To gel or not to gel: correlating molecular gelation with solvent parameters
To gel or not to gel: correlating molecular gelation with solvent parameters†

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Rational design of small molecular gelators is an elusive and herculean task, despite the rapidly growing body of literature devoted to such gels over the past decade. The process of self-assembly, in molecular gels, is intricate and must balance parameters influencing solubility and those contrasting forces that govern epitaxial growth into axially symmetric elongated aggregates. Although the gelator–gelator interactions are of paramount importance in understanding gelation, the solvent–gelator specific (i.e., H-bonding) and nonspecific (dipole–dipole, dipole-induced and instantaneous dipole induced forces) intermolecular interactions are equally important. Solvent properties mediate the self-assembly of molecular gelators into their self-assembled fibrillar networks. Herein, solubility parameters of solvents, ranging from partition coefficients (log $P$), to Henry’s law constants (HLC), to solvatochromic parameters ($\alpha$, $\beta$, and $\pi$), and to Hansen solubility parameters ($\delta_p$, $\delta_h$, $\delta_s$), are correlated with the gelation ability of numerous classes of molecular gelators. Advanced solvent clustering techniques have led to the development of a priori tools that can identify the solvents that will be gelled and not gelled by molecular gelators. These tools will greatly aid in the development of novel gelators without solely relying on serendipitous discoveries. These tools illustrate that the quest for the universal gelator should be left in the hands of Don Quixote and as researchers we must focus on identifying gelators capable of gelling classes of solvents as there is likely no one gelator capable of gelling all solvents.

What is a molecular gel?

Gels are comprised of chemically diverse systems that are frequently easier to recognize than to define.1,2 Since the introduction of a gel theory by Thomas Graham in 1861, the definition of what is a gel has been evolving.3 Among the numerous attempts made to define gels, Dr Dorothy Jordan Lloyd proposed that gels must be composed of two components with one being liquid at the temperature of consideration and the other one being solid (i.e., the gelator). As well, the system must have mechanical properties of a solid.2 This definition is useful in determining a gel but, imprecise because not all colloids are gels and not all gels are colloids.4 Over several decades, the definition of a gel evolved to the point where Hermans depicted gels as “coherent colloid disperse systems of at least two components that exhibit mechanical properties characteristic of the solid state”1 and “both the dispersed component and dispersing medium extend themselves continuously throughout the whole system”.5 Due to the exclusivity of this definition, Ferry provided a less vigorous and more descriptive one: “A gel is a substantially diluted system which exhibits no steady state flow.”6 In order to link the microscopic and macroscopic properties, a substance can be classified as a gel if it: (1) has a continuous microscopic structure with macroscopic dimensions that is permanent on the time scale of an analytical experiment and (2) is solid-like in its rheological behavior despite being mostly liquid.1 Depending on the continuous structures of gels discussed, Flory proposed a scheme and classified gels into four categories: (1) well-ordered lamellar structures, including gel mesophases; (2) covalent polymeric networks, completely disordered; (3) polymer networks formed through physical aggregation, predominantly disordered, but with regions of local order; (4) particulate, disordered structures.7 Herein, molecular gels reside in category three and perhaps four.

Unlike polymeric gels, molecular gels are comprised of low molecular weight gelators (LMOGs) (arbitrarily limited to <3000 Da). The molecules self-assemble via highly specific, non-covalent interactions that lead usually to elongated fibrillar structures resulting in the formation of an entangled self-assembled network (a SAFIN).8,9 The thermally-activated formation...
of molecular gels follows a multi-step process. The initial step of gelation involves dissolution of LMOGs under specific conditions, such as elevated temperature, to obtain a solution or sol. Upon cooling, super-saturation drives the aggregation of LMOGs via stochastic nucleation. Unlike common crystallization processes where macroscopic phase separation occurs and bulk solids and liquids are visible, the gelation here involves microscopic phase separation. The nucleation event requires highly specific interactions that promote preferential 1-dimensional (1D) growth. These interactions include hydrogen-bonding, π–π stacking, electrostatic interactions, van der Waals interactions and so on. The consequent fibers formed from 1D growth can be described in many cases as “crystal-like” and they play the same role as polymer chains in polymeric gels. The morphology of these fibers may be tubules, strands, tapes, chiral ribbons or any aggregate with a large aspect ratio. The junction zones and branching between these SAFiN strands are responsible for the rigidity of the microstructure of the gel matrixes. The junction zones serve as the “glue” that combines 1D fibers into 3-dimensional (3D) networks that pervade the whole system and entrap the liquid component macroscopically via capillary forces and surface tension.

Practically, it is sometimes difficult to determine experimentally whether a material is a gel. For the purpose of this review, it is extremely important to be as inclusive as possible when defining gelation ability. The advantage of advanced rheological techniques in defining gels is that they are able to differentiate weak and strong gels. However, these determinations are very time consuming. Given the extremely large number of materials to be screened, the authors here have elected to use a simpler
rheological technique, the inverted tube method.\textsuperscript{14} Specifically, a solid–liquid mixture in a glass vial was heated above the sol–gel transition temperature (\textit{i.e.}, where all of the solid was dissolved) and allowed to cool to room temperature. After being stored at room temperature for 24 h, it was inverted for 1 h to observe flow. If no flow was detected, it was classified as a gel.

An arising question in the expanding field of molecular gels

Rational design of small molecular gelators has remained elusive, despite the vast and rapidly growing body of literature devoted to such gels over the past decade (Fig. 1).\textsuperscript{15} Determining gelation by small molecules is still an empirical science, and the vast majority of new gelators are discovered serendipitously.\textsuperscript{16} In polymer gels, the basic elements of the 3D network are 1D objects (\textit{i.e.}, chains with covalent links between monomer units); conversely, the 3D network for molecular gels are composed of zero-dimensional (0D) objects (\textit{i.e.}, molecules that must self-assemble through non-covalent interactions initially into 1D objects and then into 3D networks).\textsuperscript{17,18} The fibrillar networks (SAFiNs) form spontaneously via aggregation-nucleation-growth pathways of the small molecule gelators.\textsuperscript{19–31}

![Histogram of citations by year to ‘organogel(s)’ in the web of science.](image)

At this time, the research community is faced with more questions than answers concerning molecular gels: (1) why do some molecules self-recognize and orient into supramolecular networks while others do not? (2) Why do these fibrillar aggregates grow without minimizing the surface area between the liquid and crystal interface, and hence have an excess of interfacial free energy? (3) Why do slight variations in molecular structure frequently have drastic consequences in the nanoscopic, microscopic and mesoscopic properties of the gels? (4) How does each molecule of these building blocks immobilize macroscopically up to one thousand (or more) liquid molecules? The intricate nature of SAFiNs lies in their hierarchical assembly process that must arise as a consequence of weak non-covalent interactions that include hydrogen-bonding, $\pi-\pi$ stacking and van der Waals interactions.\textsuperscript{32} However, the possibility of creating, \textit{a priori}, or discovering serendipitously a ‘universal’ gelator for all solvents seems impossible given the importance of the solvent–gelator interplay in directing self-assembly.

The quest for rational design criteria in creating new gelators

Each class of molecular gelators can gel a limited set of solvents. No universal law may be applied to all gelator–solvent systems. As an example, originating with the simplest reported molecular gelators, $n$-alkanes, we have incrementally increased the complexity of the structures in an attempt to identify the features that lead to the most efficient gelators.\textsuperscript{33–38} Alkanes are driven to assemble via van der Waals interactions. By inserting an oxygen at various positions along the carbon chain, other interactions such as dipole–dipole and hydrogen-bonding come into play, leading in some cases to SAFiNs and gels. Further addition of a terminal carboxylic acid to a C18 hydrocarbon chain leads to one of the most widely studied, highly efficient molecular gelators known to date, R-12-hydroxystearic acid (HSA).\textsuperscript{21,22,33,34,36,37,39–55} The efficiency of gelation in silicone oil and aromatic solvents can be improved by modifying the carboxylic acid of HSA. Thus, the efficiency of gelation is in the order – primary amide $>$ carboxylic acid $>$ secondary amides $>$ amines – while impeding gelation of more polar solvents such as water (Fig. 2).\textsuperscript{38} Therefore, the molecular features that make up an ideal gelator for silicone oil are clearly not the same as for water.

Without exception, every study has shown only select solvents are capable of being gelled by a given gelator! Thus, it is our opinion that the quest for the universal gelator should be left to Don Quixote. The diversity of attractive and repulsive forces that can operate between gelators and solvents is enormous. In that regard, a recent meta analysis of 50 individual gelators, with an extremely diverse set of molecular structures, has been conducted.\textsuperscript{56} None was capable of gelling all of the solvents examined. However, analyses of solvent parameters and the gelator structures do provide an understanding of why certain gelators form gels in specific solvent types and others do not. For example, the molecules that act as gelators of benzene have

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strong hydrogen-bonding interactions, whereas molecules that form solutions/sols tend to interact mainly through dispersive forces. Therefore, the “holy grail” of this field of science may not be to develop a universal gelator; but, instead, to create reliable \textit{a priori} methodologies to understand why molecular gels form in specific classes of solvents and not in others.

What role does solvent play in molecular gel formation?

The role of solvent chemistry on the capacity of small molecules to assemble into SAFiNs is as important as the gelator structure! The first pioneering work on the influence of both solvent and gelator structure on the formation of SAFiNs showed that the gelation number (\textit{i.e.}, the maximum number of solvent molecules gelled per gelator molecule) correlated with Hildebrand solubility parameters when the primary functional group of the solvent remained constant.\textsuperscript{57} In that study, various primary alcohols were gelled by trehalose-based gelators. The authors found that when the substituent R group was short (either a diacetate or dibutyrate) the capacity to gel solvents was inversely proportional to the Hildebrand solubility parameter of the solvent.\textsuperscript{57} Since this finding, similar observations have been reported for HSA,\textsuperscript{58} derivatives of HSA,\textsuperscript{46,59} 1,3:2,4-dibenzylidene sorbitol (DBS),\textsuperscript{60} two component dendritic gels,\textsuperscript{61} l-lysine based gelators,\textsuperscript{62} and di-peptides\textsuperscript{63} (among others).

Changing the solvent can alter the morphology of the SAFiN. In a study with dipeptide (diphenylalanine) gelators, changing the solvent from toluene to ethanol led to a change in gel morphology from fibers to microcrystals (Fig. 3). Solvent-induced morphological changes have been observed with morphs of cholesteryl 4-(2-anthryloxy)butanoate (CAB) gelators\textsuperscript{64} and HSA.\textsuperscript{55,65,66} HSA gels in various alkanes and thiols have fibrous SAFiNs, with a hexagonal sub cell spacing and a multi-lamellar morphology in which the distance between lamellae is greater than the length of 2 molecules of HSA. HSA in solvents with nitriles, aldehydes and ketones as functional groups, assemble less effectively due to the formation of sphereulitic objects with a triclinic, parallel sub cell, and interdigitation in the lamellar arrangement.\textsuperscript{55} It is clear that solvent properties are central in dictating non-covalent interactions driving self-assembly.\textsuperscript{67–74}

Correlating molecular gelation to global solvent properties

Attempts have been made to correlate solvent parameters with gelation ability.\textsuperscript{75–77} Important considerations include: (1) can the solvent be considered as a macroscopic continuum characterized only by bulk physical properties? (2) Can the outcome of all gelator–solvent systems, capable of forming a gel, be predicted by a universal set of solvent parameters? (3) What solvent parameters are most crucial in predicting the likelihood of gel formation?

In that regard, Parker established three primary classifications of solvents: protic, dipolar aprotic and apolar (low polarity) aprotic.\textsuperscript{78} Although, solvent ‘polarity’ seems intuitively simple, Katrizky \textit{et al.} stated that “…the simple concept of polarity as a universally determinable and applicable solvent characteristic is a gross oversimplification”.\textsuperscript{79} Reichardt and Welton point out that solvent polarity may be interpreted as a permanent dipole moment of a solvent, its relative permittivity, or a sum of all of the molecular properties responsible for all interaction forces between a solvent and solute (\textit{i.e.}, coulombic, directional, inductive, dispersion, and hydrogen-bonding forces).\textsuperscript{80} To assess the correlation between gelation and solvent parameters, we have conducted studies and meta-analyses covering a total of 22 molecular gelators that were obtained from a combination of literature and from our laboratories. The gelators are sub-classified into azobenzene
gelators (Scheme 1),\textsuperscript{18,81} ALS gelators (Scheme 2),\textsuperscript{82} 1-O-methyl-4,6-O-benzylidene derivatives (MBD gelators) (Scheme 3),\textsuperscript{83,84} and miscellaneous highly efficient gelators (Scheme 4).\textsuperscript{16,58,60,85} The studies were conducted in 34 to 80 solvents, including protic, dipolar aprotic and apolar aprotic solvents! Data for all 80 solvents with every gelator were not obtained.

In this section, we discuss studies that have correlated molecular gelation with global solvent parameters, \textit{i.e.}, a single parameter is used to characterize the solvent as a macroscopic continuum. These are further subdivided into (1) physical properties, (2) solvatochromic properties, and (3) thermodynamic properties.

1.1 Global physical solvent properties

\subsection{1.1.1 Dielectric constant.} The dielectric constant \(\varepsilon_r\) (\textit{i.e.}, static relative permittivity)\textsuperscript{86} is routinely used to assess the ‘polarity’ of a solvent. Studies have attempted to correlate this parameter with molecular gelation, but the results have been mixed.\textsuperscript{58,61} Only in limited cases have correlations between gel properties (\textit{e.g.}, sol–gel transition temperature\textsuperscript{61,87} or critical gelator concentration (GGC)\textsuperscript{58}) and the dielectric constant been reported, and that too is only within a class of solvents with the same functional groups (\textit{e.g.}, alcohols of differing alkyl chain length). Here, we take a closer look at the correlation between gelation and \(\varepsilon_r\) for the gelators shown in Schemes 1–4.

AZO gelators rely on \(\pi-\pi\) stacking and are unable to donate hydrogen bonds.\textsuperscript{81} In the case of these gelators, there is no differentiation of solvents capable of being gelled and those that cannot, based simply on the \(\varepsilon_r\) (Fig. 4 and Fig. S1, ESI\textsuperscript{†}). ALS gelators have similar cholesterol derivatives as the AZO gelators; however,
the more polar azobenzene group is replaced with the less polar 2-anthraquinonyl group (Scheme 2).\textsuperscript{64,88} ALS gelators are limited to gelling solvents with lower $\varepsilon_r$ compared to AZO gelators (Fig. 4 and Fig. S2, ESI†). Again, dielectric constants are insufficient to differentiate the nature of the gelator–solvent phases.

Gelators based on 1-O-methyl-4,6-O-benzylidene derivatives (MBD-1 through MBD-10) (Scheme 3) are more polar compared to ALS and AZO, and hence form gels in lower polarity solvents (typically, $\varepsilon_r < 20$). When comparing the vastly different gelators in Scheme 4, some intuitive insights may be drawn from the dielectric constants (Fig. 4 and Fig. S3, ESI†). DBS, $N,N'$-DBU and $N,N'$-DCHU have both the ability to form hydrogen bonds as well as undergo $\pi-\pi$ stacking and van der Waals interactions; thus, these gelators are able to gel a range of solvents with varying dielectric constants. HSA, on the other hand, relies mainly on hydrogen bonding and gels solvents with lower $\varepsilon_r$ values. Conversely, CAB gelators rely on $\pi-\pi$ stacking and tend to gel solvents with higher $\varepsilon_r$ values. However, differentiation of outcomes based on $\varepsilon_r$ is not possible for any of these gelators when the solvent structure varies greatly (Fig. S4, ESI†).

1.1.2 Refractive index. The refractive index $n$ of a solvent increases with both the molecular mass and polarizability, where the larger the dipole moment the more efficiently the solvents respond to an applied electric field. $n$ is related to its molar polarizability $P_m$ by the Lorenz–Lorentz equation (eqn (1)):

$$P_m = \frac{(n^2 - 1)}{(n^2 + 1)} \tag{1}$$

It has been well established that the refractive index increases with the polarizability of the molecule, where the larger the dipole moment the more efficiently the molecule responds to an applied electric field.

Correlations between gelation and $n$ or $P_m$ for the gelators shown in Schemes 1–4 are now examined. For the AZO gelators, although $n$ and $P_m$ cannot differentiate gels from sols, important insights can once again be garnered (Fig. S5, ESI†). First, it is clear that an increase in the acyl chain length on the gelator increases the range of solvents (in terms of $n$ and $P_m$) that are capable of being gelled. AZO-ME ($\sim$CH$_3$) gels solvents between 1.3 < $n$ < 1.4 while AZO-DE ($\sim$C$_{10}$H$_{21}$) gels solvents between 1.3 < $n$ < 1.65 (Fig. S5, ESI†). For ALS gelators, $n$ and $P_m$ are fairly useful in differentiating sample states (Fig. S6, ESI†). Empirically, it is shown that the linker (L) group in these gelators affects gelation.\textsuperscript{82} For example, ALS-1 gels solvents with 1.38 < $n$ < 1.45; in this case, the linker group has both an ester and an ether. In comparison, ALS-8 has a shorter linker group with only an ester, and it gels solvents with 1.41 < $n$ < 1.45. These effects are amplified when comparing the polarizability $P_m$ instead of $n$.

The relation of the MBD gelators with $n$ and $P_m$ is plotted in Fig. S7 (ESI†); here, we include the different isomers of the same sugar-based MBD gelators. For MBD-1, gels are found for 1.5 < $n$ < 1.6; while the other MBD gelators gel solvents over a much wider range of $n$, i.e., 1.4 < $n$ < 1.65. Comparing MBD-1–2, and -6, we see that even though the gelators are very similar structurally, a conformational modification of the sugar radically changes their ability to form gels. Numerous other studies have illustrated the importance of chirality and geometric isomerization on gelation ability, so albeit not a surprising finding, it points to the complexities in designing gelators.\textsuperscript{53,36,37,52–54,89–95}

Finally, Fig. 5 depicts correlations with $n$ and $P_m$ for the gelators in Scheme 4. DBS is capable of gelling the broadest range of solvents with regard to $n$; however, differentiation of sample outcomes (gel vs. sol) based on $n$ or $P_m$ does not appear to be possible. $N,N'$-DCHU and $N,N'$-DBU are both derivatives of the smallest known organogelator, $N,N'$-dimethyleurea, where the methyl groups are replaced with cyclohexyl or benzyl groups, respectively. Among the two, $N,N'$-DBU gels solvents that have relatively higher $n$, i.e., are more polarizable. Also, in the case of $N,N'$-DCHU, the gel and precipitate states can be differentiated to some extent based on $n$ and $P_m$. Overall, we conclude that $n$ and $P_m$ do not permit significant differentiation of sample outcomes across a range of solvents for any of the gelators in Schemes 1–4.

1.1.3 Octanol/water partition coefficients and Henry’s law constants. The distribution of a pure substance between two partially miscible solvents, such as 1-octanol and water,
which are in intimate contact is the basis of the partition coefficient parameter \( \log P \) (eqn (2)).\(^{96} \)

\[
\log P = \log \left( \frac{X_{\text{organic}}}{X_{\text{aqueous}}} \right)
\]

\( X \) is the mole fraction of the substance in each phase. Overall, the \( \log P \) is related to the lipophilicity of the substance. Henry’s law constants (HLCs) are influenced by the same factors associated with the \( \log P \) constants.\(^{96} \) Instead, HLCs measure the volatility of the substance or the water-to-air partition. This becomes a measure of the intermolecular interactions between molecules within the mixture.\(^{97,98} \)

We have examined the correlation between gelation of the compounds in Schemes 1–4 with \( \log P \) and HLC. Examples are shown in Fig. 6 for the AZO gelators, MBD (Fig. S8, ESI†), ALS (Fig. S9, ESI†), and other gelators (Fig. S10, ESI†). Neither of these parameters is useful in differentiating the sample states (i.e., gel vs. sol vs. precipitate). The lack of differentiation is not surprising because \( \log P \) and HLC are global parameters that cannot distinguish specific types of intermolecular interactions.

### 1.2 Global solvatochromic solvent properties

#### 1.2.1 Reichardt’s \( E_T(30) \) solvent parameter

The notion that solvatochromic dyes are sensitive measures of solvent polarity was introduced nearly 70 years ago.\(^{99} \) It was quickly realized that the solvent can alter absorption bands of a dye because of: (1) dipolar interactions between the solvent and dye; (2) changes in the dipole moment of a dye arising from an optical transition; and (3) the Franck–Condon principle.\(^{100} \) Solvatochromic techniques rely on a pronounced change in the position of UV/VIS absorption bands of a solvatochromic dye caused by the solvent in which it is dissolved.\(^{101} \) Reichardt’s

\( E_T(30) \) parameter corresponds to the molar transition energy for the long wavelength electronic transition of diphenyl-4-[2,4,6-triphenylpyridinio]phenolate dye (Dimroth–Reichardt’s betaine dye) in a solvent at 25 °C and at 0.1 MPa (Fig. 7). This parameter accounts for all possible (specific and nonspecific) intermolecular forces between solvent and solute molecules.\(^{60,102} \) The \( E_T(30) \) parameter is calculated (eqn (3)) using the shift in the longest wavelength (\( \lambda_{\text{max}} \)) absorption band of the dye.\(^{103} \)

\[
E_T(30) = \frac{hc}{\lambda_{\text{max}}}N_A
\]

Here, \( h \) is Planck’s constant, \( c \) is the speed of light and \( N_A \) is the Avogadro number.

The \( E_T(30) \) parameter has been correlated with gel properties in very small sample sets and for some simple gelators.\(^{57} \) However, our results for AZO (Fig. S11, ESI†) and ALS-based gelators (Fig. S12, ESI†) show that \( E_T(30) \) has limited utility for explaining gel formation. AZO gelators gel solvents with \( E_T(30) \) values from 30 to 55 kcal mol\(^{-1} \). For the ALS gelators, the \( E_T(30) \) values
tend to cluster for the higher polarity solvents and again with very low polarity solvents (i.e., alkanes).

For the MBD-gelators (Fig. 8), some patterns emerge. In the case of MBD-1 and MBD-6, gels, precipitates and sols are differentiated based on \( E_T(30) \). However, for MBD-9, the distinction between sols and precipitates is lost. Generally, a better correlation is seen when the gelator mainly exploits hydrogen-bonding for self-assembly. This can also be noted by comparing the results for \( N,N'\)-DCHU (with cyclohexyl groups) to those for \( N,N'\)-DBU (with benzyl groups). For \( N,N'\)-DCHU, which relies mainly on hydrogen bonding, the gel and sol states are well separated based on \( E_T(30) \) (Fig. 8). However, for \( N,N'\)-DBU, which has the capacity for both hydrogen bonding and \( \pi-\pi \) stacking, the \( E_T(30) \) parameter is insufficient to differentiate the different states of the samples.

1.2.2 Pyrene fluorescence solvent parameter. Two pyrene scales \( (I_1/I_3) \) are obtained as the ratio of \( I_1 \) and \( I_3 \) vibronic band intensities in absorbance or fluorescence spectra. \(^{104-106}\) The changes in the \( I_1 \) and \( I_3 \) band intensities are theoretically attributable to: (1) dipolar interactions between the transition moment of the pyrene and the permanent dipole moments of the solvent molecules in the immediate solvent sphere; (2) inductive interactions with the quadrupole moment of the solvent; (3) dispersive interactions; (4) hydrogen bonding interactions. \(^{104}\) The \( I_1/I_3 \) scale is much less sensitive to hydrogen bonding effects than to dipolarity-polarizability of the solvent. \(^{104}\) Also, the \( I_1/I_3 \) scale has been linearly correlated to the \( E_T(30) \) parameter. \(^{104}\) Correlations with the \( I_1/I_3 \) scale for the AZO (Fig. S11, ESI†) and ALS gelators (Fig. S12, ESI†), do not provide additional information than the \( E_T(30) \) parameter. Similarly, for the MBD gelators, the same correlations are found with the \( I_1/I_3 \) scale (Fig. 9) and the \( E_T(30) \) scale. In the case of HSA, both the \( E_T(30) \) and \( I_1/I_3 \) scales show a region of sols with gel regions flanking it.

1.3 Global thermodynamic solvent properties

1.3.1 Hildebrand solubility parameter. For a solute–solvent pair, the molar Gibbs energy of mixing \( \Delta G_m \) is related to the enthalpy of mixing \( \Delta H_m \) and an entropy term \( T \Delta S_m \). The Hildebrand solubility parameter, as proposed by Hildebrand and Scott \(^{107}\) and Scatchard, \(^{108}\) relies solely on the enthalpy of mixing \( \Delta H_m \) (eqn (4)).

\[
\Delta H_m = V \left( \frac{\Delta E_1}{V_1} \right)^{1/2} \left( \frac{\Delta E_2}{V_2} \right)^{1/2} \phi_1 \phi_2 \tag{4}
\]

Here \( V \) is the mixture volume, \( \Delta E_i^V \) is the energy of vaporization, \( V_i \) is the molar volume, and \( \phi_i \) is the volume fraction of component \( i \).
Under conditions of isothermal vaporization of saturated liquid, the cohesive energy density \( (\Delta E_i^c) \) is the negative of the energy of vaporization per cm\(^3\) of sample, corresponding to the Hildebrand parameter, \( \delta_i \) (eqn (5)).

\[
\delta_i = \left( \frac{\Delta E_i^c}{V_i} \right)^{1/2}
\]

The cohesive energy combines the dispersion forces and polar interactions (including hydrogen bonding). The cohesive energy density has been shown to be important in determining intermolecular forces required to overcome solvent–solvent interactions.\(^{111}\)

Previous studies have shown only weak correlations between gelation outcomes and \( \delta_i \); and only for small subsets of solvents.\(^{57,61,112}\) For the gelators in Schemes 1–4, the same results hold. In the vast majority of gelators examined here (i.e., HSA, DBS, \( N,N'\)-DBU, \( N,N'\)-DCHU (Fig. 10), and ALS (Fig. S13, ESIF\(^*\)), AZO (Fig. S14, ESIF\(^*\)), and MBD (Fig. S15, ESIF\(^*\)), correlations between \( \delta_i \) and gelation outcomes are very poor. The reason is that \( \delta_i \) is only useful for low polarity solvents without hydrogen bonding.\(^{113}\) CAB, a non-hydrogen-bond donating molecule, is the only gelator for which we see fairly good discrimination between sols, gels and precipitates (Fig. 10).

1.4 Conclusions derived for global solvent parameters

It is clear that global solvent parameters lack the specificity to predict gelation capabilities of a range of organogelators. While individual parameters may correlate well with individual gelators, global perspectives may not be derived from this class of solvent parameters. Considering a solvent as a macroscopic continuum characterized by a single physical constant (e.g., dipole moment, dielectric constant, refractive index, etc.), solvatochromic shift, or Hildebrand solubility parameter is insufficient in predicting gelation behavior. Next, multi-term solvent parameters are considered, which can distinguish individual types of interactions and have been reported as better predictors of gelation.

Multi-term solvent parameters

Numerous multi-term parameters have been developed to define accurately solvent properties. These parameters attempt to separate the contributions of individual interactions such as ion–dipole, dipole–dipole, dipole induced-dipole, hydrogen bonding, electron pair donor–electron pair acceptor interactions (EPD/EPA), and solvophobic interactions. Examples include Kamlet–Taft,\(^{114–119}\) Catalan\(^{120–123}\) and Swain\(^{116,124}\) solvatochromic parameters, as well as the Hansen,\(^{58,75,125–129}\) Flory Huggins,\(^{130–133}\) and Modified Separation of Cohesive Energy Density Model (MOSCED)\(^{134,135}\) thermodynamic models.

1.5 Multi-term solvatochromic solvent parameters

1.5.1 Kamlet–Taft solubility parameters. Kamlet–Taft parameters characterize a solvent with regards to its polarizability \( \pi \), hydrogen-bond donating (the acidity parameter, \( \alpha \)), and hydrogen-bond accepting (the basicity parameter, \( \beta \)) terms.\(^{79,117,136,137}\) The polarizability parameter originates from electrostatic and dispersive interactions.\(^{79,117,136,137}\) It is obtained using the spectra of either \( p \)-nitroanisole or \( N,N\)-dimethyl-\( p \)-nitroaniline (NNDN) (Fig. 11A) along with eqn (6).

\[
\pi = \frac{\nu - 28.18}{-3.52}
\]

The \( \beta \) parameter is specific to solute–solvent interactions where the solute plays the role of an electron-pair acceptor and the solvent is an electron-pair donor, while for the \( \alpha \) parameter, the solute plays the role of an electron-pair donor and the solvent is an electron-pair acceptor.\(^{79}\) The \( \beta \) parameter is determined using the difference in wavenumbers of the intensity maximum of the absorption bands between \( p \)-nitrophenol \( (\nu_2) \) (Fig. 11B) and the non-hydrogen-bond accepting molecule, NNDN \( (\nu_1) \) (eqn (7)).

\[
\beta = \frac{0.9841\nu_1 + 3.49 - \nu_2}{2.759}
\]

The constants (in cm\(^{-1}\)) arise from a standardized value of 1 for hexamethylphosphoramide.\(^{116}\) \( \nu_1 \) is the frequency maximum of the NNDN UV-Vis absorption band (cm\(^{-1}\)), and the
constants (cm\(^{-1}\)) arise from normalization of the \(\pi\) parameter between 0.0 for cyclohexane and 1.0 for DMSO.\(^{119}\) The \(\alpha\) parameter is determined by the difference in the wavenumbers for the maximum intensities of the absorption bands for the Dimroth–Reichardt betaine dye (\(\nu_3\)) (Fig. 11C), and for NNDN (\(\nu_1\)) (eqn (8)).

\[
\alpha = \frac{1.873\nu_1 - 74.58 + \nu_3}{5.47}
\]  

(8)

The constants are based upon a standardized value of 1 for methanol.\(^{115}\)

Kamlet–Taft parameters have been applied to L-lysine bis-urea gelators to understand their capacity to form molecular gels.\(^{137}\) The \(\alpha\) parameter was correlated with gelling ability, while the \(\beta\) and \(\pi\) parameters had subsidiary roles: the magnitude of \(\beta\) affected the stability of the gel and \(\pi\) indicated the influence of fiber–fiber interactions.\(^{137}\) More recently, Kamlet–Taft parameters have been used to show that 2,3-dihydroxycholestane-based steroids are incapable of gelling solvents with large \(\alpha\) parameters while \(\pi\) is less important since solvents with both high and low polarizability could be gelled.\(^{138}\)

In the present work, we have correlated the gelators from Schemes 1–4 with the Kamlet–Taft parameters. For DBS (Fig. 12), these parameters are quite effective at demarcating sol and gel regions. For example, sols fall in an intermediate range of polarizability \(\pi\) and electron-donating ability \(\beta\). Also, when the \(\alpha\) parameter of the solvents is close to 0, DBS forms a sol, whereas DBS gels the solvent when \(\alpha > 0\). Similar to DBS, \(N,N'\)-DBU forms gels in solvents whose \(\alpha\) parameters are sufficiently high (>0.7). Good demarcation is also seen for \(N,N'\)-DCHU: gels form in solvents with \(\pi\) parameters close to 0.75, intermediate \(\beta\) parameters (0.15 to 0.5) and \(\alpha\) parameters close to 0. This is stark different from DBS and \(N,N'\)-DBU, where gels form in solvents when \(\alpha > 0\).

The difference is that DBS and \(N,N'\)-DBU are capable of \(\pi\)-\(\pi\) stacking interactions, whereas \(N,N'\)-DCHU is not. ALS gelators also show strong correlations with Kamlet–Taft parameters (Fig. 13). Much like DBS, gels are found in solvents of either low \(\pi\) (<0.5) or high \(\pi\) (>0.8). For 0.5 < \(\pi\) < 0.8. For MBG gelators (Fig. S16, ESI†) and AZO gelators (Fig. S17, ESI†), outcomes were not well distinguished based on Kamlet–Taft parameters.

1.5.2 Catalan’s solubility parameters. Catalan’s multi-term solvatochromic parameters are similar to Kamlet–Taft’s \(\alpha\), \(\beta\), and \(\pi\) parameters, differing only in the use of an alternate series of probes/homomorphs.\(^{120–122}\) Here, the three solvent parameters characterize solvent polarizability (SPP), solvent acidity (SA) and solvent basicity (SB). SPP is determined using the long wavelength absorption of 2-(dimethylamino)-7-nitrofluorene (DMANF) and its homomorph, 2-fluoro-7-nitrofluorene (FNF).\(^{120}\) DMANF is sensitive to the polarizability while the homomorph FNF has a similar structure with much lower dipole moment because of the strong electron withdrawing fluorine atom. The difference between the maxima of the lowest energy absorption bands for DMANF and FNF (\(\Delta\nu\)) is determined for the solvent and with DMSO and cyclohexane as references SPP is then calculated using eqn (9).

\[
SPP = \frac{\Delta\nu(\text{solute}) - \Delta\nu(\text{cyclohexane})}{\Delta\nu(\text{DMSO}) - \Delta\nu(\text{cyclohexane})}
\]  

(9)

The SB parameter is determined using the acidic molecule, 5-nitroindoline (NI), and its ‘non-acidic’ analogue, 1-methyl-5-nitroindoline (MNI), which lacks the amino group.\(^{121}\) The SB parameter for a solvent is determined by eqn (10).

\[
SB = \frac{\Delta\nu(\text{solute}) - \Delta\nu(\text{gas})}{\Delta\nu(\text{TMG}) - \Delta\nu(\text{gas})}
\]  

(10)

Fig. 12 The capacity of DBS and urea gelators to form gels, precipitates or solutions as a function of Kamlet–Taft parameters.
Here, \( \Delta \nu_{(\text{solvent})} \) is the difference between the maxima of the lowest energy absorption bands between 350 and 400 nm for NI and MNI, \( \Delta \nu_{(\text{TMG})} \) is the difference between the lowest energy absorption bands for NI and MNI in tetramethylguanidine (TMG), and \( \Delta \nu_{(\text{gas})} \) is the lowest energy absorption band in a series of \( n \)-alkanes and extrapolating the Lorentz–Lorentz function \( f(n^2) = \frac{n^2 - 1}{(n^2 + 1)n} \). Finally, the SA parameter is measured using a basic probe (o-tert-butylstilbazolium betaine, TBSB) and its ‘non-basic’ analogue (o,o-di-tert-butylstilbazolium betaine, DTBSB). The SA scale is set at 0.2 for ethanol (eqn (11)).

\[
\text{SA} = \frac{\Delta \nu_{(\text{solvent})} \times 0.4}{1299.9} 
\]  

(Catalan’s parameters show some trends similar to those of the Kamlet–Taft parameters for the gelators in Schemes 1–4, although the demarcation into distinct sol, gel, and precipitate regions is not as good. Among the gelators in Fig. 14 (DBS, \( N,N' \)-DBU and \( N,N' \)-DCHU), the best demarcation is for \( N,N' \)-DCHU when examined with the SB parameter. That is, \( N,N' \)-DCHU gels are confined to solvents with an SB between 0.2 and 0.4; for both lower and higher SB, precipitates (or sols) form. MBD gelators do not show as convincing trends (Fig. S18, ESI†), and neither do the AZO (Fig. S19, ESI†) and ALS gelators (Fig. S20, ESI†).

1.6 Multi-term thermodynamic solvent parameters

1.6.1 Hansen solubility parameters. Hansen solubility parameters (HSPs) were devised to overcome the limitations of the Hildebrand solubility parameter that do not include the effects of specific intermolecular interactions such as polar and hydrogen-bonding. To derive the HSPs, it is assumed that the energy of vaporization of a species, which is a measure of its total cohesive energy \( E \), is the sum of three individual energetic components, i.e., due to dispersion interactions (\( E_d \)), dipole–dipole interactions (\( E_p \)) and hydrogen-bonding interactions (\( E_h \)) (eqn (12)).

\[
E = E_d + E_p + E_h 
\]  

The dispersive component \( E_d \) dominates for simple, low polarity solvents such as saturated hydrocarbons. The second component \( E_p \) accounts for interactions due to polar groups on the solvent. Nitroparaffins, propylene carbonate, and tri-\( n \)-butyl phosphate are examples of solvents that are polar, but immiscible with water. The third component \( E_h \) arises from hydrogen bonding interactions. Dividing eqn (12) by the molar volume (V) gives the square of the total HSP (which is identical to the Hildebrand solubility parameter) as the sum of squares of the individual HSP components (eqn (13) and (14)).

\[
\frac{E}{V} = \frac{E_d}{V} + \frac{E_p}{V} + \frac{E_h}{V} 
\]

\[
\delta^2 = \delta_d^2 + \delta_p^2 + \delta_h^2 
\]

HSPs have been commonly used to select solvents for dissolving polymers. However, their importance in new fields is rapidly emerging and they were first introduced to the field of molecular gels by Raynal and Bouteiller in 2011. When considering a gelator in a given solvent, the HSPs of the solvent are typically known (e.g., from Hansen’s book); however, the HSPs of the gelator are typically not known. The HSPs of the gelator may be estimated, e.g., using the group contribution method, where it is assumed that the different functional groups that affect the energy of vaporization are additive. However,
this method is seldom applicable for complex types of molecules such as most of the gelators shown in Schemes 1–4 because it does not differentiate among positional of stereo isomers.\textsuperscript{109} We now discuss the correlation between sample outcomes (sol, gel, or precipitate) for gelator–solvent pairs versus each of the HSPs. This was done for the gelators in Schemes 1–4 and the results are shown in Fig. 15 and Fig. S21–S23 (ESI\textsuperscript{†}). The dispersive HSP $\delta_d$ did not correlate with sample behavior for any gelator investigated. The correlation was better in the case of the polar HSP $\delta_p$ and the hydrogen-bonding HSP $\delta_h$, i.e., sample outcomes clustered over different regions of these parameters. For example, consider MBD-6 and MBD-9 (Fig. 15). These molecules form gels in solvents with low $\delta_p$ and precipitates in solvents with higher $\delta_p$. Similar trends were observed in $\delta_h$, also, with minor exceptions. Also, AZO-ME and AZO-BU form sols or gels in solvents with low $\delta_h$ and precipitates at high $\delta_h$ (the two points in the precipitate side refer to water and glycerol) (Fig. S21 and S22, ESI\textsuperscript{†}). Note, however, that none of the HSPs is able to distinguish sols from gels.

1.6.2 Determining Hansen space. Although the correlation with individual HSPs is marginal, their key utility is manifested when they are examined in their entirety. Note that the three HSPs can each be treated as the three axes on a 3D plot, which is referred to as Hansen space. Each solvent is represented as a point in this space. In the context of polymer–solvent miscibility, the closer a polymer and a solvent are in Hansen space, the higher the probability that they are miscible.\textsuperscript{126} To extend this approach to gelators, it is necessary to examine gelator–solvent outcomes in Hansen space. The idea was first proposed by Raynal and Bouteiller.\textsuperscript{75} They noted that when data for a given gelator

![Fig. 14](https://example.com/fig14.png) The capacity of DBS, CAB, N,N'-DBU, N,N'-DCHU gelators to form gels, precipitates or solutions as a function of Catalan’s SPP, SA, and SB parameters.
are plotted in Hansen space, one can discern clusters corresponding to solvents in which sols, gels, and precipitates are formed. The vast majority of the LMOG data sets analyzed so far have shown compact and globular domains. Therefore, the observed clusters of points can be enclosed in spheres, following the methodology proposed by Hansen. For example, the sol sphere (S) encloses all (or most of) the solvents in which the gelator forms a sol, while excluding the solvents in which the gelator gives rise to a different outcome.

The parameters of the estimated spheres, i.e., center location and radius, have been extensively used to improve the analysis and predictive ability of LMOG datasets, e.g., calculation of the distance between the center of the sphere and particular solvents to optimize solvent selection. Different approaches have been employed to obtain meaningful spheres that adequately characterize each domain. In this section the main features of the available methodologies will be discussed.

The HSPiP software, developed by Hansen, Abbott and Yamamoto, allows for the numerical determination of solution (or, alternatively, gel and precipitate) spheres and also provides 2D and 3D visualizations of the enclosed region. This software was originally envisioned to find adequate, i.e., effective and safe, solvents or solvent mixtures that solubilize specific polymers. It calculates the sphere that includes all the “good” solvents (defined in HSPiP as “inside” the sphere) and excludes all the “bad” solvents by implementing a minimization algorithm coupled with a desirability function that, in principle, reduces the likelihood of misclassifications. This specialized software does not allow the simultaneous estimation of several spheres. Therefore, estimations for different gelation outcomes have to be independently performed. For the estimation to run, the data set should include at least one outside (“bad”) solvent. It should be noted that the selection of outside (“bad”) solvent(s) from different domains during the single estimation of the sphere domain might yield different results depending on whether the

---

Fig. 15 The capacity of MBD derivatives to form gels, precipitates or solutions as a function of the dispersive Hansen solubility parameter ($\delta_d$), polar Hansen solubility parameter ($\delta_p$) and hydrogen-bonding Hansen solubility parameter ($\delta_h$).
outside solvent is selected from an overlapping or distant location. The HSPiP software allows easy identification of wrongly classified data points (a feature provided only in this software). Thus, sequential running of this software aids in the identification of “outside” data sets and in obtaining reliable results. A reference value, e.g., HSP parameters of the gelator, can be included during the estimations so that concentric spheres can be determined.

Bonnet et al.\textsuperscript{129} have implemented a similar approach to the one provided by Hansen’s HSPiP software using the Nelder–Mead optimization algorithm and a default minimization routine, the Solver function, in Microsoft Excel (Microsoft Corporation, Seattle, WA). Although the parameters of the gelation sphere can be estimated using this simple method, only a 2D rendition of the distance to the center of the sphere is automatically plotted in this application. Additionally, and as the authors have pointed out, the robustness of the estimates highly depends on the completeness of the data set.\textsuperscript{129}

Following the principles described by Hansen\textsuperscript{126} and Bonnet et al.,\textsuperscript{129} Lan et al.\textsuperscript{60} have applied a global constrained optimization procedure programmed in Mathematica 9 (Wolfram Research, Champaign, IL) to calculate minimal enclosing spheres that contained all the points pertaining to each outcome, i.e., solution, gel, and precipitate. This program uses the “NMinimize” built-in function in Mathematica to obtain the location of the center of each sphere while solving for the smallest possible radius. The NMinimize function was combined with a direct search method, namely differential evolution, to reach a numerical global optimum solution. The selection of this direct search method was based on its robustness despite being computationally more expensive. Any proficient user of this software can easily implement this program in Mathematica. It allows the simultaneous estimation of independent and concentric spheres and 2D and 3D plots can be included. Similar procedures can be programmed and implemented in alternative mathematical softwares.

The UMD Complex Fluids and Nanomaterials Laboratory’s Hansen Solubility Parameter Data Fitting Software, developed by Diehn et al.,\textsuperscript{128} is comprised of a user-friendly, self-explanatory, graphical interface that facilitates LMOG data analysis and visualization. Although this program has been coded using MATLAB (MathWorks, Natick, MA), no programing knowledge is needed and its use requires only the installation of the MATLAB Compiler Runtime (MCR – Version 8.1). The LMOG solubility data, displayed following a pre-set format, can be directly input from Excel spreadsheets, which facilitates its use. The UMD Complex Fluids HSP Program offers three fitting methods.

The “concentric sphere shell fit” method fits all categories radiating out from a center sphere (the solution sphere). This method first optimizes the fit of the solution sphere. The center of the solution sphere is then fixed as the optimal center and is used to estimate the radii of the subsequent classifications (gels, precipitates, etc.). The “non-concentric sphere shell fit” recalculates an optimal center for every data class. Finally, the “independent sphere fit” treats each sub-classification independently, optimizing the radius and center for each sphere. As in the case of the Hansen parameters, the UMD Complex Fluids HSP Program implements a desirability function that allows for the optimization of the radii of the spheres so that most of the data of a single category are included within the enclosed region at the same time that most of the data points corresponding to other categories are excluded.

Hansen’s HSPiP software, Lan et al.’s approach and the UMD Complex Fluids HSP Program, have been used to obtain the individual spheres of 4 LMOG types—DBS, AZO-ET, ALS-10, and MBD-10. It should be pointed out that the same premises were employed when performing the estimations to facilitate the comparison of the outcomes. Although similar 3D plots for all tested LMOGs were obtained using the described methods (Fig. 16), the radii estimated using the UMD program are, in most cases, smaller than the ones reported using the two other approaches (Table 1). The implementation of the desirability function narrows the scope of the sphere, providing tighter regions where prediction reliability will be high. This feature will be extremely useful to predict the behavior of constrained data sets such as solvent mixtures. However, certain regions of the HSP 3D space might be mischaracterized if the data set is not sufficiently comprehensive. In the particular case of DBS,
Continuous efforts to increase the comprehensiveness of the relevant information. Regardless of the approach selected, a priori characterized by a concentric shell approach. However, relying ion from solutions to gels to precipitates which will be well-

The behavior of an LMOG. It is reasonable to expect a progress-
magnitude but also directionality can play a role in predicting the user’s needs.

Therefore, the data analysis tool should be selected based on identification of regions with extremely high predictability. Domain to further scout solvent–gelator combinations information, such as the identification of broad regions for each
to analyze the data will provide important and equally valuable that should be excluded from the analysis. Different approaches
gel sub-class that should be further differentiated or a subset
result of the scarcity of data points in that region or a hint of a gel sphere estimated by the UMD program, which might be a
20% larger than the one estimated by the UMD Program. The

[Table 1] Hansen coordinates for the center of the solution, gel and precipitate spheres and the radii of the spheres for the independent Hansen space plots estimated using the three described approaches

<table>
<thead>
<tr>
<th>Gelator</th>
<th>Sphere</th>
<th>Method</th>
<th>$\delta_D$ (MPa$^{1/2}$)</th>
<th>$\delta_P$ (MPa$^{1/2}$)</th>
<th>$\delta_H$ (MPa$^{1/2}$)</th>
<th>Radius (MPa$^{1/2}$)</th>
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</thead>
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<td>DBS</td>
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<td>17.5</td>
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<td>5.5</td>
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<td></td>
<td></td>
<td>Diehn et al.</td>
<td>18.3</td>
<td>16.9</td>
<td>8.8</td>
<td>6.5</td>
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<tr>
<td></td>
<td></td>
<td>Lan et al.</td>
<td>18.1</td>
<td>17.5</td>
<td>7.9</td>
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<td>4.8</td>
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<td>8.6</td>
<td>4.1</td>
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<td>6.3</td>
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<td>7.7</td>
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</table>

the approach taken by Lan et al. provides a gel sphere almost 20% larger than the one estimated by the UMD Program. The points corresponding to DBS in polyols are not included in the gel sphere estimated by the UMD program, which might be a result of the scarcity of data points in that region or a hint of a gel sub-class that should be further differentiated or a subset that should be excluded from the analysis. Different approaches to analyze the data will provide important and equally valuable information, such as the identification of broad regions for each domain to further scout solvent–gelator combinations versus identification of regions with extremely high predictability. Therefore, the data analysis tool should be selected based on the user’s needs.

From the presented data, it can be inferred that not only magnitude but also directionality can play a role in predicting the behavior of an LMOG. It is reasonable to expect a progression from solutions to gels to precipitates which will be well-characterized by a concentric shell approach. However, relying a priori or solely on this assumption might result in loss of relevant information. Regardless of the approach selected, continuous efforts to increase the comprehensiveness of the data sets will result in more robust and reliable predictions. Until then, a careful and critical examination of the data sets to be fit should be conducted to obtain meaningful results.

### 1.6.3 Vector distance in Hansen distance

In turn, we can calculate $R_{ij}$, the vector distance in Hansen space from a solvent ($j$) to the center of the sol sphere ($i$). We refer to this as the Hansen distance, and it is calculated by eqn (15).

$$R_{ij} = \sqrt{4(\delta_{Dj} - \delta_{Di})^2 + (\delta_{Pj} - \delta_{Pi})^2 + (\delta_{Hj} - \delta_{Hi})^2}$$  (15)

Note that the factor of 4 for $\delta_D$ comes into play because typically the axis with the dispersive HSP is plotted as 2$\delta_D$, form order a value of $R_{ij}$. Thus, $R_{ij}$ allows us to
quantify the intermolecular interactions between a solvent and a gelator. We should point out that the gelator–solvent scenario is more complicated than that for polymers and solvents. This is because there are at least three possible outcomes in the gelator–solvent case: sol, gel, and precipitate. For the polymer–solvent case, the only outcomes considered are soluble vs. insoluble. The related complication for the gelator–solvent case arises when drawing the best-fit spheres in Hansen space for the various outcomes. The spheres corresponding to sol (S), gel (G), and precipitate (P) are typically not concentric. For the polymer–solvent case, only one sphere needs to be drawn, encompassing the soluble points, and so this complication is absent.

We now examine the sample outcomes (sol, gel, precipitate) for the gelators in Schemes 1–4. They are shown in Hansen space, i.e., as 3-D plots, and the outcomes are also plotted on 2-D plots against $R_{ij}$. They were created using the software application developed by the Rogers group; the other software packages give similar results for the most part.

An excellent confinement of solutions was found within solubility sphere for DBS, HSA and CAB (Fig. 17). For DBS, none of the gel points was found in the solubility sphere. This can also be graphically represented using $R_{ij}$. Similar phenomena were observed for HSA where only 2 gel points were found within the solubility sphere. The gelation spheres appear to be less effective in determining the ability to cluster points in Hansen space. Only in the gelation sphere of CAB were all the solution points excluded. The other two gel spheres fail to exclude the soluble points, and so this complication is absent.

From the data presented, it can be inferred that not only magnitude but also the directionality of $R_{ij}$ can play a role in predicting the behavior of an LMOG. This is clear for DBS and MBD-10 (Fig. 18) where both gelators have excellent confinement in Hansen Space using the UMD software; however, for MBD-10 this does not translate to a clear correlation with $R_{ij}$. It is reasonable to expect a progression from solutions to gels to precipitates, which will be well characterized by a concentric shell approach. However, and as it was mentioned before, relying a priori, or solely, on this assumption might result in loss of relevant information. Regardless of the approach selected, continuous efforts to increase the comprehensiveness of the data sets will result in more robust and reliable predictions.

1.6.4 Modified Separation of Cohesive Energy Density Model (MOSCED) model. MOSCED (Modified Separation of Cohesive Energy Density) is a thermodynamic model that was first proposed to predict limiting activity coefficients (γ−1’s) using only pure component parameters. Several versions of MOSCED models have been reported. The most updated revision and extension of the parameters was done in 2005 to include binary systems of low polarity, polar and hydrogen-bonding compounds. MOSCED utilizes 5 adjustable parameters.
Table 2  Hansen coordinates (MPa0.5) for the center of the solution, gel and precipitate spheres as well as the radii of the spheres for AZO and ALS gelators

<table>
<thead>
<tr>
<th>Gelator</th>
<th>2δq</th>
<th>δp</th>
<th>δh</th>
<th>Rsphere</th>
<th>2δq</th>
<th>δp</th>
<th>δh</th>
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<td>5.4</td>
<td>10.0</td>
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<td>6.3</td>
<td>11.1</td>
<td>12.7</td>
<td>32.9</td>
<td>14.1</td>
<td>35.8</td>
<td>7.1</td>
</tr>
<tr>
<td>AZO Et</td>
<td>36.9</td>
<td>7.9</td>
<td>5.1</td>
<td>9.9</td>
<td>30.4</td>
<td>6.3</td>
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<td>32.9</td>
<td>14.1</td>
<td>35.8</td>
<td>7.1</td>
</tr>
<tr>
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<td>36.9</td>
<td>7.9</td>
<td>5.1</td>
<td>9.9</td>
<td>30.4</td>
<td>6.3</td>
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<tr>
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</tr>
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<td>7.1</td>
</tr>
<tr>
<td>ALS 1</td>
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<td>7.8</td>
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<td>12.9</td>
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<td>8.7</td>
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<td>5.6</td>
<td>13.0</td>
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</tr>
<tr>
<td>ALS 6</td>
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<td>5.6</td>
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</tr>
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<td>4.2</td>
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<td>6.9</td>
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<td>11.1</td>
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<tr>
<td>ALS 9</td>
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<td>6.3</td>
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<td>30.0</td>
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<td>11.1</td>
<td>12.7</td>
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</tbody>
</table>

Fig. 18  $R_t$ and UMD software groupings in Hansen space for DBS and MBD-10.

($\lambda$, $\tau$, $q$, $x$, and $\beta$) to characterize the energy of interaction of a solute in solution. These 5 parameters are partly derived from experimental values and partly fitted to experimental data. The original formulations of MOSCED are shown below (eqn (16)–(22)). MOSCED parameters can be considered as modifications of Hansen solubility parameters, which divides the regular solubility parameters into three components—dispersion, polarity and hydrogen bonding. Analyses of sample outcomes by MOSCED improved the poor performance of the Hansen model for associated and solvating systems.\(^{134}\)

$$
\ln \gamma_\infty^T = \frac{v_2}{RT} \left( (\lambda_1 - \lambda_2)^2 + \frac{q_1^2 q_2^2 (\tau_1^T - \tau_2^T)^2}{\phi_1} + \frac{(x_1^T - x_2^T) (\beta_1^T - \beta_2^T)}{\xi_1} \right)
$$

\[(16)\]

$$
d_{12} = \ln \left( \frac{v_2}{v_1} \right)^{aa} + 1 - \left( \frac{v_2}{v_1} \right)^{aa}
$$

\[(17)\]

$$
aa = 0.953 - 0.002314 (x_1^T)^2 + x_2^T \beta_1^T
$$

\[(18)\]

The adjustable parameters for each molecule are the dispersion parameter $\lambda$, the polarity parameter $\tau$, the induction parameter $q$, the hydrogen-bond acidity $x$ and the hydrogen-bond basicity parameter $\beta$. Eqn (17) is the combinatorial Flory–Huggins expression with the empirically derived exponent $aa$ calculated from eqn (18). Eqn (19) describes the temperature dependence of $x$, $\beta$ and $\tau$. The $\phi_1$ and $\xi_1$ are empirical asymmetry terms calculated from eqn (20)–(22).\(^{134}\)

The dispersion parameter $\lambda$ describes the polarizability of a molecule. The polarity parameter $\tau$ measures the fixed dipole of a compound in a solution. The induction parameter $q$ measures the dipole-induced dipole and induced dipole-induced dipole interactions that can be found in molecules with large dispersion (polarizability) parameters, such as halogenated compounds and aromatic compounds. The strong interaction of dispersion forces tends to lessen the dipolar interaction, leading to decreased values of $q$ (i.e., $<1$). For aromatic compounds, $q$ is arbitrarily set to 0.9; for halogenated compounds, $q$ is varied for best fit. The acidity and basicity parameters $x$ and $\beta$ account for specific interactions from hydrogen bond formation through both association and solvation.\(^{134}\)

The MOSCED parameters are effective at discerning solvents that resulted in different material states (Fig. 19). For DBS, albeit the differentiation of dispersion parameter $\lambda$ is not perfect, useful trends can still be observed: solvents with relatively higher $\lambda$ values or higher polarizability are more likely to form sols. Similar conclusions were drawn from the Kamlet–Taft $\pi$ parameter and Catalan’s SPP parameter as noted previously. It appears that both the acidity and basicity are important in determining the final states formed upon cooling for DBS. Gels tended to form
in more acidic solvents, while solvents that formed sols resided at very low $z$ values ($z < 2$). There is a stark difference between the basicity parameter of solvents that formed gels vs. sols. Gels clustered at low and $\beta$ values ($\beta < 13$), while solvents that formed solutions resided at relatively higher region when $\beta > 13$. MOSCED parameters also predict the gelation behavior of HSA very well. The dispersion parameter $\lambda$ differentiated gels from precipitates but not from sols, and $z$ seems to be an important parameter. All gel points clustered at $z < 1$, where solvents have very low acidities. This behavior is interesting when compared with DBS, whose gels formed in both very low and high acidity ranges. The enhanced gelation capability of DBS is a result of its ability to form a wider variety of non-covalent interactions than can HSA (whose primary driving force for aggregation is hydrogen bonding and van der Waals interactions). In solvents with high acidity, aromatic groups promote insolubility and $\pi-\pi$ interactions aid formation of SAFIN structures. Although some correlations were also found with CAB, convincing conclusions could not be drawn due to the small number of data points. Similar trends involving acidity and basicity parameters $z$ and $\beta$ were observed for the two urea gelators (Fig. S26, ESI†). For both, gels were observed in weakly acidic solvents; precipitates formed in more acidic solvents. The basicity parameter $\beta$ seems to be more promising at differentiating material states for $N,N'$-DBU. At low ($\beta < 2.23$) and intermediate ($7.8 < \beta < 12$) regions of the $\beta$ scale, precipitates formed. All gels clustered between $2.23 < \beta < 7.8$ and solutions formed in very basic solvents (i.e., $\beta > 14$). The same trend for the basicity parameters was obtained with $N,N'$-DCHU except that no solution data was available.

For the AZO gelators, the hydrogen-bond acidity and basicity parameters $z$ and $\beta$ are most important (Fig. S27, ESI†). Solvents that resulted in solutions clustered at low $z$ and $\beta$ values.

Fig. 19 Ability of DBS, HSA and CAB to form gels, precipitates or solutions as a function of the MOSCED parameters.
Good separation between gels and solutions was observed for AZO-ME and AZO-BU based on the β scale where the cut off points are 7 and 5, respectively. When comparisons were made within AZO groups, it was found that when the length of the aliphatic side chain was increased, the homologue was able to gel a broader range of solvents based on the α and β scales. This behavior is ascribed to the decreased solubility in high acidity and basicity solvents in the presence of longer aliphatic side chains that promote epitaxial growth of fibers.

Compared to AZO gelators, the replacement of more polar azobenzene group to less polar anthraquinonyl group (ALS derivatives) limits the gelation ability of ALS derivatives, and more precipitates were also observed (Fig. S28, ESF†). MOSCED parameters showed some evidence in predicting gelation behaviors of ALS derivatives. Solvents with lower polarizability tended to form precipitates and either gels or sols formed in highly polarizable solvents. Good separation of points for precipitates from the other two material states was observed with β, but not between gels and sols. Gels also tended to cluster with low values of q when dipolar interactions in solvents are weaker. The hydrogen-bond acidity parameter α seemed able to separate gels from other material states only for ALS-10. Although the differences among the behaviors of the ALS derivatives may involve the nature of the C-17 alkyl side chains,82 the reason is not yet clear.

For MBD-based gelators, only the hydrogen-bond acidity and basicity parameters α and β have good correlations with gelation behaviors (Fig. S29, ESF†). Solvents with low acidity (α < 4.2) tended form gels for all MBD derivatives, and solvents with high acidity (α > 4.2) were more likely to form precipitates. Solvents that resulted in gels tended to cluster at low β values for gluco (MBD-1) and manno (MBD-6) isomers. The galacto isomer (MBD-10) was able to gel solvents with both low (β < 3.3) and intermediate (7 < β < 9.5) β values. The enhanced gelation ability is attributed to the geometric isomerization; however, details of the reason are unclear. From these observations, it is clear that MOSCED parameters can be useful tools to predict the gelation behaviors of different gelators, and that hydrogen-bond acidity and basicity of solvents are especially important in determining whether SAFiN structures will be formed in a particular solvent.

1.7 Conclusions derived from multi-term solvent parameters

The ability to differentiate solvents capable of being gelled by a particular gelator, remaining as solutions or forming precipitates is better when using multi-term solvent parameters than when using global solvent parameters. Although the most significant insights gleaned involve the ability to accept or donate a hydrogen bond, other important conclusions and insights are available from analyses of other parameters.

General classification of solvents using a multivariate statistical treatment of quantitative solvent parameter

Organic solvents have been classified by a Fage nonhierarchical, multivariate, statistical method using eight solvent variables that include the Kirkwood function, molecular refraction, dipole moment, the Hildebrand parameter, index of refraction, boiling point, and HOMO and LUMO energies.148 Using this approach, 83 solvents were categorized into 9 classes: aprotic dipolar, aprotic highly dipolar, aprotic highly dipolar and highly polarizable, aromatic low-polarity, aromatic relatively polar, electron pair donating, hydrogen bonding, strongly-associated hydrogen bonding, and others that do not fit into one of the categories mentioned.148 Because the solvents are classified using their physical properties rather than their chemical structure, some oddities are observed, such as CCl₄ being classified as an ‘aromatic polar solvent’. This statistical approach is not intended to develop an all-encompassing and rigid classification, but, instead, a quantitative, logical approach to solvent classification.148 Using it, we have identified solvents that were included in their analysis and with the data sets which have been the focus of this review. Thus, parameters that are insensitive to predicting self-assembly of molecular gels, when combined, become useful in grouping material states. For example, DBS is incapable of crystallizing only in aprotic highly dipolar solvents and in all other solvent categories DBS will form gels (Table 3).

In the case of HSA, similar groupings are observed: it remains in the solution state in aprotic highly dipolar, hydrogen bonding and hydrogen bonding self-associating solvents (Table 4). In comparison, it is likely that DBS relies on π-π stacking in the hydrogen bonding and hydrogen bonding self-associating solvents. HSA is capable of gelling the aromatic apolar, aromatic polar, electron pair donor and most of the aprotic dipolar solvents. In a similar manner to DBS, the only category to have overlapping results is the aprotic dipolar solvents, where both precipitates and gels are observed.

For the MBD gelators, the ability to differentiate gel states is less apparent (Table 5). It is clear that MBD-1 forms solutions in both aprotic dipolar and aprotic highly dipolar solvents and gels in aromatic apolar and aromatic polar solvents. However, in the electron pair donor, hydrogen bonding and hydrogen bonding self-associating solvents, either sols or precipitates are observed. With more advanced clustering tools or more refined solvent categories, predictability may be enhanced, allowing for

<table>
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<th>Aprotic</th>
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<th>Aprotic highly</th>
<th>Aromatic</th>
<th>Aromatic</th>
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<tr>
<td>Methanol</td>
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<tr>
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<table>
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<th>Electron</th>
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<th>Misc.</th>
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</tbody>
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very useful *a priori* tools which provide deeper insights into the structure-function mechanisms that govern the formation of fibrillar aggregates in different organic solvents.

## Cluster analysis of dual solvent parameter data to assess their combined predictive ability

The use of cluster analysis of dual solvent parameter data sets has been proposed as a means to pinpoint synergistic parameter combinations with enhanced predictive ability. This approach provides, in principle, a way to assess the relevance of pair (or higher order) parameter combinations in the design of novel molecular gels.

As has been presented in previous sections, individual solvent parameters, including physical solvent properties and thermodynamic solvent parameters can predict gelation outcomes only to a limited extent. The availability of larger data sets on organogelators will allow additional data analyses to extract important information that is difficult to attain using limited databases. Cluster analysis is an unsupervised learning technique that allows the organization of a collection of data points, in our case solvent parameter combinations, into clusters based on a distance or dissimilarity function. The capability of the combinations to predict gelator behavior can be evaluated based on their probability to group data points pertaining to a single type of outcome (solution, gel or precipitate) into distinctive clusters. This technique requires some minimal data pre-processing. For example, to avoid larger-scale parameters to dominate others, each parameter is commonly normalized by mean removal and

| Table 4 | HSA solution (blue), precipitate (green) and gel (red) states clustered using a Fage nonhierarchical multivariate statistical method using eight solvent variables |
| Table 5 | MBD-1 solution (blue), precipitate (green) and gel (red) states clustered using a Fage nonhierarchical multivariate statistical method using eight solvent variables |
| Table 6 | Classification of DBS–solvent mixtures based on cluster analyses using two solvent parameters and the degree of their predictability as % |

SA, SB and SPP are Catalan parameters; $E_f(30)$ is Reichardt’s parameter; Py corresponds to the shift in pyrene absorbance; FH is the Flory–Huggins Parameter; $R_d$ is the distance in Hansen space; $D_{total}$ is the total Hansen solubility parameter; $d$, $h$ and $p$ are the individual HSPs; $D/A$ is the hydrogen bond donor/acceptor ratio; alpha, beta and pi are Kamlet–Taft parameters; Henry is Henry’s law constant; DI is the dielectric constant; Dipole is the dipole moment of the solvent; $\log P$ is the octanol/water partitioning coefficient.
variance scaling prior to the classification stage. Cluster analysis then can be performed using, for example, a k-medoids algorithm (related to k-means algorithm). This clustering technique can be programmed in most mathematical software, such as Mathematica, which through an iterative approach finds a local optimum clustering. Several distance functions can be used to reflect dissimilarity between two data points. Although the Euclidean distance (i.e., square root of the sums of the squares of the differences between the coordinates of the points in each dimension) is often used, other distance functions such as Canberra or Chebyshev’s functions are commonly implemented. It should be noted that utilization of centroid based algorithms, such as the k-medoids, for classification requires specification in advance of the number of output clusters. In our case, the number of output clusters was set to 3, which accounts for all possible outcomes—solution, gel or precipitate for our samples.

The predictive ability of the solvent parameter pair combinations for DBS, evaluated based on the percentage of correct grouping, is presented in Table 6. As can be observed, several of the solvent parameter pairings did not cluster the solvents effectively into their appropriate group, producing values slightly above those that can be attributed to random grouping. That is the case for all the combinations of the Kamlet-Taft $\beta$ parameter with any of the Hansen solubility parameters; values are 40–43%. This analysis also corroborates findings reported in previous sections, in which the performance of $R_p$ as a classification criterion was enhanced by its pairing with additional parameters such as Flory–Huggins, HLC and dielectric constant. These pairings consistently placed >60% of the solvents into their appropriate cluster. The results also point out that parameters that do not allow an appropriate classification alone, such as most of the physical properties parameters, might produce better outcome estimations when combined with other parameters. The combinations may reflect polarity and/or the gelation mechanism more completely than individual solvent properties.

The effectiveness of this kind of statistical data analysis relies on the availability of comprehensive, well-curated data sets. However, the preliminary results presented here suggest that the applicability of particular combinations might not be universal since parameters that are better related to the mechanism of gelation will also have a better performance on the classification.

Conclusions

It is apparent that without consideration of the solvent properties, the ability to develop a priori tools for designing new and unique molecular gels, as well to gain truly fundamental understanding of the driving forces for epitaxial growth, will continue to be unsuccessful. Clearly, global solvent parameters lack the specificity to predict gelation capacity for organogelators with diverse structures. Although individual parameters may correlate well with individual gelators, global perspectives may not be derived from this approach. Considering a solvent as a macroscopic continuum, characterized by a single physical constant (e.g., dipole moment, dielectric constant, refractive index, etc.), solvachormic shift, or Hildebrand solubility parameter is insufficient in predicting gelation behavior. Although multi-term solvent parameters drastically improve the predictive abilities of solvent gelator states, they are far from perfect; however, they greatly aid in understanding the mechanism of assembly. True insights into developing a priori tools will only be garnered when developing complex relationships between numerous solvent parameters. However, the benefits of greater accuracy afforded by this kind of analysis must be considered in terms of the additional analytical procedures that must be applied and the balance will have to be determined by the information sought.

Acknowledgements

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Notes and references

7 P. J. Flory, Faraday Discuss., 1974, 57, 7–18.
Supplemental Information

To Gel or Not to Gel: Correlating Molecular Gelation with Solvent Parameters

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Figure S1: The capacity of the AZO derivatives (scheme 1) to form gels, precipitates or solutions as a function of the static relative permittivity of the solvents.
Figure S2: The capacity of the ALS gelators (Scheme 2) to form gels, precipitates or remain as solutions as a function of the static relative permittivity of the solvents.
Figure S3: The capacity of the MBD gelators (Scheme 3) to form gels, precipitates or remain as solutions as a function of the static relative permittivity of the solvents.
Figure S4: The capacity of the miscellaneous highly efficient gelators (Scheme 4) to form gels, precipitates or remain as solutions as a function of the static relative permittivity of the solvents.
Figure S5: The capacity of the AZO derivatives to form gels, precipitates or solutions as a function of the refractive index and polarizability of the solvents.
Figure S6: The capacity of the ALS gelators to form gels, precipitates or solutions as a function of the refractive index and polarizability of the solvents.
Figure S7: The capacity of the MBD derivatives to form gels, precipitates or solutions as a function of the refractive index and polarizability of the solvents.
Figure S8: The capacity of the MBD derivatives to form gels, precipitates or solutions as a function of partition coefficients and Henry’s law constants of the solvents.
Figure S9: 15: The capacity of the ALS derivatives to form gels, precipitates or solutions as a function of partition coefficients and Henry’s law constants of the solvents.
Figure S10: The capacity of DBS, HSA, N,N'-DBU and N,N'-DCHU to form gels, precipitates or solutions as a function of partition coefficients and Henry’s law constants of the solvents.
Figure S11: The capacity of the AZO derivatives to form gels, precipitates or solutions as a function of $E_T(30)$ and the Py scale of the solvents.
Figure S12: The capacity of the ALS derivatives to form gels, precipitates or solutions as a function of $E_T(30)$ and the Py scale of the solvents.
Figure S13: The capacity of the ALS derivatives to form gels, precipitates or solutions as a function of the Hildebrand solubility parameter of the solvents.
Figure S14: The capacity of the AZO derivatives to form gels, precipitates or solutions as a function of the Hildebrand solubility parameter of the solvents.
Figure S15: The capacity of the MBD derivatives to form gels, precipitates or solutions as a function of the Hildebrand solubility parameter of the solvents.
Figure S16: The capacity of the MBD gelators to form gels, precipitates or solutions as a function of Kamlet Taft Parameters, $\pi$, $\beta$, and $\alpha$, of the solvents.
Figure S17: The capacity of the AZO gelators to form gels, precipitates or solutions as a function of Kamlet Taft Parameters, $\pi$, $\beta$, and $\alpha$, of the solvents.
Figure S18: The capacity of the MBD gelators to form gels, precipitates or solutions as a function of Catalan’s SPP, SA, and SB solvent parameters.
Figure S19: The capacity of the AZO gelators to form gels, precipitates or solutions as a function of Catalan’s SPP, SA, and SB solvent parameters.
Figure S20: The capacity of the ALS gelators to form gels, precipitates or solutions as a function of Catalan’s SPP, SA, and SB solvent parameters.
Figure S21: The capacity of the AZO derivatives to form gels, precipitates or solutions as a function of the dispersive Hansen solubility parameter ($\delta_d$), polar Hansen solubility parameter ($\delta_p$) and hydrogen- bonding Hansen solubility parameter ($\delta_h$) of the solvents.
Figure S22: The capacity of the ALS derivatives to form gels, precipitates or solutions as a function of the dispersive Hansen solubility parameter ($\delta_d$), polar Hansen solubility parameter ($\delta_p$) and hydrogen-bonding Hansen solubility parameter ($\delta_h$) of the solvents.
Figure S23: The capacity of the DBS, N,N'-DBU, N,N'-DCH to form gels, precipitates or solutions as a function of the dispersive Hansen solubility parameter ($\delta_d$), polar Hansen solubility parameter ($\delta_p$) and hydrogen-bonding Hansen solubility parameter ($\delta_h$) of the solvents.
Figure S24: 3D Hansen space and distances in Hansen space to the centers of the sol and gelation spheres for ALS gelators. The blue horizontal line represents the radius of the solubility sphere, and the red line represents the radius of gel sphere. The x-axes of the gelators are comprised of arbitrary values to aid in the visualization of differences in $R_{ij}$. 
Figure S25: 3D Hanson space and distances in Hansen space to the center of the sol and gelation spheres for MBD gelators. The blue horizontal line represents the radius of the solubility sphere, and the red line represents the radius of the gel sphere. The x-axes are comprised of arbitrary values to aid in the visualization of differences in $R_\phi$. 
Figure S26: The capacity of the urea gelators to form gels, precipitates or solutions as a function of the MOSCED (MOdified Separation of Cohesive Energy Density) parameters ($\lambda$, $\tau$, $q$, $\alpha$ and $\beta$) of the solvents.
Figure S27: The capacity of the AZO gelators to form gels, precipitates or solutions as a function of the MOSCED (MOdified Separation of Cohesive Energy Density) parameters ($\lambda$, $\tau$, $q$, $\alpha$ and $\beta$) of the solvents.
Figure S28: The capacity of the ALS gelators to form gels, precipitates or solutions as a function of the MOSCED (MOdified Separation of Cohesive Energy Density) parameters ($\lambda$, $\tau$, $q$, $\alpha$ and $\beta$) of the solvents.
Figure S29: The capacity of the MBD gelators to form gels, precipitates or solutions as a function of the MOSCED (MOdified Separation of Cohesive Energy Density) parameters (λ, τ, q, α and β) of the solvents.