Isotropic, nematic, and lamellar phases in colloidal suspensions of nanosheets

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The phase diagram of colloidal suspensions of electrically charged nanosheets, such as clays, despite their many industrial uses, is not yet understood either experimentally or theoretically. When the nanosheet diameter is very large (~100 nm to 1 μm), it is quite challenging to distinguish the lamellar liquid-crystalline phase from a nematic phase with strong stacking local order, often called “columnar” nematic. We show here that newly upgraded small-angle X-ray scattering beamlines at synchrotron radiation facilities provide high-resolution measurements which allow us to identify both phases unambiguously, provided that single domains can be obtained. We investigated dilute aqueous suspensions of synthetic Sb2P2O7−n nanosheets that self-organize into two distinct liquid-crystalline phases, sometimes coexisting in the same sample. Close examination of their X-ray reflection profiles in the directions perpendicular to the director demonstrates that these two mesophases are a columnar nematic and a lamellar phase. In the latter, the domain size reaches up to ~20 μm, which means that each layer is made of >600 nanosheets. Because the lamellar phase was only rarely predicted in suspensions of charged disks, our results show that these systems should be revisited by theory or simulations. The unexpected stability of the lamellar phase also suggests that the rims and faces of Sb2P2O7−n nanosheets may have different properties, giving them a patchy particle character.

Significance

Colloidal suspensions of electrically charged nanometric sheets (nanosheets), like graphene oxide or clays, which are widely used in industry, form liquid-crystalline phases. These include the nematic phase, where all nanosheets are approximately parallel, and the lamellar phase, where they also form equidistant layers. When the particle diameter is large (100 nm to 1 μm), distinguishing these phases is quite challenging. Using newly available synchrotron small-angle X-ray scattering setups, we unambiguously identified both phases, in H3Sb2P2O14 nanosheet suspensions, by analyzing their X-ray scattering patterns. The lamellar domain size reaches 20 μm, which means each layer is made of ~1,000 nanosheets. Because the lamellar phase was rarely predicted in suspensions of charged disks, these systems should be revisited by theory or simulations.

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third-generation synchrotron radiation facilities allowed us to perform such a study. We used an upgraded small-angle X-ray scattering (SAXS) beamline to distinguish a lamellar phase (Fig. 1A) from a so-called “columnar” nematic phase (Fig. 1B). This latter phase, of nematic symmetry, is made of disks which tend to stack over short distances and form short columns that align parallel to each other. We revisited the phase diagram of colloidal suspensions of synthetic H₃Sb₂P₂O₁₄ nanosheets (13), a material of interest for solid-liquid extraction of rare earth elements (21), and discovered that in addition to the isotropic liquid phase, many samples display two different coexisting mesophases that we identified as nematic and lamellar by high-resolution X-ray scattering. Our study also demonstrates that in the lamellar phase, the positional order of the nanosheets extends over a length scale of at least ∼20 μm and involves at least 600 nanosheets per layer.

Materials and Methods

Synthesis and Characterization (XRD). H₃Sb₂P₂O₁₄ was synthesized according to a previously published method (13). The synthesis was performed in two consecutive steps: the first high-temperature reaction led to the synthesis of K₅Sb₂P₂O₁₄, and the second led to the title compound by cation exchange in aqueous solution. For the synthesis of K₅Sb₂P₂O₁₄, a stoichiometric mixture of NH₄H₂PO₄ (23 mmol, 2.65 g, synthesized from mixing stoichiometric amounts of H₂PO₄ 85 wt % in water with NH₃ 30 wt % in water), Sb₂O₃ (17 mmol, 5.06 g; Merck), and KNO₃ (34 mmol, 3.51 g; Prolabo) was placed in a platinum crucible and heated in air, first at 300 °C (ramp of 50 °C/h) for 10 h to decompose NH₄H₂PO₄, then at 1,000 °C (ramp of 50 °C/h) for 100 h, yielding 8.78 g (12 mmol; yield = 100%) of K₅Sb₂P₂O₁₄ (X-ray powder diffraction showed impurities level <1%).

For the second synthesis step, 3.4 g of K₅Sb₂P₂O₁₄ powder was thoroughly ground and stirred in a 0.33 M solution of 7.5 M nitric acid at 50 °C for 22 h. During this process, the solid was centrifuged (3,000 g) and rinsed with 18 MΩ water a few times until a swelling of the solid was noticed. This procedure was repeated three times to ensure a complete exchange of the alkaline metal cations for protons to yield H₅Sb₂P₂O₁₄ (over 98% potassium to proton exchange). It should be noted that after the last cation exchange and centrifugation, the solid was only rinsed once to avoid too much swelling.

Further purification of H₅Sb₂P₂O₁₄ was achieved according to the following procedure: the gel collected after cation exchange was placed in a regenerated cellulose tubular membrane (Cellu Sep, width 46 mm, thickness 28 μm, pore size 1 nm) and was subjected to dialysis in 18 MΩ water, replacing the water regularly, until the nitrate ion concentration decreased to less than 1 ppm (JBL, nitrate test). During this dialysis, the lamellar, protonated phosphoatoamionic acid H₅Sb₂P₂O₁₄ kept on swelling.

Size fractionation procedures were then applied by centrifuging, on a Heraeus Biofuge Primo centrifuge (Thermo Fisher Scientific), the stock suspension at 8,000 g for 86 h, in 50 mL Falcon polypropylene conical tubes, leading to five different size fractions from top to bottom, named A to E. All fractions were carefully removed one by one, from top to bottom, using a pipette. The bottom fraction E was very small and consisted of white and insoluble solid impurities; it was discarded. The top two fractions were both transparent liquids and were separated by an interface. When observed between crossed polarizers, the top fraction showed some flow birefringence when agitated, whereas the one underneath was permanently birefringent. Both fractions were concentrated by rotoevaporation, yielding fractions A and B, for the very top and second ones, respectively. The third fraction from the top, labeled C, was made of a thixotropic white gel. It was separated by two interfaces from fraction B over it and fraction D below it, the latter being a dense white gel. After a preliminary fast exploration of the phase diagram of each fraction, samples were prepared in glass test tubes by diluting the fractions, up to a dilution factor of 100, so as to span the concentration range of interest.

Thermogravimetric Analysis. The weight fractions of the suspensions were determined by thermogravimetric analysis (TGA) under N₂ (model no. TGA92 from SETARAM Inc.) by heating to 250 °C. All calcined materials collected after TGA were white, indicating that no organic contamination arose from the cellulose dialysis membranes. Weight fractions for A to D were measured to be 1.01 wt %, 1.77 wt %, 1.95 wt %, and 5.63 wt %, respectively.

Atomic Force Microscopy. Nanosheets were imaged by atomic force microscopy (AFM) to determine their size. They appeared as irregular flat objects of 1.1 ± 0.2 nm thick [in agreement with the crystallographic thickness of 1.10 nm of a single Sb₂P₂O₁₄ layer (13, 22)]. Approximating these objects as regular disks, their average equivalent diameters (and SDs), for fractions A to D, are 44 (31) nm, 660 (615) nm, 790 (395) nm, and 960 (490) nm, respectively. A typical AFM image is given in SI Appendix, Fig. S1A, and the whole size distributions for A to D are shown in SI Appendix, Fig. S1B–E.

Optical Studies. Test tubes were filled with samples of different concentrations and were first observed with the naked eye, either in the natural light of an LED hand lamp shone from below the tubes or between crossed polarizers in transmission. Reported observations were made 6 mo after the dilutions were performed, which gave time for the samples to reach equilibrium. No significant evolution has been observed since.
Polarized-light microscopy was performed with a BX51 microscope (Olympus) equipped with an Olympus digital camera. For texture observations, samples were filled in flat glass capillaries (Viroctom) of 100- or 200-μm thickness, 2-mm width, and about 80-mm length. The capillaries were flame-sealed after filling and then stored vertically. Their evolution in time was monitored over several months, and they remained stable for more than a year.

**SAXS Studies.** Samples were inserted into cylindrical Lindemann glass capillaries of 1-mm diameter (Glas-Technik & Konstruktion) that were flame-sealed and left vertical in the field of gravity. Many of the X-ray capillaries were carefully scanned by X-ray scattering to characterize the different phases and to find areas where the X-ray beam impinged on only one (or a few) domain in reflection condition. One particular capillary was filled with a sample taken from a triphasic test tube (fraction C, 0.49 wt %) after remixing. Within 3 d, this triphasic sample appeared triphasic again and remained stable.

A first set of SAXS experiments were carried out at the SWING beamline of the SOLEIL synchrotron radiation facility. Measurements were made using a fixed energy of 12.0 keV and a sample-to-detector distance of 6.56 m. The typical accessible range of scattering vector modulus q was $10^{-2}$ to $1 \text{ nm}^{-1}$ ($q = (4\pi\lambda)/\sin\theta$, where $\theta$ is the scattering angle and $\lambda = 0.1033 \text{ nm}$ is the wavelength). Scattering patterns were recorded on an Avies 170170 CCD camera formed with four detectors and placed in a vacuum detection tunnel. The pixel size was 41.7 μm, and the binning was either 2 x 2 or 4 x 4; the beam size at the detection level was about 50 μm. The scattering patterns were radially averaged to obtain the scattering curves (q). The particular (fraction C, 0.49 wt %) capillary mentioned above was scanned by SAXS on SWING every 200 μm from its very bottom up to 28 mm above (well into the isotropic phase). The lamellar period, d, was obtained by using the relation $d = 2\pi/\Delta q$, where $\Delta q$ is the scattering vector modulus at the maximum of the reflection.

High-resolution SAXS experiments were also performed at the recently upgraded high-brilliance ID02 beamline of the European Synchrotron Radiation Facility in Grenoble, France (23). The X-ray energy was 12.48 keV ($\lambda = 0.0983 \text{ nm}$), and the sample-to-detector distance was 20.007 m, which leads to an accessible q range of 0.005–0.15 nm$^{-1}$. The detector was a Frelon camera with pixel size of 24 μm. In high-resolution SAXS conditions, the instrumental resolution is mostly governed by the beam size at the detector level (120 x 120 μm$^2$). In reciprocal space, the resolution can be modeled as a Gaussian of full-width at half-maximum $2.7 \times 10^{-4} \text{ nm}^{-1}$, which gave a coherence length of ~20 μm.

**Results and Discussion**

*Observations of Samples with the Naked Eye.* For all four size distributions A to D, some of the sample tubes clearly displayed two liquid–liquid interfaces that separated the liquid samples in three regions. [The A fraction already showed three phases at its initial synthesis (maximum) weight fraction of 1.01 wt %, B, C, and D remained monophasic down to 1.04, 0.84, and 0.81 wt %, respectively.] Typical triphasic sample tubes observed with the naked eye, either in natural light or between crossed polarizers, are shown in Fig. 2. The upper regions were clear and isotropic, but they always showed a strong flow birefringence at the slightest movement, especially near the top interface. Below this phase, a usually small birefringent region (5–15% of the overall height of the sample) was observed between two interfaces. The lowest phase was always birefringent and often displayed visible light diffraction, resulting in a blueish hue, as previously observed for the lamellar phase close to its maximum swelling periodicity (~ 200 nm) (13, 24). The appearance of three coexisting phases in samples of several centimeters height left under the influence of gravity is actually due to a vertical particle concentration gradient (evidence of this gradient was obtained by X-ray scattering; SI Appendix, Fig. S2). Such a phenomenon was already reported for colloidal suspensions of sterically stabilized gibbsite nanodisks in toluene (25).

In this system of hard disks, the proportions of isotropic, nematic, and columnar phases were precisely modeled by balancing the effect of gravity with osmotic pressure. The gravitational length scale of Sb$_2$P$_2$O$_{14}$ nanosheets, $l_g = k_BT/$(gVΔρ) (where $k_B$ is the Boltzmann constant, T is the temperature, g is the gravity acceleration, V is the particle volume, and Δρ is the specific mass contrast of the particle with the solvent), is of the order of 1 mm, like that of gibbsite disks. However, detailed modeling of this system is much more complicated than for gibbsite suspensions because the equation of state of charged nanosheets is much less understood than that of hard disks. A similar phase coexistence was also reported for

![](image_url)

**Fig. 3.** Photographs of samples of H$_3$Sb$_2$P$_2$O$_{14}$ suspensions observed by polarized-light microscopy. (A) Suspension of nanosheets (size C, 0.35 wt %), showing three coexisting phases (IL, isotropic liquid; L, lamellar phase; and N, nematic phase) in a flat glass capillary. (The width of the capillary is 2 mm.) (B) Typical nematic texture of the intermediate phase (size A, the overall sample tube weight fraction is 0.51 wt %). (Scale bar, 100 μm.) (C) Texture of the bottom phase (size C, the overall sample tube weight fraction is 0.35 wt %). (Scale bar, 100 μm.) The polarizer and analyzer directions are parallel to the edges of the photographs.

**Fig. 4.** A 2-mm-diameter X-ray capillary (size C, 0.49 wt %) was scanned from bottom to top, every 0.2 mm on the SWING SAXS beamline. Representative patterns are displayed: (A) isotropic phase just above the I-N interface, nematic phase (B) just below the I-N interface and (C) just above the N-L interface, (D) lamellar phase just below the N-L interface, and (E) in the middle of the lamellar phase region. (Note that in the photograph on the left, colored spots can be seen in the lamellar phase due to diffraction of visible light.)

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Description of Microscopic Textures. The coexistence of three different phases in the same sample was also observed in many flat capillary tubes by optical microscopy (Fig. 3A). For example, a sample of size C nanosheets, at 0.35 wt %, left undisturbed for several months in the field of gravity, displays three different liquid phases separated by sharp interfaces. The top phase is optically anisotropic, whereas both the middle and bottom phases are birefringent and therefore liquid-crystalline. The middle phase is strongly homeotropically anchored on the glass surfaces so that it mostly looks dark in the polarizing microscope. Such homeotropic anchoring is expected for nematic suspensions of plate-like particles for entropic reasons (27). This alignment could easily be disturbed by applying a magnetic field of ~800 mT delivered by small permanent magnets. After a minute, the texture became distorted, and the phase birefringence was revealed (SI Appendix, Fig. S3). After field removal, full homeotropic alignment of this mesophase was recovered after about an hour. In another sample that has weaker homeotropic anchoring, the middle phase shows a typical threaded texture (Fig. 3B), which strongly suggests that it is a nematic phase. This conclusion was indeed confirmed by SAXS experiments (see below). The nematic phase of H$_3$Sb$_2$P$_2$O$_{14}$ suspensions looks quite similar to that displayed by aqueous dispersions of HSbP$_2$O$_6$ (13), bentonite (28), beidellite, and nontronite natural clays (16). In contrast, the liquid-crystalline bottom phase did not show strong homeotropic anchoring and was not affected by the magnetic field, probably because it is more viscoelastic than the middle phase. Its optical texture (Fig. 3C) is also quite different because it is slightly reminiscent of the mosaic textures of ordered smectics (29). However, the texture is not typical enough to allow unambiguous identification of this meso-phase, a task that requires X-ray scattering experiments.

Triphasic Capillary Tube Studied by SAXS. Representative SAXS patterns recorded while scanning a vertical capillary tube filled with a triphasic suspension of H$_3$Sb$_2$P$_2$O$_{14}$ are shown in Fig. 4. In the upper region, all patterns are typical of an isotropic phase, even just above the I/N interface (Fig. 4A). The next pattern in the series, just below this interface, is strongly anisotropic and presents broad peaks (d ~ 230 nm); it is typical of an oriented nematic phase (Fig. 4B). Similar patterns were observed throughout the whole intermediate region till just above the bottom interface (Fig. 4C). Just below that interface and throughout the whole bottom region, very anisotropic patterns displaying sharp and intense diffraction peaks [with up to six equidistant (00l) orders] were recorded (Fig. 4 D and E). Such diffraction patterns strongly suggest the existence in this system of a lamellar phase, somewhat reminiscent of that of charged surfactant membranes diluted in pure water. The lamellar period decreases from 230 to 190 nm from top to bottom, due to the above-mentioned gravity-induced particle concentration gradient (SI Appendix, Fig. S2). In many scattering patterns, the X-ray reflections and diffuse peaks of both liquid-crystalline phases had maxima around the horizontal direction, which reveals some alignment of the nanosheets parallel to the capillary axis. This partial alignment may be due to flow during sample filling and/or slow nanosheet sedimentation under gravity.

High-Resolution SAXS. High-resolution SAXS experiments on selected samples allowed us to probe the large-scale structures of the different liquid-crystalline phases observed between crossed polarizers. Fig. 5A presents the X-ray scattering pattern of a well-aligned region of the middle phase. This pattern only shows broad scattering features arising from liquid-like positional correlations of the nanosheets. Because the pattern is also quite anisotropic, it is typical of a nematic phase, in agreement with the microscopic texture observations. A radial scan of the scattered intensity reveals two broad peaks whose scattering vector moduli are in a ratio of 1:2; they correspond to a period of
(~240 nm (Fig. 6A). The broad peaks indicate a marked positional short-range order in the nematic phase which can then be regarded as a columnar nematic (Fig. 1B). The peak width in the perpendicular direction may be due to either the finite diameter of the nanoparticles or their orientational distribution, quantified by the nematic order parameter, S. Because the transverse width of the second-order broad peak is about twice that of the first, we conclude that the broadening of the peak is mostly due to the orientational distribution. Then, by neglecting in a first approximation the broadening due to the finite nanosheet diameter and by using classical methods, S can be estimated by fitting an azimuthal scan of the scattered intensity (SI Appendix, Fig. S4) going through the first scattering peak (30). The value obtained, $S = 0.93 \pm 0.05$, is very large but is not uncommon for colloidal nematic suspensions which usually show a strongly first-order I/N transition (3). This large value also suggests that the scattering essentially arises from a nematic single domain.

The high-resolution SAXS patterns of the other mesophases are completely different from those of the nematic phase. They usually show several very sharp dotted diffraction rings (SI Appendix, Fig. S5). Uniform rings were very rarely observed, whatever the sample treatment, so that good-quality, powder-like samples of the bottom phase could not be obtained. Nevertheless, by carefully scanning capillaries with the small X-ray beam, the scattering pattern of a single domain of this mesophase could be acquired for several samples (Fig. 5B). This pattern displays two very sharp diffraction spots that can be indexed as the (001) and (002) reflections from a one-dimensional ordered stack. A radial scan of the scattered intensity going through the (001) reflections reveals sharp peaks whose width increases with $l$, as expected for a lamellar phase. The full-width at half-maximum (FWHM) of $5.3 \times 10^{-2} \text{nm}^{-1}$ of the (001) peak in the radial direction is close to the experimental resolution ($2.7 \times 10^{-2} \text{nm}^{-1}$) of the setup. A fit of the whole scattered intensity profile along the radial direction (SI Appendix, Fig. S6) was used to estimate the size of the lamellar domain along the director, $L_\perp \sim 25 \mu m$. The (001) lamellar reflection is thus quite different from the first broad scattering peak of the nematic phase which, with FWHM of $8.4 \times 10^{-2} \text{nm}^{-1}$, is ~16 times broader (Fig. 6).

Even though the radial profiles of scattered intensity in each phase differ so much, the distinction between the two types of organization depicted in Fig. 1 can only be made through the close inspection of the scattering profile of the reflections along the orthoradial direction. As mentioned above, for the nematic phase (Fig. 1A), this profile results from both the in-plane form factor of the nanosheets, leading to a widening of the reflection inversely proportional to the nanosheet diameter, and the orientational distribution, leading to the deformation of the reflection along a small circular arc. For the lamellar phase, neglecting the line broadening due to the Landau–Peierls instability of the lamellar phase (SI Appendix, Fig. S7) in a first approximation, the reflection width should be inversely proportional to the lamellar domain size, $L_\perp$, in the plane perpendicular to the director (Fig. 1B). In fact, the transverse scan of the scattered intensity going through the first reflection, shown in Fig. 5B, is barely wider than the experimental resolution function of the beamline (Fig. 6B). By simply applying Scherrer’s formula (31), taking the experimental resolution into account, $L_\perp$ can be roughly estimated to ~20 μm. This size is much larger than the nanosheet diameter, $<D> \sim 800 \mu m$ for this sample, and leads to approximately (20 μm/800 nm)~600 nanosheets self-assembled to build up a smectic layer.

Because no other scattering feature could be detected, the nanosheets have no long-range positional correlations within the layers. This rules out the possibility of a smectic phase with solid-like layers (like smectic B, smectic E, smectic G, etc.) and allows us to identify the bottom phase as a usual lamellar phase of the same symmetry as the SmA of thermotropic liquid crystals and the $L_\perp$ phase of surfactants.

For comparison, the SAXS pattern of a beidellite natural clay suspension, recorded in the same experimental conditions, is shown in Fig. 5C. This sample displays a clear nematic schlieren texture in polarized-light microscopy (16). Its SAXS pattern is indeed quite similar to that of the nematic phase of $\text{H}_3\text{SbP}_2\text{O}_{14}$ suspensions (Fig. S4), and it is completely different from that of the lamellar phase (Fig. 5B). Clay suspensions have not yet been reported to display any other mesophase than the nematic one because they undergo a sol/gel transition at concentrations only slightly larger than that of the isotropic/nematic transition, hence forbidding any higher order to take place at larger concentrations.

Detailed experimental studies of charged gibbsite, Ni(OH)$_2$, and layered double hydroxide nanoplates, of aspect ratio ~12–15, most often show that the nematic and columnar phases are more stable than the lamellar one (32–37). There is indeed only one report in literature of a lamellar phase in suspensions of gibbsite particles, but it is not based on the detailed analysis of the scattering from a single domain (38), and no lamellar/nematic phase coexistence was reported. However, the $\text{H}_3\text{SbP}_2\text{O}_{14}$ nanosheet suspensions described here have much larger aspect ratios (>100) and therefore form liquid-crystalline phases at much smaller concentrations than these systems. Moreover, the electrostatic repulsions between nanodisks and nanosheets may strongly differ due to the higher anisotropy of the latter. There are also several reports of the existence of a lamellar phase in systems of nanosheets (14, 18, 39–42), but they all rely on the analysis of the scattering from unaligned (i.e., powder-like) samples. Indeed, none of these studies show the detailed analysis of the transverse profiles of the X-ray reflections from single domains of the lamellar phase, which is the most stringent criterion to tell the difference between a columnar nematic and a lamellar phase (Fig. 1). Moreover, $\text{H}_3\text{SbP}_2\text{O}_{14}$ nanosheet suspensions are so far the only experimental system where both the columnar nematic and lamellar phases coexist in the same samples, which confirms that these phases have different structures. Nevertheless, the (isotropic–nematic–lamellar) phase sequence, upon increasing concentration, is quite robust in this system because it is observed for all four size fractions, A to D, of
H$_2$SB$_2$P$_2$O$_4$ nanosheets. The phase transition concentrations decrease with increasing particle size (SI Appendix, Text), as already reported in previous studies (40).

Theoretical studies of colloidal suspensions of plate-like particles rarely predict the existence and stability of the lamellar phase with respect to the nematic and columnar ones (43, 44). For example, it was shown that disk polydispersity may enhance the stability of the lamellar phase. Moreover, there have only been a few theoretical studies of the phase diagram of electrically charged plate-like particles reported so far because the electrostatic interaction potential of two charged disks in suspension has only recently been calculated (45). Based on this potential, numerical simulations were made to draw rich phase diagrams that present the nematic and columnar phases, among others, but not the lamellar one (10, 46).

In this context, the observation of a lamellar phase in the suspensions of H$_2$SB$_2$P$_2$O$_4$ nanosheets is somewhat surprising, and more theoretical work is needed to revise the phase diagrams of charged nanosheets. Other microscopic details could also play a role in the relative phase stabilities. For example, the chemical properties of atoms located at the nanoparticle rim are likely to differ from those of atoms on the particle faces, leading to differences in charge density that may affect the particle interactions, even at long range. Such particle heterogeneity that is reminiscent of patchy nanoparticles might affect the phase behavior of H$_2$SB$_2$P$_2$O$_4$ suspensions (47, 48). Also, cation-mediated attractions between like-charged nanosheet rims might also take place. More experimental investigations are therefore needed to better characterize the surface properties at the particle rim and faces.

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