Solvent-Induced Helix Superstructure in Achiral Dumbbell-Shaped Hydrazine Derivatives

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Supporting Information

ABSTRACT: We studied hydrogen-bonding assemblies in a series of dumbbell-shaped hydrazine derivatives, namely oxalyl N,N'-bis(3,4-dialkoxybenzoyl)-hydrazide (BFH-n, n = 4, 6, 8, 10) and oxalyl N,N'-dibenzoxyldihydrazide (FH-0). It has been demonstrated that NH-1 protons of BFH-n precipitated from tetrahydrofuran (THF) or dimethylformamide (DMF) were involved in intramolecular H-bonding to form 6-membered rings. Meanwhile, NH-2 protons of BFH-n precipitated from THF formed intermolecular hydrogen bonds with C=O groups of neighboring molecules, while NH-2 protons of BFH-n precipitated from DMF formed intermolecular hydrogen bonds with C=O group of neighboring DMF molecules. C=O, −CH3, and −CH groups of DMF molecules participated in multiple intermolecular hydrogen bonds with the −N−H and −C=O groups of FH-0 molecules in single-crystals formed in DMF, leading to a double helix morphology with a pitch of 24.2 Å along the ε direction. Both left- and right-handed helical micrometer-length ribbons with nonuniform helical pitches were observed in an achiral BFH-10 xerogel precipitated from DMF.

INTRODUCTION

Supramolecular chirality is highly intriguing for its important role in biological phenomena, such as molecular recognition, information storage, and potential applications in materials science.1 Generally, chiral assemblies are often achieved among chiral molecules or between chiral building blocks.2 Chirality can also be induced by adding chiral molecules to achiral molecules resulting in various chiral architectures, based on the “sergeants and soldiers principle” and the “majority rules”.3 It is also possible to obtain chiral supramolecular assemblies from achiral molecules. Chiral symmetry breaking from achiral molecules is known to occur in helical supramolecular organization through coordination with metal ions or through hydrogen bonding.4 Organogels are the materials in which three-dimensional networks are formed through self-assembly of low-molecular-weight organogelators (LMOGs) through noncovalent interactions, and the network can absorb a large volume of solvent.5 The formation of an organogel is the result of both gelator−gelator and solvent−gelator interactions, and a gelator only has the ability to induce macroscopic gelation in certain solvents.6−9 Recently, several detailed studies revealed that solvents play an important role in tuning the morphology of the organogel.10 For example, Tritt-Goc and co-workers reported that the polarity of the solvent influences the hydrogen-bonding network and the microstructure of the gel network using 1,2-O-(1-ethylpropyliyldene)-α-D-glucopyranose as the gelator,10a whereas Shinkai and co-workers reported that the solvent polarity affects the gel microstructure in methyl-4,6-O-(p-nitrobenzylidene)-α-D-glucopyranose gels, which form helical fibers in water and frizzled fibers in carbon tetrachloride.10d Interestingly, helical morphologies can be obtained from achiral organogelators through specific gelator−solvent interactions.11−13 For example, we reported that both left- and right-handed helical ribbons are formed in the achiral twin-tapered dihydrazide derivatives xerogel from ethanol,12 whereas You and co-workers reported that a helical nonracemic nanotubular structure is induced by a simple achiral molecule when linked by a linear metal connecting point to form a fibrillar network thereby immobilizing a wide range of...
Although some key points have been recognized, a complete mechanism for the self-assembly of achiral molecules chiral supramolecular structures is still beyond our understanding.

Among the noncovalent interactions, hydrogen bonding is most commonly used to direct self-assembly processes because of its strength, directionality, reversibility, and selectivity. Peptide, amino acid, amide, and urea groups have been widely employed as building blocks to afford supramolecular gels. Wang and co-workers reported a hydrazide quinolinone-based quadruple hydrogen-bonded building block, which can gel dichloromethane/hexane at concentrations higher than 2 wt%. Li and co-workers reported a new series of highly stable hydrazide-based ADDA-DAAD heterodimers, which represented the first successful application of hydrazide derivatives in the self-assembly of hydrogen-mediated supramolecular systems with well-established structures. Herein, we focus on the self-assembly property of a series of achiral dumbbell-shaped molecules, namely oxalyl \( \text{N}_2 \text{N}_0 \)-bis(3,4-dialkoxybenzoyl)-hydrazide (BFH-\(n\), \(n = 4, 6, 8, 10\)) and oxalyl \( \text{N}_2 \text{N}_0 \)-dibenzyolhydrazide (FH-0) (Scheme 1), which were prepared according to a reported procedure. It was demonstrated that DMF molecules were involved in multiple intermolecular hydrogen bonds with FH-0 molecules in FH-0 single-crystals grown from DMF, leading to a double helix with a pitch of 24.2 Å along the \(c\) direction. Moreover, left- and right-handed helical micrometer-length ribbons with nonuniform helical pitches were observed in a BFH-10 xerogel precipitated from DMF.

### EXPERIMENTAL SECTION

**Characterization.** \(^1\)H NMR spectra were recorded with a Mercury-300BB 300 MHz spectrometer, using tetramethylsilane (TMS) as an internal standard. Field emission scanning electron microscopy (FE-SEM) images were taken with a JSM-6700F apparatus. Samples for FE-SEM measurement were prepared by wiping a small amount of gel onto a silicon plate followed by evaporating the solvent at ambient temperature. X-ray diffraction was carried out with a Bruker Avance D8 X-ray diffractometer. FT-IR spectra were recorded with a Perkin-Elmer spectrometer (Spectrum One B). The xerogels were obtained by freezing and pumping the organogel of BFH-\(n\) for 12 h, and then the xerogels were pressed into a tablet with KBr for FT-IR measurement.

**Gelation test:** The weighted gelator was mixed in a capped sealed test tube \(3.5 \text{ cm (height)} \times 0.5 \text{ cm (radius)}\) with an appropriate amount of solvent, and the mixture was heated until the solid dissolved. The sample vial was cooled to \(4 \degree C\) and then turned upside down. When a clear or
slightly opaque gel formed, the solvent therein was immobilized at this stage. Melting temperature ($T_m$) was determined by the "falling drop" method. An inverted gel was immersed in a water bath initially at or below room temperature. The water bath was heated slowly up to the point at which the gel fell due to the force of gravity, i.e., the $T_m$.

**RESULTS AND DISCUSSION**

**Hydrogen Bonding in BFH-$n$.** Temperature dependent $^1$H NMR spectroscopic experiments were performed to explore the hydrogen bonding motif in BFH-$n$. Generally, internally hydrogen bonded amides are expected to show a much smaller shift with temperature (<3.0 × 10^{-3} ppm K^{-1}) compared to those directed externally and accessible for hydrogen bonding to a polar solvent (>4.0 × 10^{-3} ppm K^{-1}). As shown in Figure 1(a), the NH-2 (adjacent to central bicarbonyl) of BFH-10 precipitated from THF (called BFH-10-THF) showed large shifts (7.0 × 10^{-3} ppm K^{-1}) with temperature in 20% DMSO-d$_6$/CDCl$_3$, suggesting the primary involvement of NH-2 protons in intermolecular hydrogen bonding. However, the NH-1 (closer to alkoxyphenyl) showed smaller shifts (3.5 × 10^{-3} ppm K^{-1}) with temperature, indicating the involvement of NH-1 protons in intramolecular hydrogen bonding. The packing model of BFH-10-THF was proposed in Scheme 2a, in which protons of NH-2 associated with C=O groups of neighboring molecules and protons of NH-1 were involved in intramolecular hydrogen bonding forming two 6-membered rings.
As shown in Figure 1b, NH-1 and NH-2 of BFH-10 \((1 \times 10^{-3} \text{ mol L}^{-1})\) precipitated from DMF (called BFH-10-DMF) showed similar temperature-dependent shifts \((3.9 \times 10^{-3} \text{ and } 6.7 \times 10^{-3} \text{ ppm K}^{-1}, \text{respectively})\) with those of BFH-10-THF in 20% DMSO-\(d_6/\text{CDCl}_3\) in the first heating run. Thus, the NH-1 and NH-2 protons participate in intramolecular and intermolecular hydrogen bonding, respectively, for BFH-10 precipitated either from THF or DMF. The presence of DMF in BFH-10-DMF was confirmed by the \(^1H\) NMR spectrum of BFH-10-DMF (Figure 2, in 20% DMSO-\(d_6/\text{CDCl}_3\)); \(\delta\) of protons in DMF were found at 7.92, 2.89, and 2.78 ppm, respectively.\(^{19}\) The molar ratio of BFH-10/DMF was calculated to be 1:2 according to the integrated peak area of the protons, which is consistent with the observation in FH-0 single-crystals, suggesting that similar structures might be formed in BFH-10-DMF and FH-0 single crystals. In contrast to that of BFH-10-THF, the characteristic peak of the NH-2 protons of BFH-10-DMF appeared slightly further upfield. Addition of DMF \((3.21 \times 10^{-2} \text{ mol L}^{-1})\) to BFH-10-THF in 20% DMSO-\(d_6/\text{CDCl}_3\) caused a similar upfield shift of NH-2 (Figure 1c), suggesting an interaction between DMF and BFH-10. The temperature-dependences of the chemical shifts of NH-1 and NH-2 of BFH-10 in 20% DMSO-\(d_6/\text{CDCl}_3\) are quite similar for the first cooling run and the second heating run (Figure S1). Based on the results of \(^1H\) NMR spectroscopy, as well as the poor volatility of DMF, it can be concluded that DMF is associated with BFH-\(n\) in the course of precipitation from DMF.

A packing model of BFH-10 in DMF is shown in Scheme 2b, in which NH-1 protons form two 6-membered rings via intramolecular hydrogen bonding, while NH-2 protons associate with \(\text{C}==\text{O}\) groups of neighboring DMF molecules. In addition, \(\text{C}==\text{O}\) groups of BFH-10 molecules form intermolecular hydrogen bonds with \(-\text{CH}_3\) groups of DMF molecules.

In order to validate the above hypothesis, FH-0 without terminal alkoy chains was synthesized by the same method used for BFH-\(n\). Single crystals of FH-0, suitable for X-ray analysis were obtained at room temperature from DMF by slow evaporation of the solvent. Crystal data are given in the Supporting Information. DMF molecules interacted with FH-0 molecules in a single-crystal state, as shown in Figure 3a. Protons of \(\text{N}==\text{Hs}\) of FH-0 molecules formed intermolecular hydrogen bonds with \(\text{C}==\text{O}\) groups of DMF molecules (one \(\text{C}==\text{O}\) associated with two \(\text{N}==\text{Hs}\) coming from two different FH-0 molecules). \(\text{C}==\text{O}\) (near to phenyl) groups of FH-0 molecules were involved in intermolecular hydrogen bonding with \(-\text{CH}_3\) groups of DMF molecules, while one \(\text{C}==\text{O}\) (central bicarbonyl) group of an FH-0 molecule formed an intermolecular hydrogen bond with a \(\text{C}==\text{H}\) group of a DMF molecule and the other \(\text{C}==\text{O}\) was involved in intermolecular hydrogen bonding with a \(\text{C}==\text{H}\) group of a phenyl ring of a neighboring FH-0 molecule. Single crystal analysis indicated that the molar ratio of FH-0/DMF is 1:2. In addition, double helical structures were induced by DMF molecules with a helical pitch of 24.2 Å as seen from the view of the \(\varepsilon\) direction (Figure 3b).

Combining the results of single crystal analysis of FH-0 and the temperature dependent NMR spectroscopic results of BFH-10-DMF, we propose the intermolecular hydrogen bonding motif of BFH-10 from DMF as shown in Scheme 2b. Different from that of BFH-10-THF, NH-2 protons of BFH-10-DMF are H-bonded to \(\text{C}==\text{O}\) groups of DMF resulting in intermolecular H-bonding.

### GELATION PROPERTIES

The gelation properties of BFH-\(n\) were examined for 7 different organic solvents and the minimum gel concentrations are summarized in Table 1. The length of the terminal alkoy chains clearly affected the gelation ability. Compounds BFH-6, -8, and -10 bearing longer terminal chains can gelate toluene, chloroform, benzene, and DCE, but not in DMF or DMSO.
whereas BFH-10 formed a stable gel in DMF. In contrast, compound BFH-4 with shorter terminal chains cannot form a gel in any of the 7 solvents tested. Figure 4 shows the melting temperature ($T_m$) of a BFH-10 gel in DMF as a function of concentration. $T_m$ values were determined by the falling drop method.\textsuperscript{17} The $T_m$ increases from 20 °C for the gel at 1.0 $\times$ 10^{-2} mol L^{-1} to 60 °C at 2.0 $\times$ 10^{-2} mol L^{-1}.

In order to investigate the morphology of these materials, xerogels of BFH-$n$ were prepared as described in the Experimental Section and subjected to scanning electron microscopy (SEM). As shown in Figure 5b–d, xerogels of BFH-$n$ ($n = 6, 8, 10$) prepared from toluene consist of flat ribbons 0.2–1.5 μm wide and several tens of micrometers long, indicating that self-assembly is driven by strong directional intermolecular interactions. Interestingly, both right- and left-handed helical ribbons with the width of 2–8 μm and ten to hundreds of micrometers in length were observed for a BFH-10 xerogel prepared from DMF (Figure 5a), even though it does not possess any chiral groups. The helical pitches were nonuniform, suggesting that the helical structure did not arise from molecular chirality.

Figure 6 shows X-ray diffraction patterns of BFH-10 crystal and xerogels prepared from toluene and DMF. The XRD pattern of the BFH-10 crystal showed one sharp peak (32.0 Å) and three weak peaks (15.8, 10.7, and 8.0 Å) in the low angle range, suggesting a layered structure. A similar arrangement was observed for the BFH-10 xerogel formed from toluene, indicating a similar supramolecular arrangement in both cases. In contrast, the XRD pattern of the xerogel prepared from DMF was quite different from the others, indicating that the solvent greatly affected the packing modes.

FT-IR spectra of FH-0 single-crystals grown from DMF and BFH-10 xerogels prepared from toluene and DMF were measured (Figure 7). Bands of N=O stretching vibrations of hydrazide groups appear at 3235 and 3195 cm^{-1} as well as amide I bands at 1671 and 1611 cm^{-1}, indicating that the majority of the N–Hs of the hydrazide group are associated with the C=O groups via N–H...O=C hydrogen bonding in the xerogel and BFH-10 precipitated from THF (Figure 7, panels a and b), which are involved in intramolecular and intermolecular hydrogen bonding.\textsuperscript{20–22} Two new absorption bands for the BFH-10 xerogel formed from DMF appear at 1698 and 1651 cm^{-1} (Figure 7c); these are assigned to the free and associated C=O stretching vibration of DMF in the BFH-10 xerogel, respectively. This assignment is further supported by the observation of bands of the C=O stretching vibration of DMF at 1703 cm^{-1} (free) and 1650 cm^{-1} (H-bonded) in an FH-0 single crystal grown from DMF (Figure 7d). The $^1$H NMR spectrum of the xerogel formed from DMF (Figure S4) further confirms the presence of the DMF molecules (CDCl$_3$; δ of protons in DMF at 8.01, 2.93, and 2.87 ppm, respectively).\textsuperscript{29}

Combining the results of single crystal analysis of FH-0, SEM, FT-IR spectroscopy and $^1$H NMR spectroscopy of the xerogel, a possible packing motif for BFH-10 molecules in the xerogel is illustrated in Figure 8. In this model, BFH-10 molecules rotate to form a twisting supramolecular array through intermolecular hydrogen bonding between BFH-10 and DMF, as well as van der Waals interaction between alkyl chains. N–H protons of BFH-10 molecules form intermolecular hydrogen bonds with C=O groups of neighboring DMF molecules. Meanwhile, C=O groups of BFH-10 molecules are involved in intermolecular hydrogen bonding with –CH$_3$ groups of DMF molecules. Hydrogen-bonding interactions between DMF and

Figure 7. FT-IR spectra of (a) BFH-10 xerogel formed from toluene (1.9 $\times$ 10^{-2} mol L^{-1}), (b) BFH-10 precipitated from THF, (c) BFH-10 xerogel formed from DMF (1.2 $\times$ 10^{-2} mol L^{-1}), and (d) FH-0 single-crystals grown from DMF.

Figure 6. XRD patterns of BFH-10 (a) crystal, (b) xerogel formed from toluene (2.1 $\times$ 10^{-2} mol L^{-1}), and (c) xerogel formed from dimethylformamide (1.8 $\times$ 10^{-2} mol L^{-1}).
the gelator play an important role in the formation of helical ribbons of the BFH-10 xerogel prepared from DMF. During gelation, DMF molecules penetrate into the twisted molecular arrays, which are sustained by the weak van der Waals interactions between alkyl chains in adjacent screwed arrays. Balancing the stresses in the separated mono or multiple screwed molecular arrays might then give rise to the helical ribbons. More detailed investigation of microscale helical ribbons in BFH-10 gels from DMF is currently in progress.

**CONCLUSION**

It has been demonstrated that the NH-1 protons of BFH-n precipitated from either THF or DMF form intramolecular H-bonded 6-membered rings. While the corresponding NH-2 protons of BFH-n precipitated from THF form intermolecular hydrogen bonds with C=O groups of neighboring BFH-n molecules, the NH-2 protons of BFH-n precipitated from DMF form intermolecular hydrogen bonds with C=O groups of DMF molecules. Single crystal analysis of FH-0 shows that the C=O, –CH₃, and –CH groups of DMF molecules form multiple intermolecular hydrogen bonds with the –N–H and –C=O groups of neighboring FH-0 molecules. A double helix with a pitch of 24.2 Å along the c direction is observed in the FH-0 single crystal grown from DMF. Furthermore, left- and right-handed helical micrometer-length ribbons with nonuniform helical pitches were observed in a xerogel formed by the achiral BFH-10 precipitated from DMF, which demonstrates that intermolecular hydrogen-bonding interactions between DMF and the gelator play an important role in the formation of helical ribbons of BFH-10 xerogel prepared from DMF. This new self-assembly of dumbbell-shaped molecules containing hydrazone units offer excellent opportunities for the organization of other functional groups.

**REFERENCES**


