Fluorescence-based thermal sensing with elastic organic crystals

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Operation of temperature sensors over extended temperature ranges, and particularly in extreme conditions, poses challenges with both the mechanical integrity of the sensing material and the operational range of the sensor. With an emissive bendable organic crystalline material, here we propose that organic crystals can be used as mechanically robust and compliant fluorescence-based thermal sensors with wide range of temperature coverage and complete retention of mechanical elasticity. The exemplary material described remains elastically bendable and shows highly linear correlation with the emission wavelength and intensity between 77 K to 277 K, while it also transduces its own fluorescence in active waveguiding mode. This universal new approach expands the materials available for optical thermal sensing to a vast number of organic crystals as a new class of engineering materials and opens opportunities for the design of lightweight, organic fluorescence-based thermal sensors that can operate under extreme temperature conditions such as are the ones that will be encountered in future space exploration missions.

Exploration of extreme environments, such as for example the possibility to sustain life at very low temperatures on Earth or in outer space requires robust cryogenic engineering materials that will be able to respond to a number of requirements1–3. Cooling has an inevitable, and sometimes drastic effect on the properties of both biogenic and artificial materials, as is well known with liquefaction or solidification of natural gas4, deceleration or cessation of cell metabolism5,6, and low-temperature superconductivity7,8. The oil and gas, machining, chemical industry, food, biomedicine, and aerospace sectors are in increasing demand for cryogenically stable and robust temperature sensing materials9–11. Low-temperature environments also imply the need for accurate measurement of low temperatures, which currently relies on either thermally controlled resistance9 or integrated circuit (IC) temperature sensing10. However, devices based on resistive measurement may fail due to the opening of the electrical circuit by mechanical separation of the components or as a result of materials’ aging. While the IC sensors are comparatively more robust, they typically cover only a narrow temperature range for measurement. While being effective for specific applications, a common drawback of both of these sensing technologies is that their physical components are made of stiff and brittle materials, and are therefore prone to mechanical damage in devices that are exposed to vibrations or shock over prolonged periods of time. A long sought-after alternative to these materials are fundamentally new soft, light, mechanically compliant, and robust temperature-sensing materials that can be used for accurate and reliable thermal measurement.

Organic fluorescent crystals based on π-conjugated small organic molecules with photoluminescent properties have been reported14. In a crystalline condensed state, both the intermolecular interactions and the molecular arrangement are known to affect the luminescence15–17.
Within that context, molecular organic crystals from π-conjugated molecules have been studied extensively as promising emissive materials due to the effects of molecular aggregation on the emission\(^5\). However, organic crystals have not been traditionally considered the materials of choice for temperature measurement. The recent realization of their chemical versatility, anisotropy in structure and properties, and long-range structural order has led to increased recognition of these compounds as a new materials class for organic optoelectronic components, such as resonators\(^19,20\), circuits\(^21\) and lasers\(^22,23\). A particularly important aspect of their property profile that has been highlighted recently is their pronounced elasticity\(^24,25\). This newly realized facet is central to their mechanical compliance and has been recently explored for electronics applications that require lightweight materials, such as optical waveguides for passive transduction of information in both the visible and near-infrared spectral regions\(^26,27\). Opportunities for application of the fluorescence of some flexible organic crystals have also been implicated\(^28,29\) for active signal transmission\(^30\). In line with the increasing demand for resilient cryogenic materials, we sought to develop lightweight, flexible materials that can be used for sensing or optical information transduction at low temperatures. The material that we report here combines all of these assets. Specifically, crystals of the organic crystalline material that we describe display reversible, reproducible, and strong temperature-induced shifts of their fluorescence, which translates into an opportunity for reproducible optical temperature measurement based on fluorescence. Within the broader set of available luminescence thermometry techniques\(^30\), this work brings a new class of materials that could be considered as an alternative to the currently used fluorescent thermometers.

**Results and discussion**

**Preparation and characterization**

The material described here, \(\text{C}_{1} \text{H}_{1} \text{O}_{2} \text{F}_{3} \text{N}_{2} \text{S}_{2}\), was discovered during our broader screening of materials in an effort to identify organic crystals that are both elastic and emissive. It was synthesized in two steps with a 72% yield (Supplementary Figs. 1 and 2). Long, yellow needle-like crystals of about 1.5 cm in length were readily prepared by liquid-phase diffusion (Fig. 1b). The material has a good thermal stability and does not undergo a phase transition up to its melting point at 467 K (Supplementary Fig. 3). The crystals of compound 1 are reversibly elastic; they can be bent without any visible damage (Supplementary Movie 1), as it was confirmed by the scanning electron microscopic images of bent specimens that were fixed in their bent state for inspection (Fig. 1c). No obvious defects could be observed on the crystal surface even after bending the crystals up to 5000 times (Supplementary Fig. 4). The crystals absorb light with a maximum at 415 nm, and emit bright green fluorescence with a maximum emission at about 540 nm and a quantum yield of 0.16 at 298 K (Fig. 1d). The fluorescence quantum yield is 0.30 at 77 K, which is higher compared to room temperature due to the decreased non-radiative rate at low temperature (Supplementary Table 1). Their bulk elastic modulus at 298 K was found to be 1.6 GPa by tensile measurements (Fig. 1e)\(^2\). The elasticity of the crystals is preserved even at low temperatures, and they can be bent repeatedly even when they are immersed in liquid nitrogen (Fig. 1f–i).

![Fig. 1](https://example.com/fig1.png)

**Temperature-induced change of fluorescence**

During the mechanical characterization at low temperature, we noticed that the color of emission of crystals of compound 1 changes visibly from green to orange when they are transferred from room temperature to liquid nitrogen (Fig. 1j–o; Supplementary Movie 2). The temperature-dependent fluorescence spectra show that the color change is due to a red-shift of the emission maximum upon cooling, whereupon the emission intensity gradually increases (Fig. 2a; Supplementary Fig. 5). The maximum emission wavelength changes from 540 to 580 nm upon cooling from 277 to 77 K, and the dependence is linear in the temperature range from 77 to 277 K \(\left(\frac{\lambda_{\text{max}}}{\text{nm}} = -0.21 \frac{T}{K} + 596.91, R^2 = 0.9948\right)\) (Fig. 2b). More quantitatively, when plotted on a CIE 1931 color space diagram\(^3\) this change in color of the emitted light is reflected in the change of the CIE coordinates from (0.38, 0.59) to (0.51, 0.48) (Fig. 2c). At 77 K, the emission intensity increases to 1.79 times compared to that at 277 K. The change of emission intensity with temperature is also linear within a certain temperature range \(\left(\frac{l_{0}}{l_{\text{max}}} = -0.0022 \frac{T}{K} + 1.14, R^2 = 0.9711\right)\) (Fig. 2d). This linear and strong response of the fluorescence of compound 1 with temperature favors this material as a temperature-sensing medium. The lowest detectable temperature of compound 1 is lower than that of some metal sensors\(^4,5\), however, the sensitivity (maximum emission wavelength vs. temperature) is higher than other organic materials with temperature-dependent fluorescence\(^1,6\). In order to verify whether the red-shift of compound 1 is caused by triplet luminescence, the lifetimes at 298 and 77 K were determined (Supplementary Fig. 6). The lifetime of the emission of compound 1 at 77 K is 17.99 ns, which is longer than that of 298 K (6.13 ns), confirming that the emission does not originate from a triplet state. The lifetime and quantum yield of bent crystals were also recorded and indicate that the bending does not have any significant effect on non-radiative pathways (Supplementary Fig. 6, Supplementary Table 2).
Optical-waveguiding capability

The remarkable flexibility of the compound 1 crystals at both ambient and low temperatures appears as a prospective platform for their application in thermally resistant (opto)electronics. Similar to other flexible crystals, single crystals were tested as active optical waveguides (Fig. 2d). The optical loss coefficient of a straight crystal at room temperature was found to be 0.16 dB mm⁻¹ (Supplementary Fig. 7). A bent crystal had a nearly identical optical loss, 0.17 dB mm⁻¹ (Supplementary Fig. 8). This result indicates that the crystal bending does not have a significant effect on the optical waveguiding performance, as is expected from the retention of the crystal integrity upon bending, described above. The cooling did not affect the waveguiding ability of the crystal (Fig. 2f); at 77 K, the optical loss factors were found to be 0.17 dB mm⁻¹ for a straight crystal and 0.20 dB mm⁻¹ for a bent crystal (Supplementary Figs. 9 and 10, Supplementary Movie 3).

Temperature sensing using fluorescence

We hypothesized that the combination of linear dependence of the emission wavelength on the temperature and the favorable mechanical flexibility of compound 1 at low temperature could carry some potential for the development of flexible temperature sensors. Along this line of thought, a crystal of about 1.0 cm in length was selected, and liquid nitrogen was dropped continuously on a small area at one end (Fig. 3a). The cold spot was excited with a laser, and the optical output was collected and analyzed at both ends of the crystal (Supplementary Fig. II). Consistent with the temperature difference, the cold end of the crystal emitted orange light, while the opposite end that was at higher temperature emitted green light. The temperature of the warm end was determined to be about 230 K (Supplementary Figs. 12 and 13). However, the emission wavelength maximum of the spectra recorded at both ends was 580 nm, showing that only the orange light was transduced (Fig. 3b). We note that the optical waveguide is in active mode here. The photoluminescence (PL) spectra of crystals of millimeter size at 77 and 298 K were also recorded.

Fig. 2 | Temperature dependence of the emission of compound 1. a Variable-temperature emission spectra of a crystal of compound 1. b Correlation between crystal emission wavelength and temperature in the temperature range 77–277 K. The straight broken line shows a linear fit to the data. c The CIE coordinates of crystal emission bands at different temperatures. The region in the boxed area is shown as a zoomed-in image in the inset. d Correlation between the crystal fluorescence intensity ratio and temperature in the temperature range 77–277 K. The straight broken line shows a linear fit to the data. e, f Emission spectra were measured at one end of the crystal at 298 K (e) and 77 K (f). The values 0–5 mm, corresponding to curves with different symbols, represent the distance from the excitation site to the point of measurement. The excitation was performed at 355 nm.

Fig. 3 | Temperature-dependent optical waveguiding. a Schematic diagram and real picture of a crystal waveguide at low temperature. The gray dashed box denotes the area where the liquid nitrogen was dropped (77 K). b Emission spectra recorded at the excitation end (77 K) and at the output end of the crystal (230 K). c Schematic diagram and real picture of a crystal waveguide at room temperature. d Emission spectra recorded at the excitation end (230 K) and at the output end of the crystal (77 K). e Emission spectra of the crystal optical waveguide at different temperatures. f CIE 1931 color space coordinates of crystal optical waveguide signals at different temperatures. The region in the boxed area is shown as zoomed-in image in the inset. g Correlation between the emission intensity ratio and temperature in the temperature range 77–277 K. The straight broken line shows a linear fit to the data. h Correlation between the crystal waveguide emission wavelength and temperature in the temperature range 77–277 K. The straight broken line shows a linear fit to the data.
Comparing the PL spectrum at 77 K with the optical waveguide signal at 77 K, we conclude that the peak shapes are almost identical (Supplementary Fig. 14d). This result indicates that the optical signal propagating in the waveguide is not affected during its propagation. Other two crystals that have been reported earlier (compounds 2 and 3) were also investigated to confirm this property of organic crystals (Supplementary Fig. 15). The position of laser excitation was then changed, and the crystal was excited at the higher temperature end, while it was being cooled at the opposite end (Fig. 3c). Contrary to the previous case, the warm end emitted green light and the cold end emitted orange light. The spectra of the fluorescent remained identical at both ends, with a maximum emission at 550 nm (Fig. 3d). These results demonstrate that the crystal can be indeed cooled locally, and it does not equilibrate thermally during the experiment. More importantly, when the crystal is used as a medium for optical transduction, the output signal depends only on the temperature at the point of excitation; the output is not affected even when the intermediate section of the crystal is at lower temperature.

In another experiment, a 1.5 cm-long crystal was selected, and one of its ends was brought into contact with evaporating liquid nitrogen. This cool end was excited with a laser at 355 nm, and the spectrum of the output light was recorded at the other end (Supplementary Fig. 16). As the liquid nitrogen evaporated, the excited end of the crystal gradually warmed up, and the output spectrum changed (Fig. 3e). As expected from the other experiments above, upon warming the position of the maximum emission was blueshifted while its intensity decreased, as it is illustrated by changes in the respective CIE coordinates (Fig. 3f). This is in accordance with the results obtained by cooling, as shown in Fig. 3g and h. The changes in maximum emission wavelength and intensity with temperature on warming are linear ($\lambda_{max}/nm = -0.19 T/K + 595.00$, $R^2 = 0.9921$; $I/I_{max} = -0.0015 T/K + 1.12$, $R^2 = 0.9964$). These experiments demonstrate that the crystal does not only respond to temperature changes by emitting light, but it also transduces and outputs the temperature information by acting as an optical waveguide. The measurement was also performed after the crystal was repeatedly bent, and the results show that the sensitivity was practically unaffected even after the crystal was bent 1000 times (Supplementary Fig. 17, Supplementary Table 3). The signal from the crystal is not affected by the temperature of the section through which the optical signal is transmitted. The crystal’s output reflects only the temperature of the point of excitation, and this property favors this material, and in principle also other emissive elastic organic crystals, as an active sensing medium for luminescence thermometry.

### Computational modeling of the fluorescence

To investigate the root cause of the temperature dependence of fluorescence, density functional theory (DFT) calculations were performed on a molecule of compound 1. Supplementary Fig. 18 shows the frontier molecular orbitals based on the ground-state ($S_0$) geometry of compound 1. Obviously, the LUMO, HOMO, and HOMO-1 are mostly $\pi$-orbitals, while the HOMO-2 is an n-orbital and distributed over the carbonyl group and its surroundings. The HOMO-1 is localized on the indole ring and the HOMO spreads over the indole ring and the adjacent vinyl group. In contrast, the LUMO is delocalized over the entire molecular backbone. Vertical excitations from $S_0$ to $S_1$, $S_2$, and $S_3$ are dominated by the transitions from the HOMO, HOMO-2, and HOMO-1 to LUMO, respectively. The $S_2$ and $S_3$ excitations energies are close to the $S_2$ excitation energy (Supplementary Table 4). Deeper insight into the fluorescence properties necessitated closer inspection of the electronic structures in the excited-state optimized geometries. The nature of the frontier orbitals in the excited state is hardly altered; moreover, the $S_2$ and $S_3$ excitations still correspond to the HOMO and HOMO-2 to LUMO transitions, respectively. Interestingly, in the $S_3$ geometry, besides the HOMO → LUMO transition, the HOMO → LUMO transition has a significant contribution (14.8%) to the $S_3$ excitation, with significant oscillator strength ($f = 0.30$) (Fig. 4a). Because of the relatively close energies between the $S_1$ and $S_3$ states (Supplementary Tables 4 and 5), the $S_3$ population will be increased when the temperature is elevated, which is beneficial to enhance the luminescence from $S_3$ and results in blue-shift in the emission (Fig. 4b). This result supports the experimental observations.

### Relation to the crystal structure

The change in emissive properties of any crystal is also inevitably related to its crystal structure, and therefore the structure was determined at 298 and 100 K (Supplementary Fig. 19, Supplementary Table 6, Supplementary Data 1 and 2). At 298 K, the crystal is in the monoclinic system (space group P2_1/n) with $Z = 4$ and cell parameters $a = 12.1007(9)$ Å, $b = 4.8876(4)$ Å, $c = 3.7142(2)$ Å, and $\beta = 91.548(3)^\circ$, and unit cell volume $V = 1875.0(2)$ Å³. The torsion angles around the two...
single chemical bonds between the benzene and the indole rings are 12.2° and 13° (Supplementary Fig. 19). The molecules are stacked in a parallel fashion and form columnar structures by π···π interactions with an intermolecular distance of 3.738 Å along the crystallographic [010] direction (Fig. 5a). Upon cooling to 100 K, the crystal symmetry is retained, but the cell parameters change to \(a=12.0224(7)\) Å, \(b=4.8149(3)\) Å, \(c=31.4650(18)\) Å and \(\beta=92.271(2)^\circ\), and the unit volume is \(V=1819.97(19)\) Å\(^3\). The unit cell shrinks by 0.7% along the \(a\) axis, 1.5% along the \(b\) axis, and 0.8% along the \(c\) axis. The torsion angles between the benzene ring and the indole ring remain very close, 11.0° and 14° respectively, and thus the high degree of planarity of the molecule is retained at low temperature (Supplementary Fig. 19). The stacking distance of the molecules is slightly reduced, to 3.579 Å (Fig. 5b). Since the molecular geometry is practically conserved, similar to other cases\(^6,9,10\), the change in emission can be attributed to the decreased π···π distance, which is related to the significant shrinking of the lattice along the \(b\) axis. Furthermore, the enhanced intensity of emission at low temperatures is related to the restriction of non-radiative transition pathways and enhancement of the competing radiative processes\(^8\). In addition, the interaction energy among the molecules at 298 and 100 K were both calculated. The energy frameworks were constructed by using Crystal Explorer\(^6\) and the B3LYP hybrid functional with the 6–31G(d,p) basis set, where semiempirical dispersion was included by using D2 version of Grimme’s dispersion, and they are visualized in Fig. 5c. Based on the preceding discussion, the structural changes observed upon cooling indicate that the unit cell contraction along the \(b\) axis is the most likely contributor to the appealing change in emission. The calculated interaction energy between molecule 1 and molecule 2 of \(-66.4\) kJ mol\(^{-1}\) at room temperature was found to decrease to \(-67.5\) kJ mol\(^{-1}\) at 100 K (Table 1). This decrease in energy is in accord with the strengthening of the π···π interaction and is also consistent with the observations from the crystal structures.

The crystal structure is also directly related to the elastic properties of compound 1. The thicker blue bars in Fig. 5c correspond to the π···π interactions along the [010] direction. The C–H···O interactions (\(d=2.600\) Å, \(\theta=148.8^\circ\) and \(d=2.418\) Å, \(\theta=135.6^\circ\)) and F···F interactionsLeutwyler et al. (2015) (2015) (2015) (2015) (2015) (2015) (2015) (2015). The effects of temperature on the structure and energy framework of compound 1 are illustrated in Fig. 5a. The arrows indicate the torsional angles discussed in the text. The energy bars showing the interaction energies. The blue bars represent the interactions between molecules, and their thickness indicates the value of interaction energy. The energy frameworks viewed along the \(b\) axis are consistent with the qualitative requirements for bending. Parallel packing structure, and the purported expansion and contraction of the outer and inner arcs of the crystal that are suggested to occur during the bending of the crystal.
The intermolecular interaction energy between different molecules (molecules 1–3, as shown in Fig. 5c).

### Data availability

Crystallographic data for the structures reported in this Article have been deposited at the Cambridge Crystallographic Data Centre, under deposition numbers CCDC 2155928 and 2155929. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via www.ccdc.cam.ac.uk/data_request/cif. All data are available from the corresponding authors upon request.

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Author contributions
Q.D., P.N., and H.Z. conceived the study. Q.D. prepared and performed most experiments on the crystals. Q.D. and B.L. performed a series of experiments on crystal optical waveguiding properties, X.M. and Y.Y. performed the density functional theory calculations. Q.D., X.Y., and L.F.L. revised the article under the guidance of L.L. The paper was written with contributions from Q.D., L.L., P.N., and H.Z. All authors have given their approval for the final version of this paper.

Competing interests
The authors declare no competing interests.

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