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Self-assembled Metallogels Formed From N, N', N''-tris(4-pyridyl)-trimesic amide in Aqueous Solution Induced by Fe(III)/Fe(II) Ions

Jin-Lian Zhong\textsuperscript{a,b}, Xin-Jian Jia \textsuperscript{b}, Hui-Jin Liu \textsuperscript{b}, Xu-Zhong Luo\textsuperscript{b*}, San-Guo Hong\textsuperscript{b}, Ning Zhang\textsuperscript{a}, Jian-Bin Huang\textsuperscript{c*}

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In this work, we report self-assembled metallogels formed from a ligand of trimesic amide, N,N',N''-tris(4-pyridyl)trimesic amide (TPTA), induced by Fe(III)/Fe(II) ions. TPTA is difficult to dissolve in water even in the presence of some metal ions such as Cu\textsuperscript{2+}, Co\textsuperscript{2+}, Ni\textsuperscript{2+}, K\textsuperscript{+}, Na\textsuperscript{+} and Mg\textsuperscript{2+}. Interestingly, upon heating TPTA can be dissolved easily in aqueous solution containing Fe\textsuperscript{3+}/Fe\textsuperscript{2+}, and subsequently self-assembled into metallogels after cooling. The metallogels could also be formed in aqueous solutions of mixed metal ions containing Fe\textsuperscript{3+}/Fe\textsuperscript{2+}, indicating the other metal ions do not affect the formation of Fe(III)-TPTA and Fe(II)-TPTA metallogels. The high selectivity of metallogels formation to Fe\textsuperscript{3+}/Fe\textsuperscript{2+} may be used for application in test of Fe\textsuperscript{3+}/Fe\textsuperscript{2+}. The obtained metallogels are characterized by scanning electron microscopy, Fourier transform infrared spectra, nuclear magnetic resonance spectra, rheological measurement and scanning tunneling microscope. The results indicate that TPTA can self-assemble into fibrous aggregates in Fe\textsuperscript{3+}/Fe\textsuperscript{2+} aqueous solution through the metal-ligand interactions and intermolecular hydrogen bonding. This kind of metallogels also possesses good mechanical property and thermoreversibility.

1. Introduction

Self-assembly has emerged as a powerful process for the production of well-ordered supramolecular structures at the nanometer scale. Supramolecular gels formed from small molecules (i.e., supramolecular gelators or molecular gelators) and liquids have recently triggered enormous research activities because of their wide potential applications in drug-delivery systems,\textsuperscript{1-3} organic transistors,\textsuperscript{4} soft organic nanomaterials,\textsuperscript{5,6} biofilm simulation\textsuperscript{7,9} and other fields.\textsuperscript{10-13} Recently, great efforts have been directed toward the synthesis of the supramolecular gels based on hydrogen bonding,\textsuperscript{14,15} host-guest interactions,\textsuperscript{16,18} metal-ligand coordination\textsuperscript{19,20} and acid-base interactions.\textsuperscript{21} As one kind of “smart” soft materials, the unique property of supramolecular hydrogels allows them easily to be biocompatible and biodegradable, and their resemblance to extracellular matrices have stimulated efforts to design and synthesize novel supramolecular hydrogelators as materials for biomedical applications.\textsuperscript{22,23}

One of the most rapidly developing areas of functional materials nowadays is metallogels.\textsuperscript{24,25} The incorporation of metal ions to gels brings new properties to the system. Examples are catalytic and redox activity, conductivity, luminescence, and magnetism which significantly increase as catalysts, electronic devices, sensors etc. The self-assembly of Fe(III) ions with small molecules to form metallogels has attracted increasing interest in recent years. For example, Fe\textsuperscript{3+}-based metallogels formed in organic liquids were reported by Banerjee\textsuperscript{26} and Weiss\textsuperscript{27} groups. Several metallogels based on trimesic amide derivatives and the other metal ions were also extended by Dastidar et al.\textsuperscript{27} Although there are many reports on the low molecular weight metallogels, it is very difficult to control the self-assembly of low weight molecular in the metallogels formation process. So, it still remains a great challenge to synthesize metallogels derived from the low weight molecular through a facile method.

Herein, we develop a facile method to synthesize a low molecular weight metallogels derived from Trimesic Amide. We design and synthesize compound, N,N',N''-tris(4-pyridyl)trimesic amide (TPTA), which contains pyridine N atom as coordination atom and amide group as hydrogen-bond acceptors and donors. TPTA is difficult to dissolve in water even in the presence of some metal ions such as Cu\textsuperscript{2+}, Co\textsuperscript{2+}, Ni\textsuperscript{2+}, K\textsuperscript{+}, Na\textsuperscript{+} and Mg\textsuperscript{2+} under heating, and it exhibits no gelation ability in the condition. Interestingly, upon heating TPTA can be dissolved easily in aqueous solution containing Fe\textsuperscript{3+}/Fe\textsuperscript{2+}, and subsequently self-assembled into metallogels after cooling. The high selectivity of metallogels formation to Fe\textsuperscript{3+}/Fe\textsuperscript{2+} may assign the metallogels application in test of metal ions. The obtained metallogels possess good mechanical property and thermoreversibility.
2. Experimental section

2.1 Materials
All chemicals were commercially available. Deuterium oxide and dimethyl sulfoxide-\textit{d}_6 were purchased from J&K chemical company.

N,N’,N”-tris(4-pyridyl)trimesic amide (TPTA) was synthesized according to the method reported in reference.\textsuperscript{28a} The resulting product was characterized by IR, NMR and MS (Scheme S1, Figures S1, S2, and S3 in ESI†).

2.2 Gelation experiments
The gelation ability of TPTA was investigated by a typical test tube experiment. Weighted amounts of TPTA and metal ion aqueous solution were placed into sealed scintillation vial with a cap. The scintillation vial was heated in a water bath until the solid material dissolved. The resulting solutions were slowly cooled to room temperature and the formation of the gels was confirmed by inverting the scintillation vial containing the solution.

2.3 Characterization

Scanning electron microscopy (SEM) images were performed using a FEI QUANTA 450 with an accelerating voltage of 15.0 KV. The preparation of samples for SEM involved placing a drop of hydrogel on the copper substrate. The hydrogel was subjected to shock-freezing by liquid nitrogen, followed by lyophilization for 3 h. It was then submitted to a SEM scan after gold-coating for 3 min. Fourier transform infrared (FT-IR) spectra measurements were carried out on a Nicolet iS50 FTIR spectrophotometer. CaF\textsubscript{2} substrates and KBr pellet were used for transmission spectra of xerogel and crystal, respectively. A drop of metallogel was cast onto CaF\textsubscript{2} substrates, which was dried at 60 °C by vacuumizing method and then the obtained xerogel was subjected to FT-IR spectra measurements. Small-angle X-ray Scattering (SAXS) patterns of dried hydrogel were measured on a Bruker D8 Focus diffractometer using the CuK\textalpha radiation. X-ray was generated with a Cu anode and the CuK\textalpha beam (\lambda = 1.5418 Å) was taken out via a graphite monochromator. \textsuperscript{1}H Nuclear Magnetic Resonance (\textsuperscript{1}H NMR) spectra were measured on a Bruker AVANCE 400 MHz spectrometer. Differential scanning calorimetry (DSC) was conducted on Setaram \textmu DSC7-Evo instrument (heating rate, 1°C⋅min\textsuperscript{-1}). Rheological measurement was performed using a stress-controlled rheometer (HAAKE RheoStress 6000) with parallel plate type geometry (plate diameter, 3.5 cm). A solvent trap equipped with the rheometer was used to protect the sample from evaporation. Temperature dependences of the storage shear moduli (G\textsuperscript{'} and the loss shear moduli (G\textsuperscript{''}) were carried out by heating the sample at a rate of 0.1 °C/S. The viscoelastic moduli were monitored under small-amplitude oscillatory shear at an applied frequency of 1Hz and 0.1 °C/S. The viscoelastic moduli were monitored under small-amplitude oscillatory shear at an applied frequency of 1Hz and stress of 1 Pa. Frequency sweeps at selected temperatures were carried out over a range of 0.1-100 rad⋅s\textsuperscript{-1} at a stress of 1 Pa. Scanning tunneling microscope (STM) measurements were performed by using a Nanoscope 3D (Bruker) with mechanically formed Pt/Ir (80/20) tips. A droplet (2 mL) of the 1-phenyloctane solutions was deposited on a freshly cleaved HOPG surface and immediately observed by STM in constant current mode. The specific tunneling conditions are given in the corresponding figure captions. UV absorption spectra were obtained using a UV-1800 UV-Vis spectrophotometer. In the experiments, for SEM, IR, DSC, NMR, UV and Rheological measurement, FeCl\textsubscript{2}·6H\textsubscript{2}O and FeSO\textsubscript{4}·7H\textsubscript{2}O have been used for gel and solution preparation. For STM measurements, FeCl\textsubscript{2}·4H\textsubscript{2}O and FeCl\textsubscript{3}·6H\textsubscript{2}O have been used for solution preparation.

3. Results and discussion

3.1 Gelation test
By experiments, we found that TPTA was difficult to dissolve in water even on heating. Previous report\textsuperscript{28c} revealed that TPTA could form stable thermoreversible gels in some polar organic solvent containing a certain amount of water. While some other trimesic amide derivatives could form stable gels in water\textsuperscript{29a,b} and some organic solvents\textsuperscript{29c–f} TPTA contains pyridine groups which can coordinate with metal ions. In order to understand the effect of metal ions on the gelation abilities of TPTA, the gelation test of TPTA in aqueous solution containing metal ions were performed. The results are listed in Table S1 in ESI†. TPTA was difficult to dissolve in aqueous solution containing most metal ions tested in the experiment such as Cu\textsuperscript{2+}, Co\textsuperscript{2+}, Ni\textsuperscript{2+}, K\textsuperscript{+}, Na\textsuperscript{+}, Mg\textsuperscript{2+} ions even on heating and no hydrogels were formed. Interestingly, TPTA can be dissolved easily in aqueous solution containing Fe\textsuperscript{3+}/Fe\textsuperscript{2+} upon heating and gelatinized aqueous solution efficiently after cooling to form the metallogel (Fig. 1,
denoted by Fe(III)-TPTA and Fe(II)-TPTA gel, respectively). The results indicate that the formation of TPTA metallogels has a high selectivity for Fe$^{3+}$ and Fe$^{2+}$. In order to determine the gelatinizing ability of TPTA in Fe$^{3+}$/Fe$^{2+}$ aqueous solution, the minimum gelator concentration (MGC) is measured. The minimum amounts of TPTA dissolved in 1L Fe$^{3+}$/Fe$^{2+}$ aqueous solution of different concentration necessary for gelatinizing aqueous solution efficiently was assigned to MGC of TPTA. The MGCs of TPTA in Fe$^{3+}$/Fe$^{2+}$ aqueous solution were listed in Table 1. From table 1, it can be known that the MGCs increase with the concentrations of Fe$^{3+}$/Fe$^{2+}$. For example, the MGCs of Fe(III)-TPTA are 11.2 g and 45.0 g for 0.050 mol/L and 0.20 mol/L Fe$^{3+}$ aqueous solution, respectively.

![Diagram](Image)

Fig. 1 (a) Metallogels Formation; The photographs of gel-sol transition process of (b) Fe(III)-TPTA; (c) Fe(II)-TPTA

In addition, TPTA does not dissolve in the aqueous solution of K$_2$[Fe(CN)$_6$] and K$_4$[Fe(CN)$_6$] and exhibits no gelation ability (Table S1 in ESI†), indicating that only free ions of Fe$^{3+}$/Fe$^{2+}$ could induce TPTA to self-assemble into the hydrogel. On the other hand, metallogels are also formed in aqueous solution using Fe$^{3+}$/Fe$^{2+}$ salts of Cl$^-$, NO$_3^-$ and SO$_4^{2-}$, indicating that anions can not affect the formation of hydrogels. Similar results were also observed in metallogels formed from Cu(II) ions and tetratopic ligand rctt-1,2-bis(3-pyridyl)-3,4-bis(4-pyridyl) cyclobutane.$^{29}$

The effects of other metal ions on the Fe(III)-TPTA and Fe(II)-TPTA gel formation were also studied(Fig. S4 in ESI†). TPTA does not dissolve in aqueous solution containing Na$^+$, Cu$^{2+}$, Mn$^{2+}$, Co$^{2+}$, K$^+$ and Cr$^{3+}$ ions, respectively or their mixed solution even under the condition of heating. However, TPTA exhibits good solubility in aqueous solutions of mixed metal ions containing Fe$^{3+}$/Fe$^{2+}$ upon heating. When a certain amount of TPTA was dissolved in aqueous solutions of mixed metal ions containing Fe$^{3+}$/Fe$^{2+}$ upon heating, metallogel can be formed after the solution cooled to room temperature. The results indicate that the other metal ions do not affect the formation of Fe(III)-TPTA and Fe(II)-TPTA metallogels. Therefore, Fe(III)-TPTA and Fe(II)-TPTA metallogels have a good test function for Fe$^{3+}$/Fe$^{2+}$.

Previous reports$^{28b,c}$ revealed that the addition of the polar solvent to water can enhance the solubility of TPTA significantly and thermo-reversible gels were formed on cooling the heated solutions to room temperature, although TPTA is difficult to dissolve in water. The aggregation behaviour of some asymmetric benzene-1,3,5-tricarboxamides in organic solvents was systematically studied by Meijer et al.$^{28}$ Their results revealed that even small changes in the molecular structure of substituted benzene-1,3,5-tricarboxamides would affect their self-assembly behaviour in dilute solutions. In our work, we find that the addition of HCl can also enhance the solubility of TPTA significantly in water, but no gel formation is observed. That the addition of Fe$^{3+}$/Fe$^{2+}$ enhances the solubility of TPTA significantly in water and stable gels are formed on cooling the heated solutions to room temperature may be acribled to self-assembly induced by the coordination effect of Fe$^{3+}$/Fe$^{2+}$ and TPTA, which would be confirmed further by scanning tunneling microscopy (STM) observation and the experiments of coordination number of TPTA-Fe$^{3+}$ determining (vide infra).

### Table 1 The MGCs of TPTA for Fe(III) and Fe(II) aqueous solution of different concentration

<table>
<thead>
<tr>
<th>Gelators</th>
<th>Concentration of Fe$^{3+}$ (mol/L)</th>
<th>Concentration of Fe$^{2+}$ (mol/L)</th>
<th>The MGC of hydrogel (g/L)</th>
</tr>
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<tbody>
<tr>
<td>TPTA</td>
<td>0.050</td>
<td>----</td>
<td>11.2</td>
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<tr>
<td></td>
<td>0.10</td>
<td>----</td>
<td>22.8</td>
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<td>0.15</td>
<td>----</td>
<td>33.7</td>
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<td>0.20</td>
<td>----</td>
<td>45.0</td>
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<tr>
<td>TPTA</td>
<td>----</td>
<td>0.010</td>
<td>4.4</td>
</tr>
<tr>
<td></td>
<td>----</td>
<td>0.020</td>
<td>9.1</td>
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<tr>
<td></td>
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<td>0.030</td>
<td>14.4</td>
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### 3.2 Morphologies and thermal properties

In order to gain an insight into the aggregate morphology, xerogels were studied by scanning electron microscopy. The xerogels were obtained by freeze-drying the hydrogels for avoiding damage from the high vacuum or drying of the samples. As shown in Fig. 2, the xerogels of Fe(III)-TPTA and Fe(II)-TPTA metallogels are composed of fibrous structures at different
concentration. It indicated that the concentrations of Fe$^{3+}$ and Fe$^{2+}$ can not change the aggregates morphology of gelators in metallogels. On the basis of a fibrous structure, the inter-twisted and interlocked fibers form the 3D porous network structure. Large amounts of entangled fibers and fiber bundles are clearly observed. The width of a fiber bundle is approximately 1-3 µm, and the small fiber diameter ranged from 0.1 to 0.4 µm. It indicated that the observed fiber bundles are composed of smaller fiber. It is obvious that the original fibers, aggregates formed from TPTA induced by Fe$^{3+}$/Fe$^{2+}$, proceed to assemble into small fibers, and then get entangled to form a close-knit porous network. Gelator molecules created complex three dimensional networks by entangling numerous tiny fibers and entrapped abundant water in the interspace of the networks by surface tension and capillary forces, leading to the formation of metallogels.

Facile thermoreversible gel-sol transition is a notable feature of supramolecular gels. Typical heating and cooling DSC curves of metallogels obtained from varying concentrations of TPTA in aqueous solution containing Fe(III)/Fe(II) ions are studied. The gel-sol transition temperatures ($T_{gs}$) of metallogels at different concentrations of TPTA were obtained from DSC experiments. By visual inspection, the samples were liquids above $T_{gs}$ and then became gels below $T_{gs}$. Shown in Fig. 3a, b and c are the gel-sol transition temperatures ($T_{gs}$) profiles of metallogels prepared by varying concentrations of TPTA in aqueous solution containing 0.050 mol/L Fe$^{3+}$, 0.10 mol/L Fe$^{3+}$ and 0.010 mol/L Fe$^{2+}$, respectively.

It can be known from Fig. 3 that the thermal stability of the Fe(II)-TPTA gels is greater than that of the corresponding Fe(III)-TPTA gels at the same concentrations of TPTA. The $T_{gs}$ increased gradually and eventually reached a plateau with the increasing of the concentration of TPTA when the concentration of Fe$^{3+}$/Fe$^{2+}$ remained unchanged. The temperature of plateau region was denoted as a concentration-independent $T_{gs}$. The plateau regions occurred at beginning TPTA concentrations of 19.0, 34.0 and 13.0 g/L for Fe(III)-TPTA(Fe$^{3+}$)=0.050 mol/L, Fe(III)-TPTA(Fe$^{3+}$)=0.10 mol/L and Fe(II)-TPTA(Fe$^{2+}$)=0.010 mol/L gels, respectively. The corresponding concentration-independent $T_{gs}$ are 44.6, 55.6 and 74.5 °C, respectively. These results imply that at the beginning, the intricately and densely entangled fibrous were formed and the thermal stability of each gel was gradually enhanced following the increasing of the concentration of TPTA. Thereby, a high temperature is need to break the fibrous structure. However, when the concentration increased to a certain value, the $T_{gs}$ reached a constant.

3.3 Rheological properties

The mechanical performance of a material is extremely important for its practical uses. To explore the property of gel in detail, rheological measurements were conducted on Fe(III)-TPTA and Fe(II)-TPTA gels. According to Fig. 4a, frequency sweep exhibits typical solid-like rheological behavior with the storage moduli (G') dominating the loss moduli (G'') over the investigated oscillating frequency range. Below a certain level of stress, G' and G'' slightly increase with the frequency from 0.01 to 10 Hz and the value of G' is always larger...
than that of $G''$ in the whole range (0.01-10 Hz), suggesting that the gels are fairly tolerant to the external force.

Additionally, variable-temperature studies were conducted to characterize the thermo-responsive behaviors of Fe(III)-TPTA and Fe(II)-TPTA gels.\(^{37}\) Fig. 4b shows the variation tendency of the storage shear moduli ($G'$) and loss shear moduli ($G''$) of gels with increasing temperature. As shown in Fig. 4b, both $G'$ and $G''$ of Fe(III)-TPTA gel (0.10 mol/L Fe\(^{3+}\), 28.0 g/L TPTA) almost keep constant from room temperature to 35°C, above which both $G'$ and $G''$ decrease rapidly with the increase of temperature, indicating the gradual transformation from gel to solution. When the temperature reaches 45°C, a cross-over point where $G'$ equals to $G''$ appears, marking the transition from primarily elastic to viscous properties. Upon further heating to 55°C, $G'$ exceeds $G''$, indicating that gel transforms to solution completely. The result is consistent with the $T_g$ obtained by DSC experiments. Similar results were observed for Fe(II)-TPTA gel (0.010 mol/L Fe\(^{2+}\), 6.0 g/L TPTA) (Fig. 4b).

3.4 IR and \(^1\)H NMR Spectral Studies

It is well-known that FT-IR spectroscopy is a powerful tool for investigating intermolecular hydrogen bonding interaction. In order to gain insight into the hydrogen-bonding environment of the amide C=O group, FT-IR experiments were performed. As shown in Fig. 5, the FT-IR spectrum of crystal TPTA is characterized by bands appear at 3235 and 1687 cm\(^{-1}\) attributed to the N-H and C=O stretching bands of amide, respectively. Generally, N-H and C=O stretching bands of amide for a free secondary amide group locate at 3440, 1680 cm\(^{-1}\), respectively.\(^{38,39}\) The red shift of N-H stretching band of amide for crystal TPTA compared to a free secondary amide group may be attributed to the strong hydrogen-bonding interactions. The C=O stretching band of amide is close to a free amide indicating that the C=O group is free. The analyses agree well with previous reported crystal structure of TPTA reported by our group\(^{28a}\) and Dastidar et al.\(^{28c}\) In the crystal TPTA, only the hydrogen bonds between N-H of amide and N of pyridine can form and the C=O groups of amides are free.

For Fe(III)-TPTA xerogel (Figure 5b), N-H stretching band of amide appears at 3203 cm\(^{-1}\), implying N-H of amide group takes part in the formation of intermolecular hydrogen bonds and no free N-H group exists. The C=O stretching bands of amides appear at 1702 and 1638 cm\(^{-1}\), ascribing to free and strong hydrogen bonding amide groups respectively. The result demonstrates some of the C=O groups of amides participate in formation of intermolecular hydrogen bonds and the others are in free state. Compared to IR results of crystal TPTA, it can be deduced that the intermolecular hydrogen bonds between
adjacent amide groups and the intermolecular hydrogen bonds between N-H of amide and N of pyridine have formed in the Fe(III)-TPTA gel. The shift of C=O stretching vibration of free C=O group from 1687 (for crystal TPTA) to 1702 cm⁻¹ (for Fe(III)-TPTA xerogel) may be ascribed to the influence of coordination interaction between Fe(III)/Fe(II) and TPTA. The IR spectrum of Fe(II)-TPTA xerogel (Figure 5c) is almost the same as that of Fe(III)-TPTA gel (Figure 5b), suggesting that the pattern of hydrogen bonding in Fe(II)-TPTA xerogel is close to that in Fe(III)-TPTA gel. The result indicates that all the N-H groups take part in the formation of intermolecular hydrogen bonds. Two types of intermolecular hydrogen bonds have formed. One is between adjacent amide groups and the other is between N-H of amide and N of pyridine.

It is well known that ¹H NMR techniques can give a great deal of information on self-assembly process in the gel state. Especially, ¹H NMR experiments may provide an insight into how molecules are orientated with respect to one another in a self-assembled state. In general, ¹H NMR signals of molecules in solution at moderate field strengths are sharp, while those of samples in liquid crystal, glass, and gel state broaden and also shift from their usual ¹H NMR frequency values. Thus, a correlation of the ¹H NMR spectra of the species in solution with that of gel provides an insight into the self-assembly process.

Figure 6(a) shows the ¹H NMR spectra of a solution of Fe(III)-TPTA in D₂O-DMSO and the gel of Fe(III)-TPTA in D₂O at various temperatures. As shown in Figure 6a, aromatic proton signals of Fe(III)-TPTA in solution phase appeared at 9.01 (s, H₂,₆,9 ph), 8.75 (b, H₃,₅ py) and 8.35 ppm (s, H₂,₆ py) at 20°C, while those in gels phase further broadened, to the extent that they almost disappeared, suggesting that almost all of the Fe(III)-TPTA was incorporated into the gel, rendering it more solid-like, and thus its molecular motion was restricted. Upon heating, the aromatic proton signals in the gel phase gradually appeared with downfield shift from 8.07 (b, H₃,₅ py) and 7.75 (b, H₂,₆ py) at 20°C to 8.54 (b, H₃,₅ py) and 8.20 ppm (b, H₂,₆ py) at 70°C, respectively. Recoveries of the resonance lines of aromatic proton were observed closer to the gel-melting temperature (50 °C) and there was a dramatic sharpening of these resonance lines upon complete gel melting (60 °C). The upfield shift of the aromatic protons in the gel phase may arise from π-π stacking interactions.

For understanding the information of hydrogen bond formation in the solution phase, concentration-dependent ¹H NMR studies were conducted. The ¹H NMR spectra of a solution containing different amount of Fe(III)-TPTA in D₂O-DMSO at 20 °C are shown in Figure 6(b). It is seen that the signals of the amide protons shifted gradually to lower field (from 11.84 to 12.03 (11.93) ppm) on increasing the concentration of the gelator from 1.0 to 4.0 mg/mL, suggesting that intermolecular hydrogen bonds have formed in D₂O-DMSO solution of Fe(III)-TPTA at a relatively high concentration. The signals of the amide protons at 12.03 and 11.93 ppm for solution of 4.0 mg/mL Fe(III)-TPTA can be attributed to different type intermolecular hydrogen-bonded N-H groups. The above results imply that no free N-H groups exists in a relatively high concentration solution of Fe(III)-TPTA in D₂O-DMSO. According to the above analyses, it can be deduced that in a gel of Fe(III)-TPTA, all the N-H groups take part in the formation of intermolecular hydrogen bonds because the concentration of a gels is much higher than a solution. The result is consistent with that obtained by IR experiments.
Similarly, $^1$H NMR spectroscopy of Fe(II)-TPTA gel in D$_2$O at various temperatures showed significant line broadening of the protons of the aromatic ring at 20 °C (Fig. S6a in ESI†). Upon heating to 60 °C, the resonance peaks started to become sharp with downfield shift from 8.45 (s, H$_{2,4,6}$-ph), 8.31 (b, H$_{3,5}$-py) ppm and 7.99 (b, H$_{2,6}$-py) to 9.02 (s, H$_{2,4,6}$-ph), 8.88 (b, H$_{3,5}$-py) and 8.53 ppm (b, H$_{2,6}$-py), respectively, indicating a π-π stacking interactions of Fe(II)-TPTA in the gel phase. This observation was consistent with the understanding that the TPTA molecules experience severe mobility restrictions in a gel and that gel melting would indirectly imply melting of that TPTA are now in a state of higher mobility, thereby averaging out or minimizing the dipolar interactions between $^1$H nuclei. The results of $^1$H NMR of a solution containing different amount of Fe(II)-TPTA in D$_2$O-DMSO at 20 °C (Fig. S6b in ESI†) also suggest that in a gel of Fe(II)-TPTA, all the N-H groups take part in the formation of intermolecular hydrogen bonds.

3.5 SAXS study

SAXS is a powerful technique for characterizing the morphologies of various types of colloidal dispersions. The aggregation structures of Fe(III)-TPTA and Fe(II)-TPTA gels were characterized by SAXS.

As shown in Figure 7, there is no clear diffraction peak at the concentration of 22.8 g/L, while two clear diffraction peaks appear at 5.02 ° and 24.50 °, corresponding to the d-spacing values of 1.78 nm and 0.36 nm, respectively, at concentration of 28.0 g/L. When the concentration up to 35.0 g/L, three clear diffraction peaks appear at 5.02 °, 10.95 ° and 24.50 °, corresponding to the spacings of 1.78 nm, 0.81 nm and 0.36 nm, respectively. The ratio of the three spacings suggests a lamellar structure with a layer distance of 1.78 nm. These results indicate that with increasing concentration of Fe(III)-TPTA, the aggregation structures of xerogels gradually transformed from amorphous to polycrystalline in Fe$^{3+}$ ion solution. In addition, similar results were also obtained for Fe(II)-TPTA gels, of which there are no clear diffraction peaks at low concentration, but with increasing concentration of Fe(II)-TPTA, more and more clear diffraction peaks appear.

3.6 The structure of self-assembled gel

In order to understand the effect of the Fe$^{3+}$/Fe$^{2+}$ on the enhancement of the solubility of TPTA in water and supramolecular structures at the single molecule level, scanning tunneling microscopy (STM) is used to investigate the assembly structures. STM images of TPTA and Fe(III)-TPTA assembling structure are presented in Fig. 8. Owing to the higher electronic density of states of the aromatic rings, the aromatic ring appears in higher contrast than in the image. From Fig. 8a, disordered arrangements can be observed when the pure TPTA solution is deposited on graphite surface. STM investigation on the 2D assembly of Fe(III)-TPTA is presented in Fig. 8b. The adjacent molecules in the same rows are parallel to each other with a uniform direction. The bright dots in the image can be assigned to TPTA. The parameters were measured to be $a = 1.0 \pm 0.1$ nm, $b = 1.6 \pm 0.1$ nm and $\alpha = 60 \pm 1.0^\circ$, respectively. In order to get detail information of the microstructure of the assembly structures, the coordination molar ratio and coordination stability constant of Fe$^{3+}$ to TPTA in aqueous solution were determined by Spectrophotometry (Fig. S7, in ESI†). The coordination molar ratio and coordination stability constant of Fe$^{3+}$ to TPTA obtained by the experiment are 1.0 and 2.14×10$^3$, respectively. In accordance with the STM image (Fig. 8c) and the coordination molar ratio of Fe$^{3+}$ to TPTA described above, the corresponding two dimensional molecular model is proposed in Fig. 8d.

The coordination molar ratio and coordination stability constant of Fe$^{2+}$ to TPTA in aqueous solution determined by spectrophotometry are 1.0 and 2.73×10$^3$, respectively (Fig. S8, in ESI†). STM investigation (Fig. S9, in ESI†) also reveals that order assemblies of Fe(II)-TPTA in two dimension are formed in...
solution. Considering the disordered arrangements of TPTA in solution, it can be inferred that the coordination of Fe$^{3+}$/Fe$^{2+}$ to TPTA is responsible for the formation of the order aggregates of TPTA in solution.

Based on the results of SEM, FT-IR, $^1$H NMR and SAXS, it can be deduced that the TPTA molecules in the two dimensional molecular layer (Fig. 8d) was connected by coordination and the intermolecular hydrogen bonds between N-H of amide and N of pyridine. The two dimensional molecular layer was further connected by inter-layer hydrogen bonds between part adjacent amide groups and π-π stacking interactions to form a hydrogen bond network and then develop a three dimensional network superstructure. The metal-ligand coordination interactions, hydrogen bonding and π-π stacking interaction played important roles in self-assembling process.

4. Conclusions

In summary, self-assembled metallogels have been successfully fabricated from TPTA in aqueous solution induced by Fe(III)/Fe(II) ions. The high selectivity of metallogels formation to Fe$^{3+}$/Fe$^{2+}$ may be used for the application in test of Fe$^{3+}$/Fe$^{2+}$. The experimental results reveal that the driving forces are intermolecular hydrogen bonding, π-π stacking interactions and coordination interactions between Fe$^{3+}$/Fe$^{2+}$ and N-py of TPTA. These interactions are responsible for the formation of three dimensional network superstructures, which finally gelate the water. The obtained metallogels possess good mechanical property and thermoreversibility. These gels may have potential application as biomaterials.

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Notes and references

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Self-assembled Metallogels Formed From N, N’, N”-tris(4-pyridyl)trimesic amide in Aqueous Solution Induced by Fe(III)/Fe(II) Ions

Jin-Lian Zhong\textsuperscript{a,b}, Xin-Jian Jia\textsuperscript{b}, Hui-Jin Liu\textsuperscript{b}, Xu-Zhong Luo\textsuperscript{b*}, San-Guo Hong\textsuperscript{a*}, Ning Zhang\textsuperscript{a}, Jian-Bin Huang\textsuperscript{c*}

We report the self-assembled metallogels formed from N,N’,N”-tris(4-pyridyl)trimesic amide (TPTA) in aqueous solution containing Fe\textsuperscript{3+}/Fe\textsuperscript{2+}. This high selectivity of metallogels formation to Fe\textsuperscript{3+}/Fe\textsuperscript{2+} may be used for the metallogels application in test of Fe\textsuperscript{3+}/Fe\textsuperscript{2+}. 

\[
\begin{align*}
\text{TPTA} & + \text{Fe}^{3+}/\text{Fe}^{2+} \\
\text{aqueous solution} & \xrightarrow{\text{dissolved upon heating and cooled to room temperature}}\\
\end{align*}
\]