

## Electronic Supporting Information

# Gels, Xerogels and Films of Polynuclear Iron(II)–Aminotriazole Spin-Crossover Polymeric Complexes<sup>†</sup>

**Antoni Sánchez-Ferrer<sup>‡a</sup>, Irene Bräunlich<sup>‡b</sup>, Janne Ruokolainen<sup>c</sup>, Matthias Bauer<sup>d</sup>,  
Rahel Schepper<sup>d</sup>, Paul Smith<sup>b</sup>, Walter Caseri<sup>\*b</sup> and Raffaele Mezzenga<sup>\*a</sup>**

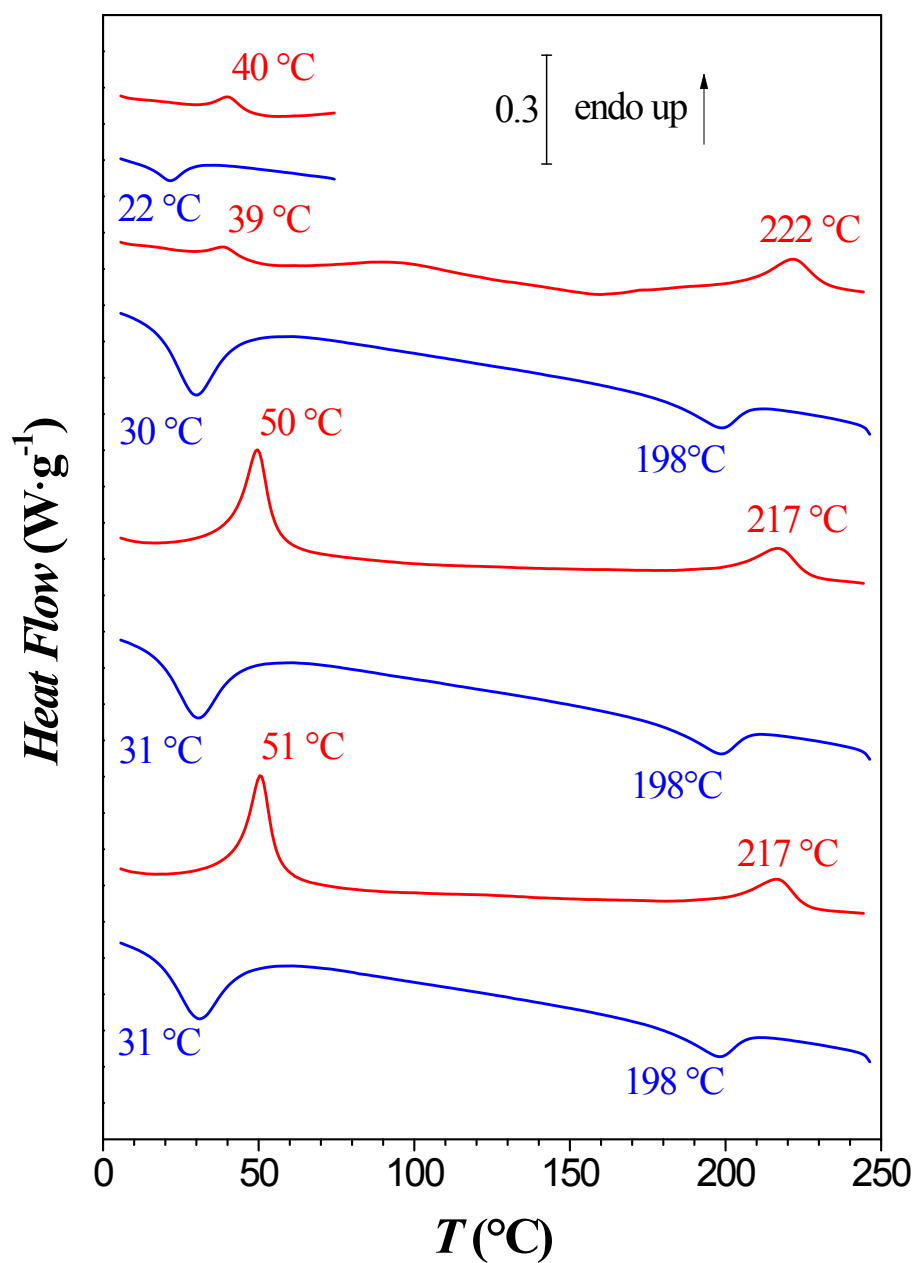
<sup>a</sup> *Department of Health Sciences and Technology, Eidgenössische Technische Hochschule (ETH) Zürich, Schmelzbergstrasse 9, 8092 Zürich, Switzerland. Fax: +41 44 632 16 03; Tel: +41 44 632 91 40; E-mail: [raffaele.mezzenga@hest.ethz.ch](mailto:raffaele.mezzenga@hest.ethz.ch)*

<sup>b</sup> *Department of Materials, Eidgenössische Technische Hochschule (ETH) Zürich, Vladimir-Prelog-Weg 5, 8093 Zürich, Switzerland. Fax: +41 44 632 11 78; Tel: +41 44 632 22 18; E-mail: [walter.caseri@mat.ethz.ch](mailto:walter.caseri@mat.ethz.ch)*

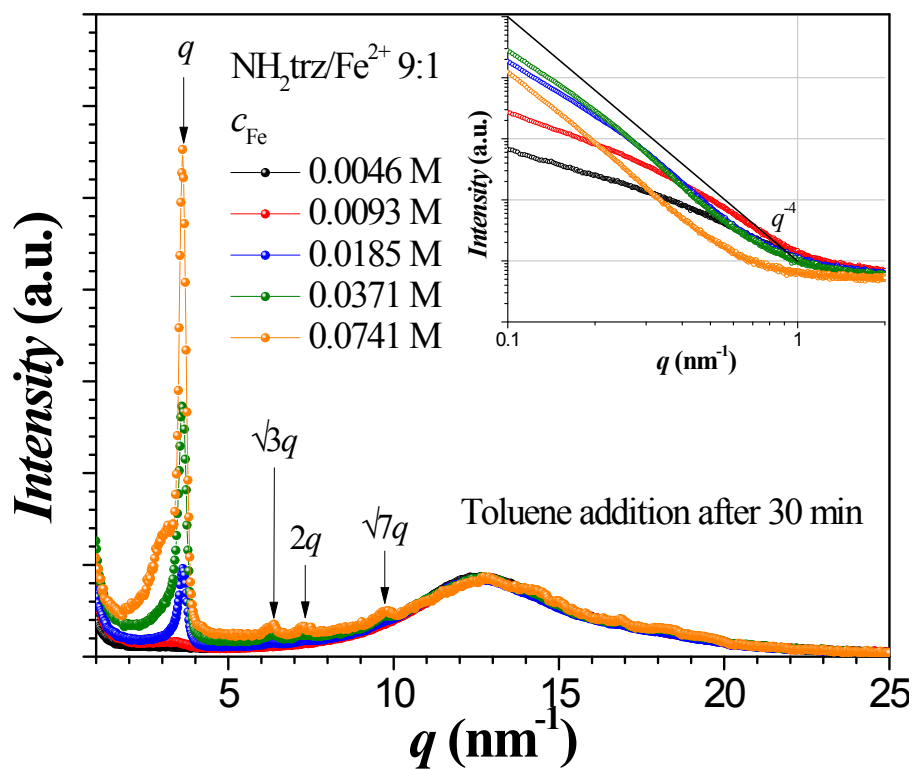
<sup>c</sup> *Department of Applied Physics, Aalto University School of Science, Puumiehenkuja 2, 00076 Aalto, Finland.*

<sup>d</sup> *Fachbereich Chemie, Universität Paderborn, Warburger Straße 100, 33098 Paderborn, Germany.*

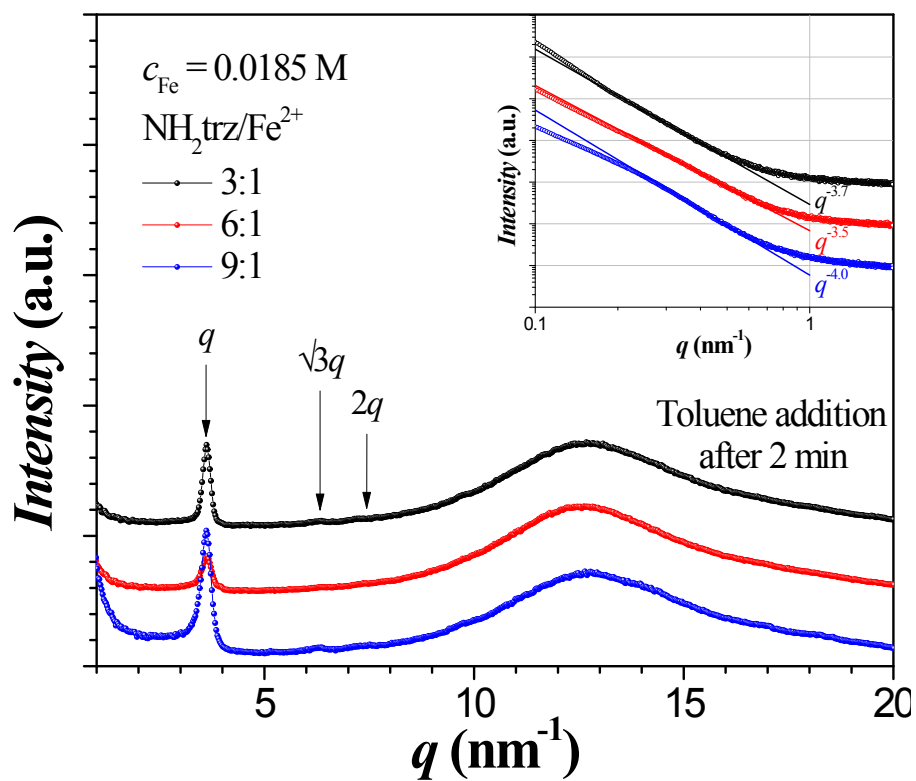
<sup>‡</sup> The first two authors contributed equally to this study.



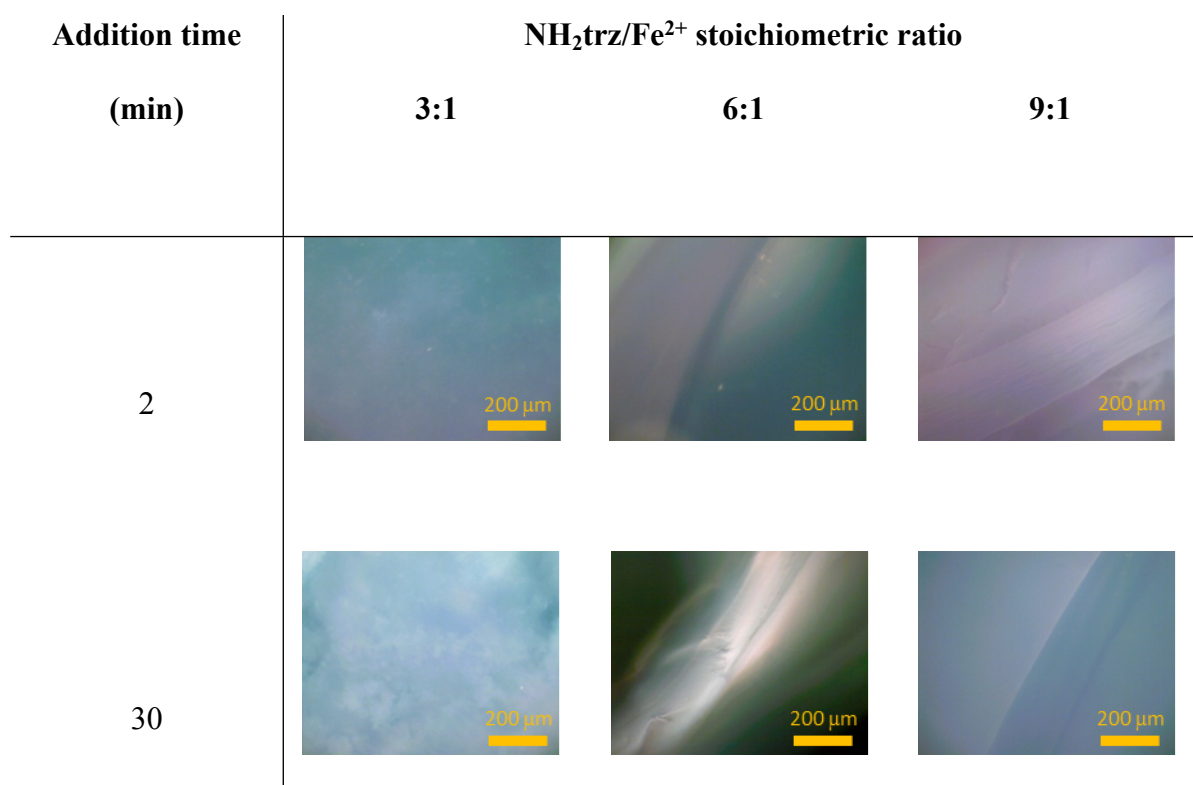
**Figure ESI-1:** Differential scanning calorimetry thermograms of  $[\text{Fe}(\text{NH}_2\text{trz})_3](2\text{ns})_2$  during 4 heating and cooling cycles (from top to bottom) at heating and cooling rates of  $10\text{ }^\circ\text{C}\cdot\text{min}^{-1}$ .



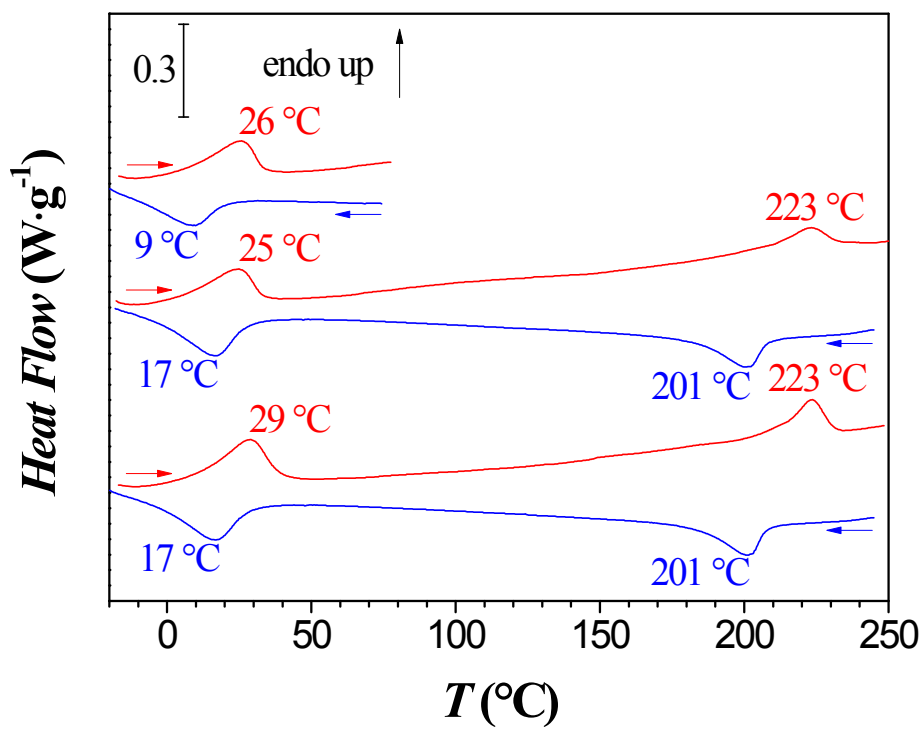
**Figure ESI-2:** WAXS 1D scattering profile for gels of *type B* at different iron concentration ( $c_{\text{Fe}}$  between 0.0046 M and 0.0741 M) and at constant stoichiometric ratio ( $\text{NH}_2\text{trz}/\text{Fe}^{2+}$  9:1) by adding toluene after 30 min of sample preparation. The peaks show a columnar hexagonal packing of the rigid rods with a lattice parameter of  $a = 2.00$  nm and correlation length of  $\xi = 25$  nm. Inset is the SAXS 1D scattering profile for the gels of *type B* with slopes close to -4 for the well-formed gels ( $c_{\text{Fe}} \geq 0.0185\text{M}$ ).



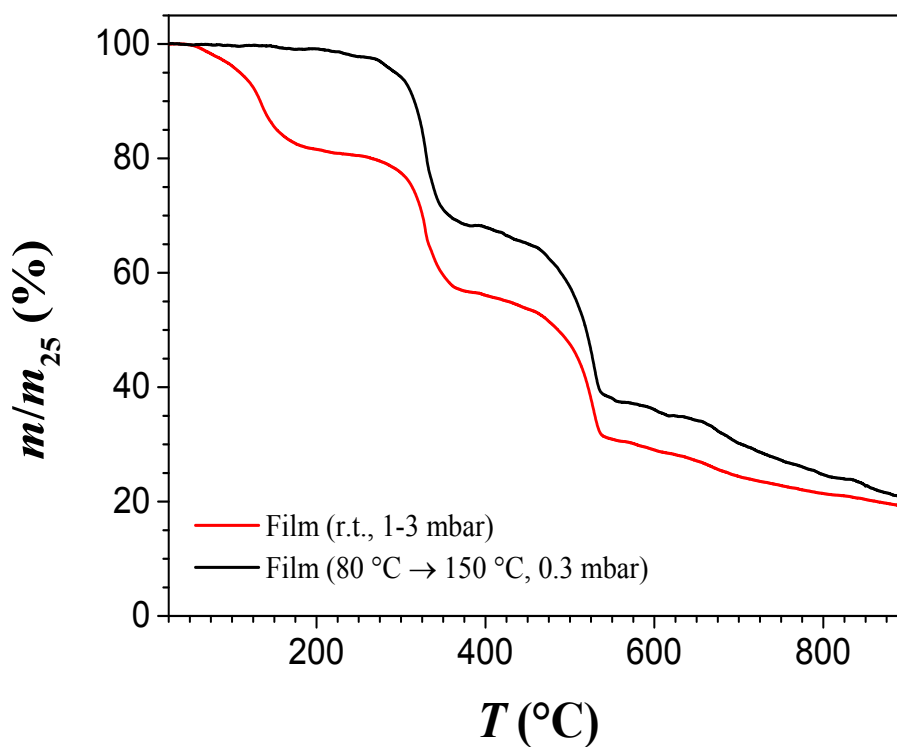
**Figure ESI-3:** WAXS 1D scattering profile for the gels of *type B* different stoichiometric ratio ( $\text{NH}_2\text{trz}/\text{Fe}^{2+}$  3:1, 6:1 and 9:1) and at constant iron concentration ( $c_{\text{Fe}} = 0.0185 \text{ M}$ ) and by adding toluene after 2 min of sample preparation. The peaks show a columnar hexagonal packing of the rigid rods with a lattice parameter of  $a = 2.00 \text{ nm}$  and correlation length of  $\xi = 29 \text{ nm}$ . Inset is the SAXS 1D scattering profile for the gels *type B* with slopes close to -4.



**Figure ESI-4:** Polarized optical microscopy (POM) images taken with crossed polarizers of gels of *type B* at  $c_{\text{Fe}} = 0.0185 \text{ M}$  and  $\text{NH}_2\text{trz}/\text{Fe}^{2+}$  3:1, 6:1 and 9:1, and at different addition time of toluene after sample preparation.



**Figure ESI-5:** Differential scanning calorimetry thermograms of the film during 3 heating and cooling cycles (from top to bottom) at heating and cooling rates of 10 °C·min<sup>-1</sup>.



**Figure ESI-6:** Thermogravimetric analysis (TGA) of a film dried at 1-3 mbar at room temperature for 24 h (red), and after subsequent drying at 0.3 mbar at 80 °C for 8 h and 150 °C for 1 h (black). The curve of the sample dried at r.t., 1-3 mbar (red curve) shows a maximum mass loss around 150 °C which corresponds to the loss of residual DMF. Indeed, the solvent could be completely removed by drying the film at elevated temperature and reduced pressure (150 °C, 0.3 mbar), as shown in the corresponding TGA curve (black curve), in which the film does not show a significant loss of weight up to ca 250 °C.