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Luminescent (metallo-supramolecular) cross-linked lanthanide hydrogels from a btp (2,3-bis(1,2,3-triazol-4-yl)picolinamide) monomer give rise to strong Tb(III) and Eu(III) centred emissions†

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The development of the 2,3-bis(1,2,3-triazol-4-yl)picolinamide (**btp**) ligand **2**, from the amine **1** (which was also structurally characterised) possessing a methacrylamide unit that facilitated the copolymerisation of this **btp** ligand into polymeric hydrogels, using 2-hydroxyethyl methacrylate (HEMA), methylmethacrylate (MMA) and ethylene glycol dimethacrylate (EGDMA), is described, resulting in the formation of two polymers with two different concentrations (cast as polymeric films), **F1** (0.03 wt%) and **F2** (0.3 wt%), respectively. These polymers were then used to form luminescent soft-materials using lanthanide ions, which can coordinate to the **btp** ligands; the focus herein on the latter polymer was shown to be more luminescent. The non-polymerisable ligand **3**, possessing an acetamide moiety was also developed as a model compound. We also present the formation of a soft material (a supramolecular hydrogel) from the precursor **1**, which was shown to be thermoreversibly formed, and its morphological and rheological properties were probed. We also show that upon addition of Tb(III), to this supramolecular gel, we have interrupted the soft-material features upon coordination of the ion to the **btp** ligands. While the focus herein is on the **F2**, in both cases the metal ion coordination (which was achieved upon hydration of the polymers in solutions of these ions) results not only in the formation of a luminescent material (Tb(III) is known to give rise to highly luminescent green emission upon coordination of the ion to **btp** ligands), but also in the instantaneous formation of metal mediated crosslinking within the polymeric matrix. The photophysical properties of both **2** and **3** are presented, both in solution, and as in the case of **2**, after the polymerisation within the hydrogel; which in the case of Tb(III) results in the formation of green emitting polymers, clearly visible to the naked eye. Similarly, we demonstrate that upon using Eu(III), red emission is observed from polymer **F2**.

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Introduction

The development of responsive supramolecular systems is currently a highly topical area of research,^{1–3} with significant developments occurring in the developments of coordination polymers, MOFs, and COFs in recent times.^{3–6} Self-assembly supramolecular polymers have emerged as highly attractive

targets for the formation of hierarchical materials.^{7–9} The use of low molecular weight gelators (LMWG),¹⁰ or more recently, the use of low molecular mass ionic organogelators (LMIOGs),¹¹ as developed by Pfeffer *et al.*, has been particularly studied with this in mind. The incorporation of coordination ligands for such self-assembly formations has also been explored, resulting in the generation of metallo-supramolecular polymers and gels,^{12–14} some of which have properties beyond that of the ligand or metal-complex alone,¹⁵ as we have recently demonstrated using bio-inspired ligand design and Eu(III) as the f-metal ion of choice,¹⁶ but the lanthanides have both physical and coordination properties that are highly desirable for the generation of responsive luminescent materials.

The lanthanides are strong Lewis-acids, have diverse coordination requirements, are paramagnetic and some also display delayed metal-centred luminescence, which is both line-like, occurring at long wavelengths (visible to NIR) and long

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lived.¹⁷ As such, lanthanide complexes are highly desirable to be employed in the development of higher order supramolecular and responsive structures and materials.^{18,19} Unfortunately, in most cases, due to their Laporte-forbidden transitions, it is necessary to employ sensitizing antennae to take full advantage of these aforementioned photophysical properties. We have strong interest in harvesting these properties within supramolecular and material chemistries.^{20–22}

Recently we have developed 2,3-bis(1,2,3-triazol-4-yl)picolinamide (**btp**) derivatives for use in various supramolecular chemistry applications.²³ This includes their use in the formation of luminescent coordinating polymers and metallo-gels.^{24,25} Gratifyingly, the **btp** based ligands can sensitise the excited states of various lanthanides, including the ⁵D₄ state of Tb(III) (resulting in green emission) resulting in the formation of complexes possessing high Tb-centered quantum yield (up to 90%), and to a lesser extent, the ⁵D₀ state Eu(III), which results in red emission (but much lower quantum yield) in solution.²⁶ We also recently incorporated **btp** based lanthanide complexes into polymer matrixes in a non-covalent manner.²⁷ While this work shows promise, the ligand system is found to be susceptible to leaching. To overcome this issue, covalently tethering the **btp** ligand to the polymer matrix was considered, but we have shown in related work, that ligands and Ln-complexes made from naphthyl-dipicolinic amide (**dpa**)²⁸ ligands, can be functionalised in two different ways to allow for such incorporation into polymers, and that this modification influences their photophysical properties within the polymers. Consequently, we set out to develop the **btp** monomer **2**, Fig. 1, from amine **1**, as well as forming **3**, as a model complex for studying the solution properties of the ligand and the corresponding Ln-complexes.

In the case of **2**, the methacrylamide unit would facilitate the co-polymerisation of the **btp** ligand into polymeric hydrogels. This modification would (i) allow for the generation of luminescent responsive systems, and (ii) facilitate the generation of lanthanide cross-linked hydrogels, the functionalization at the 4th position of the pyridine facilitating 1:3 (metal to ligand) complex formation. This second point is very important, as due to the high-coordination requirement of the lanthanides, we would expect three **btp** ligands to coordinate to a single Ln-ion, thus resulting in the potential to modulate both the 'hard' and 'soft' nature of the materials. At the same time, the luminescent properties of the lanthanides, would

furnish them with rich spectroscopic properties (*i.e.* point *i*). Here, we give full account of this work.

Results and discussion

Synthesis and characterisation of 1–3 and polymer films F1 and F2

The synthesis of the monomer **2** was achieved from the amine **1**, which was formed in a seven-step synthetic procedure (see ESI†), starting from the commercial starting material 2,6-dibromopyridine, using synthetic procedures developed in our laboratory.^{24–27} ‡ All the intermediates were fully characterised by NMR spectroscopies, IR spectroscopy and HRMS. The final two steps in the formation of **1** involved the use of click chemistry under CuAAC conditions using CuSO₄·5H₂O in a DMF:H₂O mixture, followed by removal of a Boc protecting group to give the free amine **1**, being formed in a moderate yield of 52%, as a brown precipitate, and TFA salt (The structure of **1** was further confirmed by X-ray crystallography as will be discussed further below).

The **btp** monomer **2** was formed by reacting the amine **1** with methacryloyl chloride for three hours in the presence of triethylamine. After concentrating the reaction mixture, the crude product was extracted into CHCl₃, and purified by column chromatography to yield **2** as a white solid in 63% yield. Formation of the desired product was confirmed by ¹H NMR spectroscopy (see ESI† for **1**, **2** and **3**). Two singlet peaks at 5.68 and 5.34 ppm correspond to the alkyne protons, and a peak at 1.85 ppm assigned to the CH₃ of the methacrylate moiety, confirm the presence of monomer **2**. As for **1**, compound **2** was fully characterised as outlined in the Experimental section.

As our main objective was to develop a lanthanide luminescent material, it was essential to photophysically characterise the resulting material, and compare its photophysical properties with a reference complex. As compound **1** has a primary amine at the backbone of this ligand, we anticipated that the luminescent properties of the neutral ligand might be compromised by an active photoinduced electron transfer (PET) from the amine to the excited state of the ligand, resulting in so called PET quenching.²⁹ Because of this it could not be used as a model compound for these studies, and hence, we synthesised compound **3**. The acetamide derivative **3** is not PET active, as the amide increases the oxidation potential of the 'previous amine', which prevents any PET activity. This structure also provided us with a 'structural amide model compound' to investigate the effect of the covalent attachment of **2** to a polymeric backbone, and for studying the photophysical behaviour of this system upon cross-linking by lanthanide ions,

‡ One of the intermediates from this synthesis, 2,6-dibromo-4-nitropyridine, crystallised during the workup and the resulting crystals were found to be suitable for X-ray crystallographic analysis. The resulting structure showed the presence of extended intermolecular halogen bonding interactions between the two bromine atoms of the pyridine and that of and the pyridine nitrogen, as well as the nitro group. These interactions are presented in the ESI.†

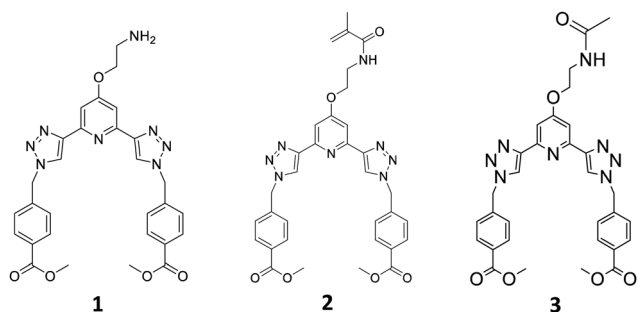


Fig. 1 The three main ligands **1**(as free base)–**3** developed for this project.





Scheme 1 Synthesis and structure of monomer **2**, formed from ligand **1**, and the two polymeric films **F1** and **F2**. Reagents and conditions: (i) NEt_3 , CH_2Cl_2 (ii) 80°C 6 hours. **F1** (0.03 wt%) and **F2** (0.3 wt%), respectively. Pictures show the films formed before and after being hydrated.

such as $\text{Tb}(\text{III})$. To achieve this, the primary amine of **1** was simply protected with an acetyl group, by reacting **1** with acetic anhydride in the presence of triethylamine. This generated **3** as a white solid in 83% yield, after workup; **3** being fully characterised by NMR and IR spectroscopy and HRMS (see Experimental).

The general synthesis of the polymeric films **F1** and **F2** is shown in Scheme 1. This simply involved stirring **2** in a solution of 2-hydroxyethyl methacrylate (HEMA), methylmethacrylate (MMA) and ethylene glycol dimethacrylate (EGDMA,) at room temperature until **2** was completely dissolved (0.3 and 3 mg for **F1** and **F2**, respectively). AIBN was then added and the clear solution was injected into a non-stick mould (see Fig. in ESI[†]). It was heated in the oven at 80°C for 6 hours and after cooling to room temperature the material was released from the mould and cut into sections. The material was soaked in H_2O overnight, followed by CH_3OH overnight to remove any remaining inhibitor or unreacted monomer. As seen in Scheme 1, the obtained film was stiff (and could easily be broken) but upon hydration gave rise to an easily handled soft, flexible film. To ensure that the film remained soft, the films were stored in a deionised aqueous solution. Having these three **btp** systems at hand as well as **F1** and **F2**, we started exploring their physical and photophysical properties, commencing with **3** and its $\text{Tb}(\text{III})$ complex **3.Tb**.

Synthesis and characterisation of the $\text{Tb}(\text{III})$ complex **3.Tb**

As mentioned above, we have shown that **btp** ligands upon complexation with lanthanide ions, give rise to up to 30-fold higher quantum yields for complexes formed from $\text{Tb}(\text{III})$ than $\text{Eu}(\text{III})$.^{26a} With this in mind, we commenced our work with exploring the photophysical properties of $\text{Tb}(\text{III})$, and the $\text{Tb}(\text{III})$ complex of **3**, **3.Tb**, was formed, structurally characterised and photophysically analysed. This was achieved by reacting **3** with $\text{Tb}(\text{CF}_3\text{SO}_3)_3$ in a 1 : 3 M : L stoichiometric ratio in CH_3OH under microwave irradiation for 15 min at 70°C . Thereafter, the solvent was concentrated *in vacuo*, and the desired product isolated by vapour diffusion of diethyl ether into the reaction mixture, to yield a white solid in 79% yield.

The complex was characterised by ^1H NMR spectroscopy, IR spectroscopy and HRMS. The ^1H NMR spectrum (400 MHz, $\text{CD}_3\text{CN}-d_3$) can be seen in Fig. 2a, and shows both broadening and significant resonance shifts when compared to the free



Fig. 2 (a) The ^1H NMR spectrum (400 MHz, $\text{CD}_3\text{CN}-d_3$) of $[\text{Tb}(\mathbf{3})_3](\text{CF}_3\text{SO}_3)_3$ showing significant shift and broadening of the signals relative to the free ligand **3** (see ESI[†] for NMR). (b) The bright green $\text{Tb}(\text{III})$ emission arising from the **3.Tb** complex in CH_3CN under UV-light ($\lambda_{\text{ex}} = 254$ nm).

ligand **3**, due to the paramagnetic nature of the $\text{Tb}(\text{III})$ ion. Elemental analysis was used to confirm the formation of this tris complex, as well as HRMS analysis, though only the characteristic isotopic distribution pattern for the $[\text{Tb}(\mathbf{3})_3(\text{CF}_3\text{SO}_3)_2]^+$ complex was observed in the MALDI HRMS with a $m/z = 1677.2877$ (see ESI[†]). The photophysical investigation of this ligand (as well as that of the methacryloyl derivative **2**) and the resulting $\text{Tb}(\text{III})$ complex is outlined in next two sections.

Photophysical evaluations of the $\text{Tb}(\text{III})$ complex **3.Tb**

Having successfully synthesised **3.Tb** we next investigated the photophysical properties of the complex in CH_3CN , and in MeOH , some key results are shown in Fig. 3. In CH_3CN , the UV-visible absorption spectrum of **3.Tb** was dominated by a high-energy band at $\lambda_{\text{max}} = 236$ nm and a lower energy band at $\lambda = 298$ nm, with a shoulder at $\lambda = 309$ nm (Fig. 3a). Excitation of the complex at $\lambda = 240$ nm gave rise to $\text{Tb}(\text{III})$ -centred emission with characteristic line-like transitions at $\lambda = 490, 545, 585, 622, 648, 667$ and 678 nm, Fig. 3b. These transitions are indicative of $\text{Tb}(\text{III})$ -sensitisation by the ligand antenna **3**, and can be assigned to the $^5\text{D}_4 \rightarrow ^7\text{F}_j$ transitions ($J = 6-0$). The complex was found to be stable over 48 hours with no loss of luminescence.

The excitation spectrum, recorded for the $\text{Tb}(\text{III})$ -centred luminescence with $\lambda_{\text{em}} = 545$ nm, also provided evidence for the sensitisation of the $\text{Tb}(\text{III})$ ion from the ligand-centred excited states. The structural features of the excitation spectrum closely resemble the absorption spectrum of **3**, which indicates indirect excitation of the metal centre from the antenna moiety (Fig. 2a). Furthermore, green emission was also observed from both the isolated solid complex, and from the complex dissolved in solution, such as CH_3CN , upon





Fig. 3 (a) The UV-vis absorption (red) and excitation spectra ($\lambda_{\text{em}} = 545$ nm) of the **btp** complex **3.Tb** in CH_3CN showing sensitisation arising from ligand-centred energy transfer (left), and time-gated emission spectrum of **3.Tb** showing the characteristic $\text{Tb}(\text{III})$ centred-emission.

irradiation with UV light ($\lambda_{\text{exc}} = 254$ nm), as demonstrated in Fig. 2b, which was clearly visible to the naked eye.

We also evaluated the hydration state (the so-called q value) of the complex, which indicates the number of associated metal-coordinated solvent molecules, and indicated the coordination number of the complex.³⁰ This was achieved by measuring the $\text{Tb}(\text{III})$ excited state lifetimes in CH_3OH and CD_3OD , respectively. These lifetimes were best fitted to a monoexponential decay function, indicating the presence of a single luminescent species in both solvents, with $\tau = 1.88 \pm 0.1$ ms and 1.98 ± 0.2 ms, in CH_3OH and CD_3OD , respectively. From these measurements a q value of ≈ 0 was determined, demonstrating that the $\text{Tb}(\text{III})$ inner coordination sphere was fully saturated by the three terdentate **3** ligands, and that under thermodynamic control, **3.Tb** excited as 1:3 (metal to ligand) stoichiometry in solution.

The photoluminescence quantum yield (Φ_{Tot} , %) of **3.Tb** was also determined in CH_3CN as 72% which is very close to relative quantum yields reported for other such complexes as mentioned above.^{26a,31}

Photophysical evaluations of **3** in presence of $\text{Tb}(\text{III})$ *in situ*

Before investigating the photophysical properties of the **btp** Ln-cross-linked polymers **F1** and **F2**, it was necessary to probe the photophysical properties of the lanthanide complexes formed from **3** under kinetic control. This mimics the condition of the free ligands within the polymers upon their hydration with solutions of lanthanide ions. To achieve this, both the ground and the excited state properties were probed *in situ* upon titrating the ligand with $\text{Tb}(\text{III})$ triflate solutions in CH_3CN solution. From this we were able to determine both the stoichiometry and the binding constants for the formation of the $\text{Tb}(\text{III})$ complex(es) with **3**.^{32,33}

The UV-visible absorption spectrum of compound **3** consists of two main bands appearing at $\lambda_{\text{max}} = 231$ nm and $\lambda_{\text{max}} = 300$ nm (see ESI[†]), from which $\epsilon = 54487 \text{ cm}^{-1} \text{ M}^{-1}$ ($\lambda_{\text{abs}} = 231$ nm) was determined. Significant changes were observed in the UV-visible absorption spectrum upon addition of $\text{Tb}(\text{III})$ (see ESI[†]); the high energy band underwent a hypochromic shift, up to the addition of 0.15 equivalents of $\text{Tb}(\text{III})$, which was followed by a redshift to $\lambda = 236$ nm and a concomitant hyperchromic shift up to the addition of 0.3 equivalents of $\text{Tb}(\text{CF}_3\text{SO}_3)_3$, indicating the successful

formation of the aforementioned 1:3 stoichiometry (see ESI[†]). Three isosbestic points at 232, 247 and 286 nm were observed during these titrations. Subsequent additions of $\text{Tb}(\text{III})$ did not result in any further changes to the band at 236 nm. The lower energy band at $\lambda = 298$ nm also experienced a hyperchromic shift up to the addition of 0.3 equivalents of $\text{Tb}(\text{III})$.

Upon excitation of the ligand **3** at $\lambda = 237$ nm, ligand centred fluorescence emission was observed, with a band centred at $\lambda = 319$ nm (see ESI[†]). This ligand luminescence was affected by the self-assembly process, which was evident by quenching of the ligand fluorescence upon addition of $\text{Tb}(\text{III})$. Upon addition of 0.3 equivalents of $\text{Tb}(\text{III})$, a 96% reduction in the emission was observed, which can be attributed to the energy transfer, and hence, sensitisation of the $^5\text{D}_4$ excited state of $\text{Tb}(\text{III})$ by the ligand **3**.

The delayed $\text{Tb}(\text{III})$ -centred luminescence was also recorded upon excitation at 237 nm, reflecting the appearance of the $\text{Tb}(\text{III})$ emission due to the effective sensitisation by the **btp** ligand, Fig. 4a. A gradual enhancement in the $\text{Tb}(\text{III})$ luminescence was observed from 0 \rightarrow 0.3 equivalents of $\text{Tb}(\text{III})$, with characteristic $\text{Tb}(\text{III})$ -centred emission bands appearing at $\lambda = 490, 545, 583, 620, 646, 665,$ and 675 nm, corresponding to the deactivation of $^5\text{D}_4 \rightarrow ^7\text{F}_j$ states ($J = 6-0$). The $^5\text{D}_4 \rightarrow ^7\text{F}_{2-0}$ bands are typically weak and not always seen in such titrations, but they can be clearly observed in the formation of **3.Tb** *in situ*. Subsequent additions of $\text{Tb}(\text{III})$, from 0.3 \rightarrow 3.0 equivalents, resulted in quenching of the luminescence intensity by approximately 20%.

As can be seen from Fig. 4a, The most emissive state in solution was seen at approximately 0.3 equivalents of $\text{Tb}(\text{III})$, which is consistent with previously reported **btp**- $\text{Tb}(\text{III})$ systems, as at this stoichiometry, the $\text{Tb}(\text{III})$ ion is coordinatively saturated, and thus completely shielded from any solvent quenching. Lifetime measurements obtained at the addition of 0.3 equivalents of $\text{Tb}(\text{III})$ gave $\tau = 1.85$ ms, which is very close to the lifetimes observed above for the thermodynamically formed **3.Tb**. These titrations demonstrate that the 1:3 stoichiometry is the most favourable one formed at low concentration and is quite stable to dissociation upon increasing $\text{Tb}(\text{III})$ concentration.

With the view of quantifying the above stability, the changes observed in the ground and the excited state titrations were

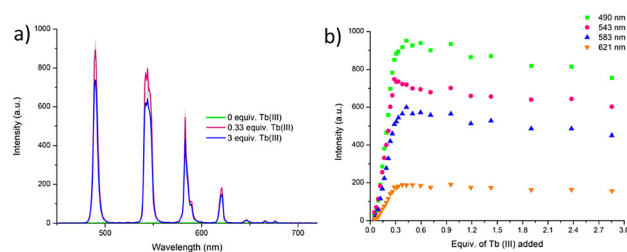


Fig. 4 (a) The overall changes to the $\text{Tb}(\text{III})$ -centred emission upon titrating **3** (1×10^{-5} M) against $\text{Tb}(\text{CF}_3\text{SO}_3)_3$ (0 \rightarrow 3 equiv.) in CH_3CN at 22 °C. (b) The corresponding experimental binding isotherms for the $\text{Tb}(\text{III})$ emission at $\lambda = 490, 545, 583$ and 621 nm. These results were fully reproducible (see enlarged version in ESI[†]).



fitted using non-linear regression analysis program ReactLab EQUILIBRIA[®]. Analysis of the UV-visible absorption data of **3** titrated with Tb(CF₃SO₃)₃ in CH₃CN solution suggests the presence of three main species in solution (**3**, 3 : 1 metal : ligand and 1 : 1 metal : ligand ratios) with the distribution of these species estimated from this analysis (see ESI[†]); with the ML₃ species being the most dominant in solution, reaching 97% abundance at 0.3 equivalents of Tb(III). The 1 : 3 metal : ligand assembly formed with $\log \beta_{1:3} = 23.1 \pm 0.6$, which is in good agreement with binding constants reported for similar **btp** systems in the literature with Tb(III).^{22,23} The second species (1 : 1 metal : ligand) had a calculated binding constant of $\log \beta_{1:1} = 8.3 \pm 0.2$, which again, is in good agreement with previously reported Tb-**btp** assemblies.[§]

Similar analyses of the changes in the Tb(III)-centred emission in Fig. 4a were also carried out and indicated the presence of four main species in solution, (**3**, 3 : 1 metal : ligand, 2 : 1 metal : ligand and 1 : 1 metal : ligand species). Again, at 0.3 equivalents of Tb(III), the Tb₃ species was the most predominant in solution, reaching an abundance of 96%, Fig. 4b. Upon subsequent additions of Tb(III), the 1 : 2 species was observed, which reaches 20% abundance at the addition of 1.5 equivalents of Tb(III) before reaching a plateau. The evolution of the 1 : 1 species occurred almost simultaneously with the 1 : 2 species; however, it is less predominant until the addition of 1.8 equivalents of Tb(III), after which point it continues to increase with a concomitant decreasing of the 1 : 3 species (see ESI[†]). From these luminescent changes, the binding constants were determined as $\log \beta_{1:3} = 22.7$, $\log \beta_{1:2} = 14.5 \pm 0.1$, and $\log \beta_{1:1} = 7.0$. The stability constants are roughly additive, suggesting that the binding process of each ligand to the metal centre was subject to a similar formation constant.

In addition to these measurements, the self-assembly of compound **2** with Tb(CF₃SO₃)₃ was also investigated in CH₃CN solution, in an identical manner to that described above for compound **3**. The changes to the UV-visible absorption, fluorescence, and time-gated emission spectra were very similar to those shown for compound **3** with Tb(III). The data was fitted using non-linear regression analysis, and four species were again estimated to be present in solution for the Tb(III) complex formation of **2**. The global analysis of these changes was found to be in good agreement with the analysis for compound **3** with Tb(III).

Supramolecular gelation studies of **1**

In the past, we have demonstrated that **btp** ligands possessing carboxylates at both the 4th position of the pyridine, as well as at the two aryl groups on the appended arms (which in the case

[§] The changes in the UV-visible absorption spectra were not large and the resulting speciation therefore may not be fully representative of the speciation in solution. No acceptable fit was obtained when the Tb₃ species was included in the UV-visible absorption data model. The reliability of this fit should therefore be considered with caution. However, the speciation obtained from the Tb(III)-centred emission spectral data, can be considered a more accurate representation of the likely speciation in solution. These titrations were fully reproducible as was the fitting of the data.

of **1–3**, are protected as methyl esters) are excellent LMWG, and give rise to the formation of transparent or opaque self-healing soft materials, which upon interactions/complexation with lanthanides, can result in the formation of luminescent supramolecular gels.²⁴ Because of this, compound **1** was investigated as well as a potential LMWG.

Ligand **1** was suspended in a variety of solvents; CH₃CN, CH₃OH, CH₃OH:H₂O, C₂H₅OH, C₂H₅OH:H₂O, and subject to different gelation conditions including heating and sonication. Of these, the suspension of **1** in CH₃CN, followed by sonication, resulted in the formation of an off-white soft material almost immediately. Following this result, the optimum concentration for gelation was determined. Compound **1** (16 mg) was suspended in CH₃CN (1 mL) and sonicated for 60 seconds to yield an opaque gel that was robust to the naked eye. The inversion test was performed, and the gel supported its own weight over several hours, Fig. 5c. The thermoreversibility of the gel was investigated by heating the material to up to 60 °C. Upon heating, the gel dissolved and, after standing at room temperature for 15 minutes, the gel reformed, indicating that the material was formed in a thermoreversible manner.

In order to investigate the LMWG wt% at which the gel forms, TGA was carried out on the material. TGA was carried out in the temperature range of 15 to 500 °C, and a mass loss of 96.2% was observed at 100 °C, indicating the gel (as shown in Fig. 5) was formed at 3.8 wt% of **1** (see ESI[†]). However, as the gel was prepared at 1.6 wt%, this experiment indicates that the gel does not retain all of the solvent in the sample. While inverting the gel upon formation did not result in any loss of residual solvent (*i.e.* the gel did not collapse, as shown in Fig. 5c).



Fig. 5 (a) Amplitude (strain) sweep experiment of **1** gel showing the linear viscoelastic region (γ_L) and confirming the behaviour of the material as a gel. (b) Amplitude (stress) sweep showing yield point (τ_y) and flow point (τ_f) of the material. (c) Image of the gel during inversion test. (d) Frequency sweep experiment showing the long-term stability of the gel.



In order to examine the mechanical properties of the gel, rheological experiments were conducted using 3.8 wt% of **1**, and an amplitude sweep was conducted in order to confirm the gel-like behaviour of the material. The strain on the sample was increased logarithmically, while the frequency was kept constant. In Fig. 5 it can be seen that the storage modulus, G' was larger than the loss modulus, G'' , which indicates that the material can be classified as a gel or a viscoelastic solid material ($G' = 1.09 \times 10^4$ Pa and $G'' = 1.46 \times 10^3$ Pa). The limit of the linear viscoelastic region (LVE, labelled γ_L in Fig. 5), the range in which further tests could be carried out on the sample without destruction of the sample, was determined to be 0.3% for this sample. After this point, G' and G'' begin to converge and the material starts to flow.

From the graph of G' and G'' vs. shear stress (τ), in Fig. 5, the yield point and flow point of the material can be determined. The yield point, τ_y , is value of shear stress at the limit of the LVE region. For this material its value was determined to be 28 Pa, and the flow point, τ_f , was observed to be 253 Pa. The rise in G'' to a maximum which begins to drop again prior to the cross-over point as seen in Fig. 5a and b suggests the breakdown of the gel microstructure prior to formation of macro cracks leading to material flow. A frequency sweep experiment was also conducted to gain an insight into the long-term stability of the gel. Having determined that the limit of the LVE region was 0.3%, the amplitude was set at 0.1%. As seen in Fig. 5 $G' > G''$ over the entire frequency range, which indicates that the material has a cross-linked structure. Rheological tests were repeated three times with three gel samples to ensure reproducibility.

A xerogel of compound **1** (as its salt form) in CH_3CN was also generated by drying the sample under vacuum in order to study the morphology of the system by SEM. The SEM images, Fig. 6a, show the fibrous, entangled nature of the gel structure. This morphology is typical of many supramolecular gels in which non-covalent interactions are involved in forming cross-links within the structure immobilising the solvent, such as hydrogen and ionic interactions. At higher wt%, Fig. 6b, a precipitate appeared on the fibres, seen in the SEM images as bright spots. These are very likely due to salt formation, however, the nature of these was not analysed further.

Having formed organogels from **1**, the possibility of forming a supramolecular metallogel with Tb(III) was next investigated. To do this, a solution of $\text{Tb}(\text{CF}_3\text{SO}_3)_3$ was added to the

pre-formed organogel, by layering a solution of known concentration of the ion on the top of the gel and allowing it to diffuse through the gel over period of time. However, unlike previous observations for **btp** based ligands (*cf.* above discussion) upon standing overnight, the material lost its soft-material properties and the gel dissolved, giving a solution, which was shown to emit green light under UV-irradiation ($\lambda_{\text{exc}} = 254$ nm) (see ESI†). We have previously seen this behaviour for **dpa** based gels.³² This strongly indicates the formation of a Tb(III) complex, which likely breaks-down the network of interactions which otherwise generate the supramolecular soft-material upon formation (*e.g.* such as hydrogen bonding and π - π stacking).

Crystallographic characterisation of $[\text{H}_1\text{1}^+][\text{CF}_3\text{SO}_3^-] \cdot \text{MeCN}$

As outlined above, the gelation studies were carried out at different wt% of **1**. When the organogel was formed at low, or less than 1.5 wt%, a gel-to-sol transition occurred overnight, and yellow crystals were formed in a colourless solution. These crystals were examined by X-ray diffraction and were found to be crystals of the TFA salt of compound **1**, $[\text{H}_1\text{1}^+][\text{CF}_3\text{SO}_3^-] \cdot \text{MeCN}$, as illustrated in Fig. 7 (see also ESI†).³⁴

The structure was solved in the monoclinic $C2/c$ space group, with the asymmetric unit containing one complete protonated molecule of **1**, as well as one counterbalancing TFA anion, and a solvent molecule of CH_3CN in the lattice. The primary intermolecular interactions in the structure of **1** are hydrogen bonds, involving the oxygen atom of the TFA anion, **O6** and the NH_3^+ of compound **1**, with an $\text{N} \cdots \text{O}$ distance of 2.793(5) Å and an $\text{N-H} \cdots \text{O}$ angle of 175.3(3)°. Unexpectedly, the triazoles in ligand **1** adopt a *syn-syn* conformation. Previous **btp** ligands have been most commonly reported in the *anti-anti* conformation, but in the presence of a cation such as a metal ion or a proton, the triazoles can adopt the *syn-syn* conformation.^{23,35,36} In this case, the RNH_3^+ of one ligand molecule, and the triazole N14 and N31 nitrogen atoms of the adjacent ligand molecule, form a hydrogen bonding interaction with an $\text{N} \cdots \text{N}$ distance of 2.998(6) and 2.885(6) Å, respectively, which causes the triazoles to adopt the *syn-syn* conformation.

Crosslinking polymers **F1** and **F2** with Ln(III) ions

Having confirmed the incorporation of **2** in the polymer matrix, the ability of the hard dehydrated polymer to uptake Ln(III) ions

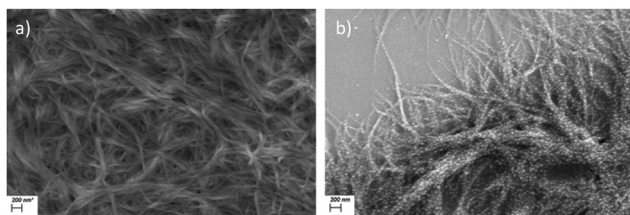


Fig. 6 (a) SEM images of **1** gel (scale bar 200 nm) showing fibrous structure of the material. (b) The formation of salt particles on the surface of the fibres upon gelation at higher concentrations.

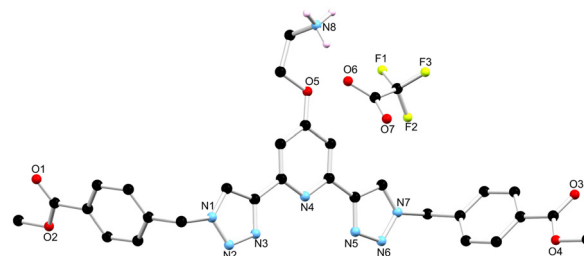


Fig. 7 The solid state X-ray structure of compound **1** with heteroatom labelling scheme; lattice acetonitrile molecule and selected hydrogen atoms omitted for clarity.



was investigated. Initially, the dry polymer **F2** (which had a higher concentration of the **btp** ligand) was soaked in an aqueous solution of $\text{Tb}(\text{CF}_3\text{SO}_3)_3$ and after 10 minutes the resulting soft, flexible polymer was rinsed with water and examined under UV-irradiation at $\lambda_{\text{exc}} = 254 \text{ nm}$, in water at room temperature. The film exhibited bright green emission under UV-light, which was clearly visible to the naked eye, indicating that crosslinking had occurred with $\text{Tb}(\text{III})$ and that the covalently bound **btp** ligand was sensitising the $\text{Tb}(\text{III})$ excited within the polymer film, as is evident from the Fig. 8a. The changes in the $\text{Tb}(\text{III})$ centred emission for **F1** was similarly enhanced upon soaking in $\text{Tb}(\text{III})$ solution over a longer period of time, with all the expected transitions being clearly visible.

Examining **F2** further, showed that upon excitation of the film (on a quartz slide) at $\lambda = 285 \text{ nm}$ in water, the delayed $\text{Tb}(\text{III})$ -centred luminescence could be recorded in water and the spectrum can be seen in Fig. 8a. Characteristic $\text{Tb}(\text{III})$ -centred luminescence was evident with $\text{Tb}(\text{III})$ -centred transitions at $\lambda = 490, 545, 583, 620, 646, 665$ and 675 nm , corresponding to deactivation from the $^5\text{D}_4$ excited states to the $^7\text{F}_j$ states (where $J = 6-0$). **F2** was found to be stable in water; with no evidence of changes in the intensity of the $\text{Tb}(\text{III})$ -luminescence spectra seen over several hours. This suggests that the $\text{Tb}(\text{III})$ ions underwent diffusion through the polymer film, and did not simply attach to the surface of the film. Thus, it is likely that $\text{Tb}(\text{III})$ mediated self-assembly occurs within the polymer matrix, leading to the green luminescence observed, and that the resulting cross-linked **btp-Tb** complexes within the film are stable to dissociation under such competitive media. This is a very important observation that demonstrates that the grafting of **2** to the backbone of the polymer indeed adds stability to the **btp-Ln** interaction, which we have not previously seen when **btp-Tb** complexes were incorporated into such matrixes in a non-covalent manner.²²

As was mentioned above, the emission of the $\text{Eu}(\text{III})$ is significantly less than for the $\text{Tb}(\text{III})$ complexes for the **btp**

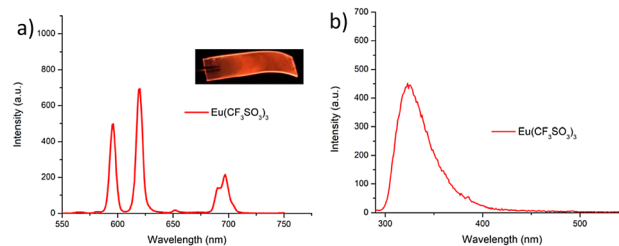


Fig. 9 (a) The $\text{Eu}(\text{III})$ -centred emission observed for **F2** when measured in H_2O . (b) The corresponding fluorescence spectrum of **F2** measured in H_2O . Inset (a) photograph of polymer following soaking for 10 min in $\text{Eu}(\text{CF}_3\text{SO}_3)_3$ solution under UV-irradiation showing the characteristic red luminescence from $\text{Eu}(\text{III})$.

based ligands. Nevertheless, to demonstrate that other lanthanides such as $\text{Eu}(\text{III})$ could also be used in the formation of luminescent cross-linked polymers was also briefly investigated. As for the $\text{Tb}(\text{III})$ above, the polymer film **F2** was soaked in a $\text{Eu}(\text{CF}_3\text{SO}_3)_3$ solution for 10 min at room temperature and the spectra were recorded in H_2O resulting in the characteristic red emission of $\text{Eu}(\text{III})$ under UV-irradiation, which again, was clearly visible to the naked eye, as shown in Fig. 9a, and demonstrates that $\text{Eu}(\text{III})$ can be equally employed in such luminescent material, even though it has (as we have demonstrated in solution on many occasions) much lower Ln-centred quantum yields. As for $\text{Tb}(\text{III})$, the delayed $\text{Eu}(\text{III})$ -centred luminescence was also recorded upon excitation at $\lambda = 285 \text{ nm}$ and the spectrum is shown in Fig. 9a, for **F2**, $\text{Eu}(\text{III})$ -centred transitions appeared at $\lambda = 595, 620, 652$ and 695 nm corresponding to the $^5\text{D}_0 \rightarrow ^7\text{F}_j$ ($J = 0-3$) states characteristic of $\text{Eu}(\text{III})$ emission. The fluorescence emission is shown in Fig. 9b, mirroring that seen for $\text{Tb}(\text{III})$ above in Fig. 8b for $\text{Tb}(\text{III})$.

Conclusion

Herein, we have developed the **btp** ligands **1**, **2**, and **3** as coordinating ligands for metal ions, and explored the



Fig. 8 (a) The $\text{Tb}(\text{III})$ -centred emission of **F2** when measured in H_2O after having soaked the polymeric film in a solution of $\text{Tb}(\text{III})$ triflate at different times (given in minutes). Inset (a): Photograph of **F2** film following soaked in $\text{Tb}(\text{CF}_3\text{SO}_3)_3$ under UV-irradiation ($\lambda_{\text{exc}} = 245 \text{ nm}$), showing the characteristic green $\text{Tb}(\text{III})$ luminescence visible to the naked eye. (b) The fluorescence emission spectrum of **F2** in H_2O at different times.



luminescent properties of these ligands upon formation of Tb(III) complexes both in solution and as, in the case of **2**, upon co-polymerising the ligand with HEMA, MMA and EGDMA. The model compound **3** was employed to form a Tb(III) complex, which was formed in 1 : 3 (M : L) stoichiometry. This complex was luminescent, possessing high quantum yield of luminescence, and when formed *in situ* was produced as the main species in solution at 96%, upon addition of 0.3 equivalents of Tb(III). We also demonstrated that a supramolecular gel with fibrous morphology, can be produced from the precursor **1**, which upon interactions with Tb(III) dissolves, as weak supramolecular interactions that produce the gel are replaced with stronger coordination bonds.

The **btp** based polymers **F1** and **F2**, were formed and both were shown to be luminescent, and metal ion mediated cross-linked, upon interaction with Tb(III) (and Eu(III) in the case of **F2**). We further demonstrated that unlike the incorporation of **btp**-based Tb(III) complexes non-covalently into such polymers (such as into a blank HEMA-co-MMA-co-EGDMA polymer), no leakages were observed from the polymeric film upon prolonged soaking in water. Moreover, the Tb(III) centred emission was shown to be stable over a period of time. Furthermore, we show that in competitive media, *e.g.* water, the polymer is highly luminescent indicating that no dissociation occurs in the cross-linked **btp**-based Tb(III) complexes within the hydrogel. This is highly important, as it demonstrates significant improvement on the use of non-covalently used **btp**-based Tb(III) complexes, which indeed showed rapid dissociation and subsequent loss of metal centred luminescence.

The lanthanide cross-linked polymers offer a desirable means of forming functional polymeric systems, where both conventional covalent and coordination metal ion bonds are employed, and we are currently investigating the advantage of such features. This includes the use of ligands that have conjugated receptor sites within the ligand structures of these and related systems, with the view of forming stimulated responding polymers that can be used in luminescent sensing.

Experimental

Synthesis

Please refer to Scheme S1 in ESI.†

Dimethyl 4,4'-(((4-(2-aminoethoxy)pyridine-2,6-diyl)bis(1H-1,2,3-triazole-4,1-diyl))bis(methylene)dibenzoate (1). Ligand **10** (see ESI†) was deprotected by stirring in neat TFA at room temperature for 30 min. The resulting solution was neutralised with Na₂CO₃ and the product was filtered to yield a brown solid (0.493 g, 0.87 mmol, 80%). The product decomposes over 182 °C. HRMS (*m/z*) (ESI+): calculated for C₂₉H₂₉N₈O₅H⁺ *m/z* = 569.2261 [M + H]⁺. Found *m/z* = 569.2266; ¹H NMR (400 MHz, DMSO): δ = 8.70 (s, 2H, triazole H), 7.98 (d, *J* = 8.2 Hz, 4H, Ar H-COOME), 7.54 (s, 2H, pyridine H), 7.46 (d, *J* = 8.2 Hz, 4H, Ar H-CH₂), 5.81 (s, 4H, CH₂), 4.31 (t, *J* = 4.3 Hz, 2H, O-CH₂), 3.84 (s, 6H, OCH₃), 3.15 (q, *J* = 4.3 Hz 2H, N-CH₂), ¹³C NMR (150 MHz, DMSO): δ = 165.8, 151.5, 147.2, 141.1, 129.7, 129.4, 128.5, 128.1, 124.1, 104.9, 64.7, 52.6, 52.2, 38.2; IR

*ν*_{max} (cm⁻¹): 3112, 3088, 3086, 2955, 2478, 2203, 2091, 1933, 1865, 1796, 1714, 1685, 1610, 1281, 1174, 1041, 801.

Dimethyl 4,4'-(((4-(2-methacrylamidoethoxy)pyridine-2,6-diyl)bis(1H-1,2,3-triazole-4,1-diyl))bis(methylene)dibenzoate (2). Ligand **1** (0.07 g, 1.23 mmol) was dissolved in a solution of CH₂Cl₂ (3 mL) with NEt₃ (0.017 mL, 1.29 mmol) in an ice bath. To this solution, methacryloyl chloride (0.012 mL, 1.29 mmol) was added dropwise in CH₂Cl₂ (5 mL). The mixture was stirred for 30 min on ice and then allowed to stir at room temperature for two hours. The solution was then diluted in CH₂Cl₂, washed with dilute NaHCO₃ and H₂O and then dried over MgSO₄. The solution was dried under reduced pressure at 30 °C to yield colourless oil. The compound was purified by flash chromatography (CH₂Cl₂:CH₃OH gradient up to 5% CH₃OH) to yield a white solid (0.071 g, 64%); m.p. 148–154 °C; HRMS (*m/z*) (ESI+): calculated for C₃₃H₃₂N₈O₆H⁺ *m/z* = 637.2523 [M + H]⁺. Found *m/z* = 637.2541. ¹H NMR (400 MHz, DMSO): δ = 8.69 (s, 2H, triazole H), 8.17 (s, *J* = 5.7 Hz, 1H, NH), 7.98 (d, *J* = 8.4 Hz, 4H, Ar H-COOME), 7.51 (s, 2H, pyridine H), 7.45 (d, *J* = 8.4 Hz, 4H, Ar H-CH₂), 5.80 (s, 4H, CH₂), 5.68 (s, 1H, alkene H), 5.34 (s, 1H, alkene H), 4.29 (s, *J* = 11.3, 2H, O-CH₂), 3.84 (s, 3H, 2CH₃), 3.56 (q, *J* = 11.3, 5.7 Hz, 2H, CH₂-NH), 1.85 (s, 3H, CH₃). ¹³C NMR (150 MHz, DMSO): δ = 151.9, 147.7, 141.6, 140.2, 130.2, 129.9, 128.5, 124.6, 119.8, 105.3, 66.9, 53.1, 52.7, 38.8, 19.1; IR *ν*_{max} (cm⁻¹): 3434, 3289, 2514, 2094, 1711, 1612, 1588, 1493, 1279, 1203, 1184, 1010, 1000, 799, 726, 563.

Dimethyl 4,4'-(((4-(2-acetamidoethoxy)pyridine-2,6-diyl)bis(1H-1,2,3-triazole-4,1-diyl))bis(methylene)dibenzoate (3). Ligand **1** (0.13 g, 0.02 mmol) was dissolved in dry CH₂Cl₂ and triethylamine (0.01 mL, 0.07 mmol) was added. Acetic anhydride (0.003 mL, 0.03 mmol) was added to the mixture, which was cooled over ice and the reaction was stirred at room temperature overnight. The solvent was removed under reduced pressure and the resulting solid was taken up in EtOAc and washed with brine. The solution was dried over MgSO₄ and the solvent was removed under reduced pressure to yield a brown solid. (0.077 g, 0.01 mmol, 55%). m.p. 155–158 °C; HRMS (*m/z*) (ESI+): calculated for C₃₁H₃₀N₈O₆H⁺ *m/z* = 611.2367 [M + H]⁺. Found *m/z* = 611.2370. ¹H NMR (400 MHz, DMSO): δ = 8.69 (s, 2H, triazole H), 8.13 (s, 1H, NH), 7.98 (d, *J* = 8.3 Hz, 4H, Ar H-COOME), 7.50 (s, 2H, pyridine H), 7.46 (d, *J* = 8.3 Hz, 4H, Ar H-CH₂), 5.80 (s, 4H, CH₂), 4.23 (s, 2H, O-CH₂), 3.48 (d, *J* = 5.4 Hz, 2H, CH₂-NH), 1.83 (s, 3H, CH₃). ¹³C NMR (150 MHz, DMSO): δ = 169.6, 165.9, 135.9, 151.4, 147.3, 141.2, 129.7, 129.4, 128.1, 124.1, 104.9, 66.9, 52.6, 52.3, 38.0, 22.6; IR *ν*_{max} (cm⁻¹): 3292, 3082, 2954, 1714, 1643, 1609, 1517, 1548, 1462, 1433, 1381, 1277, 1227, 1177, 1106, 1044, 1015, 982.

Synthesis of Tb.3₃. Ligand **3** (0.01 g, 0.016 mmol) was added to Tb(CF₃SO₃)₃ (0.004 g, 0.0055 mmol) in CH₃OH (5 mL) and heated at 70 °C under microwave irradiation for 15 minutes. The resultant solution was dried under vacuum to yield colourless oil that was taken up in CH₃OH and dropped slowly into a large excess of swirling Et₂O to afford Tb.3₃ as a white solid in 79% yield. m.p. (decomp) > 280 °C; Elemental analysis for C₉₃H₉₃N₂₄O₃₆Tb·3H₂O Calculated: C 46.27 H 3.88 N 13.48; Found: C 46.19 H 3.62 N 14.02%; HRMS (*m/z*) (ESI+): C₆₄H₆₀N₁₆O₁₈F₆S₂Tb *m/z* = 1677.2871. Found *m/z* = 1677.2877; IR *ν*_{max} (cm⁻¹): 3292, 2955, 1735, 1630, 1609, 1580, 1381, 1277, 1245, 1106, 982, 939, 744, 668.



Synthesis of F2. Polymer gels were generated by a modified procedure previously reported by McCoy and co-workers.²² Ligand 2 (3 mg) was stirred in 2-hydroxyethyl methacrylate (HEMA, 7.5 mL), methylmethacrylate (MMA, 0.25 mL) and ethylene glycol dimethacrylate (EGDMA, 0.1 mL) at RT until compound 2 was completely dissolved. AIBN (100 mg) was added and the clear solution was injected into a non-stick mould. It was heated in the oven at 80 °C for 6 hours and after cooling to room temperature the material was released from the mould and cut into sections. The material was soaked in H₂O overnight followed by CH₃OH overnight to remove any remaining inhibitor or unreacted monomer.

Synthesis of F1. Same as for F2, using: 2 (0.3 mg).

General experimentals and methods

Photophysical measurements. All photophysical measurements were taken using spectroscopic grade solvent and measured in quartz cells with path length 10 mm. UV-visible absorption spectra were recorded using a Varian Cary 50 spectrophotometer in a spectroscopic window of 400–200 nm. The blank used was a sample of the solvent in which the titration was carried out. Luminescence spectra were measured using a Varian Cary Eclipse spectrometer and reported in arbitrary units and the luminescence data was collected between 450 and 700 nm for Tb(III) emission and 450 to 720 nm for Eu(III) emission. Time-gated emission spectra were recorded over an average integration time of 0.1 seconds (delay time: 0.1000 ms; gate time: 5.000 ms). Luminescence lifetime measurements of Tb(III)-centered emission was recorded using Varian Cary Eclipse spectrophotometer as a time-resolved measurements at 298 K. All lifetimes were averaged from at least five measurements at different gate times between 0.02–0.045 ms. The decay curves were fitted to mono- or bi-exponential decay functions.

SEM studies. Microscopy analysis of samples by Scanning Electron Microscopy (SEM) was carried out using the facilities of the Advanced Microscopy Laboratory (AML) in Trinity College Dublin. Samples were prepared by drop-casting the sample onto clean silicon wafers. The manually drop cast samples were dried overnight in ambient conditions and under high vacuum for at least 2 hours prior immediately to their imaging. In some cases, samples were coated with a conductive Pd/Au layer using a Cressington 208Hr high-resolution sputter coater, to improve contrast where static charging interfered with the imaging. Low kV SEM was carried out using the Zeiss ULTRA Plus using an SE2 detector.

Rheological studies. All the rheological measurements were carried out on 1.0 mL gel samples prepared 12 h before the measurement, using an Anton Paar Rheometer: MCR 302 equipped with serrated parallel plates with a diameter of 25 mm at a gap size of 1 mm and a temperature of 20 °C. Excess gel was trimmed just before the final gap height was reached. To avoid evaporation of solvent during the measurements a solvent trap was used and refilled with solvent if necessary. The strain sweep measurements were performed at a frequency of 10 rad s⁻¹ and a strain rate of 0.01–100%. The frequency sweep

measurements were performed at a constant strain rate of 0.1% and a frequency range of 100–0.1 rad s⁻¹.

Conflicts of interest

There are no conflicts to declare.

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