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Pyridine coupled mono and bisbenzimidazoles as supramolecular gelators: selective metal ion sensing and ionic conductivity†

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Pyridine coupled mono and bisbenzimidazoles **1–6** are synthesized and their gelation properties are examined in various polar as well as nonpolar solvents. Among the structures, compounds **1** and **2** form instant gels from DMSO/H₂O and MeOH/H₂O solvents. The gel states of both **1** and **2** are responsive to Ag⁺ and Cu²⁺ ions whereas the gel to sol transition of **2** in DMSO–H₂O is additionally caused by Hg²⁺ ions. On the other hand, compounds **3** and **4** produce gels only in the presence of Ag⁺ ions under similar conditions and validate their visual readout. In relation to this, the nongelation behavior of *p*-isomers **5** and **6** (structural isomers) under identical conditions emphasizes the positional role of the pyridyl ring nitrogen with respect to the imidazole ring. Furthermore, metallogels of **3** and **4** exhibit thermally activated ionic conductivity due to movement of Ag⁺ ions within the gel network.

Introduction

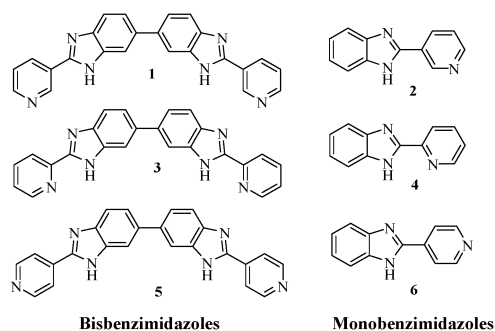
Molecular architectures that are capable of forming dimensionally controlled aggregates in solution *via* non-covalent interactions (hydrogen bonds, π – π stacking, electrostatic and van der Waals interactions, metal coordination, *etc.*) draw attention in the area of supramolecular chemistry.^{1–6} Emphasis has been given to such a special class of compounds not only to correlate the fundamental processes of gelation with structural diversities^{3,5,7–9} but also to develop new materials with wide applications in biomedicine, drug delivery, reaction catalysis, tissue engineering, water purifiers, photophysics, *etc.*^{10–14} In addition, gelators that exhibit a gel to sol transition in the presence of different external stimuli such as light, redox, pH, ions, *etc.*, have gained attention with a view to development of new materials.^{4,15–19} In this endeavor, metal and anion-induced phase transitions of gelators are worth mentioning. Many research groups in this field have shown interest especially to develop metal responsive supramolecular gels owing to their wide applications in catalysis, sensing, nanoparticle synthesis, optics and magnetic fields, *etc.*^{18–20} In this regard, a number of organogelators are known which contain pyridine-based ureas and amides in the structural backbones as they form directed and strong hydrogen bonds.^{21–26} In contrast, pyridine coupled

benzimidazoles that are capable of forming self-aggregates are less reported in the literature^{27–29} and thus there is a surge of interest to examine these compounds in gelation with possible applications.

As part of our ongoing research on the development of stimuli responsive supramolecular gelators,^{30–33} herein we report metal responsive behaviours of some pyridine coupled mono and bisbenzimidazoles **1–6** (Fig. 1).

Importantly, the synthesis of compounds **1–6** and DNA binding studies of some of them are reported by Neidle *et al.* and other research groups.^{34–36} But no attempt on examining their gelation characteristics, which is the prime focus of this study, has been made yet.

Analysis of structures **1–6** reveals that the compounds are different due to the change in the position of the pyridine ring nitrogens. This change is deeply associated with their different gelation behaviours. As the gels are formed, they behave

Fig. 1 Structures of compounds **1–6**.^a Department of Chemistry, University of Kalyani, Kalyani-741235, India.

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^b Department of Physics, University of Kalyani, Kalyani-741235, India† Electronic supplementary information (ESI) available: Gelation results, emission, absorption, FTIR and ¹H NMR spectra, Job plots, binding curves, detection limit and copies of ¹H, ¹³C NMR and HRMS. See DOI: 10.1039/c7qm00505a

differently to metal ions. While the gel states of **1** and **2** are converted to sols in the presence of Ag^+ and Cu^{2+} ions, the gel state of **2** additionally exhibits strong affinity for Hg^{2+} ions. The structural isomers **3** and **4**, in comparison, exhibit gelation selectively in the presence of Ag^+ ions under similar conditions. Along this direction, the nongelation behavior of **5** and **6** (*p*-derivatives) under identical conditions emphasized the positional role of the pyridyl ring nitrogen with respect to the imidazole ring.

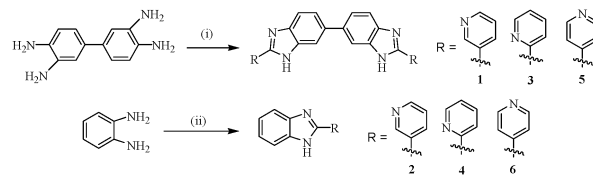
Compounds **1–4** in solution also showed measurable interaction with the said metal ions as verified by fluorescence. Fluorescence sensing of metal ions draws attention due to their environmental and physiological significance.³⁷ Of various metal ions, Ag^+ , Cu^{2+} and Hg^{2+} ions are of considerable importance. An Ag^+ ion is associated with bioaccumulation and toxicity.^{38,39} Similarly, a Cu^{2+} ion is one of the essential elements in our life and is linked with a variety of fundamental physiological processes in organisms.^{40,41} It serves as a catalytic cofactor for a variety of metalloenzymes like superoxide dismutase, tyrosinase and cytochrome oxidase. Exposure to high levels of copper can also cause gastrointestinal disturbance and liver or kidney damage. This metal ion also causes environmental pollution.⁴¹ Mercury is another important analyte because of its toxicity to the ecosystem and damage to health,⁴² as demonstrated by Minamata disease.⁴³ Mercury exposure can have a number of severe adverse effects on health, such as brain damage, kidney failure, and various cognitive and motion disorders.⁴⁴ Therefore, selective recognition of these metal ions is of utmost interest. In spite of reasonable progress in the detection/sensing of these ions in solutions, the use of organogelators for the same is less explored and thus requires attention.

In addition to the metal sensing behavior in sol–gel medium, the metallogels obtained from **3** and **4** exhibited a good response in terms of ionic conductivity, because of the movement of the metal ions within the gel networks under an applied electrical field. Conducting gels are widely used in the simulation of electrochemical devices such as super capacitors, fuel cells, rechargeable batteries, solar cells, *etc.*^{20,45} In general, long range delocalization of charges is necessary for a gel medium to illustrate conducting properties. However, most of the examples found in the literature are based on polymer gels and the methods that are abruptly used in the fabrication of conducting gels basically involve either direct production of gels from conducting polymers or blending of conducting polymers into non-conducting gel networks.⁴⁶ Not enough data are found regarding the fabrication of ion conducting supramolecular gels derived from simple metallo-organic architectures.⁴⁷

Results and discussion

Synthesis

Compounds **1–6** were achieved according to Scheme 1.³⁶ Oxidative condensation between 3,3'-diaminobenzidine and 2-, 3- and 4-pyridine carboxaldehyde in dry nitrobenzene at 140 °C resulted in one step synthesis of compounds **1**, **3** and **5**, respectively. On the other hand, condensation of *o*-phenylenediamine with the



Scheme 1 (i) 2-, 3-, or 4-pyridine carboxaldehyde, nitrobenzene, 140 °C, 20–24 h; (ii) 2-, 3-, or 4-pyridine carboxaldehyde, DMSO, 90 °C, 6–10 h.

said aldehydes in dry DMSO at 90 °C produced compounds **2**, **4** and **6** in appreciable yields. All the compounds were characterized by usual spectroscopic methods.

Gelation, thermal stability, morphology, rheology and metal ion sensing behaviors of the gels

Compounds 1 and 2. Structural analysis reveals that compounds **1**, **3** and **5** possess bispyridyl benzimidazole units which may exhibit different conformations^{36,48–50} during hydrogen bonding interactions. Compounds **2**, **4** and **6**, which are the monomeric forms of **1**, **3** and **5**, can acquire more directional hydrogen bonds involving imidazole –NH and pyridine ring nitrogen atoms. The hydrogen bonding may enhance the π – π stacking interactions of benzimidazole units.^{27–29,48–52} Considering such different attributes, initially the gelation abilities of compounds **1** and **2** were examined in a wide range of protic and aprotic solvents with different polarities. Table S1 (ESI†) summarizes the results of the gelation test of compounds **1** and **2** in different solvents. It was observed that the presence of an increasing amount of water improved the gelation propensities of both the compounds. In DMSO:H₂O (1:3, v/v), compound **1** formed a brown colored gel at a minimum concentration of 3.8 mg mL^{–1} whereas at least 5 mg compound was required for **2** to gelate 1 mL of the DMSO:H₂O (1:6, v/v) solvent mixture to form a white colored gel. The gels were thermo-reversible and upon heating were transformed into clear solution which upon cooling again to room temperature reappeared as gels (Fig. S1, ESI†). The thermal stability of the gels derived from aqueous DMSO was studied. A sharp increase in the gel melting temperature was observed with an increase in the gelator concentration (Fig. S2, ESI†).

The morphology of the gels was derived from SEM images that predicted a fibrous network (Fig. 2A). Structural difference of the two compounds tuned the morphology of the gels. Gel of **1** is composed of very thin nanofibers which are intertwined in a regular shape demonstrating well-organized molecular packing. In contrast, the gel state of **2** exhibited thin discrete particles.

To examine the mechanical properties of the gels, rheology experiments (Fig. 2B) were carried out. For both the gels the elastic storage modulus (G') was higher than the corresponding loss modulus (G''). With a gradual increase in the applied stress at a fixed frequency ($\omega = 10 \text{ rad s}^{-1}$), nonlinear characteristics were observed for both G' and G'' although no crossover point was found within the experimental stress region. However, in the case of **1**, the values of G' and G'' were almost independent of stress compared to the case of **2**, which demonstrated that gel **1** is stronger than gel **2**. The plot of G' and G'' against the angular frequency ω at a constant strain of 1% indicated that

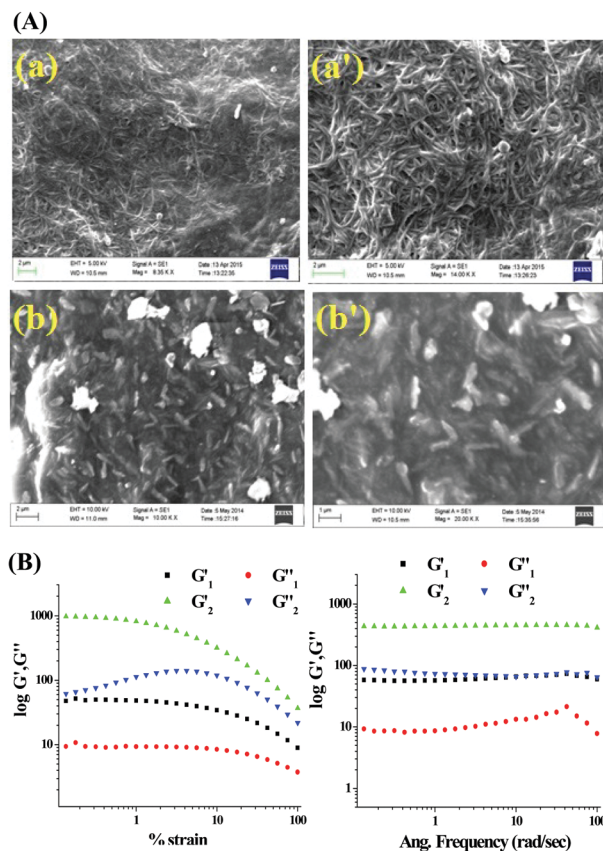


Fig. 2 (A) SEM images of xerogel of **1** (a and a') and **2** (b and b') prepared from aqueous DMSO at different scale bars; (B) rheology experiments of the DMSO : H₂O gels of **1** and **2**.

both G' and G'' were frequency invariant over the entire frequency range and the values of G' were almost 4 to 6 times higher than those of G'' , indicating their true gel behaviour.

Out of different possibilities, we believe that the imidazole ring nitrogen, pyridine nitrogen and the N-H groups of **1** and **2** are involved in intermolecular hydrogen bonding according to the modes described in Fig. 3 to form the aggregates. Water presumably acts as a bridging entity involving hydrogen bonding with the pyridine ring nitrogens and helps in establishing the cross-linked network in solution.³¹ In FTIR of **1** and **2**, the stretching frequency for -NH in the amorphous states was higher than the gel states which explained the hydrogen bond formation during gelation (Fig. S3, ESI[†]). Additionally, the appearance of a new signal in the gel states of both in the region 2090 cm⁻¹ to 2101 cm⁻¹ is likely to be due to the involvement of pyridine nitrogen in the formation of strong hydrogen bonds in the network. In addition, the stretching of C=N of benzimidazoles in **1** and **2** which appeared at 1577 cm⁻¹ and 1575 cm⁻¹, respectively underwent blue-shift.

To gain more insight into the aggregate formation, absorption spectra of **1** and **2** in sol and gel states were compared. A shoulder at 388 nm in the UV-vis spectrum of **1** in the gel state corroborated stacking induced aggregation (Fig. 3c). Additionally, a bathochromic shift of 14 nm of the absorbance at 330 nm is possibly due to planarization of the biphenyl backbone which in turn enhanced the

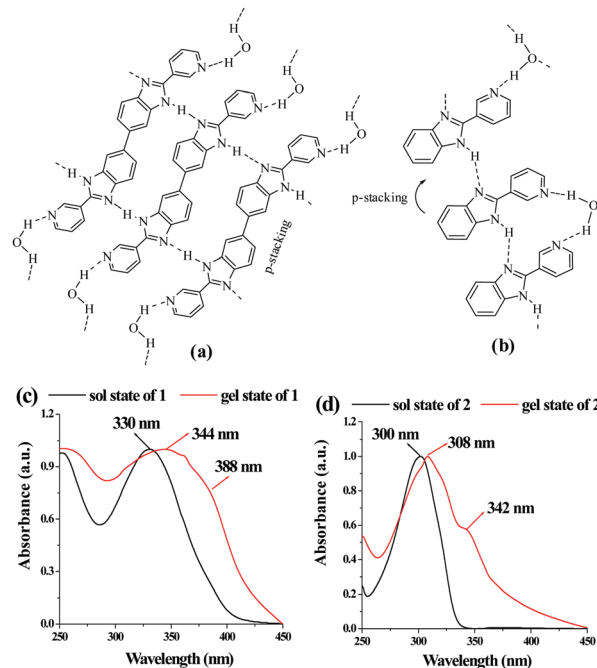


Fig. 3 Suggested modes of interaction of (a) **1** and (b) **2** to form the networks during gelation. Comparison of absorption spectra of **1** (c) and **2** (d) in sol and gel states.

effective π -conjugation length.^{53,54} The stacking-induced aggregation was much efficient in the case of gel of **2** that exhibited a new absorption at 342 nm (Fig. 3d). Such aggregation resulted in quenching of emission with red shifts of 33 nm and 28 nm in the case of **1** and **2**, respectively (Fig. S4, ESI[†]).

The gel phases of **1** and **2** exhibited a sharp response to the metal ions, leading to the gel-to-sol transition. For this, we studied the gelation propensity of the compounds in the presence of a series of metal ions in aqueous DMSO. In the case of **1**, the presence of 4 equiv. amounts of Ag⁺ and Cu²⁺ inhibited the gel formation (Fig. 4A) whereas other metal ions taken in the study left a marginal effect on gelation. We also investigated, on the other hand, the stimuli responsive nature of the gel by adding different metal ions to the gel state of **1**. The sustainability of the gel state was checked upon addition of 2 equiv. amounts of Ag⁺, Cu²⁺, Hg²⁺, Cd²⁺, Zn²⁺, Co²⁺, Ni²⁺, Pb²⁺, Mg²⁺, and Fe²⁺ (as perchlorate salts) in DMSO : H₂O (1 : 3, v/v). Only the gel phase of **1** in the presence of 2 equiv. amounts of Ag⁺ and Cu²⁺ ions was disrupted to the sol (resulting in precipitation) within 30 mins. Other metal ions, under similar conditions, did not show any effect on phase transition of **1** suggesting that gelator **1** is highly sensitive and selective towards Ag⁺ and Cu²⁺ ions over the other metal ions tested. The selectivity of **1** towards Ag⁺ and Cu²⁺ is presumed to be due to the positional role of pyridine nitrogen with respect to the imidazole nitrogens. Pyridine nitrogen away from the imidazole nitrogens is incapable of forming a metal chelate involving any collaborative effect. As a result, pyridine and imidazole will likely be involved in metal ion coordination individually. Although it is true for all the ions taken in the study, the pronounced effect of

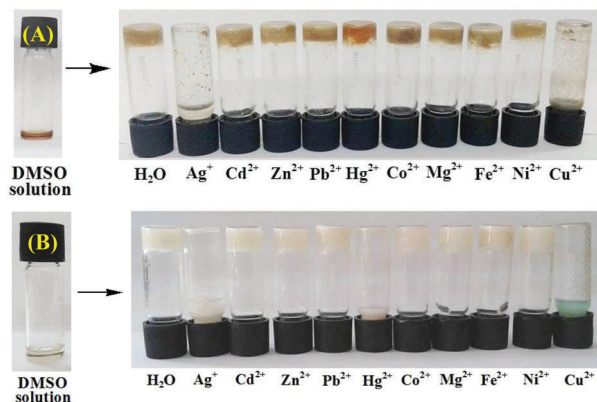


Fig. 4 Photographs showing the changes of the gels of (A) **1** (8 mg mL^{-1} , DMSO : $\text{H}_2\text{O} = 1 : 3 \text{ v/v}$) and (B) **2** (8 mg mL^{-1} , DMSO : $\text{H}_2\text{O} = 1 : 6 \text{ v/v}$) when the gels were prepared with 4 equiv. amounts of different metal ions [$c = 0.2 \text{ M}$ in H_2O as perchlorate salt] after 2 h.

Ag^+ and Cu^{2+} ions is due to their high affinity towards nitrogen atoms of either pyridine or imidazole rings to attain favourable geometries which, in turn, involve disruption of self-assemblies.⁵⁵

The metal-induced broken gels were found to be chemically reversible as tested by externally added chelating species (Fig. S5, ESI[†]). The Ag^+ -induced disrupted gel was recovered in the presence of excess KCl solution after 2 h. The scavenging of Ag^+ ions through the formation of AgCl led to ligand **1** free to attain its original structural feature for which gelation took place. Likewise, the Cu^{2+} -induced broken gel was recovered upon addition of ethylene diamine (in excess) that resulted in a blue color solution which ultimately became thick after 6 h. In this case, the color of the recovered gel was changed to blue rather than brown due to the presence of a Cu-ethylene diamine complex within the gel matrix.

Compound **2**, which formed a gel in DMSO : H_2O (1 : 6, v/v) in a similar study, was incapable of forming gels in the presence of Ag^+ , Cu^{2+} and Hg^{2+} ions (Fig. 4B). This highlighted the sensitivity of **2** towards Ag^+ , Cu^{2+} and Hg^{2+} ions. In the presence of 1 equiv. amount of Ag^+ and Cu^{2+} ions individually, the gel state of **2** was disrupted at a faster rate (within 1 h) than the case with Hg^{2+} which almost took 2.5 h to transform the gel into a sol. The Ag^+ , Cu^{2+} and Hg^{2+} -broken gels were not recovered in the presence of external chelators (e.g. Cl^- , ethylene diamine and I^- ions) and thereby revealed their chemical irreversibility. The nonappearance of the gel or disruption of the gel of gelator **2** in the presence of Ag^+ , Cu^{2+} and Hg^{2+} is presumably due to the same reason as given for gelator **1**. Like **1**, the pyridine and imidazole nitrogens in **2** interact strongly with Ag^+ and Cu^{2+} ions to attain their favourable geometries. The mercury ion exhibits weak affinity in comparison to Ag^+ and Cu^{2+} ions and therefore, takes longer time to break the self-assembly of **2**.

Due to the presence of pyridyl and benzimidazole nitrogens as the basic centre³⁶ we wished to study the effect of pH on the gel states of both **1** and **2** (Fig. S6, ESI[†]). The gels were stable within the pH range 6 to 9. At pH below 6 the gel state derived from **1** started to disintegrate, while at pH 5 there was partial gelation, and the gel was completely converted to a brown

colored sol at pH 4. However, upon further change of pH to 6, the gel reappeared instantly. At above pH 9, the gel was ruptured to the brown colored solution presumably due to the deprotonation of benzimidazole -NH. Similar observations were noted in the case of gel **2**. In this case, gel **2** was initially found to be stable at pH 5 for 30 h and then was converted to the sol slowly. These experimental observations underline the fact that the gels of **1** and **2** are stable within the pH window 6 to 9. Below pH 6 or above pH 9, the nonexistence of the gel states is due to either protonation or deprotonation phenomenon, respectively.

Compounds 3 and 4. As we move to compounds **3** and **4** (isomers of **1** and **2**, respectively), there was a change in gelation behaviour. Interestingly, while compounds **3** and **4** resulted in either precipitation or solution in DMSO : H_2O solvent (Table S1, ESI[†]), they formed thick gels instantly only in the presence of Ag^+ ions. It is mentionable that in the presence of 2 equiv. amounts of Ag^+ ions, compound **3** at a minimum concentration of 4.9 mg mL^{-1} formed a yellowish brown-colored gel in DMSO : H_2O (1 : 1, v/v) whereas at least 4.5 mg compound was necessary for **4** to gelate 1 mL of DMSO : H_2O (1 : 2, v/v) solvent to produce a white colored gel. The presence of less than 2 equiv. amounts of Ag^+ ions resulted in partial gelation in both cases. We also investigated the effect of other metal ions during gelation of **3** and **4** separately (Fig. 5). Interestingly except for Ag^+ , other metal ions taken in the study were unable to convert sol states of **3** and **4** into their gel states. This indicated the selectivity of the gelators to Ag^+ ions.

The morphologies of the gels were characterised by SEM images. While close packed globular microstructures were noted for **3**, intertwined tape like fibers were observed for the gel of **4** (Fig. 6).

The Ag^+ -induced gels of **3** and **4** exhibited a temperature dependent sol-gel reversible transition (Fig. S7, ESI[†]). At $56 \text{ }^\circ\text{C}$, the gel state of **3** (at mge) was converted to the sol state with brown color. However, the gel state of **4** at mge was transformed into a clear solution at $64 \text{ }^\circ\text{C}$. In both cases, T_{gel} varies directly with the gelator concentration (Fig. S8, ESI[†]). Such kind of gel

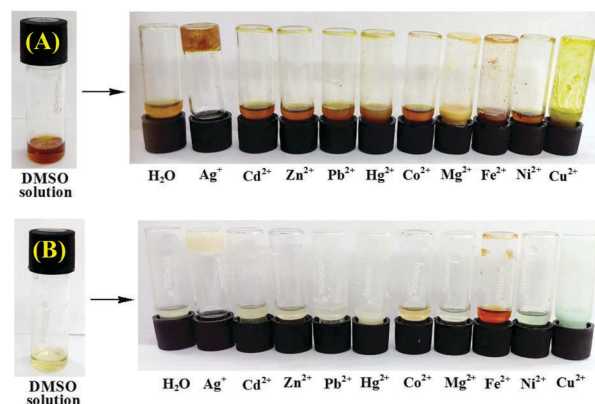


Fig. 5 Photographs showing the phase changes of the gels of (A) **3** (4.9 mg mL^{-1} , DMSO : $\text{H}_2\text{O} = 1 : 1$, v/v) and (B) **4** (4.5 mg mL^{-1} , DMSO : $\text{H}_2\text{O} = 1 : 2$, v/v) in the presence of 2 equiv. amounts of different metal ions [$c = 0.2 \text{ M}$ in H_2O as perchlorate salt] after 2 h.

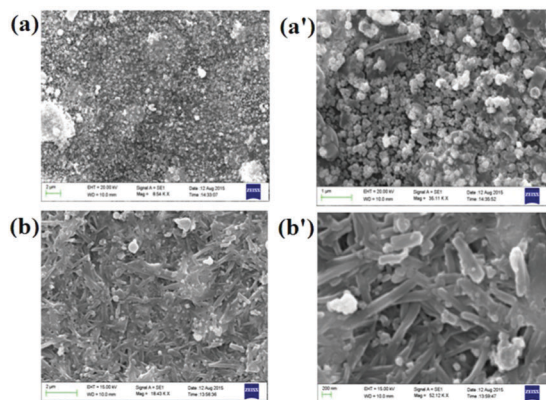


Fig. 6 SEM images of xerogels derived from **3** (a and a') and **4** (b and b') with Ag^+ from aqueous DMSO at different scale bars.

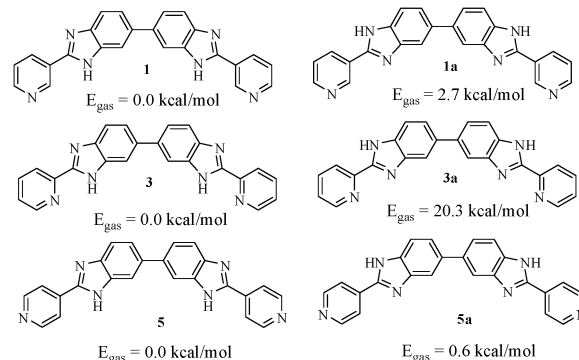


Fig. 7 Two relative conformations of the isomeric compounds based on the protons attached to the benzimidazole N face and their relative gas phase energies as ascertained by Bhattacharya *et al.*,³⁶ using the B3LYP/6-31G* level of theory.

to sol transition was also observed in the presence of Cl^- ions at room temperature. Addition of 2 equiv. amounts of Cl^- ions caused rapid destruction of the gels within 1 h giving a white precipitate of AgCl . Further addition of Ag^+ ions assembles ligands **3** and **4** to attain the required structural features, responsible for gelation (Fig. S9, ESI[†]).

Fig. S10 (ESI[†]) represents the probable binding modes of **3** and **4** out of several possibilities during gel formation. The strong coordination of Ag^+ ions involving the pyridyl and benzimidazole ring nitrogens and the hydrogen bonding interplaying involving water are presumably the directives to attain a cross-linked structure in the solution to afford the gels. In FTIR spectra of **3** and **4**, from amorphous to their gel state there was a shifting of the $-\text{NH}$ stretching to higher wavenumber (Fig. S11, ESI[†]). Moreover, the stretching of $\text{C}=\text{N}$ of the imidazole rings appearing at 1589 cm^{-1} and 1588 cm^{-1} in **3** and **4**, respectively underwent a blue-shift and moved to 1595 cm^{-1} and 1592 cm^{-1} , respectively. The blue-shift essentially reflects the strengthening of the $\text{C}=\text{N}$ bond in the presence of Ag^+ and it is ascribed to the inhibition of electronic delocalization around the benzimidazolic unit.⁵⁶

Moreover, due to aggregation, the gel states of **3** and **4** exhibited strong quenching of emission with redshifts of 40 nm and 28 nm of the emission bands, respectively (Fig. S12 and S13, ESI[†]).

Compounds 5 and 6. It is important to mention that compounds **5** and **6** (*p*-isomers) like the other isomers (*ortho* and *meta*) were insensitive to gelation either alone or in the presence of any other metal ions examined. This highlighted the positional role of pyridine ring nitrogen with respect to the benzimidazole ring which controlled the self-aggregation of the molecules either alone or in the presence of metal ions.

Positional role of pyridine ring nitrogen with respect to benzimidazole in gelation

A detailed theoretical study on the molecules (**1**, **3** and **5**) was put forward by Bhattacharya and co-workers.³⁶ In their study, they have reported the conformational behaviour of the molecules in gas and solvent phases in detail. In **3** (*ortho* isomer), the intramolecular hydrogen bonding between benzimidazole $-\text{NH}$

and pyridine ring nitrogen led to coplanarity between rings and thereby ruled out the possibility of intermolecular aggregation through hydrogen bonding. On moving from **3** to **1** (*meta*-isomer) and **5** (*para*-isomer), the case of intramolecular hydrogen bonding between benzimidazole $-\text{NH}$ and pyridine ring nitrogen is ruled out. Consequently, the energy barrier related to the orientation of the pyridine ring with respect to benzimidazole is reduced to a significant extent and thus isomers **1** and **5** are conformationally mobile (Fig. 7).³⁶

Therefore, in our opinion, isomers **1** and **5** may undergo aggregation through intermolecular hydrogen bonding in several ways and may form gels. But the experimental observation reveals that only isomer **1** exhibits aggregation-induced gelation from DMSO in the presence of water as cosolvent which most likely helps in cross-linking involving pyridine to establish a network for gelation. In **5**, the involvement of water in setting up of a cross-linked network involving pyridine as that of **1** is likely to be less feasible and therefore, gelation does not occur.

The case of intramolecular hydrogen bonding in **3** and the role of water in linking the molecules were realized by considering the MMX calculation⁵⁷ of the dimeric forms of the monomeric analogues **2**, **4** and **6** (Fig. S14, ESI[†]). While in **2** the dimeric form is associated with water to form the network, structure **6** is assumed to be less efficient in producing suitable network for gelation. In comparison, the pyridine in dimeric form of **3** is unavailable for water coordination to form the network for entrapping solvent.

Metal ion binding in solution with compounds 1–4

Pyridyl benzimidazoles are established motifs in the formation of stable metal–ligand complexes.^{27–29,58–60} In order to verify the solution phase interactions of the pyridyl benzimidazoles of the present study, we performed UV-vis and fluorescence titrations of **1** and **2** ($c = 2.5 \times 10^{-5}\text{ M}$) in $\text{DMSO}:\text{H}_2\text{O}$ (1:9, v/v). In the fluorescence study, compound **1** showed strong emission at 442 nm upon excitation at 330 nm. Fig. S15 (ESI[†]) represents the change in fluorescence ratio of **1** at 442 nm in the presence of 10 equiv. amounts of metal ions. The result indicated the sensitivity of **1** towards Ag^+ ions over the other metal ions considered in the study. During titration of **1**, the intensity of

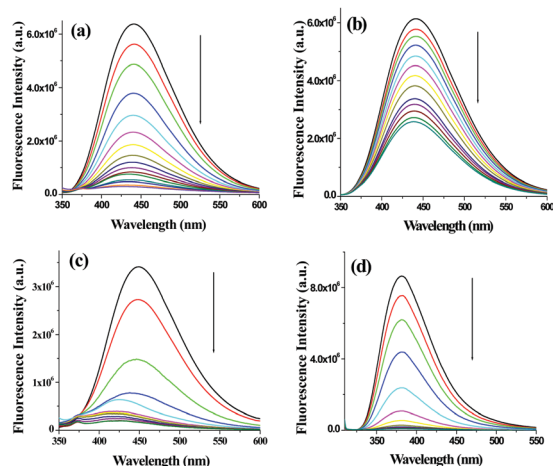


Fig. 8 Change in emission of **1** ($c = 2.5 \times 10^{-5}$ M) upon addition of 10 equiv. amount of (a) Ag^+ and (b) Cu^{2+} ($c = 1.0 \times 10^{-3}$ M) in $\text{DMSO}:\text{H}_2\text{O}$ (1:9, v/v) respectively, Change in emission of (c) **3** and (d) **4** ($c = 2.5 \times 10^{-5}$ M) upon addition of 1 equiv. and 5 equiv. amounts of Ag^+ ($c = 1.0 \times 10^{-3}$ M) in $\text{DMSO}:\text{H}_2\text{O}$ (1:9, v/v), respectively.

the emission at 442 nm underwent significant quenching only in the presence of Ag^+ ions (Fig. 8a) with a blue shift of 16 nm in emission spectra. Copper ion perturbed the emission moderately resulting in a blue shift of 5 nm (Fig. 8b), while other metal ions showed negligible effect on the emission of **1** (Fig. S16, ESI[†]). The binding constant values⁶¹ were found to be $(2.78 \pm 0.45) \times 10^4 \text{ M}^{-1}$ and $(1.88 \pm 0.28) \times 10^4 \text{ M}^{-1}$ for Ag^+ and Cu^{2+} , respectively with 1:1 stoichiometries⁶² (Fig. S17 and S18, ESI[†]). In this sensing process, the detection limits⁶³ for Ag^+ and Cu^{2+} were determined to be 3.69×10^{-6} M and 9.51×10^{-6} M, respectively (Fig. S19, ESI[†]). Ag^+ -Induced greater quenching of emission at 442 nm even in the presence of other metal ions corroborated the selectivity of **1** toward Ag^+ ions (Fig. S20, ESI[†]) thereby indicating the noninterference of other metal ions.

On the other hand, compound **2** in $\text{DMSO}:\text{H}_2\text{O}$ (1:9, v/v) upon excitation at 300 nm exhibited emission at 382 nm which was perturbed (quenching) almost equally upon interaction with metal ions such as Ag^+ , Cu^{2+} and Hg^{2+} (Fig. S21, ESI[†]). Upon gradual addition of these metal ions to the solution of **2**, the emission intensity was gradually decreased without any other change in the emission spectra. Other metal ions taken in the study did not introduce any noticeable effect on the emission of **2** (Fig. S22, ESI[†]). In solution, compound **2** showed 1:2 (H:G) stoichiometric interactions with Ag^+ , Cu^{2+} and Hg^{2+} ions (Fig. S23, ESI[†]). From fluorescence titration data, the binding constant values which described the stronger complexation of Ag^+ [$K_1 = (1.05 \pm 0.15) \times 10^4 \text{ M}^{-1}$ and $K_2 = (9.75 \pm 4.01) \times 10^2 \text{ M}^{-1}$] than Cu^{2+} [$K_1 = (9.02 \pm 2.44) \times 10^3 \text{ M}^{-1}$ and $K_2 = (6.57 \pm 1.58) \times 10^3 \text{ M}^{-1}$] and Hg^{2+} [$K_1 = (6.01 \pm 0.28) \times 10^3 \text{ M}^{-1}$ and $K_2 = (2.07 \pm 0.04) \times 10^3 \text{ M}^{-1}$] ions (Fig. S24, ESI[†]). The detection limits of **2** for Ag^{2+} , Cu^{2+} and Hg^{2+} were determined to be 3.34×10^{-6} M, 5.06×10^{-6} M and 2.02×10^{-6} M, respectively (Fig. S25, ESI[†]).

The solution phase interactions of **3** and **4** ($c = 2.5 \times 10^{-5}$ M) were also investigated with Ag^+ ($c = 1.0 \times 10^{-3}$ M) ions under similar conditions. In the fluorescence study, the presence of

only 1 equiv. amount of Ag^+ resulted in almost complete quenching of emission of **3** with a 30 nm blue shift (Fig. 8c). In comparison, compound **4** exhibited complete fluorescence quenching in the presence of 5 equiv. amounts of Ag^+ (Fig. 8d) with no spectral shift like **3**. The binding constant values were calculated to be $(1.30 \pm 0.31) \times 10^6 \text{ M}^{-1}$ and $(4.98 \pm 0.98) \times 10^4 \text{ M}^{-1}$ for 1:1 complexation of Ag^+ ions with **3** and **4**, respectively (Fig. S26–S29, ESI[†]). From the fluorescence titration data, the detection limits for Ag^+ with **3** and **4** were determined to be 1.93×10^{-7} M and 1.28×10^{-6} M, respectively (Fig. S30 and S31, ESI[†]).

In UV-vis titration of **1**, in the presence of both Ag^+ and Cu^{2+} ions, the intensity of the absorption at 330 nm was decreased with the simultaneous increase of a peak at 260 nm (Fig. 9a and b). In the case of Ag^+ , the ratiometric nature of the absorption spectra was observed only up to the addition of 2 equiv. amounts of Ag^+ . Further addition of silver ions caused a gradual increase in absorption at 330 nm. In the ground state, compound **1** interacted with both Ag^+ and Cu^{2+} ions in a 1:1 stoichiometric manner with binding constant values of $(2.13 \pm 0.48) \times 10^3 \text{ M}^{-1}$ and $(1.04 \pm 0.15) \times 10^4 \text{ M}^{-1}$, respectively (Fig. S32–S34, ESI[†]). Compound **2** interacted more strongly with Ag^+ in comparison to **1** in the ground state, showing a steady decrease in the absorption at 300 nm with a binding constant value of $(3.60 \pm 0.57) \times 10^3 \text{ M}^{-1}$ involving 1:1 complexation (Fig. S35–S37, ESI[†]). However, a red shift of 16 nm in the absorption spectra was noted during titration of **3** with Ag^+ ions (Fig. 9c). Interestingly, in the presence of Ag^+ , a ratiometric change with a stable isosbestic point at 333 nm was observed in the absorption spectra of **4** (Fig. 9d). The Job method corroborated 1:1 complexation of Ag^+ ions with both **3** and **4** with the binding constant values of $(9.47 \pm 1.07) \times 10^4 \text{ M}^{-1}$ and $(2.46 \pm 0.66) \times 10^4 \text{ M}^{-1}$, respectively (Fig. S26–S29, ESI[†]). Table S2 (ESI[†]) summarises the metal–ligand interaction profile in detail and the results

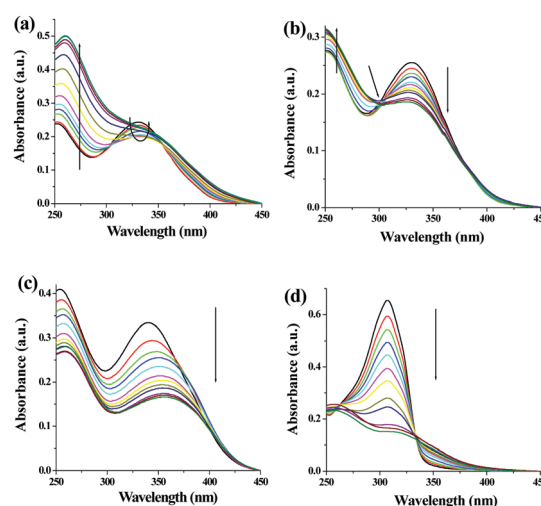


Fig. 9 Change in absorbance of **1** ($c = 2.5 \times 10^{-5}$ M) upon addition of 6 equiv. amounts of (a) Ag^+ and (b) Cu^{2+} ($c = 1.0 \times 10^{-3}$ M) in $\text{DMSO}:\text{H}_2\text{O}$ (1:9, v/v) respectively, Change in absorbance of (c) **3** and (d) **4** ($c = 2.5 \times 10^{-5}$ M) upon addition of 2 equiv. amounts of Ag^+ ($c = 1.0 \times 10^{-3}$ M) in $\text{DMSO}:\text{H}_2\text{O}$ (1:9, v/v), respectively.

indicate that the *o*-derivatives **3** and **4** chelate Ag^+ ions more strongly both in ground and excited states probably due to the formation of a more stable five membered ring during complexation.

In order to understand the metal ligand interactions, we performed ^1H NMR experiments of **1**–**4** in the absence and presence of Ag^+ in respective solvents. In ^1H NMR of **1** and **2** with an equivalent amount of Ag^+ , the downfield chemical shifts of the imidazole $-\text{NH}_a$ ($\Delta\delta$ for **1** = 0.37 ppm and **2** = 0.54 ppm in d_6 -DMSO) indicated strong involvement of the imidazole ring nitrogen in complexation with Ag^+ ions (Fig. S38 (ESI †) and Fig. 10A). Due to complexation, the benzimidazole ring proton H_c moved downfield by 0.67 ppm and 0.17 ppm for **1** (Fig. S38, ESI †) and **2** (Fig. 10A), respectively. Moreover, the downfield chemical shift of pyridyl ring protons in **1** and **2** indicated the simultaneous involvement of the pyridine rings in complexation (Fig. 10A). Such kind of dinuclear Ag^+ -coordination in M_2L_2 fashion is known in the literature.^{58,59} A similar characteristic feature in ^1H NMR spectra was observed with **3** and **4**. In ^1H NMR spectra of 1 : 1 mixture of **3** and Ag^+ in d_6 -DMSO, the aromatic protons became too broad to detect their exact positions except the imidazole- NH_a that exhibited a downfield chemical shift of 0.79 ppm (Fig. S39, ESI †). However in the case of **4**, in the presence of an equivalent amount of Ag^+ in CDCl_3 , no significant shift of the pyridyl protons of type H_b and H_c was observed.

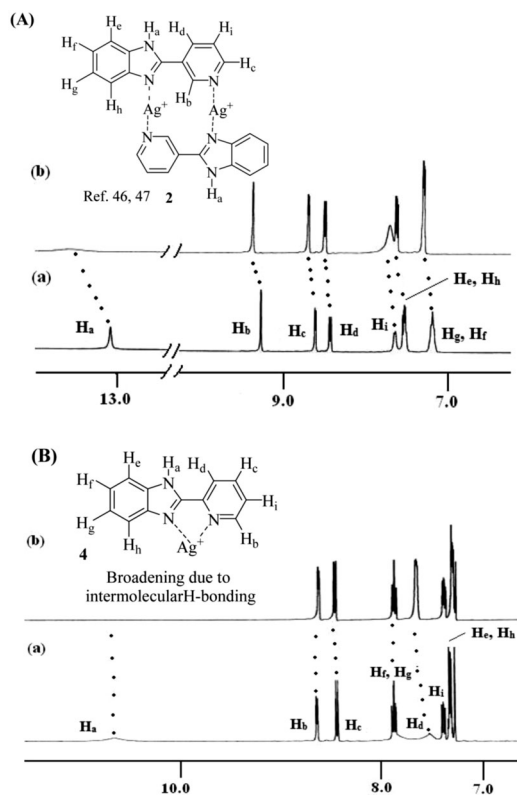


Fig. 10 (A) Probable binding mode of **2** with Ag^+ in solution along with ^1H NMR (400 MHz, d_6 -DMSO) changes of (a) compound **2** ($c = 2.56 \times 10^{-2}$ M) and (b) **2** with 1 equiv. amount of AgClO_4 . (B) Probable binding mode of **4** with Ag^+ in solution along with ^1H NMR (400 MHz, CDCl_3) changes of (a) compound **4** ($c = 3.41 \times 10^{-2}$ M) and (b) **4** with 1 equiv. amount of AgClO_4 .

Moreover, the disappearance of the signal for imidazole $-\text{NH}_a$ was attributed to the hydrogen bond mediated broadening (Fig. 10B).

Ion conductivity

As application the metallogels were explored in an ion conductivity study. The Ag^+ -induced metallogels of **3** and **4** gave a good response in terms of electrical conductivity. The dielectric properties of a charge containing medium are usually measured by means of the impedance spectroscopy technique.⁶⁴ Fig. 11a and b, in this aspect, show the frequency and temperature dependence of the real part of complex total ionic conductivity of the gels of **3** and **4** (8 mg mL^{-1}) respectively, which arises due to the migration of Ag^+ ions within the gel matrices. It is observed from both the figures that the frequency dependence of ionic conduction within the gels can be characterized by two mutually correlated phenomena *viz.* low frequency electrode polarization and high frequency independent macroscopic charge conduction. For both the gels, it is worth mentioning that below an onset frequency (ω_{on}) the conductivity $\sigma_t(\omega)$ slowly decreases with decreasing frequency. This corresponds to the electrode polarization due to the accumulation of charges at the electrolyte-electrode interface, commonly designated as the frequency dependent double layer capacitive effect (C_{dl}).^{65,66} Above the onset frequency of electrode polarizations σ_t remains frequency independent, which, in accordance with other reports,^{67,68} can be attributed to the macroscopic dc-conductivity of the materials obtained from the intersection of the extrapolated straight lines with the y-axis as depicted in the figures. It is observed from the

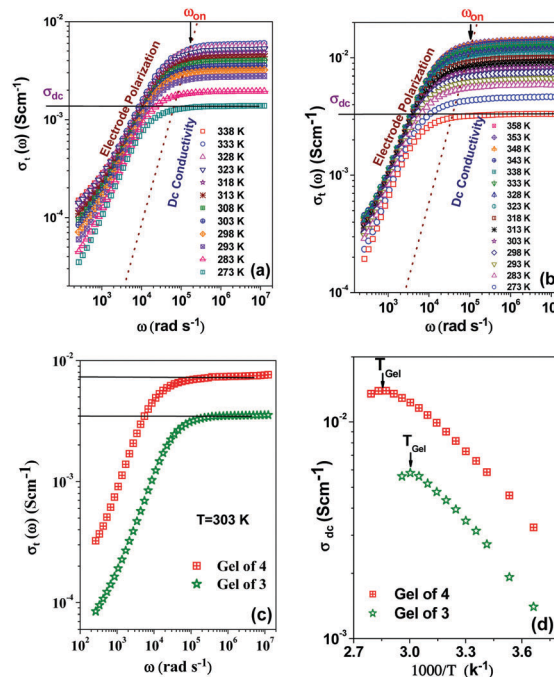


Fig. 11 Frequency dependent real part of total ionic conductivity of (a) gel of **3**, (b) gel of **4**, (c) comparison of the frequency dependent real part of total ionic conductivity of the gels at room temperature, (d) temperature dependence of the dc conductivity of the gels.

figures that as the temperature decreases, the overall conductivity decreases and the onset of electrode polarization (ω_{on}) shifts toward lower frequencies with reduction in the frequency range of electrode polarization. Fig. 11c depicts the comparison of the frequency dependent real part of ac-conductivity of the gels of **3** and **4** at room temperature (303 K). Fig. 11d displays the temperature dependence of the dc-conductivity obtained from Fig. 11a and b, respectively. It is interesting to note that σ_{dc} in both the gels shows thermally activated nature up to the respective T_{gel} . Also, as the T_{gel} approaches, the gels gradually lose their thermally activated conductive nature and conductivity tends to saturate. Above T_{gel} the conductivity decreases from the saturated value. The above nature of temperature dependence of conductivity for both the gels clearly indicates that the gel matrix favours Ag^+ ion conduction to a greater extent than the sol due to the increase in defect states which actually enhances the hopping conductivity. In the vicinity of T_{gel} the gel structure is on the way of rupturing and simultaneously the defect states are diminished which hinders further increase of conductivity. Above T_{gel} the hopping conductivity is destroyed and the uncorrelated movement of cations and anions within the sol contributes to the conductivity which is less than the hopping conductivity. Both Fig. 11c and d clearly indicate that the gel matrix of **4** favours the Ag^+ ion conduction more than the gel of **3** at each measured temperature. The hindrance to Ag^+ ion conduction in the gel of **3** is likely to be due to the greater entanglement of Ag^+ in the gel network of **3** for participation of more number of pyridyl benzimidazole units.

Conclusions

In conclusion, we have explored the possibilities of pyridine-coupled mono and bisbenzimidazoles **1–6** in gelation and their subsequent use in detection of metal ions. Of the benzimidazoles **1–6**, compounds **1** and **2** formed instant gels from DMSO/ H_2O and MeOH/ H_2O solvents. The gels of **1** and **2** are pH responsive and thermoreversible. Moreover, the gel phases of **1** and **2** are rapidly transformed into the sol states in the presence of Ag^+ and Cu^{2+} ions. Additionally, compound **2** exhibited a strong affinity for Hg^{2+} and recognized it *via* a gel to sol transition.

On the other hand, the structural isomers **3** and **4** exhibited gelation tendency in the presence of Ag^+ ions selectively from similar solvent systems and thus validated their visual sensing over a series of other metal ions. In relation to this, nongelation behavior of *p*-derivatives **5** and **6** (structural isomers) under identical conditions emphasized the positional role of the pyridyl ring nitrogen with respect to the imidazole ring. Thus the uniqueness of pyridyl benzimidazoles in gel chemistry is admirable. While they are easy-to-make, the presence of three N-atoms and imine $-\text{NH}$ groups with extended π backbone is of great use in supramolecular self-aggregation either alone or in the presence of metal ions.

As an application, the metallo gels derived from **3** and **4** were explored as conducting electrolytes as they responded well in terms of thermally activated ionic conductivity due to the movement of Ag^+ ions within the gel matrix. This kind of

conductivity study with pyridyl benzimidazoles is unexplored in the literature. Thus the present study in this report draws attention and is a new addendum to the existing merely explored pyridyl benzimidazole gelators for metal ion sensing (Table S3, ESI[†]).

Experimental

Materials

3-Amino pyridine was purchased from Spectrochem. 3,3'-Diaminobenzidine was obtained from Sigma-Aldrich. Metal salts used in the study were purchased from Sigma-Aldrich and were carefully handled. All solvents used in the synthesis were purified, dried and distilled as required. Solvents used in NMR experiments were obtained from Aldrich. Thin layer chromatography was performed on Merck precoated silica gel 60-F₂₅₄ plates. ^1H and ^{13}C NMR spectra were recorded using a Bruker 400 MHz instrument using TMS as internal standard. High resolution mass data were acquired by the electron spray ionization (ESI) technique on a XEVO GS-2 QTOF Waters mass spectrometer. FTIR measurements of all the compounds and dried gels (xerogels) were carried out using a Perkin-Elmer L120-00A spectrometer (ν_{max} in cm^{-1}) using a KBr cell and KBr pellets, respectively. Scanning electron microscopy (SEM) images were obtained on a EVO LS-10 ZEISS instrument. Fluorescence and UV-Vis studies were performed using a Horiba Fluoromax 4C spectrofluorimeter and a Shimadzu UV-2450 spectrophotometer, respectively.

Syntheses

2,2'-Di-pyridin-3-yl-3H,3'H-[5,5']bibenzoimidazolyl (1). A mixture of 3,3'-diaminobenzidine (0.200 g, 0.933 mmol) and 3-pyridinecarboxaldehyde (0.299 g, 2.800 mmol) in nitrobenzene (10 mL) was allowed to heat at 140 °C for 20 h. The progress of the reaction was monitored by TLC. After completion of the reaction compound **1** was purified by column chromatography using 10% MeOH in ethyl acetate as eluent in 72% yield (0.261 g, decomposition temperature 186 °C). ^1H NMR (d_6 -DMSO, 400 MHz): δ 13.27 (s, 2H), 9.36 (s, 2H), 8.69 (d, 2H, $J = 4$ Hz), 8.51 (d, 2H, $J = 8$ Hz), 7.62–7.59 (m, 8H); ^{13}C NMR (100 MHz, d_6 -DMSO): δ 153.9, 147.9, 145.1, 144.8, 141.2, 134.1, 127.1, 124.6 (signals for four carbons were not resolved); FTIR (KBr) ν cm^{-1} : 3350, 2926, 1632, 1577, 1429, 1384; HRMS (TOF MS ES⁺): required 389.1515 ($M + H$)⁺, found 389.1518 ($M + H$)⁺.

2-Pyridin-3-yl-1H-benzoimidazole (2). *ortho*-Phenylenediamine (0.300 g, 2.774 mmol) and 3-pyridinecarboxaldehyde (0.442 g, 4.161 mmol) were mixed in dry DMSO (5 mL) and the reaction mixture was then allowed to heat at 90 °C for 6 h. Then water was added to the reaction mixture and it was stirred for 30 minutes. The resulting precipitate was filtered to get the crude product. The crude product was purified by column chromatography using 70% ethyl acetate in petroleum ether as eluent to afford the pure compound **2** in 78% yield (0.422 g, mp 248 °C). ^1H NMR (d_6 -DMSO, 400 MHz): δ 13.07 (s, 1H), 9.28 (s, 1H), 8.61 (d, 1H, $J = 4$ Hz), 8.43 (d, 1H, $J = 8$ Hz), 7.63 (d, 1H, $J = 4$ Hz), 7.54–7.50 (m, 2H), 7.18 (brt, 2H); ^{13}C NMR (100 MHz, d_6 -DMSO): δ 150.9, 149.3, 147.9, 134.2, 126.5, 124.5, 122.9 (signals for two carbons are

unresolved); FTIR (KBr) ν cm^{-1} : 3422, 2883, 1575, 1488, 1447, 1315; HRMS (TOF MS ES⁺): required 196.0875 (M + H)⁺, found 196.0875 (M + H)⁺.

2,2'-Di-pyridin-2-yl-3H,3'-H-[5,5']bibenzoimidazolyl (3). 3,3'-Diaminobenzidine (0.200 g, 0.933 mmol) and 2-pyridine-carboxaldehyde (0.299 g, 2.800 mmol) were taken in nitrobenzene (10 mL) and the reaction mixture was then allowed to heat at 140 °C for 24 h. The progress of the reaction was monitored by TLC. Purification of crude mass by column chromatography using ethyl acetate as eluent afforded compound 3 in 52% yield (0.188 g, mp 226 °C). ¹H NMR (d₆-DMSO, 400 MHz): δ 13.11 (s, 2H), 8.68 (d, 2H, J = 4 Hz), 8.29 (d, 2H, J = 8 Hz), 7.97–7.91 (m, 2H), 7.72 (s, 2H), 7.56–7.46 (m, 6H); ¹³C NMR (100 MHz, d₆-DMSO): δ 151.7, 149.8, 148.8, 138.0, 136.9, 136.1, 125.2, 123.6, 122.4, 121.9, 120.0, 117.8; FTIR (KBr) ν cm^{-1} : 3185, 1636, 1589, 1443, 1305, 1262; HRMS (TOF MS ES⁺): required 389.1515 (M + H)⁺, found 389.1511 (M + H)⁺.

2-Pyridin-2-yl-1H-benzoimidazole (4). *ortho*-Phenylenediamine (0.300 g, 2.774 mmol) and 2-pyridinecarboxaldehyde (0.442 g, 4.161 mmol) were dissolved in dry DMSO (5 mL) and the reaction mixture was allowed to heat at 90 °C for 10 h. Then water was added to the reaction mixture and it was stirred for 30 minutes. The brown precipitate was filtered to get the crude mass which was purified by column chromatography using 40% ethyl acetate in petroleum ether as eluent. Compound 4 was obtained in 68% yield (0.368 g, mp 218 °C). ¹H NMR (CDCl₃, 400 MHz): δ 10.65 (s, 1H), 8.63 (d, 1H, J = 4 Hz), 8.44 (d, 1H, J = 8 Hz), 7.92–7.85 (m, 2H), 7.51 (brs, 1H, J = 8 Hz), 7.39–7.36 (m, 1H), 7.33–7.28 (m, 2H); ¹³C NMR (100 MHz, CDCl₃): δ 150.8, 148.9, 148.4, 144.2, 137.5, 134.2, 124.6, 123.8, 122.8, 122.0, 119.9, 111.4; FTIR (KBr) ν cm^{-1} : 3180, 1634, 1588, 1442, 1305, 1262; HRMS (TOF MS ES⁺): required 196.0875 (M + H)⁺, found 196.0878 (M + H)⁺.

2,2'-Di-pyridin-4-yl-3H,3'-H-[5,5']bibenzoimidazolyl (5). 3,3'-Diaminobenzidine (0.200 g, 0.933 mmol) and 4-pyridine-carboxaldehyde (0.299 g, 2.800 mmol) were taken in nitrobenzene (10 mL) and the reaction mixture was allowed to heat at 140 °C for 20 h. The progress of the reaction was monitored by TLC. Crude compound 5 was purified by column chromatography using 15% MeOH in ethyl acetate as eluent to yield the pure compound 5 in 76% yield (0.275 g, decomposition temperature 186 °C). ¹H NMR (d₆-DMSO, 400 MHz): δ 13.44 (s, 2H), 8.71 (d, 4H, J = 8 Hz), 8.07 (d, 4H, J = 8 Hz), 7.87 (brs, 2H), 7.73 (d, 2H, J = 8 Hz), 7.59 (d, 2H, J = 8 Hz); ¹³C NMR (100 MHz, d₆-DMSO): δ 150.9, 150.4, 149.5, 149.3, 137.0, 136.2, 124.1, 122.84, 122.82, 120.3; FTIR (KBr) ν cm^{-1} : 3180, 1634, 1593, 1443, 1306; HRMS (TOF MS ES⁺): required 389.1515 (M + H)⁺, found 389.1515 (M + H)⁺.

2-Pyridin-4-yl-1H-benzoimidazole (6). *ortho*-Phenylenediamine (0.300 g, 2.774 mmol) and 4-pyridinecarboxaldehyde (0.442 g, 4.161 mmol) were dissolved in 3 mL of dry DMSO and the reaction mixture was then allowed to heat at 90 °C for 10 h. Then water was added to the reaction mixture and it was stirred for 30 minutes, resulting in a brown precipitate which was filtered to get the crude mass and then purified by column chromatography using 80% ethyl acetate in petroleum ether as eluent which afforded the pure compound 6 in 82% yield (0.444 g, m.p. 212 °C). ¹H NMR (d₆-DMSO, 400 MHz): δ 13.24

(s, 1H), 8.69 (dd, 1H, J_1 = 8 Hz, J_2 = 4 Hz), 8.02 (dd, 1H, J_1 = 4 Hz, J_2 = 4 Hz), 7.59 (s, 1H), 7.20 (dd, 1H, J_1 = 8 Hz, J_2 = 4 Hz); ¹³C NMR (100 MHz, d₆-DMSO): δ 149.9, 148.1, 138.8, 136.5, 122.3, 119.7, 115.1; FTIR (KBr) ν cm^{-1} : 3163, 1648, 1595, 1440, 1316, 1258; HRMS (TOF MS ES⁺): required 196.0875 (M + H)⁺, found 196.0880 (M + H)⁺.

Gelation test and SEM imaging

Respective amounts of compounds of 1–6 were dissolved in 1 mL of desired solvents, slightly warmed to form a homogeneous solution and then allowed to cool slowly to room temperature to form a gel. All the gels were tested by an inversion of vial method. Samples of gels for SEM imaging were dried under vacuum and then coated with a thin layer of gold metal.

Determination of gel-sol transition temperature (T_g)

The gel-to-sol transition temperature (T_g) was measured by the dropping ball method. The T_{gel} is defined as the temperature at which the gel melted and started to flow. In this test, a small glass ball was carefully placed on the top of the gel to be tested, which was present in a test tube. The tube was slowly heated in a thermostated oil bath until the ball fell to the bottom of the test tube. The temperature at which the ball reaches the bottom of the test tube is taken as T_g of that system.

General procedure for fluorescence and UV-vis titrations

Stock solutions of the compounds were prepared in DMSO: H₂O (1:9, v/v) at a concentration of 2.50×10^{-5} M. Stock solutions of metal ions were also prepared in the same solvents at a concentration of 1.0×10^{-3} M. A solution of each compound (2 mL) was taken in the cuvette and to this solution different metal ions were individually added in different amounts. Upon addition of metal ions, the change in emission of the compound was recorded. The same stock solutions were used to perform the UV-vis titration experiment in the same way.

Binding constant determination⁶¹

Binding constant values for receptors with the ionic guests were determined by the nonlinear curve fitting procedure. The 1:1 [H:G] nonlinear fit was done using the equation $I = I_0 + (I_{lim} - I_0)/2C_H\{C_H + C_G + 1/K_a - [(C_H + C_G + 1/K_a)^2 - 4C_H C_G]^{1/2}\}$, where I represents the intensity; I_0 represents the intensity of pure host; C_H and C_G are the corresponding concentrations of the host and ionic guest; K_a is the binding constant. The binding constant (K_a) and correlation coefficients (R) were obtained from a non-linear least-squares analysis of I versus C_H and C_G .

Binding constant values for receptors with the ionic guests in 1:2 complexations were determined by the nonlinear curve fitting procedure using the equation: $I = [I_0 + (K_1 C_G I_m) + (K_1 K_2 A X^2)] / (1 + K_1 X + K_1 K_2 X^2)$, where I represents the intensity; I_0 represents the intensity of pure host; I_m represents the intensity of the host in the presence of an equiv. amount of guest; C_G is the corresponding concentrations of the guest; K_1 and K_2 are the binding constant values. The binding constant values and correlation coefficients (R) were obtained from a non-linear least-squares analysis of I versus C_G .

Method for the Job plot⁶²

The stoichiometry was determined by the continuous variation method (Job plot). In this method, solutions of host and guests of equal concentrations were prepared in respective solvents. Then host and guest solutions were mixed in different proportions maintaining a total volume of 3 mL of the mixture. The related compositions for host:guest (v/v) were 3:0, 2.8:0.2; 2.5:0.5, 2.2:0.8, 2:1, 1.8:1.2, 1.5:1.5, 1:2, 0.8:2.2, 0.5:2.5, 0.2:2.8. All the prepared solutions were kept for 1 h with occasional shaking at room temperature. Then emission and absorbance of the solutions of different compositions were recorded. The concentration of the complex, *i.e.*, [HG] was calculated using the equation $[HG] = \frac{\Delta I/I_0}{[H]}$ or $[HG] = \frac{\Delta A/A_0}{[H]}$ where $\Delta I/I_0$ and $\Delta A/A_0$ indicate the relative emission and absorbance intensities, respectively. [H] corresponds to the concentration of pure host. Mole fraction of the host (X_H) was plotted against the concentration of the complex [HG]. In the plot, the mole fraction of the host at which the concentration of the host–guest complex [HG] is maximum gives the stoichiometry of the complex.

Calculation of the detection limit⁶³

The detection limit was calculated using the fluorescence titration data. The emission of compounds was measured 5 times, and the standard deviation of the blank measurement was achieved. To obtain the slope, fluorescence intensities were plotted as concentrations of metal ions. The detection limits were calculated using the equation: detection limit = $3\sigma/k$, where σ is the standard deviation of the blank measurement, and k is the slope.

Ionic conductivity measurements

The ionic conductivity of the gels of **3** and **4** was measured by means of impedance spectroscopy (Hioki 3532-50 LCR Hi-Tester from 42 Hz–2 MHz (signal amplitude, 0.1 V)). For measurements, the sample was placed between two parallel steel electrodes with a Teflon ring spacer in a constant-volume cylindrical cell. The measurements of the gels were taken isothermally from 273 K to 358 K in temperature steps of 5 K. The temperature was controlled using a Eurotherm temperature controller and temperature constancy of ± 0.2 K was achieved in the entire range of measurements. At each temperature, the gels were thermally equilibrated for 30 min before the measurement. To examine the arbitrary errors related to the changes in the environmental conditions and contact *etc.* the experiments were repeated several times under identical experimental conditions.

Conflicts of interest

There are no conflicts to declare.

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