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Quantum electric-dipole liquid on a triangular lattice

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Geometric frustration and quantum fluctuations may prohibit the formation of long-range ordering even at the lowest temperature, and therefore liquid-like ground states could be expected. A good example is the quantum spin liquid in frustrated magnets. Geometric frustration and quantum fluctuations can happen beyond magnetic systems. Here we propose that quantum electric-dipole liquids, analogues of quantum spin liquids, could emerge in frustrated dielectrics where antiferroelectrically coupled electric dipoles reside on a triangular lattice. The quantum paraelectric hexaferrite BaFe₁₂O₁₉ with geometric frustration represents a promising candidate for the proposed electric-dipole liquid. We present a series of experimental lines of evidence, including dielectric permittivity, heat capacity and thermal conductivity measured down to 66 mK, to reveal the existence of an unusual liquid-like quantum phase in BaFe₁₂O₁₉, characterized by itinerant low-energy excitations with a small gap. The possible quantum liquids of electric dipoles in frustrated dielectrics open up a fresh playground for fundamental physics.

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eometric frustration arises on various triangle-based lattices such as one-dimensional (1D) trestle lattice, two-dimensional (2D) triangular and kagome lattices and three-dimensional (3D) B-site spinel and pyrochlore lattices, and are typically investigated in spin systems¹⁻³. It has become well known that the introduction of quantum fluctuations in geometrically frustrated magnets gives rise to a rich variety of interesting quantum phases³⁻⁵, as discussed with both the transverse Ising models⁶⁻⁹ and Heisenberg models^{10,11}. Especially, exotic quantum spin liquids (QSLs), characterized by either gapped or gapless itinerant excitations^{3,4}, have been theoretically predicted to show extremely intriguing phenomena. Compared with the impressive progress and diversity in theory, nevertheless, a clear identification of QSLs in real materials has proved challenging, with a very limited number of candidates reported so far¹²⁻¹⁶.

Similar to the situation of spin lattices in magnets, geometric frustration can occur in lattices made of electric dipoles in dielectrics. In the case of small electric dipoles with significant quantum fluctuations persisting down to $T=0 \,\mathrm{K}$, exotic disordered quantum phases, such as a quantum electric-dipole liquid (QEL), could emerge in certain conditions. In fact, some theoretical models proposed for ultracold dipolar particles trapped on 2D frustrated optical lattices have predicted topological quantum phases with fractional excitations¹⁷. In a QEL, the electric dipoles are highly entangled with one another in a form of quantum dimers (pairs of antiparallel dipoles) and continue to fluctuate in the resonating valence bond state, a picture qualitatively similar to a QSL. However, we must emphasize that the QEL should have distinctive features from QSLs because electric dipole and spin have important differences¹⁸. For instance, electric dipole neither has intrinsic angular momentum nor exhibits quantum precession as magnetic dipole (spin) does. Moreover, the nature of short- and long-range interactions between electric dipoles is very different from that of spins¹⁹. This could lead to a very different phase diagram between QSLs and QELs.

Frustration in dielectrics has been previously studied in materials with competing ferroelectric (FE) and antiferroelectric (AFE) constituents such as the KH₂PO₄/NH₄H₂PO₄ (KDP-ADP) family or containing random-site impurities such as KTaO3:Li, which usually result in electric-dipole glasses similar to spin glasses^{20,21}. However, the geometric origin of frustration and cooperative liquid-like quantum phases has been largely ignored in the studies of dielectrics. On the other hand, the role of quantum fluctuations in dielectrics has been noticed since 1970s when people were studying the abnormal dielectric behaviour of SrTiO₃ (ref. 22). It was proposed that quantum fluctuations in SrTiO₃ prevent the onset of long-range FE order so that a quantum paraelectric state persists down to zero temperature²³. Since then, quantum paraelectricity has been reported in a number of perovskite oxides with similar structures to SrTiO₃, such as CaTiO₃, EuTiO₃ and KTaO₃. The quantum paraelectrics provide a new playground for the study of quantum critical phenomena¹⁸; however, it seems hopeless to search for the QELs in those perovskite quantum paraelectrics because their crystalline structures and FE interactions usually do not introduce geometric frustration.

In this work, we demonstrate that both geometric frustration and strong quantum fluctuations can be simultaneously achieved in a unique quantum paraelectric hexaferrite $BaFe_{12}O_{19}$ in which small electric dipoles that originated from the off-centre displacement of Fe^{3+} in the FeO_{5} bipyramids constitute a 2D triangular lattice. Our experiments including dielectric permittivity, heat capacity and thermal conductivity measured down to 66 mK suggest the existence of a very unusual liquid-like ground

state, characterized by itinerant low-energy excitations with a small gap. We consider this nontrivial quantum phase as a possible candidate of the QELs. The quantum liquids of electric dipoles in frustrated dielectrics provide a new playground for fundamental physics and may find applications in quantum information and computation as well.

Result

Geometric frustration in BaFe₁₂O₁₉. Recently, we have discovered that the M-type hexaferrites, such as BaFe₁₂O₁₉, belong to a completely new family of quantum paraelectrics²⁴. Other hexaferrites containing the FeO₅ bipyramids in their crystal structures, such as the W-, Z-, X- and U-type hexaferrites, are also likely candidates of quantum paraelectrics²⁴. The M-type hexaferrite BaFe₁₂O₁₉ is one of the most popular magnetic materials with a wide use in magnetic credit cards, bar codes, small motors and low-loss microwave devices²⁵ because of its superior properties of ferrimagnetic ordering with a strong ferromagnetic moment and a very high Néel temperature (\sim 720 K), high resistivity, as well as low cost of synthesis. The crystal structure of BaFe₁₂O₁₉ is shown in Fig. 1a. It can be described by a periodically stacking sequence of two basic building blocks—S block and R block along the c axis. The Fe³⁺ ions occupy three different kinds of sites: octahedral, tetrahedral and bipyramidal sites. In particular, the FeO₅ bipyramids only exist in the middle of the R/R^* blocks and form a triangular lattice in the ab plane (Fig. 1b).

Previous experiments including Mössbauer spectroscopy²⁶, X-ray diffraction²⁷ and neutron diffraction²⁸ have revealed the existence of off-equatorial displacements for Fe³⁺ at Wyckoff position of 2b site inside the FeO₅ bipyramids to minimize the total energy, which results in two adjacent Wyckoff positions of 4e sites with a lowered symmetry (Fig. 1c). The off-equatorial displacement (4e–4e distances are 0.176(5) Å at 4.2 K and 0.369(5) Å at room temperature)²⁶ would induce a small local electric dipole P along the c axis in each FeO₅ bipyramid (Fig. 1c). A dynamic displacement persists down to the lowest temperature because of the significant quantum tunnelling between two 4e sites and the weak dipole–dipole coupling along the c axis. Consequently, a quantum paraelectric behaviour without long-range electric ordering has been observed in BaFe₁₂O₁₉ (ref. 24).

More importantly, these electric dipoles associated with the FeO₅ bipyramids reside on a triangular lattice in each R/R^* block. Because the R/R^* blocks are well separated by the S/S^* blocks, this triangular lattice thus has a 2D feature. Consequently, a dielectric system with uniaxial (Ising-type) electric dipoles on a 2D triangular lattice is practically achieved in BaFe₁₂O₁₉ (Fig. 1c). the neighbouring dipole-dipole interaction favours anti-alignment, the system confronts frustration and has a very large degeneracy of ground states. In this sense, BaFe₁₂O₁₉ is a very unique quantum paraelectric system compared with those previously known perovskite quantum paraelectrics: first, it has uniaxial electric dipoles, whereas perovskite SrTiO₃ is pseudocubic with multiple easy axis; second, it experiences geometrical frustration, whereas there is no evidence of geometrical frustration in SrTiO₃ and other perovskite quantum paraelectrics. Thus, the quantum paraelectric BaFe₁₂O₁₉ sets up a promising candidate to search the proposed QELs, where an assembly of quantum dimers (pairs of dipoles) with long-range entanglement continues to fluctuate (Fig. 1d). We then employ a series of experimental techniques to resolve the ground state of BaFe₁₂O₁₉.

Dielectric permittivity. A prerequisite of a QEL is the AFE interaction between neighbouring dipoles. To confirm the AFE

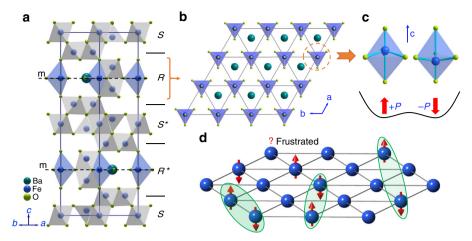


Figure 1 | Uniaxial electric dipoles on a triangular lattice in BaFe₁₂O₁₉. (a) Crystal structure of the M-type hexaferrite BaFe₁₂O₁₉. It consists of alternate stacks of *S* and *R* blocks along the *c* axis. The asterisk symbols indicate that the corresponding blocks rotate about the *c* axis by 180°. The Fe³⁺ ions occupy three different sites: octahedral, tetrahedral and bipyramidal (blue) sites. A mirror plane (m) bisects equally the bipyramids in the R/R^* blocks. (b) The 2D triangular lattice of FeO₅ bipyramids in each R/R^* block. (c) Illustration of Fe³⁺ off-equator displacements in the FeO₅ bypyramid. The upward or downward displacements at two 4e sites give rise to small electric dipoles along the *c* axis. Quantum fluctuations between two 4e sites persist to T = 0 K. (d) Frustrated electric dipoles on a triangular lattice. Each site contains an Ising-type electric dipole (red arrow), while the neighbouring interactions favour anti-alignment. Quantum dimers (marked by green ovals) with either short-range or long-range entanglement continue to fluctuate and result in a OEL.

coupling in BaFe₁₂O₁₉, we have made a careful analysis on the low-temperature dielectric permittivity. As shown in Fig. 2a, the dielectric permittivity along the c axis (ε_c) of BaFe₁₂O₁₉ increases steadily with decreasing temperature but remains nearly constant below ~ 5.5 K. No dielectric phase transition is observed down to 1.5 K. This dielectric behaviour evidences a quantum paraelectricity, similar to that in SrTiO₃. The quantum paraelectric behaviour can be well described by the mean-field Barrett formula²⁹:

$$\varepsilon = A + \frac{M}{\left(\frac{1}{2}T_1\right)\coth\left(\frac{T_1}{2T}\right) - T_0},\tag{1}$$

where A is a constant, T_0 is proportional to the effective dipole-dipole coupling constant and the positive and negative values correspond to FE and AFE interactions, respectively. T_1 represents the tunnelling integral and is a dividing temperature between the low-temperature region where quantum fluctuation is important and the high-temperature region where quantum effect is negligible. $M = n\mu^2/k_{\rm B}$, where n is the density of dipoles, μ denotes the local dipolar moment and $k_{\rm B}$ is the Boltzmann constant.

After fitting the ε_c below 160 K to the Barrett formula, we obtained $T_0 = -22.9(1)\,\mathrm{K}$ and $T_1 = 47.3(1)\,\mathrm{K}$. The negative T_0 confirms the AFE coupling between electric dipoles. We note that recent first-principle calculations 30 also predicted the AFE interaction with frustration in BaFe₁₂O₁₉. The relative strength of quantum fluctuations can be estimated by $\sim |T_1/T_0| = 2.06$, which is likely high enough to favour a liquid ground state rather than an ordered or glass phase. The uniaxial anisotropy is evidenced by comparing the dielectric permittivity along the c axis with that in the ab plane. As seen in the inset of Fig. 2a, the in-plane ε decreases slowly with decreasing temperature (less than 1 for a temperature interval of 250 K). The absence of a paraelectric behaviour in the ab plane is consistent with the uniaxial electric dipoles along the c axis.

Further lines of evidence of the AFE coupling in BaFe₁₂O₁₉ are presented in Fig. 2b. For those perovskite quantum paraelectrics with FE coupling, such as SrTiO₃, a moderated electric field is able to drive the quantum paraelectric state into a long-range

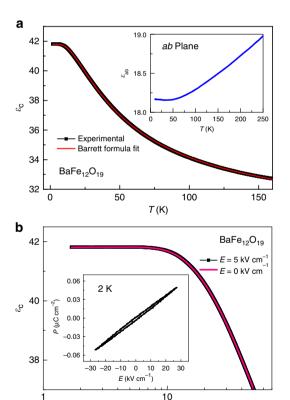


Figure 2 | **Dielectric permittivity of BaFe**₁₂**O**₁₉. (a) The temperature dependence of *c* axis dielectric permittivity ε_c . The red solid line is the fitting curve to the Barrett formula. The negative fitting parameter $T_0 = -22.9(1)$ K suggests the AFE interaction. The inset shows the dielectric permittivity measured along the [100] direction in the *ab* plane. (b) The temperature dependence of dielectric permittivity ε_c measured with d.c. bias electric fields. A bias electric field of 5 KV cm⁻¹ has no detectable influence on the quantum paraelectric behaviour. The inset shows the *P-E* loop at 2 K. The nearly linear response and the low values of *P* imply the AFE coupling.

T(K)

ordered FE state³¹. In strong contrast, for BaFe₁₂O₁₉, an external electric field of 5 KV cm⁻¹ applied along the c axis has no detectable influence on the dielectric permittivity. This inertness to external electric fields may indicate the AFE interaction in BaFe₁₂O₁₉. Moreover, the P-E loop at 2 K (the inset of Fig. 2b) shows a nearly linear response with quite small polarization up to a high electric field of 30 kV cm⁻¹, further implying the AFE coupling. It should be clarified that the magnetic moments of Fe³⁺ at the bipyramidal sites are all parallel along the c axis in the R/R* blocks (see Supplementary Fig. 1) so that there is no magnetic frustration on the triangular lattice.

Heat capacity and thermal conductivity. The thermodynamic studies at temperatures as low as possible are crucial to identify the conjectured quantum liquid state, as they provide the key information on the spectrum of low-energy elementary excitations. Heat capacity and thermal transport measurements can probe the low-energy density of states as well as determine whether these low-energy excitations are localized or itinerant, and have been indispensably employed in the study of QSLs³²⁻³⁴.

Since ${\rm BaFe_{12}O_{19}}$ is a good insulator (see Supplementary Fig. 2) with long-range collinear ferrimagnetic ordering ($T_{\rm N}=720\,{\rm K}$), both the electronic and magnon contributions to the thermal dynamics become negligible at very low temperatures³⁵. Therefore, its thermodynamics at low enough temperatures should be dominated by the lattice contribution only, and the well-known T^3 dependence would be expected for both the heat capacity and thermal conductivity. Figure 3a shows the heat capacity ($C_{\rm P}$) of ${\rm BaFe_{12}O_{19}}$ at low temperatures ($T<10\,{\rm K}$). No sharp anomaly due to a phase transition could be detected down to 0.4 K, in accordance with the quantum paraelectric behaviour. Unfortunately, the heat capacity data become scattered and noisy below $\sim 1\,{\rm K}$, possibly because of the very small values that reach the resolution limit of our equipment. Thus, a quantitative analysis of the heat capacity data is not possible.

The thermal conductivity provides more reliable and critical information on the low-lying elementary excitations because it is sensitive exclusively to itinerant excitations and is totally insensitive to localized entities that may cause the nuclear Schottky contribution and plague the heat capacity measurements at low temperatures 33,34 . For example, although the heat capacity study 32 suggested a gapless QSL in the frustrated triangular magnet $\kappa\text{-}(\text{BEDT-TTF})_2\text{Cu}_2(\text{CN})_3$, the thermal conductivity measurements 33 carried out down to 80 mK clarified instead a gapped QSL in the same material. We then have devoted a great effort to measure precisely the thermal conductivity of $\text{BaFe}_{12}\text{O}_{19}$ down to 66 mK.

Figure 3b shows the thermal conductivity κ measured in the *ab* plane as a function of temperature below $\sim 1 \text{ K}$. κ decreases rapidly with cooling, with a change more than 2 orders from 1 to 100 mK. No anomaly due to a phase transition is observed down to 66 mK. The thermal conductivity divided by temperature as a function of T^2 is plotted in Fig. 3c. The data between 0.65 and 1 K exactly follow a linear relation with an extrapolation to the origin, in a good agreement with what expected for the phonon thermal conductivity, $\kappa = \beta T^3$, with $\beta = 0.098 \, \text{WK}^{-4} \, \text{m}^{-1}$. Nevertheless, there is apparently an extra contribution below ~650 mK in addition to the normal phonon term, strongly suggesting the existence of abundant itinerant low-energy excitations other than phonons. Moreover, the thermal transport behaviour at the zero temperature limit provides the key information on the nature of these low-lying excitations. As seen in Fig. 3d, κT^{-1} in the $T \rightarrow 0$ K limit tends to vanish rather than having a finite residual value, immediately implying the absence of gapless excitations.

Instead, the data at the lowest temperature regime (T<125 mK) can be fitted to

$$\kappa = \alpha \exp(-\Delta/k_B T) + \beta T^3. \tag{2}$$

The inset of Fig. 3 shows an Arrhenius plot of $\kappa^* = \kappa - \beta T^3$ in the lowest temperature region. The good linearity confirms the validity of equation (2). The best fit gives $\Delta = 0.16(1)$ K, which is much smaller than the effective dipole–dipole interaction constant T_0 (~ 23 K). The exponential behaviour of thermal conductivity at the zero temperature limit is very similar to that observed in the frustrated triangular magnet κ -(BEDT-TTF)₂Cu₂(CN)₃, where a QSL with gapped excitations ($\Delta = 0.46$ K) was identified³². Therefore, the thermal transport behaviour excludes a frozen dipole glass or a classical paraelectric phase but strongly suggests an exotic liquid-like ground state.

To exclude the possibility that these itinerant low-lying excitations may have a magnetic origin, we further studied the influence of magnetic field on the thermal conductivity behaviour. As seen in Fig. 3d, a high magnetic field of $14\,\mathrm{T}$ applied along the easy c axis has no influence on the in-plane thermal conductivity in the lowest temperature range. The inertness of these low-lying excitations to external magnetic fields supports our argument that they stem from electric dipoles rather than spins.

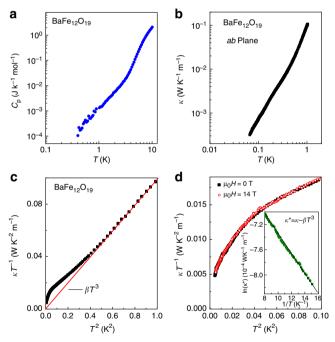


Figure 3 | **Heat capacity and thermal conductivity of BaFe**₁₂O₁₉. (a) Heat capacity C_p as a function of temperature. No phase transition is detected down to 0.4 K. (b) Thermal conductivity κ measured in the ab plane as a function of temperature. No anomaly due to a phase transition is observed down to 66 mK. (c) The ab plane thermal conductivity divided by temperature plotted as a function of T^2 below \sim 1K. The red solid line represents the expected thermal conductivity of phonons, $\kappa = \beta T^3$, with $\beta = 0.098$ WK $^{-4}$ m $^{-1}$. Apparently, there is an extra contribution beside the phonon thermal conductivity below \sim 650 mK. (d) The κT^{-1} versus T^2 plot in the lowest temperature region. κ/T tends to vanish at the $T \rightarrow 0$ K limit. An applied magnetic field of 14 T along the c axis has no influence on the ab plane thermal conductivity in this low-temperature region. The inset shows the Arrhenius plot of $\kappa^* = \kappa - \beta T^3$ below \sim 125 mK. The good linearity suggests gapped excitations with a small gap \sim 0.16 K.

Discussion

We think that the above experimental observations, especially the thermal conductivity at the zero temperature limit, strongly indicate the existence of a nontrivial ground state of electric dipoles in BaFe₁₂O₁₉ that is a very unique dielectric system because both geometrical frustration and strong quantum fluctuations play an important role, a situation similar to frustrated antiferromagnets where people are looking for QSLs. This unusual ground state is characterized by itinerant lowenergy excitations with a tiny gap ($\Delta = 0.16 \,\mathrm{K}$) that is 140 times smaller than the effective dipole-dipole interaction constant T_0 (~23 K), a feature against a dipole glass or a classical dipole liquid. Thus, we consider it as a possible candidate of an exotic QEL. As current theoretical models proposed for frustrated spin systems, such as the transverse Ising models on a triangular lattice, are inadequate for the frustrated electric dipoles because the long-range dipole-dipole interactions are quite different from the short-range spin exchange interactions, a quantitative comparison between experiments and theories is not available at this stage. We expect that our experimental findings will stimulate theoretical efforts towards this interesting subject in the future. The present work serves as a start point to draw attention on frustrated quantum electric dipoles in dielectric materials where an abundance of exotic phenomena could be awaiting ahead.

Methods

Sample preparation. The single-crystal samples of $BaFe_{12}O_{19}$ were prepared with the flux method and were characterized with X-ray diffraction, as shown in Supplementary Fig. 3. Powders of $BaCO_3$, Fe_2O_3 and fluxing agent Na_2CO_3 were weighed to a molar ratio of 1:1:1, and then were mixed and well ground. The ground raw powder was put in a Pt crucible and heated to 1,250 °C for 24 h in the air, and then cooled down to 1,100 °C at a rate of 3 °C min $^{-1}$ and finally quenched to room temperature.

Dielectric measurements. The dielectric measurements were carried out in a Cryogen-free Superconducting Magnet System (Oxford Instruments, TeslatronPT) down to 1.5 K. To measure the dielectric permittivity, silver paste was painted on the surfaces of a thin plate of crystal and annealed at 150 °C for \sim 30 min to make good electrodes. An Agilent 4980A LCR meter was used to measure the dielectric permittivity with the frequency $f\!=\!1\,\mathrm{MHz}.$

Heat capacity and thermal transport measurements. Heat capacity measurements were performed down to $0.4\,\mathrm{K}$ in a commercial Physical Properties Measurement System (Quantum Design) using a $^3\mathrm{He}$ refrigerator. A thin-plate-shaped sample with mass of 12.7 mg was used for this measurement. The contribution of attendant was measured separately and subtracted from the raw data. Thermal conductivity was measured between 60 mK and 1 K using the conventional steady-state 'one heater, two thermometers' technique in a $^3\mathrm{He}^{-4}\mathrm{He}$ dilution refrigerator 36,37 . A parallelepiped-shaped sample with size of $2.0\times0.57\times0.11\,\mathrm{mm}^3$ was cut from the as-grown crystals for thermal conductivity measurements. A chip heater and two RuO2 chip sensors are attached to the sample with gold wires. The temperature difference between the two thermometers was controlled to be typically 3% of the sample temperature. To minimize heat leak, superconducting NbTi wires with 15 μ m diameter are used as the leads of the chip sensors.

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Author contributions

Y.S. conceived the study and designed experiments with X.F.S. S.-P.S. prepared the sample and carried out dielectric and magnetic measurements. J.-C.W. and J.-D.S. performed heat capacity and thermal conductivity measurements. Y.-F.Y., Y.-S.C., D.-S.S., S.-G.W. and J.F.S. contributed to the data analysis and discussions. Y.S. and Y.-F.Y. wrote the paper, and all authors reviewed the paper.

Additional information

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