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The role of anionic design in modulating material properties: from SHG switching to ferroelastic switching†

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Crystallographic engineering and chemical synthesis modification play crucial roles in modulating material properties and designing multifunctional materials, such as those with second-harmonic generation (SHG) switching and ferroelastic behaviors. Herein, we present the role of anion-based design in crystal-line materials through anion H/F substitution. Specifically, the material $[\text{NMe}_4][\text{CH}_3\text{SO}_3]$ exhibits a high phase transition temperature ($T_c = 462$ K) and excellent SHG switching performance. The introduction of polyfluorinated substitutions results in a new organic ferroelastic material, $[\text{NMe}_4][\text{CF}_3\text{SO}_3]$, which exhibits ferroelastic phase transition at 309 K. The transition from the non-centrosymmetric $[\text{NMe}_4][\text{CH}_3\text{SO}_3]$ to the centrosymmetric $[\text{NMe}_4][\text{CF}_3\text{SO}_3]$ leads to a shift in material properties, transitioning from SHG switching to ferroelastic switching. Our findings underscore the importance of anion-based design in the development of advanced multifunctional phase-transition materials.

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Introduction

Phase transition materials undergo alterations in their physical properties with temperature variations, exhibiting multiple functional characteristics, such as switchable second-order nonlinear optical (NLO), dielectric, ferroelastic, ferroelectric, *etc.*^{1–10} Especially in second-harmonic generation (SHG) switching and ferroelastic switching materials, phase transitions play a crucial role. SHG switching materials driven by symmetry-breaking structural phase transitions have garnered considerable attention due to their unique combination of tunable optical properties and robust switching performance.^{11–14} The mechanism relies on reversible transitions between centrosymmetric and non-centrosymmetric states, where symmetry changes during phase transitions induce sharp modulation of SHG responses, enabling reproducible ‘on-off’ switching.^{15,16} Notably, organic molecular phase transition materials exhibit superior chemical tunability and dynamically adjustable nonlinear optical properties during phase transitions, resulting in excellent SHG switching performance.^{17–20} During phase transitions, the material’s crystal structure undergoes significant changes, leading to

symmetry breaking that can induce ferroelastic behavior.^{21,22} However, designing ferroelastic phase transitions necessitates adherence to one of the 94 ferroelastic phase transition types outlined in Aizu’s rule.²³

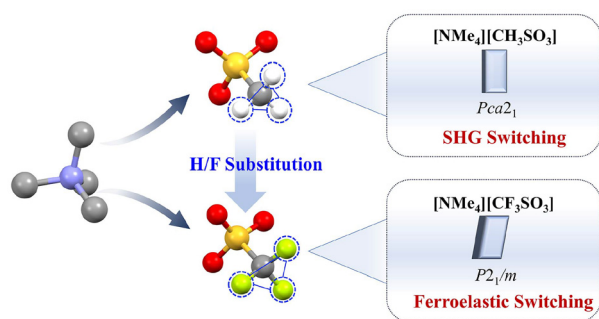
The low phase transition temperatures have significantly hindered the application and development of phase transition materials. Crystallographic engineering, particularly through fluorine substitution, plays a pivotal role in enhancing phase transition temperatures and modulating material properties.²⁴ The pronounced electronegativity difference between hydrogen and fluorine atoms induces substantial variations in molecular polarizability within H/F-substituted systems, potentially leading to significant alterations in macroscopic physical properties.²⁵ For instance, monofluorination leads to significant T_c enhancement in those hybrid molecular materials, such as (fluoropyrrolidinium) MnCl_3 ,²⁶ $[(\text{CH}_3)_3\text{PCH}_2\text{F}][\text{Cd}(\text{SCN})_3]$,²⁷ *N,N*-dimethyl-3-fluoropyrrolidinium CdCl_3 ,²⁸ $(4\text{FHQ})_2\text{RbEu}(\text{NO}_3)_6$ ²⁹ and so on.^{30–33}

Furthermore, multiple fluorine substitutions on the cations demonstrate remarkable efficacy in modifying the transition temperature, polarity, photochromism, and crystal structures, as evidenced by those hybrid compounds, such as $[4,4\text{-DFPD}]_2\text{PbI}_4$,³⁴ $[4,4\text{-DFHHA}]_2\text{PbI}_4$,³⁵ $[3,3\text{-Difluorocyclobutylammonium}]_2\text{CuCl}_4$,³⁶ $[\text{DFCBA}]_2\text{CrCl}_4$.³⁷ Besides, multiple fluorine substitutions can also induce the weaker intermolecular interactions within amantadine-pentafluorobenzoic acid and lead to a ferroelastic phase transition.³⁸ In addition to modulating material properties through H/F substitution in the cationic part of mole-

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Scheme 1 The design strategy for SHG switching CH-NMe₄ and ferroelastic switching CF-NMe₄.

cular crystals, H/F substitution in the anionic component can also be used. Fu *et al.* reported the flexible organic ferroelectric material PEA-TFMS using an anionic fluorination strategy, showing the feasibility of inducing plastic ferroelectricity.³⁹ Sugata Ray achieved multiferroicity in fluorinated inorganic oxides, Ca₂Mn₂O_{5-x}F_y, by engineering anionic networks through fluorination and vacancy ordering.²⁴

In this work, we explore the effects of H/F substitution on the anionic components to modulate the phase transition and crystalline symmetry. Herein, we choose quasi-spherical tetramethylammonium molecules as cation, which is prone to order-disorder phase transitions,^{40–42} successfully constructed two multifunctional materials, namely [NMe₄][CH₃SO₃] (CH-NMe₄) and [NMe₄][CF₃SO₃] (CF-NMe₄). As shown in Scheme 1, the organic compound CH-NMe₄ exhibits high *T_c* of 462 K and SHG switching behaviours. CH-NMe₄ crystallizes in the non-centrosymmetric *Pca2*₁ space group at 293 K, demonstrating significant nonlinear optical switching with a large on-off ratio and high stability. In contrast, the H/F substituted compound CF-NMe₄ crystallizes in the centrosymmetric *P2*₁/*m* space group at 260 K. Reversible ferroelastic domain patterns confirm its ferroelastic phase transition. This work highlights the synergy between crystallographic engineering and H/F substitution in tailoring material functionalities, offering valuable insights for advancing nonlinear optical materials and ferroelastics.

Results and discussion

Materials

Crystals of CH-NMe₄ and CF-NMe₄ are grown in ethanol solution, slowly evaporated by the solvent at a constant temperature of 30 °C, and colourless, clear crystals for single crystal measurement were obtained. Thermogravimetric analysis (TGA) confirms exceptional thermal stability, with decomposition onsets occurring at 634 K for CH-NMe₄ and 705 K for CF-NMe₄ (Fig. S1, ESI†), indicating their robust thermal stability.

Single crystal structures

At 293 K, the CH-NMe₄ crystallized in the orthorhombic *Pca2*₁ space group with cell parameter of *a* = 11.994(2) Å, *b* = 6.1359(13) Å, *c* = 11.923(3) Å, $\alpha = \beta = \gamma = 90^\circ$. The non-centro-

symmetric crystalline structure of CH-NMe₄ demonstrated intrinsic piezoelectric behavior. Quasi-static measurements revealed a *d*₃₃ coefficient of 11 pC N⁻¹, confirming its piezoelectric responsiveness (Fig. S2, ESI†). The CF-NMe₄ crystallized in the monoclinic *P2*₁/*m* space group at 260 K. The parameters are *a* = 10.3722(3) Å, *b* = 8.6225(2) Å, *c* = 11.6086(3) Å, $\alpha = \gamma = 90^\circ$, $\beta = 103.824^\circ$ for CF-NMe₄. The detailed cell parameters of CH-NMe₄ and CF-NMe₄ are listed in Table S1, ESI†. Subsequently, we performed powder X-ray diffraction (PXRD) to verify the phase purity. The experimental patterns show excellent agreement with simulated profiles (Fig. S3, ESI†), with all observed peaks aligning precisely with simulated reflections, confirming the phase purity of both CH-NMe₄ and CF-NMe₄. As depicted in Fig. 1a, the asymmetric unit of the CH-NMe₄ consists of one NMe₄⁺ cation and one CH₃SO₃⁻ anion. The CH₃SO₃⁻ parameters for CH-NMe₄ are comparable to those found in the ordered phases of other known compounds with methanesulfonic acid,⁴³ with S–O bond lengths ranging from 1.446 to 1.458 Å. The CH₃SO₃⁻ anion is involved in four non-classical hydrogen bonds C–H...O. The strongest hydrogen bonds in the structure exist between the O atom on the methanesulfonic acid and the N atom on the tetramethylamine, which are at the closest distance: H...A = 2.40 Å, D...A = 3.349 Å, D–H–A = 170° (Fig. 1b, Fig. S4a and Table S2, ESI†). Ions of opposite charges are alternately arranged along *c*-axis, forming a two-dimensional structure perpendicular to the *b*-axis under the weak interaction force. For CF-NMe₄, the asymmetric unit of the crystal structure consists of two NMe₄⁺ cations and two CH₃SO₃⁻ anions, as depicted in Fig. 1c. There exists C–H...O hydrogen-bonding interactions in the structure, forming infinite linear hydrogen-bonded chains along the *b*-axis, as shown in Fig. 1d and Fig. S4b, ESI†. The non-classical hydrogen bonds of the CH-NMe₄ and CF-NMe₄ crystals are summarized in Tables S2 and S3, ESI†. At 260 K, CF-NMe₄ exhibits six types of C–H...O hydrogen bonds, with bond lengths ranging from 3.373 Å to 3.442 Å, and corresponding bond angles ranging from 148° to 153°. Unlike most studies in which the introduction of monofluoro substitution brings

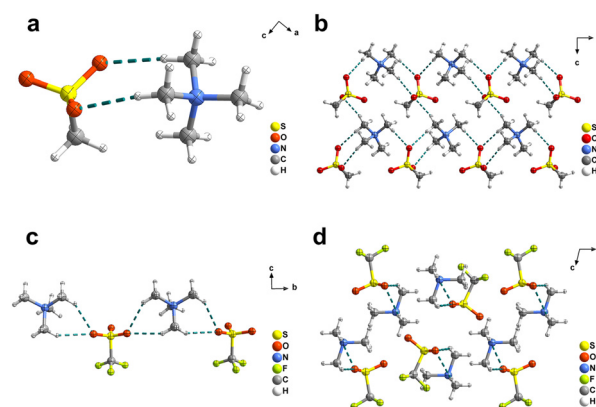


Fig. 1 The asymmetric unit of (a) CH-NMe₄ at 293 K and (c) CF-NMe₄ at 260 K. Packing structure of (b) CH-NMe₄ at 293 K and (d) CF-NMe₄ at 260 K. The dash lines denote the C–H...O hydrogen bond.

about polarity,^{26,28,33} the multiple fluorine substitution on the anions induces a notable reduction in crystal symmetry, leading to the formation of a centrosymmetric structure in CF-NMe₄.

Hirshfeld surfaces analysis

The 2D fingerprint plots enable detailed analysis and highlight the interaction forces between specific atoms, allowing for separation based on the varying contribution levels of different interactions depicted in the Fig. 2. To further explore the influence of anion fluorination on molecular dynamics, Hirshfeld surfaces and 2D fingerprint plots (Fig. 2) were generated using CrystalExplorer software, leveraging their CIF files for detailed analysis. In Fig. 2a, six distinct red spots on the Hirshfeld surface of CH-NMe₄ signify the presence of C-H...O hydrogen bonds. A comprehensive analysis of the full 2D fingerprint plot in Fig. S5, ESI† further elucidates the relative contributions of various hydrogen bond types. Detailed analysis of the fingerprint plot reveals that H...H interactions dominate, contributing 65.2% of the total Hirshfeld surface area, while H...O interactions account for 34.7%. Additionally, the H...S

Interaction represents only 0.1% of the Hirshfeld surface. In Fig. 2b, the Hirshfeld surface of CF-NMe₄ exhibits five prominent red spots, signifying the presence of C-H...O hydrogen bond interactions. Analyzing the fingerprint plot reveals that O...H interactions occupy 49.0% of the entire Hirshfeld surface area, while F...H interactions account for 42.4%. Further examination of the complete 2D fingerprint plot in Fig. S6, ESI† provides additional details about the contributions of various hydrogen bond types. Specifically, Fig. S6e, ESI† represents the F...F interaction, contributing 6.9% to the Hirshfeld surface, while Fig. S6f and S6g, ESI† depict the F...O/S...H interactions, accounting for 1.3% and 0.3%, respectively. Anionic fluorination markedly influences the spacing between H atoms on cations and O atoms on anions, thereby reshaping structural packing by introducing steric effects. Simultaneously, it reorganizes hydrogen-bonding network formation pathways through enhanced electrostatic

interactions, highlighting the profound impact of fluorination on molecular arrangements.

Phase transitions

Differential Scanning Calorimetry (DSC) serves as a robust analytical tool for precisely identifying phase transitions triggered by external thermal stimuli. Then, DSC measurements were conducted to explore the phase transition behaviors of CH-NMe₄ and CF-NMe₄. During the heating and cooling cycle of CH-NMe₄, which ranged from 425 K to 490 K, a pair of endothermic and exothermic peaks were observed at 462 K and 453 K, respectively, as depicted in Fig. 3a. This observation indicates a reversible phase transition. Notably, the significant temperature hysteresis effect during the heating and cooling processes is characteristic of first-order phase transitions.⁴⁴ It is noteworthy that CH-NMe₄ exhibits an exceptionally high phase transition temperature (462 K), which significantly exceeds that most of organic materials,^{45–54} underscoring its unique thermal stability. For CF-NMe₄, within the temperature range of 250–330 K, during the heating/cooling cycle, the DSC curve exhibits a pair of distinct thermal anomaly peaks at 309 K/283 K (Fig. 3b). The corresponding entropy changes (ΔS) for the endothermic peaks are approximately 2.98 and 17.9 J mol⁻¹ K⁻¹ for CH-NMe₄ and CF-NMe₄, respectively. Upon fluorination of the methanesulfonic acid anions, the phase transition temperature changes from 462 K to 309 K.

Phase transitions are typically accompanied by dielectric anomalies. Temperature-dependent dielectric measurements were performed at 1 MHz during a heating-cooling thermal cycle. For CH-NMe₄, a step-like dielectric anomaly is observed near 460 K (Fig. 3c), marked by a discontinuous increase in the real permittivity (ϵ') from 10.8 (low dielectric state) to 22.7 (high dielectric state), representing a 2.1-fold enhancement. Similarly, CF-NMe₄ demonstrates distinct dielectric behavior: during heating, ϵ' gradually increases from 4.5 at 307 K to 7.8

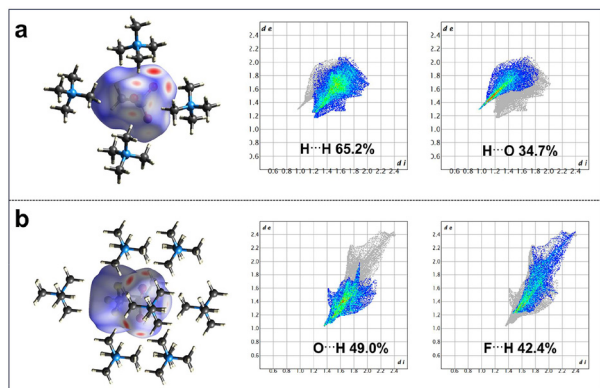


Fig. 2 Visualization map of the distribution of the interactions and the two-dimensional fingerprint plots of (a) CH-NMe₄, and (b) CF-NMe₄, show contributions from H_{outside}–O_{inside}.

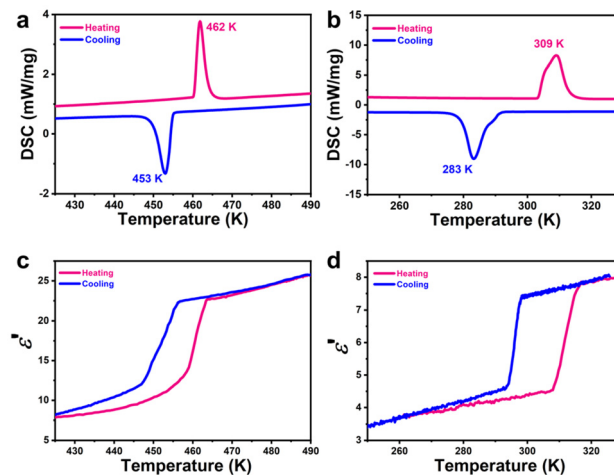


Fig. 3 The DSC curves for (a) CH-NMe₄ and (b) CF-NMe₄. The temperature-dependent real part (ϵ') of the dielectric constant was measured at 1 MHz for (c) CH-NMe₄ and (d) CF-NMe₄.

at 317 K, followed by stabilization ($\Delta\epsilon' < 0.5$) at elevated temperatures (Fig. 3d). The value of ϵ' in the high dielectric state is about 1.73 times that in the low dielectric state. The permittivity-temperature profiles were also measured at 1, 5, 10, and 100 kHz (Fig. S7, ESI†). As shown in Fig. S7,† the dielectric constant decreased with increasing frequency. Furthermore, the T_c deduced from permittivity-temperature curves is frequency-independent. Both compounds display thermal hysteresis (>5 K) between heating/cooling cycles, consistent with first-order transition characteristics observed in DSC.

To elucidate the structural phase transition, we attempted to measure the high-temperature crystal structures of CH-NMe₄ and CF-NMe₄. However, weak diffraction and severe ionic disorder at elevated temperatures hindered the successful collection of high-temperature phase data. Variable-temperature PXRD measurements were conducted for further analysis. A marked change in peak positions and intensities was observed upon heating to the phase transition point (Fig. S8, ESI†), confirming the occurrence of phase transition behavior leading to symmetry breaking. Critical evidence emerges from the progressive simplification of diffraction patterns, where specific reflections vanish above the transition temperature alongside reduced peak multiplicity. Rietveld refinement of high-temperature PXRD data reveals distinct structural transformations: CH-NMe₄ transitions from an orthorhombic *mm2* system to a tetragonal system, while CF-NMe₄ undergoes monoclinic *2/m* to tetragonal symmetry elevation (Fig. S9 and S10, ESI†). Although CH-NMe₄ exhibits a polar crystal structure that satisfies the symmetry requirements for ferroelectricity, the piezoresponse force microscopy analysis showed no evidence of domain switching or polarization hysteresis under various experimental conditions.

SHG switching

The temperature-dependent second-harmonic generation (SHG) serves as a sensitive probe for identifying structural transformations between centrosymmetric and non-centrosymmetric states.^{16,55} To further investigate the symmetry changes associated with the phase transition, we conducted SHG measurements on CH-NMe₄. At room temperature, CH-NMe₄ demonstrates a moderate SHG intensity, measuring 0.8 times that of KDP (Fig. 4a), which aligns with its non-centrosymmetric packing in the *Pca2*₁ space group. Temperature-dependent SHG measurements demonstrate a distinct bistable response, as illustrated in Fig. 4b. During heating, strong SHG signals persist below 460 K (high-NLO state) but abruptly vanish above 480 K as the material transitions to a centrosymmetric phase (low-NLO state). Conversely, upon cooling, the SHG response of CH-NMe₄ remains zero until 477 K before sharply recovering at 466 K with a 5.6 V signal, demonstrating a 9 K thermal hysteresis. The complete suppression of SHG intensity in the high-temperature phase confirms symmetry elevation to a centrosymmetric space group. Remarkably, CH-NMe₄ retains over 95% of its initial SHG intensity across 10 thermal cycles (Fig. 4c), achieving an exceptional switching contrast ratio ($I_{on}/I_{off} \approx 13$) that outper-

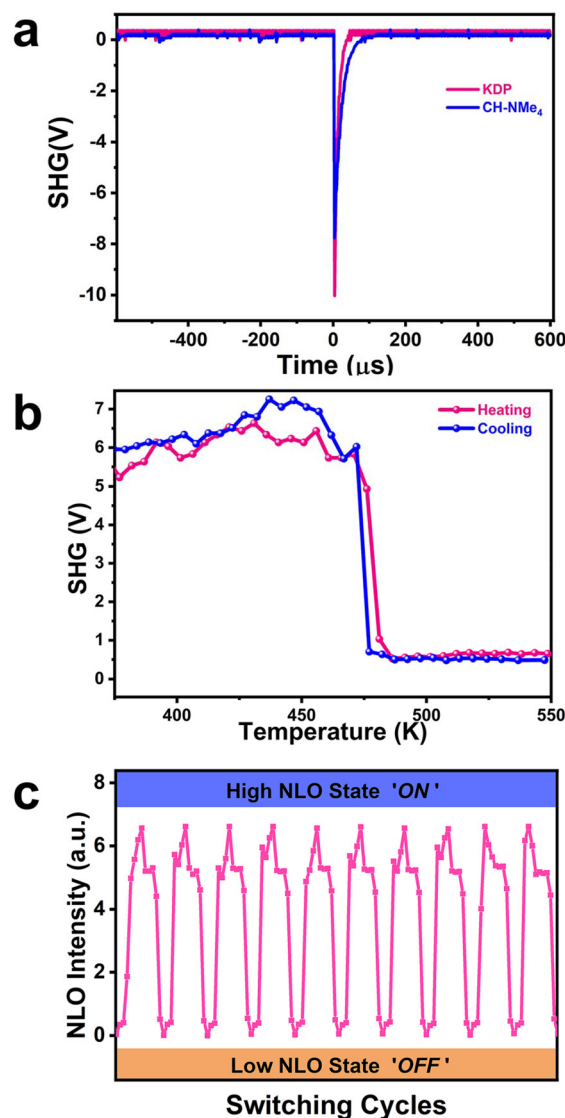


Fig. 4 (a) SHG signals of CH-NMe₄ and KDP at room temperature. (b) The temperature dependencies of SHG signals under heating and cooling run of CH-NMe₄. (c) Cycles of switching high and low NLO states of SHG signals of CH-NMe₄.

forms many existing SHG-switching materials.^{55–58} This unique combination of robust bistability, high switching contrast, and exceptional durability establishes CH-NMe₄ as a highly promising candidate for nonlinear optical switching devices that demand precise state control and long-term operational stability.⁵⁹ In contrast, the fluorine-substituted analogue, CF-NMe₄, crystallizes in a centrosymmetric space group and thus shows no SHG signal.

Ferroelastic switching

The evolution of ferroelastic domains is a sign of ferroelastic phase transitions, which we investigated through temperature-dependent polarized light microscopy. For CF-NMe₄, the appearance of stripe-like patterns in polarized microscopy

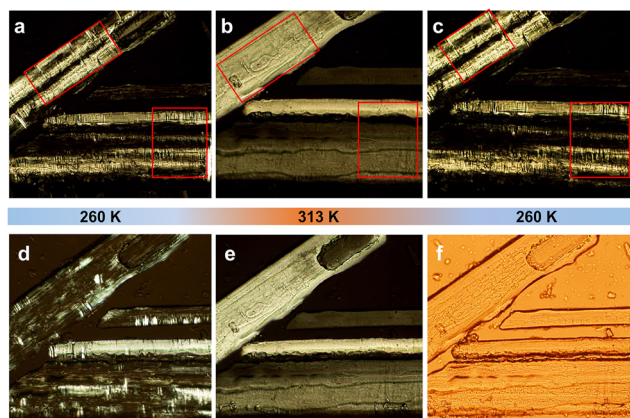


Fig. 5 The ferroelastic domain walls evolution of CF-NMe₄. (a–e) evolutions of the ferroelastic domain structure for CF-NMe₄ in the cooling-heating runs, natural optical microscope images for (f) morphology of CF-NMe₄. Ferroelastic domain evolution is indicated by the red rectangular frame.

images (at 260 K, Fig. 5a) serves as a definitive indicator of ferroelastic domain formation, which is distinct from material morphology (Fig. 5f). These domains exhibit alternating light-dark contrast, originating from the coexistence of multiple spontaneous strain orientations during the paraelastic-to-ferroelastic transition. Upon gradual heating, the ferroelastic domain structure progressively diminishes and fully vanishes near 313 K (Fig. 5b and e), signaling the transition to a paraelastic phase. The domain patterns reversibly reappear with restored stripe-like domain walls upon cooling back to 260 K (Fig. 5c and d), demonstrating robust thermal cyclability and highlighting the material's structural reversibility. DSC and temperature-dependent PXRD analyses confirm a reversible phase transition near 309 K. The temperature-dependent domain switching dynamics, coupled with the hysteretic recovery of domain patterns, provide unequivocal evidence of ferroelasticity in CF-NMe₄. However, no significant ferroelastic domain evolution was observed in CH-NMe₄, leading us to conclude that it lacks ferroelasticity. This structural transformation profoundly influences the material's properties, highlighting the critical role of chemical modifications in tailoring functional behaviors.

Conclusions

In conclusion, we reported an organic nonlinear optical switching material CH-NMe₄, and a molecular ferroelastic CF-NMe₄. Structural analyses reveal that the polyfluorinated substitution on the anion markedly alters intermolecular interactions and packing arrangements, inducing a transformation from a non-centrosymmetric (space group *Pca2*₁) to a centrosymmetric (space group *P2*₁/*m*) crystal structure at room temperature. Through anion modification, both symmetry breaking and lattice flexibility are simultaneously regulated, resulting in the disappearance of polarity and triggering ferroelasticity.

This work realizes the functional switching of phase transition materials through anion fluorination, demonstrating an effective chemical synthesis strategy and providing ideas for the development of multifunctional materials.

Conflicts of interest

There are no conflicts to declare.

Data availability

The data supporting this article have been included as part of the ESI.† Crystallographic data has been deposited at the Cambridge Crystallographic Data Centre (CCDC) as a supplementary publication no. 2449758 and 2449759.

Acknowledgements

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