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# Molecular ferroelectric pyridin-2-ylmethanaminium perchlorate with phase transition induced by disorder of perchlorate†

Xuecheng Gan, Zheng Tang, Wenjun Zhang, Cong Xu, Xiaofan Sun, Zhangran Gao, Yizhang Wu, Hong-Ling Cai \* and X. S. Wu\*

Molecular ferroelectrics are a kind of functional materials that have large application prospects. Comparing with the traditional inorganic ferroelectrics, they have some advantages, such as easy design, environment-friendliness, nontoxicity, and flexibility. In this study, a new molecular ferroelectric pyridin-2-ylmethanaminium perchlorate was discovered, which undergoes a ferroelectric phase transition at around 264.8 K. The spontaneous polarization reaches  $0.22 \mu\text{C cm}^{-2}$ , and the coercive fields are as small as  $1.1 \text{ kV cm}^{-1}$  at 202 K. The single-crystal X-ray diffraction analysis reveals that the space group transforms from a polar space group of  $P2_1$  at 173 K to a centrosymmetric space group of  $P2_1/c$  at 293 K. By analyzing the crystal structure changes below and above the phase transition temperature, it can be concluded that the phase transition is induced by the disorder of perchlorate. Its ferroelectricity was confirmed by the measurements of differential scanning calorimetry, dielectric and hysteresis loop.

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## Introduction

Ferroelectrics are a kind of polar material that has a spontaneous polarization, and the spontaneous polarization can be reversed by an external electric field.<sup>1</sup> Since its discovery in 1921, ferroelectrics have been widely applied in electric random access memories (FeRAM), infrared detectors, pyroelectric vidicons, remote sensing, nonlinear optical devices, capacitors, and laser frequency converters.<sup>2–15</sup> Molecular ferroelectrics, as one class of ferroelectric materials, compared with the traditional inorganic ferroelectric materials, have some advantages, such as easy design, nontoxicity, and flexibility.<sup>16–20</sup> The study of ferroelectricity began in 1920 with the research on the special dielectric properties of Rochelle salt by Valasek,<sup>21</sup> which was nearly 100 years ago. Thereafter, a lot of ferroelectrics have been discovered, such as  $\text{BaTiO}_3$  (BTO),<sup>21,22</sup>  $\text{KH}_2\text{PO}_4$  (KDP),<sup>23</sup> and  $\text{LiNbO}_3$ . The discoveries of these excellent inorganic ferroelectrics have greatly promoted the research process of ferroelectrics and the practical applications of ferroelectric materials. However, the development of organic ferroelectric is lagging behind. With the enhancement in people's environmental awareness, the advantages of organic ferroelectrics with respect to environmental protection have been highlighted, which has made it one of the research hotspots in recent years.

Molecular ferroelectric croconic acid, which has a spontaneous polarization at about  $20 \mu\text{C cm}^{-2}$  was discovered by Horiuchi.<sup>24</sup> The molecular ferroelectric diisopropylammonium perchlorate with spontaneous polarization ( $\sim 23 \mu\text{C cm}^{-2}$ ), which is close to BTO ( $\sim 26 \mu\text{C cm}^{-2}$ ), has been reported.<sup>25,26</sup> In addition, there are many molecular ferroelectrics that have superior ferroelectric properties, such as diisopropylammonium bromide,<sup>27,28</sup> 4-(cyanomethyl)anilinium perchlorate,<sup>29</sup> and 4-methoxyanilinium tetrafluoroborate 18-crown-6.<sup>30</sup> Recently, a number of pyridyl-based molecular ferroelectrics have been discovered, such as  $\text{PyX}$  ( $\text{Py} = \text{pyridinium}$  and  $\text{X} = \text{ClO}_4$ ,<sup>31</sup>  $\text{BF}_4$ ,<sup>32</sup>  $\text{ReO}_4$ ,<sup>33</sup>  $\text{IO}_4$  (ref. 34)), (4-amino-2-bromopyridinium)(4-amino-2-bromopyridine) tetrafluoroborate,<sup>18</sup> and pyridin-4-ylmethanaminium perchlorate.<sup>35</sup>

Herein, we discovered a new molecular ferroelectric pyridin-2-ylmethanaminium perchlorate (Fig. 1) with the chemical formula of  $\text{H}^+\text{yl-pyH}^+(\text{ClO}_4^-)_2$ , [ $\text{yl} = 2\text{-ylmethanamine}$ ,  $\text{py} = \text{pyridin}$ ] (compound 1) based on pyridin-2-ylmethanamine and perchlorate acid, which undergoes a ferroelectric phase transition at around  $T_c = 264.8 \text{ K}$ . The spontaneous polarization reaches  $0.22 \mu\text{C cm}^{-2}$ , and the coercive field is  $1.1 \text{ kV cm}^{-1}$  at 202 K. The single-crystal X-ray diffraction reveals that the space group at 173 K is a polar space group  $P2_1$ , and it is a centrosymmetric space group  $P2_1/c$  at 293 K. In addition, the ferroelectricity of compound 1 was confirmed by other measurements, which includes differential scanning calorimetry, dielectric and P–E hysteresis loop.

## Experimental Methods

All reagents and solvents were of reagent grade and used without further purification. Concentrated perchloric acid

Collaborative Innovation Center of Advanced Microstructures, Laboratory of Solid State Microstructures, School of Physics, Nanjing University, Nanjing 210093, P. R. China. E-mail: hlcai@nju.edu.cn; xswu@nju.edu.cn

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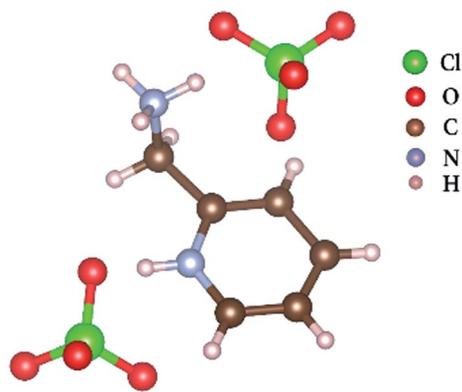


Fig. 1 Molecular structure of compound 1.

(71 wt%, 5.66 g, 0.04 mol) was added dropwise to pyridin-2-ylmethanamine (99 wt%, 2.18 g, 0.02 mol) in methanol (50 mL). Continuous stirring and resettling was done to obtain a colorless solution. A yellow block of compound was obtained after 2 weeks through the slow evaporation of aqueous solution at room temperature. Differential scanning calorimetry (DSC) and heat capacity measurements of the single crystals were performed on NETZSCH DSC 200 F3 under nitrogen protection in aluminum crucibles with a heating or cooling rate of 10 K min<sup>-1</sup>. The variable-temperature single-crystal X-ray diffraction analysis was carried out using a Rigaku Saturn 724<sup>+</sup> CCD diffractometer equipped with Mo K $\alpha$  radiation ( $\lambda = 0.71073 \text{ \AA}$ ). Data collection, cell refinement, and data reduction were performed using Rigaku Crystalclear 1.3.5. The structures of crystals were solved by direct methods and refined by the full matrix method based on  $F^2$ , using the SHELXLTL software package. All non-hydrogen atoms were refined anisotropically, and the positions of all hydrogen atoms were generated geometrically. Raman spectra (Horiba Jobin Yvon HR800) were performed using the 632.8 nm laser line of a He-Ne laser and were recorded in the back scattering geometry with a charge-coupled device detector. A 50 $\times$  microscope objective was used to focus the incident laser beam to a spot of about 3 mm in diameter. The sample was placed in a variable temperature cryostat, which can control the temperature from 293 K to 213 K. For dielectric, P-E hysteresis loops measurement, the single crystals were cut into thin plates. Silver conductive paste deposited on the plate surfaces was used as the electrodes. The complex permittivity measurement was performed using Tonghui TH2828A LCR meter at the frequency from 20 Hz to 1 MHz with an applied electric field of 0.5 V. The pyroelectric current was measured using Keithley 6517B electrometer. The P-E hysteresis loops were recorded on a Precision Premier II (Radiant Technologies, Inc.).

## Results and discussion

DSC is a commonly used thermal analysis method, which can be used to observe the latent heat of ferroelectric phase transition. The temperature-dependent DSC curve (Fig. 2) shows an endothermic peak around  $T_c = 264.8 \text{ K}$  in the heating process and an exothermic peak around  $T_c' = 246.2 \text{ K}$  in the cooling

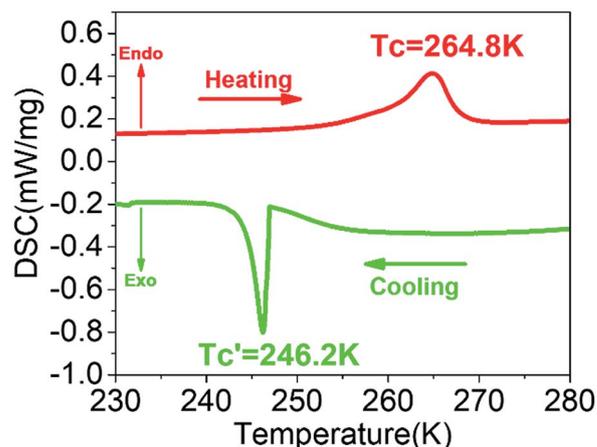


Fig. 2 Temperature dependence of DSC curve of the sample at a cooling/heating rate of 10 K min<sup>-1</sup>.

process, which reveals a reversible phase transition. Here, the exothermic peak is negative, and the endothermic peak is positive. The DSC curve shows a thermal hysteresis of about 18.6 K, which indicates a relaxation phase transition and a large thermal potential barrier. As a matter of convenience, the phases above/below  $T_c$  are labeled as high-temperature phase (HTP)/low-temperature phase (LTP), respectively. The entropy changes of phase transitions can be calculated as  $\Delta S = \Delta H/T$ , where the value of enthalpy change  $\Delta H$  can be integral from the  $T_c'$  or  $T_c$  peak area. The entropy change ( $\Delta S$ ) at around  $T_c$  is 8.28 J mol<sup>-1</sup> K<sup>-1</sup>. According to the Boltzmann equation ( $\Delta S = R \ln N$ , where  $R$  is the universal gas constant and  $N$  represents the ratio of possible configurations), it is found that  $N = 2.71$ , manifesting that the phase transition at  $T_c$  is of an order-disorder type.

Ferroelectric crystals always undergo structural transitions such as ferroelectric-paraelectric or ferroelectric-ferroelectric phase transitions. These structural phase transitions are induced by pressure, temperature, electric field, and shock wave. The large anomalies in dielectric behaviors will be engendered, which is an essential characteristic of the ferroelectrics. The dielectric behavior can be expressed by measuring the temperature-dependent permittivity. Fig. 3a shows the dielectric behavior of compound 1 measured along  $b$ -axis in the heating process with frequency from 500 Hz to 1 MHz, demonstrating that the dielectric behaviour is frequency dependent. The dielectric constant ( $\epsilon'$ ) has a peak at the temperature of  $T_c = 265.8 \text{ K}$  in the heating process. For the ferroelectric phase transition, the dielectric response should follow the Curie-Weiss law,  $\epsilon' = \epsilon_0 + C_{\text{para}}/(T - T_c)$  ( $T > T_c$ ) or  $\epsilon' = \epsilon_0 + C_{\text{ferro}}/(T_c - T)$  ( $T < T_c$ ), where  $C_{\text{para}}$  and  $C_{\text{ferro}}$  are the Curie-Weiss constant and the  $\epsilon_0$  is the optical frequency limit of the dielectric constant. Fig. 3b shows the temperature-dependent  $1/\epsilon'$  values at a frequency of 500 Hz in the heating process. The fitted Curie-Weiss constant  $C_{\text{para}}$  in HTP and the  $C_{\text{ferro}}$  in LTP are 68 and 250 K, respectively (other frequencies are listed in Table S2<sup>†</sup>). These results fit well with Curie-Weiss law.

The variable temperature single-crystal X-ray diffraction (Fig. 4) can well reveal the origin of the ferroelectric properties of



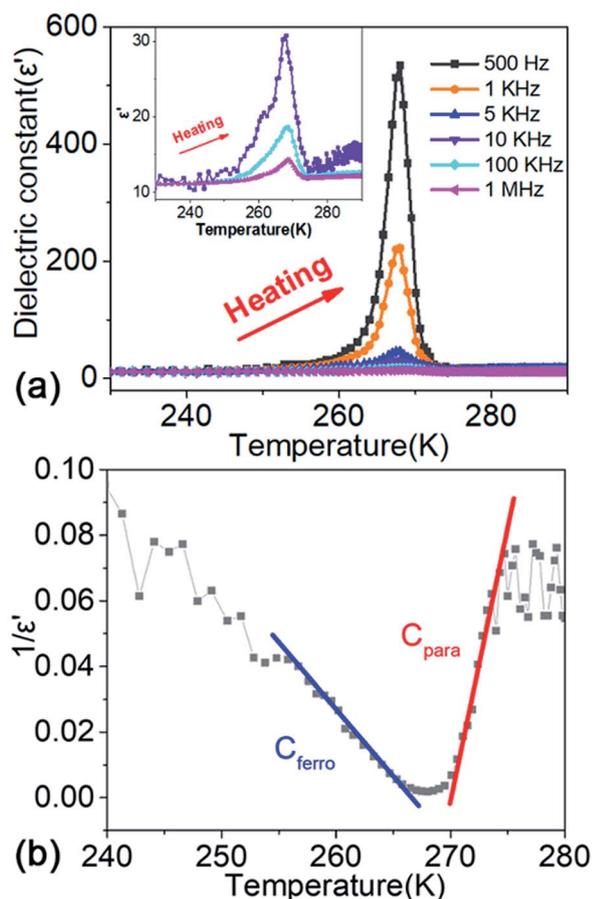


Fig. 3 (a) Temperature-dependence of real part ( $\epsilon'$ ) of complex dielectric constant of the compound in heating process. (b) The reciprocal real part of dielectric constant  $1/\epsilon'$  at 500 Hz as a function of temperature in heating process.

compound 1. Compound 1 crystallizes in the monoclinic crystal system with a centrosymmetric space group  $P2_1/c$  (point group:  $C_{2h}$ , symmetric operation elements:  $E, C_2, \sigma_h, i$ ) at 293 K (CCDC number: 1857985). Both nitrogen atoms on pyridin-2-

ylmethanaminium are protonated. Since the space group in the HTP is centrosymmetric, the dipole moment from  $\text{ClO}_4^-$  anions to  $-\text{NH}_3^+$  cations in the cell are equivalent and antiparallel, there is no net polarization. It is noteworthy that the oxygen atoms in  $\text{ClO}_4^-$  anions have multiple equilibrium positions in the HTP. A quarter of the  $\text{ClO}_4^-$  anions in each cell have disordered oxygen atoms. Compound 1 transforms to a monoclinic crystal system with a polar space group  $P2_1$  (point group:  $C_2$ ; symmetric operation elements:  $E, C_2$ ) at 173 K (CCDC number: 1857987). The two hydrogen atoms on the  $-\text{NH}_3$  attract two  $\text{ClO}_4^-$  anions to form hydrogen bonds, respectively. It is known from the direction of each dipole that the direction of polarization is along the  $b$ -axis. There is no disorder of  $\text{ClO}_4^-$  anions in each cell, which indicates that the disorder is reduced from HTP to LTP. The vibration at  $628\text{ cm}^{-1}$  comes from the inplane deformation of the perchlorate anion.<sup>16,36</sup> By varying temperature, Raman peak position shifts before and after the phase transition at  $628\text{ cm}^{-1}$  (Fig. S2†), it can also be seen that the perchloric acid disorder changes before and after the phase transition. This further indicates that the ferroelectric phase transition at  $T_c$  belongs to the order–disorder type ferroelectric. It can be concluded from the crystal structure analysis that the HTP has more slip symmetry than the LTP. In addition, the reduction of symmetric operation elements indicates that the essence of ferroelectric phase transition in compound 1 is symmetry breaking.

The spontaneous polarization of ferroelectrics can be reversed with the reversal of the external electric field, exhibited by the typical polarization–electric field (P–E) hysteresis loops below the  $T_c$ . Fig. 5 shows the P–E hysteresis loops of compound 1 at different temperatures with a frequency of 50 Hz. The P–E hysteresis loops indicate that the spontaneous polarization increases slightly with the increase in the temperature and the coercive field decreases gradually. The spontaneous polarization ( $P_s$ ) is  $0.22\text{ }\mu\text{C cm}^{-2}$ , and the coercive field ( $E_c$ ) is  $1.1\text{ kV cm}^{-1}$  as obtained from the ferroelectric hysteresis loop at 202 K. The value of the spontaneous polarization is slightly larger than that of the Rochelle salt ( $0.2\text{ }\mu\text{C cm}^{-2}$ ), which is less

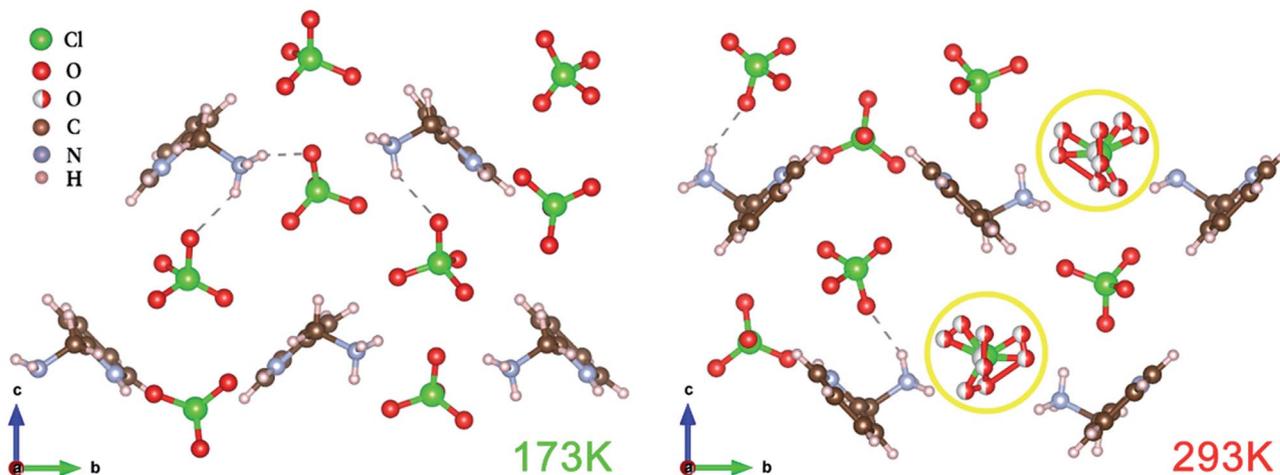


Fig. 4 Crystal structure diagrams of the compound at 173 K and 293 K.



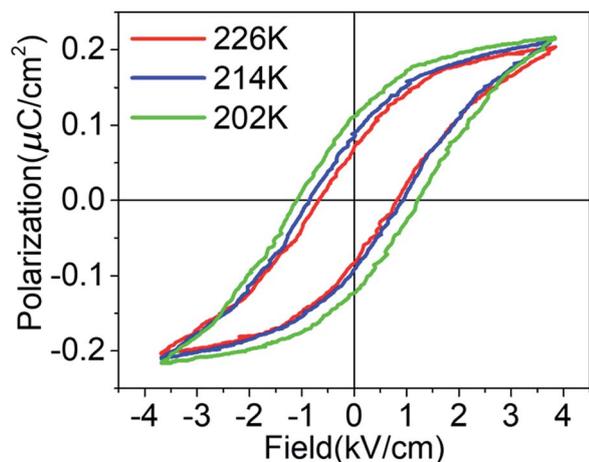


Fig. 5 P–E hysteresis loops of compound 1 at different temperatures with a frequency of 50 Hz.

than the recently reported pyridin-4-ylmethanaminium perchlorate ( $1.25 \mu\text{C cm}^{-2}$ ). The coercive field is much smaller than most organic ferroelectrics, which is much less than that of the pyridin-4-ylmethanaminium perchlorate ( $2.6 \text{ kV cm}^{-1}$ ). The hysteresis loops further demonstrate the ferroelectricity of pyridin-2-ylmethanaminium perchlorate below the Curie temperature.

## Conclusions

In summary, we have discovered a new molecular ferroelectric pyridin-2-ylmethanaminium perchlorate, which is ferroelectric below  $T_c = 264.8 \text{ K}$  and undergoes a ferroelectric phase transition at  $T_c$ . The spontaneous polarization of compound 1 is  $0.22 \mu\text{C cm}^{-2}$ , and the coercive field is  $1.1 \text{ kV cm}^{-1}$  at a temperature of  $202 \text{ K}$ . X-ray diffraction analysis reveals that the space group from LTP to HTP changes from  $P2_1$  to  $P2_1/c$ . The X-ray diffraction analysis also shows that it belongs to the order–disorder type ferroelectrics.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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## References

- 1 S. Horiuchi and Y. Tokura, *Nat. Mater.*, 2008, 7, 357–366.
- 2 M. D. Aggarwal, *Ferroelectrics*, 2008, 366, 84–121.

- 3 M. E. Lines, A. M. Glass and G. Burns, *Principles and Applications of Ferroelectrics and Related Materials*, Clarendon Press, 2001.
- 4 S. Mathews, R. Ramesh, T. Venkatesan and J. Benedetto, *Science*, 1997, 276, 238–240.
- 5 S. L. Miller and P. J. McWhorter, *J. Appl. Phys.*, 1992, 72, 5999–6010.
- 6 Q. Ye, Y. Song, G. Wang, K. Chen, D. Fu, P. Wai, H. Chan, J. Zhu, S. D. Huang and R. Xiong, *J. Am. Chem. Soc.*, 2006, 128, 6554–6555.
- 7 J. F. Scott, *Science*, 2007, 315, 954–959.
- 8 G. A. Smolenskii, V. A. Bokov, V. A. Isupov, N. N. Krainik, R. E. Pasynkov and A. I. Sokolov, *Ferroelectrics and Related Materials*, Gordon and Breach Science Publishers, 1987.
- 9 D. L. Staebler, *Ferroelectric crystals*, Pergamon, 1962.
- 10 L. Chen, Q. Ji, Q. J. Pan, X. X. Cao and G. C. Xu, *CrystEngComm*, 2017, 19, 5907–5914.
- 11 I. Collings, M. Bykov, E. Bykova, M. Hanfland, S. V. Smaalen, L. Dubrovinsky and N. Dubrovinskaia, *CrystEngComm*, 2018, 20(25), 3512–3521.
- 12 S. Q. Lu, X. G. Chen, J. X. Gao, Y. Lu, X. N. Hua and W. Q. Liao, *CrystEngComm*, 2017, 20(4), 454–459.
- 13 Y. Lu, Z. Wang, H. P. Chen and J. Z. Ge, *CrystEngComm*, 2017, 19(14), 1896–1901.
- 14 R. Mu, G. C. Xu, Y. Y. Zhang, L. Zhang and D. Z. Jia, *CrystEngComm*, 2017, 19(31), 4629–4633.
- 15 Y. Y. Zhang, J. T. Yu, B. Li, D. J. Li, Z. G. Gu, X. F. Sun, H. L. Cai, G. E. Kostakis and G. Peng, *CrystEngComm*, 2018, 20, 4582–4589.
- 16 K. Gao, Z. Cui, C. Liu, J. Zhu, H. L. Cai and X. Wu, *RSC Adv.*, 2015, 5, 62647–62651.
- 17 K. Gao, M. Gu, X. Qiu, X. N. Ying, H. Y. Ye, Y. Zhang, J. Sun, X. Meng, F. M. Zhang and D. Wu, *J. Mater. Chem. C*, 2014, 2, 9957–9963.
- 18 C. Liu, K. Gao, Z. Cui, L. Gao, D. W. Fu, H. L. Cai and X. Wu, *J. Phys. Chem. Lett.*, 2016, 7(10), 1756–1762.
- 19 C. Xu, W. Zhang, L. Gao, X. Gan, X. Sun, Z. Cui, H. L. Cai and X. S. Wu, *RSC Adv.*, 2017, 7, 47933–47937.
- 20 W. Zhang and R. G. Xiong, *Chem. Rev.*, 2012, 112, 1163–1195.
- 21 J. Valasek, *Phys. Rev.*, 1921, 17, 475–481.
- 22 L. E. Cross and R. E. Newnham, *J. Am. Ceram. Soc.*, 1986, 11.
- 23 G. Busch and P. Scherrer, *Naturwissenschaften*, 1935, 23, 737.
- 24 S. Horiuchi, Y. Tokunaga, G. Giovannetti, S. Picozzi, H. Itoh, R. Shimano, R. Kumai and Y. Tokura, *Nature*, 2010, 463, 789.
- 25 D. W. Fu, W. Zhang, H. L. Cai, J. Z. Ge, Y. Zhang and R. G. Xiong, *Adv. Mater.*, 2011, 23, 5658–5662.
- 26 D. W. Fu, H. L. Cai, Y. Liu, Q. Ye, W. Zhang, Y. Zhang, X. Y. Chen, G. Giovannetti, M. Capone and J. Li, *Science*, 2013, 339, 425–428.
- 27 K. Gao, C. Liu, Z. Cui, J. Zhu, H. L. Cai and X. S. Wu, *CrystEngComm*, 2015, 17, 2429–2432.
- 28 K. Gao, C. Xu, Z. Cui, C. Liu, L. Gao, C. Li, D. Wu, H. L. Cai and X. S. Wu, *Phys. Chem. Chem. Phys.*, 2016, 18, 7626–7631.
- 29 H. L. Cai, W. Zhang, J. Z. Ge, Y. Zhang, K. Awaga, T. Nakamura and R. G. Xiong, *Phys. Rev. Lett.*, 2011, 107, 147601.



- 30 D. W. Fu, H. L. Cai, S. H. Li, Q. Ye, L. Zhou, W. Zhang, Y. Zhang, F. Deng and R. G. Xiong, *Phys. Rev. Lett.*, 2013, **110**, 257601.
- 31 P. Czarnecki, W. Nawrocik, Z. Pajak and J. Wasicki, *J. Phys.: Condens. Matter*, 1994, **6**, 4955.
- 32 P. Czarnecki, W. Nawrocik, Z. Pajak and J. Wasicki, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **49**, 1511–1512.
- 33 J. Wąsicki, P. Czarnecki, Z. Pajak, W. Nawrocik and W. Szczepański, *J. Chem. Phys.*, 1997, **107**, 576–579.
- 34 Z. Pajak, P. Czarnecki, J. Wąsicki and W. Nawrocik, *J. Chem. Phys.*, 1998, **109**, 6420–6423.
- 35 Z. Cui, K. Gao, C. Liu, Y. Yin, D. W. Fu, H. L. Cai and X. S. Wu, *J. Phys. Chem. C*, 2016, 120.
- 36 B. A. Zakharov, B. A. Kolesov and E. V. Boldyreva, *Phys. Chem. Chem. Phys.*, 2011, **13**, 13106–13116.

