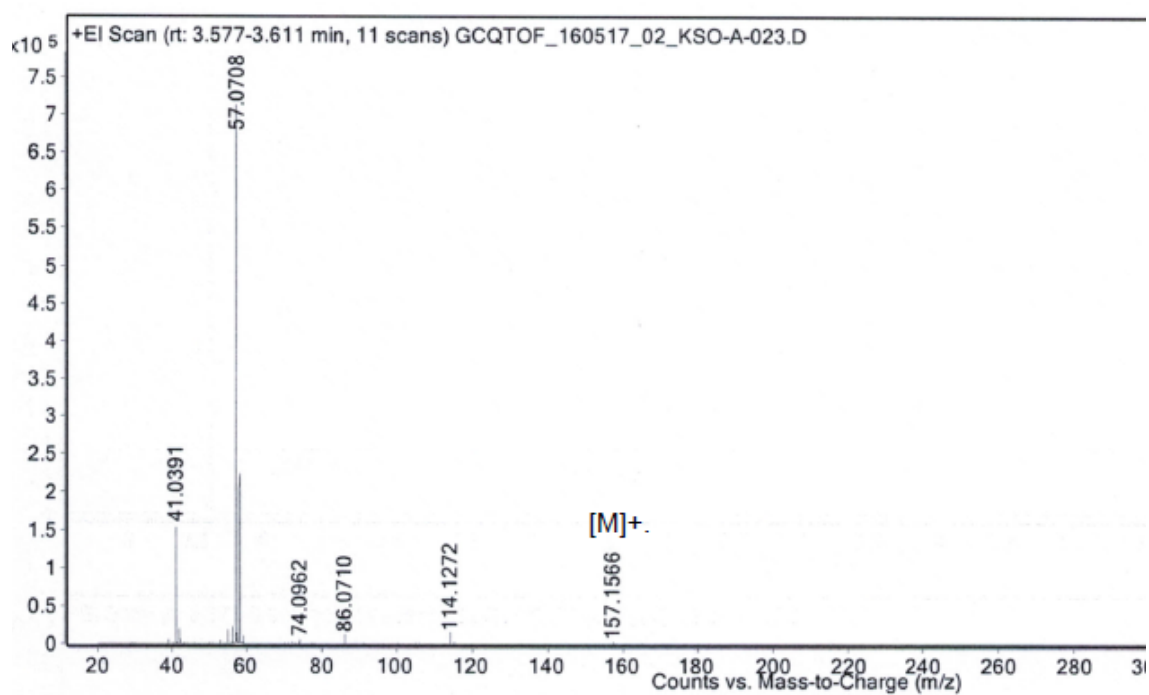


Figure S9: Mass Spectrum of **HL⁴**.

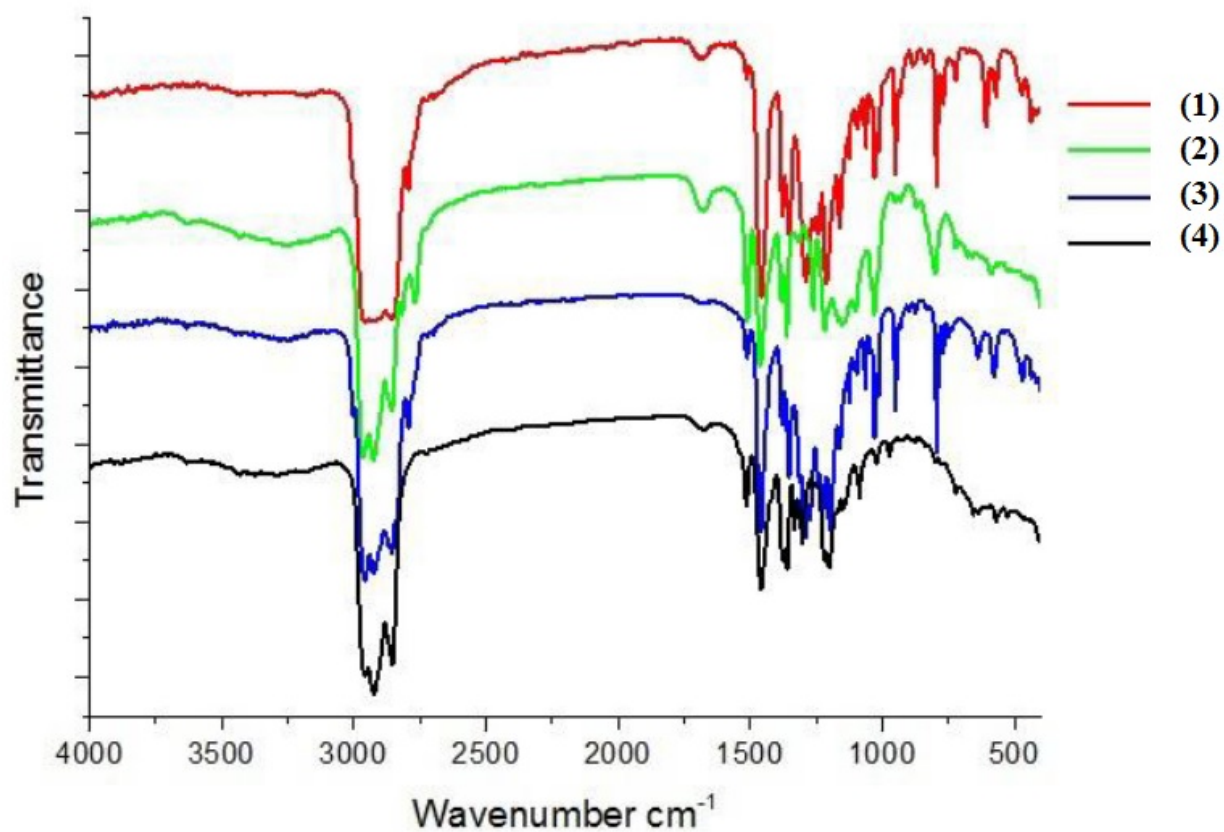


Figure S10: FT-IR spectra of the new Fe(II) complexes **1-4**.

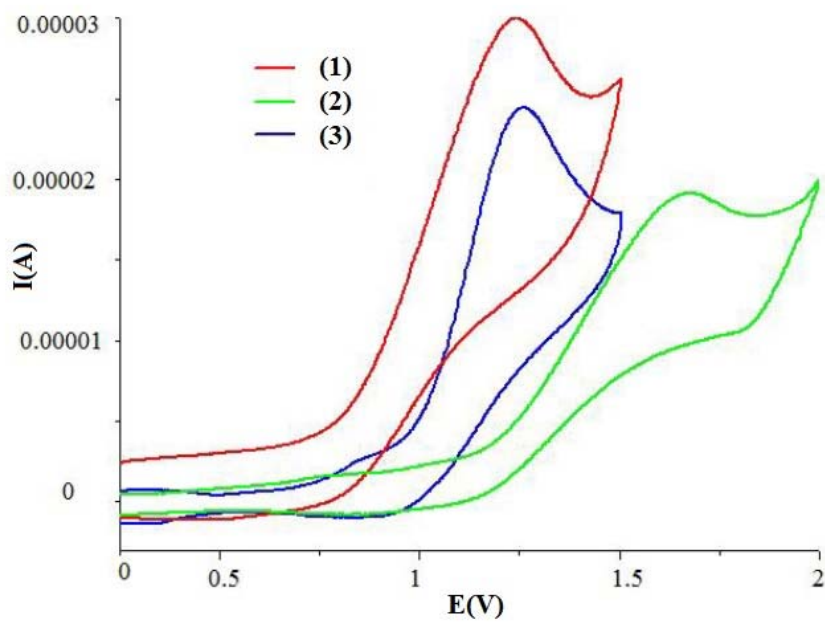


Figure S11. Voltammogram of the new Fe(II) complexes **(1)-(3)**.

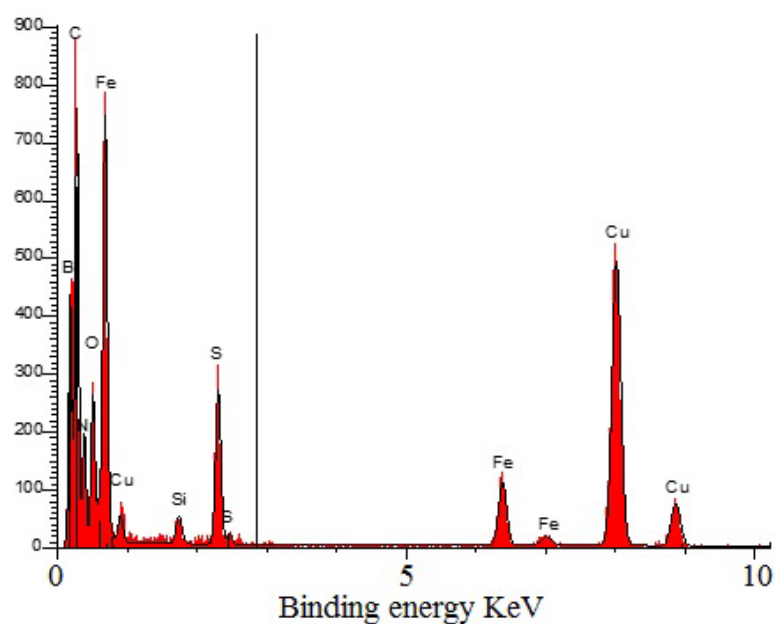


Figure S12. EDX analysis on iron nanoparticles prepared from solution reduction in toluene. The presence of N and S are due to the use of the ionic liquid as medium while Cu and C impurities come from the grid.

Table S1: % atomic concentration of species present on the least oxidized part A and in the most oxidized one B in the sample of $\text{Al}_{13}\text{Fe}_4$ nanoparticles.

% atomic concentration	Fe	Al	O	Si	Cl
A	11.3	54.16	32.06	1.4	1.08
B	8.16	3.25	80.25	-	-