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Discovery of a Tetracontinuous, Aqueous Lyotropic Network Phase with Unusual 3D–Hexagonal Symmetry

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Network phase aqueous lyotropic liquid crystals (LLCs) are technologically useful materials with myriad applications in chemistry, biology, and materials science, which stem from their structurally periodic aqueous and hydrophobic nanodomains (~0.7–5.0 nm in diameter) that are lined with well-defined chemical functionalities. To date, the exclusive observation of bicontinuous cubic network phase LLCs (e.g., double gyroid, double diamond, and primitive phases) has fueled speculations that all stable LLC network phases must exhibit cubic symmetry. Herein, we describe the self-assembly behavior of a simple aliphatic gemini surfactant that forms the first example of a triply periodic network phase LLC with the 3D–hexagonal symmetry $P6_3/mcm$ (space group #193). This normal, tetracontinuous 3D–hexagonal network LLC phase $H_{193}$ partitions space into four continuous and interpenetrating, yet non-intersecting volumes. This discovery directly demonstrates that the gemini amphiphile platform furnishes a rational strategy for discovering and stabilizing new, three-dimensionally periodic multiply continuous network phase LLCs with variable symmetries and potentially new applications.

Introduction

The concentration-dependent self-assembly of small molecule amphiphiles into lyotropic liquid crystals (LLCs), comprising structurally periodic aqueous and hydrophobic nanodomains (~0.7–5 nm in diameter) lined with well-defined chemical functionalities, has wide-ranging implications in chemistry, materials science, and biology. Common LLC morphologies include cubic micellar (I), hexagonally packed cylinders (C), and lamellar (L$_{\alpha}$) phases. In typically narrow temperature and amphiphile concentration windows situated between the C and L$_{\alpha}$ phases, hydrated surfactants form complex ordered network phase LLCs. These three-dimensionally periodic structures are comprised of two or more continuous and interpenetrating networks of hydrophobic and aqueous nanodomains. Three highly symmetric, tricontinuous network phase LLCs with cubic symmetry (Q-phases) are ubiquitous (Fig. 1): double gyroid (G; $Ia\bar{3}d$ symmetry), double diamond (D; $Pn\bar{3}m$ symmetry), and primitive (P; $Im\bar{3}m$ symmetry). These triply periodic Q-phase LLCs are a focal point for the development of membranes for molecular separations and selective ion transport in fuel cells and batteries, and as templates for mesoporous inorganic materials. Bicontinuous “lipidic cubic phase” LLCs have also enabled the crystallization and X-ray
structure determination of important integral transmembrane proteins, including G–protein coupled receptors (GPCRs). Recent work has further established the utility of Q–phase LLCs in the delivery of both therapeutic siRNAs and small molecules.

Given that many surfactants form aqueous Q–phase LLCs, conventional wisdom suggests that cubic symmetries must underlie all stable triply periodic multiply continuous aqueous LLCs. This notion stems from the fact that G, D, and P LLCs maybe derived from arranging surfactant molecules along the high symmetry Schön G, Schwarz D, and Schwarz P triply periodic minimal surfaces (TPMS), respectively, to yield LLC morphologies exhibiting negative Gaussian (“saddle splay”) curvature. These molecular arrangements minimize packing frustration in the resulting assemblies that exhibit mean interfacial curvatures in between those of the C and L phases. In spite of theoretical predictions of the existence of stable, non–cubic LLC network phases derived from non–cubic TPMS, no such network phases in simple lipid/water mixtures have been observed.

Herein, we describe the first example of a triply periodic network phase aqueous LLC with the 3D-hexagonal P6/3/mcm symmetry (space group #193) derived from a simple, aliphatic dicarboxylate gemini surfactant. This tetacontinuous hexagonal network phase H193 partitions space into four continuous and interpenetrating, yet non–intersecting volumes. This unusual thermodynamic equilibrium nanostructure exhibits an interfacial curvature intermediate between that of the C and G phases, consistent with previous theoretical predictions by Schröeder–Turk et al. The discovery that this simple gemini (“twin tail”) surfactant architecture enables access to the H193 phase suggests a powerful molecular design strategy for synthesizing amphiphiles that stabilize new families of non–cubic LLCs with potentially useful properties.

Experimental

Materials

All materials and reagent grade solvents were purchased from the Sigma–Aldrich Chemical Co. (Milwaukee, WI) and used as received unless otherwise noted. 1,4-dibromobutane and 1,6-dibromohexane were distilled and stored under nitrogen. Diisopropylamine (Sigma–Aldrich) was distilled from CaH2 and stored under nitrogen. Hexamethylphosphoramide (HMPA) was distilled from CaH2 and stored over 4 Å molecular sieves. n–Butyllithium in hexanes was titrated using diphenylacetic acid in THF. Anhydrous and anaerobic THF was obtained by nitrogen degassing of analytical grade solvent for 30 min, followed by cycling through a column of activated alumina in a Vacuum Atmospheres Solvent purification system. The synthesis of disodium docosane–9,14–dicarboxylate (Na–74) was previously reported.

Molecular characterization

1H NMR and 13C NMR spectra were recorded in DMSO–d6 or methanol–d4 at 25 °C on both Varian MercuryPlus 300 and Bruker AC+ 300 spectrometers; spectra were referenced to the residual protiated solvent peaks in the samples. Mass spectrometry was performed using a Waters (Micromass) LCT® electrospray ionization time–of–flight mass spectrometer operating in negative ion detection mode. Samples dissolved in CH3OH were sprayed with a sample cone voltage of 20 V. IR spectra were recorded on a Bruker Tensor FTIR from 4000–500 cm–1, using a diamond crystal attenuated total reflectance (ATR) stage. Elemental analysis (C, H, N) were performed by Columbia Analytical Services (Tuscon, AZ), using Combustion/TC and IR detection.

Hexacosane–11,16–dicarboxylic acid

A 500 mL 2–necked round bottom flask fitted with an addition funnel was charged with a stirbar, THF (200 mL), and disisopropylamine (14.7 mL, 105 mmol) under N2(g). Upon cooling this solution to −15 °C in an ethanol/H2O/dry ice bath, n–BuLi (36.8 mL of 2.78 M solution in hexanes, 102 mmol) was added dropwise via addition funnel. The reaction mixture was then added for 30 min, after which a solution of dodecanoic acid (10.01 g, 49.97 mmol) in THF (50.0 mL) was added dropwise via addition funnel. The reaction mixture was warmed to 22 °C for 30 min. The reaction mixture was again cooled to −15 °C, and 1,4-dibromobutane (2.90 mL, 24.4 mmol) was added dropwise. The reaction was then allowed to warm to 22 °C and stirred for 14 h. The reaction was quenched by the addition of cold 10% HCl (aq) (100 mL), transferred to a separatory funnel, and the aqueous and organic layers were separated. The aqueous layer was extracted with ether (3 x 50 mL) and the combined organic layers were washed with 10% HCl (3 x 50 mL), water (50 mL), and saturated NaCl (aq) (50 mL). After drying over MgSO4, all volatiles were removed under vacuum. The crude solid was purified by recrystallization from EtOH. Yield: 9.29 g (84%); 1H NMR (300 MHz, DMSO–d6): δ 11.96 (2H, s, COOH), 2.15 (2H, m, CH–COOH), 1.56–1.08 (44H, m, CH2), 0.848 (6H, t, JH,H = 7.0 Hz, CH3–CH3). 13C NMR (75.4 MHz, DMSO–d6): δ 177.3 (C=O), 45.14 (CH), 32.21 (CH2), 32.17 (CH2), 32.04 (CH3), 31.72 (CH2), 29.39 (CH3x2), 29.31 (CH2), 29.13 (CH2), 27.24 (CH2), 27.18 (CH3), 22.50 (CH3), 14.28 (CH3). MS (ESI–TOF) calcd. m/z for C24H45O4 453.4, found: 453.4.

Disodium hexacosane–11,16–dicarboxylate (Na–94)

In a flask equipped with a stirbar, hexacosane–11,16–dicarboxylic acid (2.505 g, 5.508 mmol) and Na2CO3 (0.584 g, 5.51 mmol) were suspended in CH3OH (55.0 mL). The mixture was stirred at 22 °C until it became homogeneous, and then stirring was continued for 1 h. All volatiles were removed under vacuum, and residual water and methanol were azeotropically distilled using CaH2 three times under vacuum. Yield: 2.75 g (quant.). 1H NMR (300 MHz, CD3OD): δ 2.14
Lyotropic Phase Sample Preparation

Aqueous lyotropic liquid crystal (LLC) samples were prepared for X-ray scattering analysis by weighing appropriate amounts of rigorously dried Gemini salt and doubly distilled water into a vial and centrifuging the tightly capped mixture for 10 min, manual mixing using a spatula, and an additional 10 min of centrifugation. After centrifugation, the samples were allowed to rest at 22 °C for at least 12 h prior to X-ray analysis. Samples were stored in tightly sealed vials whenever possible during sample preparation, in order to minimize water loss through evaporation.

Small–angle X–Ray Scattering

Synchrotron small–angle X–ray scattering (SAXS) measurements were performed at the 12-ID–B beamline at the Advanced Photon Source (Argonne, IL). Experiments employed a beam energy of 12 keV (λ = 1.034 Å) and a 2.550 mm–pixel resolution. Mounted in aluminium DSC pans, LLC samples were heated to a desired temperature in a Linkam DSC hot stage and allowed to equilibrate for 5 min prior to data collection (exposure time ~1 s). 2D patterns were azimuthally integrated to obtain Log(Intensity(q) v. q) plots using the DataSqueeze software package (http://www.datasqueezesoftware.com/).

Detailed analysis and pattern indexing of synchrotron SAXS data employed the Jade 5 software package (Materials Data, Inc., Livermore, CA). Rietveld refinement was conducted using the JANA2006 software package to extract structure factor intensities and peak full–width at half maxima associated with each allowed SAXS peak for LLC phases of interest. From these data, we generated a text file input for use with the charge–flipping algorithm SUPERFLIP to obtain an electron density reconstruction based on the SAXS data.

Rheological Measurements

Frequency–dependent storage (G’) and loss (G”) moduli of LLCs were measured using a TA Instruments ARES–LS2 rheometer in strain–controlled mode at 25 and 55 °C. All measurements were performed in a parallel–plate geometry with a 50 mm Peltier temperature–controlled bottom plate and a 8 mm top plate. To minimize water loss from the samples, a collar with a water–saturated sponge was fitted onto the bottom plate to surround the sample without making contact with the upper plate fixture or the sample. Dynamic strain sweep measurements were performed over the range |γ| = 0.03–1% with ω = 0.1 Hz at 12 logarithmically spaced values in order to identify the linear viscoelastic regime of the material. Dynamic frequency sweep measurements were then performed at |γ| = 0.1% at 19 logarithmically spaced frequencies with 0.01 ≤ ω ≤ 50 Hz.

Results & Discussion

Gemini amphiphiles, derived from linking two single–tail surfactants at or near their hydrophilic headgroups, exhibit an increased propensity to form G–phase LLCs in water.20–22 Recently, we reported that anionic gemini dicarboxylate amphiphiles (Scheme 1) form unusually stable, normal double gyroid (G1) LLC phases over large amphiphile concentration windows up to 20 wt% wide with unexpected thermal stabilities over the range T = 25–100 °C.17 Our studies of decanoic acid–derived surfactants such as Na–74 highlighted the sensitive dependence of the cubic G1–phase stability on the identity of the hydrophilic headgroup, the nature of the headgroup–counterion interaction, and the structure of the aliphatic surfactant scaffold. Subsequent molecular dynamics simulations of these self–assembled phases have demonstrated that these aliphatic gemini dicarboxylates adopt splayed chain molecular conformations, the subtle supramolecular packings of which are electrostatically controlled by the extent of counterion–headgroup dissociation at a given hydration level.23

In the course of our extended studies of the LLC phase behavior of gemini dicarboxylate surfactants, we were led to investigate the effects of hydrophobic tail length on the unusual supramolecular self–assembly of these molecules. Consequently, we synthesized the gemini surfactant Na–94 derived from dodecanoic acid and 1,4–dibromobutane in high yields according to Scheme 1. Using temperature–dependent small–angle X–ray scattering (SAXS), we investigated the LLC phases formed by variably hydrated samples of Na–94 over the concentration range 30–100 wt% amphiphile in H2O between T = 22–100 °C. We assigned the resulting LLC morphologies based on the positions of the observed SAXS peaks according to well–documented methods.4,17 For example, the azimuthally–integrated SAXS patterns shown in Fig. 2a and 2d at 50 and 65 wt% Na–94 correspond to H and G phase LLCs, respectively. The results of our SAXS analyses are summarized in the temperature versus amphiphile concentration LLC phase diagram shown in Fig. 3. Below 40 wt% Na–94 in H2O, micellar solutions with no periodic long–range order form. Upon systematically decreasing the water content, we noted...
that Na–94 forms an unusual sequence of composition–
dependent LLC phases: C → unknown → G → Lα. Since the
C, G, and unknown phases occur at higher water contents than
the zero–curvature Lα phase formed by Na–94, we deduce that
interfacial curvature toward the lipid domains is favored. Thus,
these are “normal” (Type I) LLC phases in which water is the
majority phase.

In the composition range 52–60 wt% Na–94 in H2O, we
recorded a series of SAXS patterns that exhibit peak positions
and a scattering intensity envelope inconsistent with any known
phase coexistence or any previously reported LLC structure.
The composition–dependent progression of SAXS patterns
shown in Fig. 2 illustrates the formation of a new, unidentified
phase in nearly pure form near 60 wt% Na–94 in water at 25
°C. A sample of 57.5 wt% Na–94 in water at 40 °C forms a
pure phase that exhibits the SAXS pattern shown in Fig. 4, in
which the relative positions of the observed peaks occur at
approximately \( q/q^* \approx \sqrt{4/3}, \sqrt{4}, \sqrt{5}, \sqrt{47/9}, \sqrt{16/3}, \sqrt{25/4}, \sqrt{9}, \sqrt{28/3}, \sqrt{95/9}, \sqrt{12}, \sqrt{16} \) with \( q^* = 0.0856 \text{ Å}^{-1} \) (see Table
S1 for detailed listing of the observed peak positions).
Qualitatively, the apparently homogenous optical birefringence
of this new phase implies its underlying non–cubic symmetry.
Near 80 °C, SAXS analyses also indicate that this new LLC
phase undergoes a thermoreversible order–order phase
transition to the cubic G1–phase (Fig. S1), which suggests the
thermodynamic equilibrium nature of this new LLC phase.

The observed SAXS peak positions for this new phase are
consistent with the three–dimensional hexagonal space group
symmetry \( P6_3/mcm \) (space group \#193), denoted \( H1^{91} \) wherein
the subscript designates a Type I phase, with unit cell
dimensions \( c = 6.63 \text{ nm} \) and \( a = 7.37 \text{ nm} \) (\( c/a = 0.900 \)). Table
S1 contains a complete list of 20 observed scattering peak
positions, their corresponding Miller indices, and the calculated
residuals that are in exceptional agreement with the expected
values (≤ 0.004° deviation). While the \( P3c1, P3c1, P6_3/mcm, \) and
\( P6c2 \) symmetries also produce the same scattering signature,
we favor the highest symmetry $P6_3/mcm$ space group per crystallographic convention.\textsuperscript{24} The Miller indices associated with the peaks in the azimuthally-integrated one-dimensional scattering intensity profile are given in Fig. 4 (see also Fig. S2).

SAXS patterns associated with LLC samples comprising 55.0 and 60 wt% Na–94 in water over the temperature range 30–65 °C are consistent with H$_i^{193}$ phase coexistence with either of the adjacent C$_i$ or G$_i$ phases (Fig. 2). The notable lack of C$_i$/G$_i$ phase coexistence below 70 °C at any concentration implies that the H$_i^{193}$ structure is a new, equilibrium LLC mesophase. In a LLC composition window similarly situated between the C$_i$ and G$_i$ phases, we note that the structurally homologous gemini surfactant Na–74 exhibits a similar previously unassigned phase coexistence of G$_i$ and H$_i^{193}$ phases near 55 wt% surfactant in H$_2$O (Fig. S3).\textsuperscript{17} For the Na–74 samples, we also deduce a similar unit cell ratio $c/a = 0.91$.

The percolating aqueous and hydrophobic domain structures associated with triply periodic network phase LLCs manifest in their solid–like linear viscoelastic rheological properties. Mezzenga \textit{et al.} demonstrated that inverse (Type II) bicontinuous cubic G$_{III}$ and D$_{III}$ LLCs typically exhibit large and nearly frequency–independent dynamic elastic storage shear moduli G'(o) ≈ 0.5–1 MPa over broad frequency ranges.\textsuperscript{25} Consistent with this notion, the G$_i$–phase formed by 60 wt% Na–94 in H$_2$O exhibits a frequency–independent G'(o) ≈ 10 MPa at 25 °C (Fig. S4). However, the discontinuous C$_i$–phase formed at 50 wt% Na–94 exhibits a much lower G'(o) ≈ 0.1 MPa at 25 °C, which is notably frequency dependent (Fig. S5). Dynamic elasticity measurements reveal that the H$_i^{193}$ phase exhibits a substantial G'(o) ≈ 1 MPa, which is relatively insensitive to frequency between 0.01 Hz ≤ $\omega$ ≤ 50 Hz at $T = 25$ and 55 °C (Fig. S6). We note that the elastic moduli G'(o) for the H$_i^{193}$ and G$_i$ phases differ by one order of magnitude, and that there are significant differences in the frequency-dependent viscous modulus G''(o) for these phases. We speculate that these differences arise from differences in water contents of these LLC phases and the preferred deformation axes within these Maxwellian solids; however, the shear rheology of LLCs is quite poorly understood. While this analysis is not necessarily definitive, the frequency-independent nature of G'(o) strongly suggests that the H$_i^{193}$ phase is triply periodic.

Since the nanoscale feature sizes and non–covalent, water–laden nature of the new H$_i^{193}$ LLC phase prohibited its direct, real–space imaging by transmission electron microscopy (TEM), we used the high–resolution SAXS pattern in Fig. 4 to reconstruct an electron density map of this phase. Rietveld refinement of this SAXS pattern using the $P6_3/mcm$ space group symmetry in the \textit{JANA2006} software package enabled extraction of the observed structure factor intensities (Fig. S7 and Table S2). Using these data as inputs for the charge–flipping algorithm \textit{SUPERFLIP},\textsuperscript{19, 26} we obtained an excellent fit of the experimental diffraction pattern from which we reconstructed the electron density contrast map shown in Fig. 5a (for details see Supporting Information Fig. S10). This map illustrates the labyrinthine nature of the H$_i^{193}$ phase, which confirms its triply periodic network morphology. An idealized wireframe drawing of the tectocontinuous H$_i^{193}$ structure in Fig. 5b depicts the three identical, non–intersecting and interpenetrating lipid networks within a matrix of water as the fourth network. Each of the three lipid networks in the H$_i^{193}$ phase is an [8,3]–net with $P6_3/mcm$ symmetry (space group $P6_3/mcm$), fashioned from three–fold connectors with either 0° or 90° dihedral angles between them (Fig. 5c). The [8,3]–net designation in Wells’ notation in refers to the fact that each symmetric network is comprised of 8–membered rings of three–fold connectors. This unusual structure is also referred to as the 3–etc net.\textsuperscript{14} Note that the G$_i$–phase formed upon heating the H$_i^{193}$ network above 80 °C comprises two interpenetrating lipid networks in a matrix of water, wherein each network is a [10,3]–net comprised of three–fold connectors with a 70.5° dihedral angle between them.\textsuperscript{27} The formation of three–fold connectors in these network LLCs is consistent with their lower degrees of packing frustration and smaller deviations from constant mean interfacial curvature, as compared to higher valency four– and six–fold connectors.\textsuperscript{28}

While true epitaxial relationships between the H$_i^{193}$ phase and the adjacent C$_i$ and G$_i$ phases cannot be rigorously deduced from our SAXS data, the H$_i^{193}$ model suggests some structural features that correlate with quantitative crystallographic

**Figure 5.** (a) Structural models for the tectocontinuous H$_i^{193}$ phase LLC. (A) Electron density reconstruction derived from the SAXS pattern in Fig. 2b, derived from Rietveld refinement of the data and subsequent use of the charge flipping algorithm \textit{SUPERFLIP}. (b) An idealized wire-frame model of the three interpenetrating and non-intersecting lipidic [8,3]-nets (3-etc) with overall $P6_3/mcm$ symmetry, in which each network is colored for visual clarity. (c) A wire-
Fig. 6. Schematic depiction of the structural relationships between the C$_2$ and H$_4$$^{193}$I phases. (Top) A unit cell projection along the C$_2$ phase c-axis, with an embedded black rhombus that outlines the unit cell, shows that the intercylinder distance is half of the unit cell dimension a for the tetracontinuous H$_4$$^{193}$I network phase. Arrows indicate (10) and (100) lattice vectors of the C$_2$ and H$_4$$^{193}$I phases respectively. (Bottom) A unit cell projection along the a-axis of the C$_2$-phase, dehydration of the results in the regular buckling and pinching off of the cylinders to form the three networks of three-fold connectors in the H$_4$$^{193}$I phase.

relationships apparent in the SAXS data in Fig. 2. The (100) reflection at $q = 0.0908$ Å$^{-1}$ of the H$_4$$^{193}$I network that corresponds to the unit cell $a$ parameter occurs at nearly one half of the value of the (10) reflection of the coexisting C$_2$ phase ($q = 0.186$ Å$^{-1}$) in the LLC composed of 55.0 wt% Na–94 (Fig. 2b). Thus, the $a$-axis dimension of the H$_4$$^{193}$I phase is nearly double that of the intercylinder distance in the C$_2$-phase. This relationship suggests that removing water from the C$_2$-phase results in: (1) regular buckling of every other lipid cylinder to form the zig–zag pattern of connectors running parallel to the c-axis (0° dihedral angle between connectors), while (2) the remaining lipid cylinders pinch off to form the stacks of staggered, three–fold connectors that connect the buckled cylinders (Fig. 6). This morphological relationship only holds if the H$_4$$^{193}$I phase is comprised of three interpenetrating lipid networks. The staggered stacking of three–fold connectors along the c-axis of the H$_4$$^{193}$I LLC is also reminiscent of the stacking of staggered connectors along the (111) axis of the G$_1$–phase. The distance between these stacks in the H$_4$$^{193}$I phase is $a = 7.34$ nm, which is similar to the unit cell parameter $a = 7.60$ nm for the coexisting G$_1$–phase at 60.0 wt% Na–94 (Fig. 2c). The difference in these values probably arises from the substantial differences in network connectivity and symmetry between these two phases. Efforts to prepare highly–oriented pseudo–“single crystal” samples of the H$_4$$^{193}$I phase for careful studies of these proposed morphological relationships have been thus far unsuccessful.

Aside from the ubiquity of C-phase LLCs with 2D–hexagonal P6/mmm symmetry (space group #183), LLCs with 3D-hexagonal symmetries are quite rare. Impéror-Clerc and co-workers previously reported the observation of a discontinuous micellar Type I LLC phase with P6$_3$/mm symmetry (space group #194), comprising hexagonally-closest packed spherical micelles derived from a non-ionic surfactant. In a related report, Seddon and co-workers more recently described the discovery the corresponding Type II LLC in which reverse spherical micelles form a hexagonally-closest packed lattice. Thus, our observation of the H$_4$$^{193}$I LLC constitutes only the third report of an LLC with 3D-hexagonal symmetry and the first hexagonal network phase.

Schröder–Turk and Hyde posited the existence of a variety of triply periodic multiply continuous network LLCs with mean interfacial curvatures in between those of the L$_a$ and the C–phases. However, non–cubic, network phases in self–assembling soft materials are exceedingly rare, having been only recently observed in block copolymers. Only the cubic G, D, and P network phase LLCs have been previously observed in simple lipid/water mixtures, leading to speculations that the ubiquity of cubic LLCs stems from their intrinsically low degrees of packing frustration. Schröder–Turk combined geometric packing arguments with calculations of the bending rigidity and stretching frustrations associated with lipids packed on the Schwarz H minimal surface to predict the stability of an inverse (Type II) network LLC with P6$_3$/mm symmetry. More sophisticated calculations later revealed that the inverse P6$_3$/mmc structure with mean interfacial curvature between that of the G$_a$ and H$_4$ phases is a more likely LLC candidate in H$_2$O. However, these analyses are only indirectly relevant to the normal H$_4$$^{193}$I phase reported here.

In 2009, Ying and co–workers used electron diffraction TEM studies of a few well–ordered particles of their newly synthesized mesoporous silicate IBN–9 to identify its underlying P6$_3$/mmc symmetry. Formed from a complex mixture of a silicate precursor, NH$_4$OH, H$_2$O, and a 4° ammonium surfactant likely under non–equilibrium conditions, IBN–9 consists of a single silicate sheet that partitions space into three equivalent [8,3]–networks. Thus, IBN–9 is structurally analogous to the aqueous matrix of our H$_4$$^{193}$I LLC phase. While the formation of the IBN–9 structure indirectly suggests the formation of the normal H$_4$$^{193}$I, our data constitutes the first direct observation of the H$_4$$^{193}$I phase as an equilibrium structure in simple lipid/water mixtures.

Conclusions

The discovery of the tetracontinuous H$_4$$^{193}$I network phase in gemini dicarboxylate surfactant LLCs establishes that the gemini architecture is a powerful molecular design motif for stabilizing both known and new triply periodic multiply continuous morphologies in self–assembled soft materials. Careful comparison of the previously reported Na–74 and Na–76 LLC phase diagrams with that of Na–94 strongly suggests that: (1) longer aliphatic linkers between the surfactant tails destabilize constant mean interfacial curvature phases such as L$_a$ and H$_4$; and (2) longer aliphatic surfactant tails drive formation of constant mean curvature phases. Therefore, tuning the molecular aspect ratio of a gemini amphiphile specified by
the relative lengths of the surfactant linker and tails provides enticing opportunities for subtly adjusting LLC interfacial curvature to discover new, triply periodic multiply continuous LLC morphologies with percolating and chemically well-defined nanopores. Initial atomistic molecular dynamics simulations of gemini surfactant self-assembly have revealed the complex interplay of intramolecular headgroup repulsions in defining preferred, anisotropic surfactant conformations that undergo supramolecular packing into non-constant curvature LLCs as specified by the extent of counterion-headgroup association. We anticipate that the “gemini” molecular design principle may extend to the formation of new technologically useful triply periodic network phases in other classes of self- assembling soft materials, including oligomeric and polymeric surfactants as well as block copolymers.

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Notes and References

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An aliphatic gemini dicarboxylate surfactant is shown to form a new 3D-hexagonal lyotropic liquid crystalline phase with $P6_3/mcm$ symmetry.
57.5 wt% in H$_2$O

Na-94