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Received 22nd May 2015

Accepted 29th July 2015

DOI: 10.1039/c5sc01859e

www.rsc.org/chemicalscience

Fluorescent carbon dot–molecular salt hydrogels†

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The incorporation of functionalised carbon nanodots within a novel low molecular weight salt hydrogel derived from 5-aminosalicylic acid is reported. The carbon dots result in markedly enhanced gelation properties, while inclusion within the hydrophobic gel results in a dramatic fluorescence enhancement for the carbon nanomaterials. The resulting hybrid CD gels exhibit a useful sensor response for heavy metal ions, particularly Pb²⁺.

Introduction

Impressive progress has been made in photoluminescent (PL) carbon-based dots (CDs) since their discovery in 2004,¹ in many applications such as bioimaging, drug delivery and analytical sensing. Particular applications include engineering bright nanoprobe towards pollutants and metal ions^{2–7} because of their stability, high water solubility, biocompatibility low toxicity, easy surface modification and excellent PL properties. To date CDs have been used in aqueous solution, however incorporating them into a gel medium offers particular promise in stabilising and immobilising the nanoparticles, and may result in interesting effects on the properties of the gel–nanomaterial hybrid itself. Moreover, shielding the CDs within a more hydrophobic environment may result in considerable enhancement of the CD fluorescence. Recently, a few researchers have incorporated carbon nanomaterials in ionic liquids or into agarose hydrogels to modify their electrochemical properties,⁸ as part of a sensing system for heavy metal ions⁹ or as a vehicle for drug delivery.¹⁰

Gels are flexible, soft materials that behave as solids on the analytical timescale, but are composed predominantly of a fluid phase.^{11,12} There have been a number of interesting recent reports concerning the incorporation of nanomaterials into gel phase media.^{13–19} In particular, introduction of carbon nanomaterials such as carbon nanotubes and graphene has been shown to result in gel stabilization and modification of the carbon nanomaterial properties.¹³ There has been considerable

recent interest in small-molecule supramolecular gels composed of self-assembled fibrillar networks (SAFINs) derived from relatively well-defined interactions between low molecular weight gelators (LMWG).^{11,20–29} These small-molecule SAFINs offer the opportunity to design desirable gel properties. Gelator preparation is generally straightforward allowing considerable synthetic versatility and variability, and the gels often exhibit readily reversible gelation behaviour.¹² The range of structure types of LMWG is very broad, comprising amides, nucleobases, fatty acids and dendrimers, for example.¹² Within this context, bis(urea) derivatives^{30–34} are particularly versatile because the gel-forming tendency of the urea groups is tolerant of variations in the spacer between them and the peripheral substituents, allowing the design of tailored gelators for targeted applications such as separations,³⁵ drug delivery,^{36–38} polymer templating^{39,40} and as media to control crystal growth.^{41–44}

In this work we report the preparation, properties and ion sensing applications of novel carbon dot–hydrogel hybrid nanomaterials using LMWGs designed to provide a hydrophobic environment suitable for fluorescent ion sensing within an aqueous medium. While polymer gels have been used in conjunction with CDs,^{9,18,19} to the best of our knowledge this work represents the first incorporation of CDs within an LMWG-based gel medium.

Results and discussion

Synthesis and characterization of bis(urea) gels

The bis(urea) gelators of type **1** are based on a bolaamphiphile design⁴⁵ that incorporates a hydrophobic diphenylmethane-derived central spacer (which we have found commonly results in robust gelation characteristics in bis(urea) systems⁴⁶) in conjunction with ionisable salicylic acid based peripheral substituents capable of imparting water solubility. Related urea and amide carboxylic acid systems have been studied previously.^{47–54} The bis(2-ureido-2-propyl)-*m*-phenylene analogues of type **2** represent comparator examples that are expected to be poor gelators as a result of the steric hindrance around the urea

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† Electronic supplementary information (ESI) available: Fluorescent emission spectra of CD systems and tables of gelation experiments and single crystal structure determination in CIF format. CCDC 1402495. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c5sc01859e

carbonyl group. As a result compounds with this spacer group are frequently more crystalline than most bis(ureas). In previous work we have found they act as useful model compounds allowing structural insight into the system because they are easier to isolate in a form suitable for single crystal X-ray diffraction studies.

The synthesis of the salt gelators **1a** and **2a** was carried out in a one-step reaction between the desired isocyanate and 5-aminosalicylic acid in the presence of triethylamine base. The presence of the triethylammonium cations and the loss of the acid proton in salts **1a** and **2a** was clearly evident in the ^1H NMR spectra and analytical data of the materials, and was confirmed by X-ray crystallography in the case of **2a** (*vide infra*), Fig. 1. The analogous free acid **1b** was prepared by sonicating **1a** in 1 M hydrochloric acid for 30 minutes. Compound **2b** was prepared by dissolving **2a** in water followed by the addition of 1 M hydrochloric acid upon which the compound precipitated.

Compounds of type **1** and **2** as both free acids and triethylammonium salts were screened for gelation and crystallisation in a wide range of solvents. All gelation experiments were carried out at 1 wt% using 5 mg of **1a** in 0.5 mL of solvent. Samples were sonicated for 30 seconds followed by careful heating in a sealed vial until the solid dissolved. Gelation was observed on cooling to ambient temperature.

Compound **1** proved to be a versatile gelator in both salt (**1a**) and neutral (**1b**) forms. Salt **1a** forms gels in a variety of solvents outlined in Table S1 (ESI †). Gels formed upon heating and subsequent cooling in a total of 26 out of 50 solvents studied and partial gels were also observed in 4 solvents. The compound

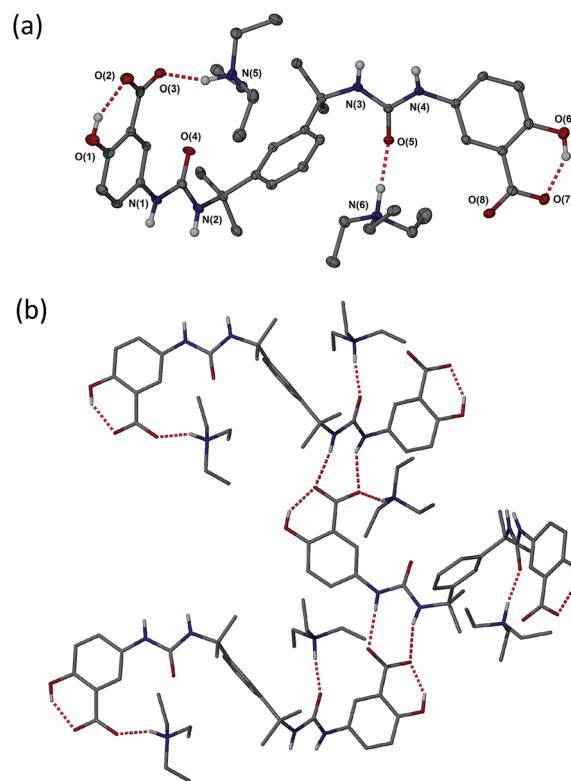


Fig. 2 (a) Molecular structure of **2a**·propan-1-ol (disordered propan-1-ol and CH hydrogen atoms omitted). (b) Hydrogen bonding in the structure of **2a**·propan-1-ol. Selected hydrogen bonding distances (Å): O1...O2 2.577(5), N1...O8 2.811(6), N4...O3 2.832(5), N5...O3 2.700(6), O6...O7 2.505(5) and N6...O5 2.775(5).

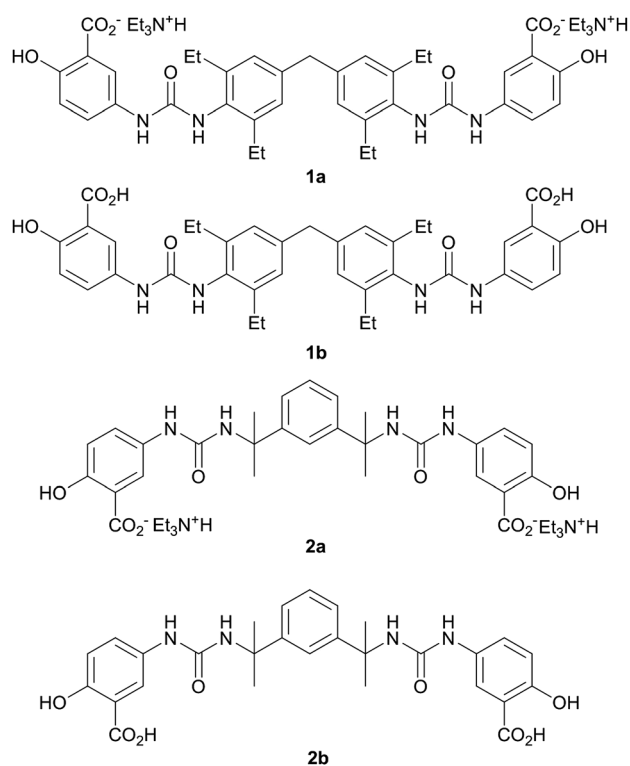


Fig. 1 The triethylammonium salt gelators **1a** and **2a**, and the analogous free acids **1b** and **2b**.

gels a range of polar solvents with both hydrogen bond donor (e.g. alcohols) and acceptor character (e.g. pyridine derivatives). The neutral acid form **1b** gels in 12 of the 50 solvents studied and partial gels were also observed in 10 solvents outlined in Table S2 (ESI †). Interestingly, room temperature gelation of **1a** on sonication was also observed in a number of instances (Table S3 †).^{55,56} Comparison between room temperature sonogels and gels produced by a combination of ultrasound and thermal methods revealed that the thermally treated materials are more robust and longer lived, although some minor degradation was evident in the colouration of some gels (Tables S1 and S3,† Fig. 3). Some of the gels formed by the sonication-only process appeared to collapse more readily and in the case of ethanol, only persist for around one hour.

While compound **1b** gels only organic solvents, the behaviour of salt **1a** in water is particularly intriguing with no gelation

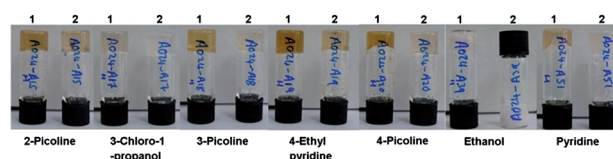


Fig. 3 Comparison between gels of **1a** formed by a combination of ultrasound and heat treatment (labelled 1) and room temperature sonogels (2) at 1 wt%.

observed for several days after dissolution at 1 wt%. However from 72 hours onwards partial gels began to form and after 10 days robust, transparent self-supporting gels form implying very slow fibre assembly within the competitive aqueous medium (ESI, Fig. S5†). Increasing the concentration to 2 wt% results in gelation in less than one day. Gelation is also markedly accelerated by lowering the pH (gelation at 1 wt% is rapid in the presence of 1 μL of concentrated sulphuric acid). Subsequent addition of base in the form of NEt_3 results in collapse of the gel.

As anticipated, compound **2** with its bulky spacer did not form gels in any solvents (Tables S4 and S5†), an outcome attributed to steric hindrance of the urea carbonyl group preventing urea tape formation (in comparison the aromatic groups of **1** may rotate to be orthogonal to the urea plane). Crystals of **2a** suitable for single crystal X-ray structure determination were obtained by slow evaporation of a solution in propan-1-ol. While not from a gel, the crystal structure serves to provide insight into the kinds of interactions possible between gelators bearing these functional groups, the triethyl ammonium counter cation and the solvent. The crystals proved to be a 1 : 1 solvate with two crystallographically unique HNEt_3^+ cations hydrogen bonded to one carboxylate group and one urea carbonyl group of a single unique bis(urea) molecule in the presence of a highly disordered molecule of propan-1-ol. The solvent molecule hydrogen bonds to a phenolic hydroxyl group. Typically gels of bis-ureas arise from the formation of α -tape hydrogen bonding interactions between the urea groups in neighbouring molecules based on a six-membered hydrogen bonded ring.^{34,57,58} In the present structure however, the Et_3NH^+ cations effectively inhibit urea tape formation by hydrogen bonding to one of the urea groups and sterically blocking the other. Instead, the strong hydrogen bond acceptor nature of the carboxylate groups results in the formation of an alternative, twisted non-planar 8-membered hydrogen-bonded ring between the urea functionalities and the carboxylate groups (Fig. 2b). The phenolic OH groups are both involved in intramolecular hydrogen to the carboxylate acceptors.

Carbon dot hybrid gels

Despite the slow assembly of hydrogels of **1a** the final gels proved to be robust and transparent and hence were investigated as host media for fluorescent carbon dots. Four different types of carbon dot were investigated. High luminescence CDs (c-CDs) were prepared from microcrystalline cellulose (MCC) whereas low luminescence CDs were obtained from multi-walled carbon nanotubes (MWCNTs). The nanotube-derived CDs were passivated with acetone (p-CDs) and in some cases subsequently functionalised to give thiol rich (t-CDs) or amine rich surfaces (a-CDs) to improve their PL properties and ability to interact with the gel and with potential analytes. CD-gel hybrids were prepared by sonicating 0.5 to 4 wt% of **1a** in aqueous solutions of the carbon dots ranging from 1–60 mg mL^{-1} . The critical gelation concentration proved to be 1 wt% with solutions of 0.5 wt% not forming self-supporting gels.

Remarkably, unlike the hydrogels which form over a period of 10 days at 1 wt%, gelation of all of the CD-gel hybrids occurred

in a matter of minutes. This fortuitous result renders the hybrid gels much more convenient to prepare and study than the pure hydrogels and hence is of marked benefit in a practical sensing context. While the mechanism for gelation is not known the CDs may well act as a hydrophobic nucleation and cross-linking node for SAFIN assembly interacting with the fibres by hydrophobic and π - π stacking interactions. The effect is not related to the increased gelation propensity on acidification since the CD solutions are slightly basic, pH 8–9. Gels containing high concentrations of CDs proved to be opaque because of the very intense absorption of the CDs, however dilution to 1 wt% solutions containing 1 mg mL^{-1} CDs gave transparent gels suitable for fluorescence study, although at this low CD concentration gelation was less readily reproducible than at higher concentrations. We postulate that hydrogen bonding to the solubilising Et_3NH^+ cations may interfere with the gel assembly process and accordingly we investigated counter cation metathesis by adding a water soluble ionic liquid 1-butyl-3-methylimidazolium tetrafluoroborate (BMIM- BF_4). The BMIM cation does not possess hydrogen bond donor functionality and hence should not interfere with gelation. Pure ionic liquids such as BMIM- BF_4 are of interest as ionogels for hosting magnetic nanoparticles, for example, and silicon-functionalised carbon dots have previously been incorporated into ionic liquid based ionogels.^{59–61} Compound **1a** does not gel pure BMIM- BF_4 ; however, addition of 1–40 wt% of the ionic liquid to the gel-CD mixture gave robust, reproducible gels for all types of CDs. The metathesis with BMIM cations proved highly effective with the resulting hybrid **1a**-CD-BMIM hydrogels forming rapidly at an optimum concentration of 2 wt% BMIM- BF_4 to give robust, transparent hybrid nanomaterials suitable for fluorescence study.

The optimised nanomaterials of composition 1 wt% **1a** in 500 μL of CD solution at 1 mg mL^{-1} containing 2% BMIM- BF_4 are shown in Fig. 4. The hybrid gels were characterised by frequency and stress sweep rheometry and by SEM. The rheological properties of the four different CD hybrid materials were similar to one another with all exhibiting an elastic modulus G' greater than the viscous modulus G'' , with G' invariant with frequency, Fig. 5. At the concentration used the gels are relatively weak with G' values of 100–300 Pa and yield stress around 1–10 Pa. The gels containing the amine-functionalised CDs proved generally to be considerably more robust than the unfunctionalised c-CDs which may indicate a specific hydrogen bonding interaction between the amine groups and the carboxylate functionalities of the gelators. The SEM images of the dried gels showed a fibrous structure consistent with similar SAFIN-based hydrogels⁶² with small fibres of average diameter of less than 30 nm consistent with the transparent appearance of the gels. In comparison the hydrogels before introduction of CDs give slightly larger, more tape-like fibres (ESI Fig. S4†).

CD-gel fluorescence

All the CDs in aqueous solution were fluorescent with a full width at half maximum (FWHM) about 100 nm which indicates their homogeneity. The bottom-up synthesised c-CDs proved to be the most fluorescent and are known to possess a higher





Fig. 4 Optimised ionic-liquid CD–Gel hybrid materials of composition 1 wt% **1a** in 500 μL of CD solution at 1 mg mL^{-1} containing 2% BMIM-BF₄ (left-to-right c-CD, p-CD, t-CD and a-CD).

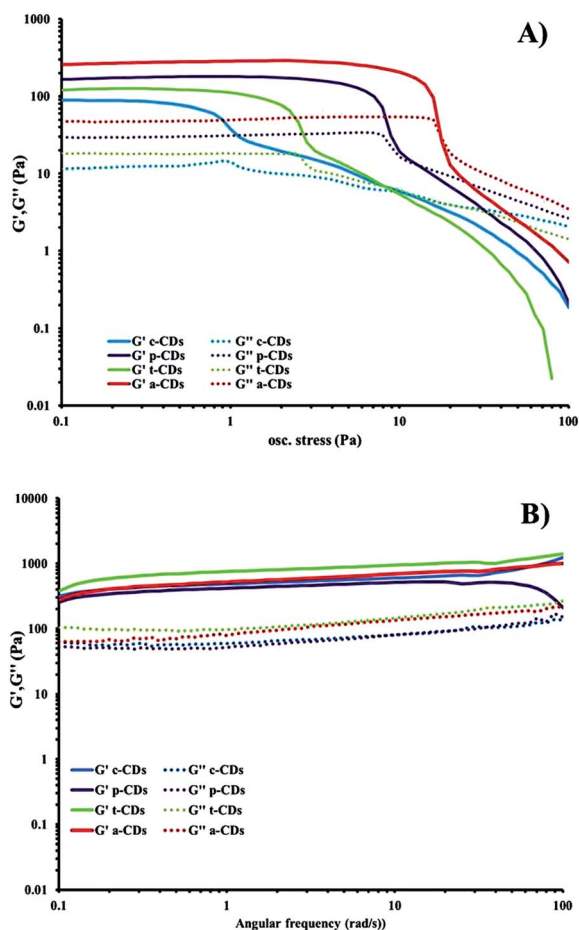


Fig. 5 (a) Stress sweep and (b) frequency-sweep rheology of dilute ionic-liquid CD–gel hybrid materials of composition 1 wt% **1a** in 500 μL of CD solution at 1 mg mL^{-1} containing 2% BMIM-BF₄.

quantum yield.⁶³ The maximum excitation wavelength was 370 nm, shifting to 365 nm for the p-CDs, t-CDs and a-CDs. Fig. S1† shows the fluorescent spectra of all the CD solutions. Consistent with previous reports,^{64–66} the functionalised CDs were more fluorescent than the passivated p-CDs (Fig. 6).

Incorporation of the CDs into the gels resulted in a striking enhancement of the CD emission intensity by at least an order of magnitude for all types of CDs studied. It is well-known that the PL intensity varies significantly with the size and core structure due to the synthetic route and precursor used, the

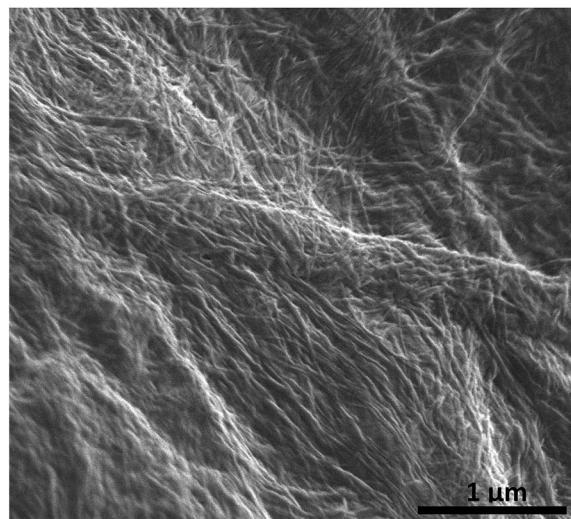


Fig. 6 SEM micrograph of the chromium coated dried (xerogel) ionic-liquid a-CD–gel hybrid materials based on **1a**. The carbon dots of average radius 3 nm are too small to be observed.

surface functionalities of CDs, as well as with the environment. As the nanodots in this case are all of similar size, the strong emission enhancement of CDs comes from the interaction of the CDs with the gelator instead of a quantum size effect. Fluorescence spectra for 1 wt% gels of 1 mg mL^{-1} CD solutions in the absence of the ionic liquid are shown in Fig. 7. The ionic liquid itself is also fluorescent, complicating the spectra, however the marked emission enhancement is also evident in the ionic liquid CD–gel hybrid materials, Fig. S2 (ESI†). The hydrophobic nature of the gels is likely to significantly reduce solvent quenching and the fact that the emission enhancement is so marked suggests significant interaction between the gel fibres and the CDs in a synergistic way, consistent with the marked effect the CDs have on the gelation properties.

The high intensity fluorescence of these novel CD gel hybrids and the possibility of tuning the properties of the materials by modification of the gelator, imidazolium cation and the size and surface functionality of the CDs offers interesting possibilities for their application as luminescent sensors, *e.g.* as environmental nanoprobe. CDs have been shown to be an excellent sensing platform for heavy metals,^{63,64} and the interaction of metal ions with the CD surface functionality may well be significantly influenced by the inclusion of the CDs within a LMWG hydrogel medium. In order to establish the feasibility of heavy metal sensing using hybrid CD gels the effect of added lead(II) and mercury(II) on the PL of the new hybrid gel system was investigated. The fluorescence of aqueous p-CD, t-CD and a-CDs solutions is moderately quenched by 10 $\mu\text{g mL}^{-1}$ of Hg²⁺ and Pb²⁺ ($I_0/I = 1.1–1.25$) while the luminescence of c-CDs is not affected. In the hybrid CD gel system, in addition to the marked increase in overall PL intensity, the response to Hg²⁺ and Pb²⁺ changes differentially. The response to Pb²⁺ of the gels containing p-CDs and t-CDs is markedly attenuated such that the luminescence is essentially unaffected, whereas the system containing the a-CDs exhibits a marked enhancement in its



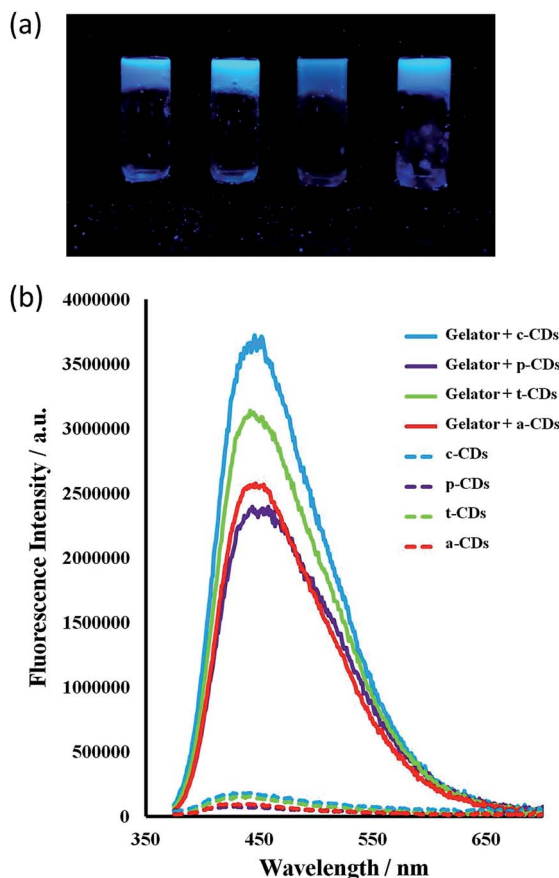


Fig. 7 (a) a-, c-, p- and t-CD gels 1 wt% under 365 nm UV irradiation. (b) Fluorescence emission spectra of aqueous CD solutions (1 mg mL^{-1} , $\lambda_{\text{ex}} = 365 \text{ nm}$ for c-CDs and 370 nm for the others) and for 1 wt% CD-gels with **1a** at the same CD concentration. Slits at 2 nm .

Pb^{2+} response in the gel phase (I_0/I increases from 1.13 to 1.32), Fig. 8. In contrast the a-CD gel exhibits essentially no response to Hg^{2+} . This differential behaviour suggests that the CD gels may provide a basis for discriminating closely related heavy metal ions in the environment. As a control the metal ion response was also examined in CD solution in the presence of the gelator without gel formation. The presence of the gelator in solution had no effect on the t-CD response, attenuated the response of the a-CD system and enhanced the response of the p-CDs. The significant and differential difference in sensor behaviour in the gel state is an interesting phenomenon. To further probe selectivity the response of a-CD-containing hydrogels (a-CDGs) as sensing probe towards a range of metal ions, namely Pb^{2+} , Hg^{2+} , Co^{2+} , Fe^{3+} and Mg^{2+} was examined (ESI, Fig. S3†). The system proved to be clearly Pb^{2+} selective in the hydrogel state. The a-CDGs thus represent a new format of selective sensor for the detection of lead(II). In future work we will systematically explore the fluorescent response of these novel hybrid nanomaterials to a broader range of analytes. A further interesting application platform could also involve the use of freeze dried hybrid CD hydrogels coated with TiO_2 allowing them to float on water as reporter medium.⁶⁷

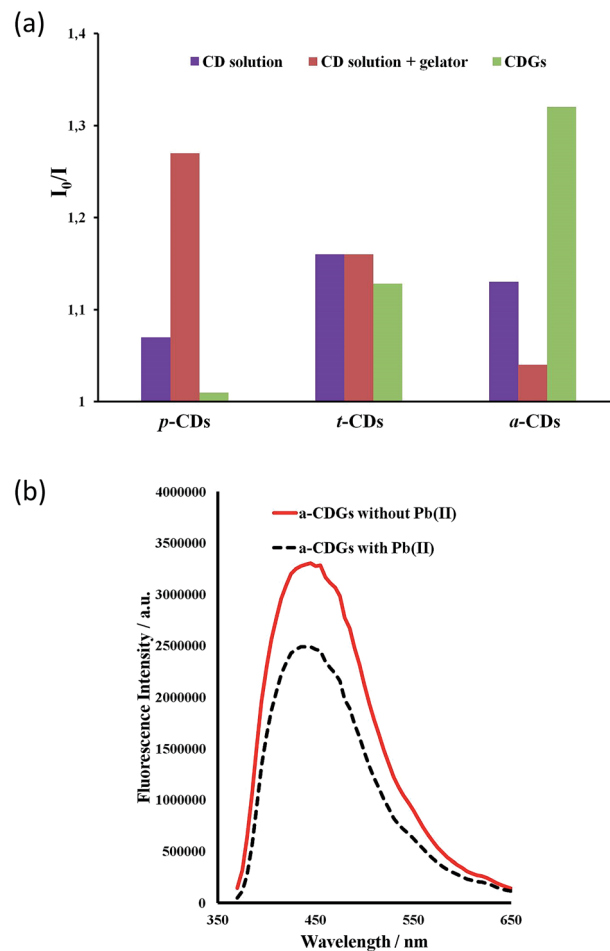


Fig. 8 (a) Photoluminescent response to $10 \mu\text{g mL}^{-1}$ of Pb^{2+} to CD solutions, CD-containing hydrogels (CDGs) and CD solutions in the presence of dissolved gelator (1 mg mL^{-1} , $\lambda_{\text{ex}} = 370 \text{ nm}$) and for 1 wt% CD-gels with **1a** at the same CD concentration. (b) Fluorescence emission spectra for a-CD gel samples with and without Pb^{2+} ($10 \mu\text{g mL}^{-1}$).

Conclusions

Inclusion of luminescent carbon dots within a hydrophobic low molecular weight hydrogel results in a dramatic enhancement of their luminescence intensity as well as an unexpected strengthening of the gels and marked increase in their rate of self-assembly. In the gel phase the CDs exhibit a differential photoluminescent response to the presence of heavy metal ions. The novel hybrid CD-gel nanostructures reported herein offer interesting potential as highly tunable luminescent media with possible applications in environmental and biomedical heavy metal ion sensing. The potential biocompatibility of these nanostructures which are derived from non-toxic precursors may open a window for their use in medical applications such as diagnosis, drug activation or slow release.

Experimental

Reagents and instrumental

Sulphuric acid (95–98%), nitric acid (69%), ethanol (99%) and chloroform were purchased from PANREAC, S.A.U. (Barcelona,



Carbon dot solutions

Four different types of CDs were synthesised as described previously.^{22–25} Briefly, CDs were prepared by acid thermohydrolyses using microcrystalline cellulose (MCC) or multi-walled carbon nanotubes (MWCNTs) as carbon precursors. Highly luminescent dots (c-CDs) were obtained using MCC, while less fluorescent CDs were obtained from MWCNTs. Later, the last CDs were passivated with acetone (p-CDs) to boost their PL properties. The p-CDs were then surface functionalised either with cysteamine to give a thiol-rich surface (t-CDs) or *N*-Boc-ethylenediamine to give an amine rich surface (a-CDs). All the CDs synthesised were neutralised and purified by precipitation with ethanol at low temperatures.

Preparation and optimization of the CD–salt hydrogels

Gels were prepared by dissolving a carefully weighed amount of bis(urea) **1** into an aqueous solution of the relevant CD heating until the gelator dissolved and sonicating the as-prepared mixture three times for few seconds each time. This procedure resulted in gel formation over the course of a few minutes, however at low concentrations of the gelator (≤ 1 wt%) in the presence of 1 mg mL^{-1} CD solution the gels proved relatively weak. Addition of BMIM-BF₄ (2 wt%) to metathesise the hydrogen bonding triethyl ammonium cation resulted in stable, robust and transparent gels.

Fluorescence studies

The differences in the fluorescence intensity and the maximum excitation wavelength between liquid and solid were studied for: c-CD, p-CD, t-CD and a-CD containing gels in comparison with aqueous solutions. Each measurement was carried out three times, $\lambda_{\text{ex}} = 365$ and 370 nm , 2 nm excitation and emission slit widths. Measurements were performed with a PTI QuantaMaster™ Spectrofluorometer.

Analyses of metal ions

Aliquots of standards (Pb²⁺, Hg²⁺, Co²⁺, Fe³⁺ and Mg²⁺ at $10 \mu\text{g mL}^{-1}$) were dropped onto the CD solution (before gel formation) and shaken for 1 min. In the case of CD gels as fluorescent probes, formation of the gels was performed as previously described.

Rheology

Rheological measurements were carried out with an AR2000 rheometer at 10°C . Frequency sweep experiments were performed at a constant oscillation stress of 1 for the angular frequency of 0.1–100, while the stress sweep was performed for the oscillation stress of 0.1–100.

Scanning electron microscopy

SEM samples were dried in air at room temperature for 2 days, coated with 3 nm of chromium using a Cressington 328 Ultra High Resolution EM Coating System, and imaged using an FEI

Helios NanoLab DualBeam microscope in immersion mode, with typical beam settings of 1.5 kV and 0.17 nA.

Acknowledgements

We thank the Engineering and Physical Sciences Research Council and the Spanish Ministry of Innovation and Science for funding projects EP/J013021/1 and CTQ2011-23790, respectively. This work was also supported by the CEIa3 (Agrifood Campus of International Excellence).

Notes and references

- 1 X. Xu, R. Ray, Y. Gu, H. J. Ploehn, L. Gearheart, K. Raker and W. A. Scrivens, *J. Am. Chem. Soc.*, 2004, **126**, 12736–12737.
- 2 H. M. R. Gonçalves, A. J. Duarte and J. C. G. Esteves da Silva, *Biosens. Bioelectron.*, 2010, **26**, 1302–1306.
- 3 W. Lu, X. Qin, S. Liu, G. Chang, Y. Zhang, Y. Luo, A. M. Asiri, A. O. Al-Youbi and X. Sun, *Anal. Chem.*, 2012, **84**, 5351–5357.
- 4 F. Du, F. Zeng, Y. Ming and S. Wu, *Microchim. Acta*, 2013, **180**, 453–460.
- 5 J. Wei, J. Ren, J. Liu, X. Meng, X. Ren, Z. Chen and F. Tang, *Biosens. Bioelectron.*, 2014, **52**, 304–309.
- 6 H. Dai, Y. Shi, Y. Wang, Y. Sun, J. Hu, P. Ni and Z. Li, *Sens. Actuators, B*, 2014, **202**, 201–208.
- 7 X. Cui, L. Zhu, J. Wu, Y. Hou, P. Wang, Z. Wang and M. Yang, *Biosens. Bioelectron.*, 2015, **63**, 506–512.
- 8 C. T. Nguyen, Y. Zhu, X. Chen, G. A. Sotzing, S. Granados-Focil and R. M. Kasi, *J. Mater. Chem. C*, 2015, **3**, 399–408.
- 9 N. Gogoi, M. Barooah, G. Majumdar and D. Chowdhury, *ACS Appl. Mater. Interfaces*, 2015, **7**, 3058–3067.
- 10 C. T. Nguyen, T. H. Tran, X. Lu and R. M. Kasi, *Polym. Chem.*, 2014, **5**, 2774–2783.
- 11 R. G. Weiss, *J. Am. Chem. Soc.*, 2014, **136**, 7519–7530.
- 12 P. Terech and R. G. Weiss, *Chem. Rev.*, 1997, **97**, 3133–3160.
- 13 S. K. Samanta, K. S. Subrahmanyam, S. Bhattacharya and C. N. R. Rao, *Chem.–Eur. J.*, 2012, **18**, 2890–2901.
- 14 S. K. Samanta, A. Pal, S. Bhattacharya and C. N. R. Rao, *J. Mater. Chem.*, 2010, **20**, 6881–6890.
- 15 B. Adhikari and A. Banerjee, *Soft Matter*, 2011, **7**, 9259–9266.
- 16 Z. Xie, F. Wang and C.-y. Liu, *Adv. Mater.*, 2012, **24**, 1716–1721.
- 17 E. Sotelo-Gonzalez, A. M. Coto-Garcia, M. T. Fernandez-Argüelles, J. M. Costa-Fernandez and A. Sanz-Medel, *Sens. Actuators, B*, 2012, **174**, 102–108.
- 18 S. Hu, Q. Zhao, Y. Dong, J. Yang, J. Liu and Q. Chang, *Langmuir*, 2013, **29**, 12615–12621.
- 19 A. Quaranta, S. Carturan, A. Campagnaro, M. Dalla Palma, M. Giarola, N. Daldosso, G. Maggioni and G. Mariotto, *Thin Solid Films*, 2014, **553**, 188–192.
- 20 V. A. Mallia, P. D. Butler, B. Sarkar, K. T. Holman and R. G. Weiss, *J. Am. Chem. Soc.*, 2011, **133**, 15045–15054.
- 21 J. A. Foster, R. M. Edkins, G. J. Cameron, N. Colgin, K. Fucke, S. Ridgeway, A. G. Crawford, T. B. Marder, A. Beeby, S. L. Cobb and J. W. Steed, *Chem.–Eur. J.*, 2014, **20**, 279–291.



- 22 K. L. Morris, L. Chen, J. Raeburn, O. R. Sellick, P. Cotanda, A. Paul, P. C. Griffiths, S. M. King, R. K. O'Reilly, L. C. Serpell and D. J. Adams, *Nat. Commun.*, 2013, **4**, 1480.
- 23 X. Yang, G. Zhang and D. Zhang, *J. Mater. Chem.*, 2012, **22**, 38–50.
- 24 G. Yu, X. Yan, C. Han and F. Huang, *Chem. Soc. Rev.*, 2013, **42**, 6697–6722.
- 25 M. George and R. G. Weiss, *Acc. Chem. Res.*, 2006, **39**, 489–497.
- 26 J. W. Steed, *Chem. Commun.*, 2011, **47**, 1379–1383.
- 27 D. K. Smith, *Nat. Chem.*, 2010, **2**, 162–163.
- 28 A. Y.-Y. Tam and V. W.-W. Yam, *Chem. Soc. Rev.*, 2013, **42**, 1540–1567.
- 29 P. Dastidar, *Chem. Soc. Rev.*, 2008, **37**, 2699–2715.
- 30 M. de Loos, A. Friggeri, J. van Esch, R. M. Kellogg and B. L. Feringa, *Org. Biomol. Chem.*, 2005, **3**, 1631–1639.
- 31 M. de Loos, B. L. Feringa and J. H. van Esch, *Eur. J. Org. Chem.*, 2005, 3615–3631.
- 32 J. H. van Esch, F. Schoonbeek, M. de Loos, H. Kooijman, A. L. Spek, R. M. Kellogg and B. L. Feringa, *Chem.-Eur. J.*, 1999, **5**, 937–950.
- 33 J. H. van Esch, S. DeFeyter, R. M. Kellogg, F. DeSchryver and B. L. Feringa, *Chem.-Eur. J.*, 1997, **3**, 1238–1243.
- 34 J. W. Steed, *Chem. Soc. Rev.*, 2010, **39**, 3686–3699.
- 35 E. Krieg, H. Weissman, E. Shirman, E. Shimon and B. Rybtchinski, *Nat. Nanotechnol.*, 2011, **6**, 141–146.
- 36 S. Sutton, N. L. Campbell, A. I. Cooper, M. Kirkland, W. J. Frith and D. J. Adams, *Langmuir*, 2009, **25**, 10285–10291.
- 37 P. D. Thornton, R. J. Mart, S. J. Webb and R. V. Ulijn, *Soft Matter*, 2008, **4**, 821–827.
- 38 K. J. C. van Bommel, M. C. A. Stuart, B. L. Feringa and J. van Esch, *Org. Biomol. Chem.*, 2005, **3**, 2917–2920.
- 39 B. Li, L. Tang, L. Qiang and K. Chen, *Soft Matter*, 2011, **7**, 963–969.
- 40 Q. Wei and S. L. James, *Chem. Commun.*, 2005, 1555–1556.
- 41 J. A. Foster, M.-O. M. Piepenbrock, G. O. Lloyd, N. Clarke, J. A. K. Howard and J. W. Steed, *Nat. Chem.*, 2010, **2**, 1037–1043.
- 42 D. K. Kumar and J. W. Steed, *Chem. Soc. Rev.*, 2014, **43**, 2080–2088.
- 43 L. A. Estroff, L. Addadi, S. Weiner and A. D. Hamilton, *Org. Biomol. Chem.*, 2004, **2**, 137–141.
- 44 F. Aparicio, E. Matesanz and L. Sánchez, *Chem. Commun.*, 2012, **48**, 5757–5759.
- 45 C. Dolle, P. Magrone, S. Riva, M. Ambrosi, E. Fratini, N. Peruzzi and P. Lo Nostro, *J. Phys. Chem. B*, 2011, **115**, 11638–11649.
- 46 L. Meazza, J. A. Foster, K. Fucke, P. Metrangolo, G. Resnati and J. W. Steed, *Nat. Chem.*, 2013, **5**, 42–47.
- 47 L. A. Estroff and A. D. Hamilton, *Angew. Chem., Int. Ed.*, 2000, **39**, 3447–3450.
- 48 D. J. Adams and P. D. Topham, *Soft Matter*, 2010, **6**, 3707–3721.
- 49 D. J. Adams, M. F. Butler, W. J. Frith, M. Kirkland, L. Mullen and P. Sanderson, *Soft Matter*, 2009, **5**, 1856–1862.
- 50 A. Bernet, R. Q. Albuquerque, M. Behr, S. T. Hoffmann and H.-W. Schmidt, *Soft Matter*, 2012, **8**, 66–69.
- 51 R. C. T. Howe, A. P. Smalley, A. P. M. Guttenplan, M. W. R. Doggett, M. D. Eddleston, J. C. Tan and G. O. Lloyd, *Chem. Commun.*, 2013, **49**, 4268–4270.
- 52 J. Li, Y. Kuang, Y. Gao, X. Du, J. Shi and B. Xu, *J. Am. Chem. Soc.*, 2013, **135**, 542–545.
- 53 Z. Yang, G. Liang and B. Xu, *Chem. Commun.*, 2006, 738–740.
- 54 S. Fleming and R. V. Ulijn, *Chem. Soc. Rev.*, 2014, **43**, 8150–8177.
- 55 G. Cravotto and P. Cintas, *Chem. Soc. Rev.*, 2009, **38**, 2684–2697.
- 56 C. Deng, R. Fang, Y. Guan, J. Jiang, C. Lin and L. Wang, *Chem. Commun.*, 2012, **48**, 7973–7975.
- 57 L. S. Reddy, S. Basavoju, V. R. Vangala and A. Nangia, *Cryst. Growth Des.*, 2006, **6**, 161–173.
- 58 M. C. Etter, Z. Urbanczyk-Lipkowska, M. Zia-Ebrahimi and T. W. Panunto, *J. Am. Chem. Soc.*, 1990, **112**, 8415–8426.
- 59 J. Yan, J. Liu, P. Jing, C. Xu, J. Wu, D. Gao and Y. Fang, *Soft Matter*, 2012, **8**, 11697–11703.
- 60 Y. Feng, H. Li, Q. Gan, Y. Wang, B. Liu and H. Zhang, *J. Mater. Chem.*, 2010, **20**, 972–975.
- 61 Z.-L. Xie, H.-B. Xu, A. Geßner, M. U. Kumke, M. Priebe, K. M. Fromm and A. Taubert, *J. Mater. Chem.*, 2012, **22**, 8110–8116.
- 62 L. A. Estroff and A. D. Hamilton, *Chem. Rev.*, 2004, **104**, 1201–1217.
- 63 A. Cayuela, M. L. Soriano and M. Valcárcel, *Sens. Actuators, B*, 2015, **207**, 596–601.
- 64 A. Cayuela, M. Laura Soriano and M. Valcárcel, *Anal. Chim. Acta*, 2013, **804**, 246–251.
- 65 A. Cayuela, M. L. Soriano, M. C. Carrión and M. Valcárcel, *Anal. Chim. Acta*, 2014, **820**, 133–138.
- 66 A. Cayuela, M. L. Soriano and M. Valcárcel, *Anal. Chim. Acta*, 2015, **872**, 70–76.
- 67 J. T. Korhonen, M. Kettunen, R. H. A. Ras and O. Ikkala, *ACS Appl. Mater. Interfaces*, 2011, **3**, 1813–1816.

