Helix-constructed polar rare-earth iodate fluoride as a laser nonlinear optical multifunctional material†

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The first trivalent rare-earth iodate fluoride nonlinear optical (NLO) crystal, Y(IO3)2F (YIF), was successfully designed and synthesized, featuring polarization-favorable helical chains constructed from trans-YO2F2 polyhedra and IO3 groups. It exhibited a suitable balance of a wide transparency range of 0.26–10.0 μm, high laser damage threshold (LDT) of 39.6 × AgGaS2, and moderate second harmonic generation (SHG) effect of 2 × KDP. A series of doped RE:YIF (RE = Pr, Nd, Dy, Ho, Er, Tm, and Yb) crystals were easily synthesized benefiting from the spring-shaped helix structure, which possess wide absorption and emission peaks as well as long lifetime, especially in the visible and near-infrared regions. Particularly, the remarkable fluorescence properties of Nd and Yb doped YIF crystals are comparable to and even better than those of traditional self-frequency doubling (SFD) crystals such as YAB, YCOB, and GdCOB. Thus, these RE-doped YIF crystals are promising laser SFD crystals. This work also indicated that constructing helical chains should be an effective strategy for the design of inorganic polar materials.

A laser self-frequency doubling (SFD) crystal is a type of excellent multifunctional crystal, which has both the simulated emission function of a laser crystal and the second harmonic generation (SHG) properties of a nonlinear optical (NLO) crystal, so that it can achieve functions such as self-activation and SFD.1 By using laser SFD crystals, all-solid-state lasers can achieve advantages such as a small size and compact structure, low cost, easy adjustability, high stability, and so on. Generally, a laser SFD crystal is composed of a host crystal and one or more types of active ions which mainly include trivalent lanthanide rare-earth ions except those with a full-shell or semi-full-shell (Sc3+, Y3+, La3+, Lu3+, and Gd3+) and some transition ions.2 However, only a few host crystals are practically available at present, mainly including LiNbO3 (LN), YAlO3 (YAB), YCa4O(BO3)3 (YCOB), GdCa4O(BO3)3 (GdCOB), etc.3 Therefore, there is still an urgent demand to explore new laser NLO multifunctional crystals.

A laser SFD host crystal must be a NLO crystal with a noncentrosymmetric (NCS) or polar structure and easily doped with active ions. However, it is known that minerals and synthetic crystals tend to crystallize into centrosymmetric (CS) structures. It is still of great technical interest but challenging to effectively synthesize polar crystals.

To accelerate the discovery of new NLO materials, the introduction of the largest electronegative element F into oxysalts such as borates, carbonates, nitrates, and phosphates has been recognized as an effective strategy.4 Besides, F can usually improve crystals’ optical properties such as broadening the band gaps and enlarging laser damage thresholds (LDTs). In recent years, metal iodate fluorides have become a research hotspot. Several metal iodate fluoride NLO crystals have been reported, mainly including Bi(IO3)2F5, BiF3(IO3)4, ABF3 (A = K, Rb, Cs, NH4), β-Ba[VFO3(IO3)]2, z/β-Ba2[VO2-F2(IO3)2]IO3, K5[W2O7F2(IO3)3], CsVO2F(IO3)4, Cs(IO3)2F2·H2O, Sn(IO3)2[IO3]2, etc. These metal iodate fluoride NLO crystals generally exhibit a wide transparent range covering the visible to mid-IR region, large SHG effects, and a high laser damage threshold (LDT). Notably, a trivalent rare-earth iodate fluoride has not been reported yet. Considering the varied and flexible coordination patterns of rare-earth ions, and that crystals containing trivalent rare-earth ions are often easily doped with other rare-earth active ions for excitation luminescence, it is meaningful to explore the potential

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trivalent rare-earth iodate fluoride photoelectric functional materials.

In this study, we have successfully designed and synthesized the first trivalent rare-earth iodate fluoride, Y(IO₃)₂F (YIF), which exhibited a good balance of a wide UV transmission range, high LDT and relatively large SHG effect. In particular, active ion doped RE:YIF (RE = Pr, Nd, Dy, Ho, Er, Tm, and Yb) crystals were also synthesized easily, showing wide absorption and emission peaks as well as long lifetime, especially in the visible and near-infrared regions. Specifically, the remarkable fluorescence properties of Nd and Yb doped YIF crystals are superior to those of several popular commercial laser SFD crystals such as YAB, YCOB, and GdCOB, indicating their potential application as laser SFD crystals. Most importantly, the novel helical structure will provide important guidance for the design of polar inorganic NLO crystal materials. Herein, we will report on the synthesis, crystal structure, and linear and nonlinear optical properties of YIF, as well as the fluorescence performance of RE doped YIF.

The YIF and RE-doped YIF crystals (Fig. 1a) were all synthesized by a mild hydrothermal method at a temperature of 240 °C. As plotted in Fig. S1 in the ESI† powder X-ray diffraction (XRD) analysis confirmed the purity of YIF and the structure-nondestructive doping of RE doped YIF. The energy dispersive X-ray spectroscopy (EDS) analysis confirmed the existence of Y, I, O, and F elements in YIF (Fig. S2†). The synthesized crystals exhibited good chemical stability. Thermogravimetric analysis (TGA) showed that the YIF was thermally stable up to about 380 °C (Fig. S3†). The weight loss of 73.5% (cal. 72.9%) at around 380–1100 °C is attributed to the release of I₂O₅, and the final residue at 1100 °C was confirmed to be YOF by powder XRD (Fig. S4†).

YIF crystallizes in the hexagonal polar space group P6₃ (the detailed crystallographic data are listed in Tables S1–S4 in the ESI†), and it has one Y atom, one F atom, two I atoms, and six O atoms in an asymmetric unit. The bond valence calculations revealed the valence states of 5.13 or 5.46 for Y⁵⁺ and 3.08 for Y³⁺, being consistent with their ideal valence. Unlike most reported metal iodate fluorides or other Y-containing oxysalt fluorides, the YIF features a novel spring-shaped one-dimensional (1D) helical chains constructed from F-bridged trans-YO₆F₂ polyhedra with IO₃ groups hung on (Fig. 1b and c). The bond lengths of Y–O, Y–F, and I–O bonds are varied in the region of 2.33(2)–2.55(2), 2.14(2)–2.22(2), and 1.77(2)–1.84(2) Å, respectively. These 1D helical chains are further connected to each other into a 3D structure by sharing IO₃, and the denging bonds on IO₃ groups are therefore eliminated, which is favorable for the enlargement of the band gap. Benefitting from the polarization-induction of the helical chains, the IO₃ groups were configured to a polar arrangement. The local dipole moment of an individual IO₃ group was calculated to be 14.62–14.90 D (Debye), which is consistent with other iodates.³–⁵ The x and y components from the twelve IO₃ groups within one unit cell cancel each other out completely, while the z component add up to a net dipole moment of −3.13 D, indicating the formation of a polar arrangement.

Notably, the helical structure has received extensive attention in the field of metal organic coordination chemistry but rarely been mentioned in the field of inorganic NLO crystals.¹⁵ Therefore, it will be meaningful and interesting to study the helical chain formation of YIF by comparing with Y(SO₄)F (Pna2₁), Y(SeO₃)F (P2₁/c)¹⁷ and NaYF(PO₄) (C2/m).¹⁸ All three crystals possess YO₆F₂ polyhedra and CS space groups. The YO₆F₂ polyhedra in NaYF(PO₄) are cis which hinder the formation of single chains. As for Y(SO₄)F and Y(SeO₃)F, their trans-YO₆F₂ polyhedra promote the formation of single YF(SO₄) and YF(SeO₃) chains. Unfortunately these chains are all centric linear chains, instead of helical, which then induce SHG-groups (SO₄ and SeO₃) packed with inversion centers (Fig. 1c and d). It can be inferred that the trans-YO₆F₂ polyhredra, rather than the cis-YO₆F₂ polyhedra, are essential for the formation of single chains. In YIF, the density of Y (6.78 × 10⁻³ Å³) is much smaller than those of Y(SO₄)F (10.87 × 10⁻³ Å³) and Y(SeO₃)F (12.45 × 10⁻³ Å³). The lower density means that the space to be filled between Y⁺ is larger, and the formation of the helical chains are conducive to tight filling. Meanwhile, the IO₃ groups with a lower valence are more favorable to achieve the charge balance of the framework with fewer positive charges but more lacunae. The molar ratio of the SHG-groups to Y may be a simple but effective parameter to guide the synthesis of polar crystals with helical chains. Obviously, the value of the IO₃/Y molar ratio in YIF is 2, which is twice that of Y(SO₄)F and Y(SeO₃)F. Hence, we deem that trans-YO₆F₂ polyhedra, SHG-groups with a lower valence, and a large molar ratio of SHG-groups to Y are all positive factors for helical chain formation. This rule may be applied to other cationic systems.

The polar structure of YIF prompted us to study its SHG properties under 1064 nm laser radiation via the Kurtz–Perry method.¹⁹ As shown in Fig. 2, the YIF exhibited a moderate SHG response of 2.0 × KDP (particle size 150–212 μm) and a type-I phase-matching behavior. Because the SHG response is proportional to the square of the effective nonlinear coefficient,
the real-space atom-cutting technique were adopted to further intuitively and quantificationally evaluate the SHG-contribution of IO$_3$ groups and YO$_6$F$_2$ polyhedra. The SHG-weighted electron densities of the occupied and unoccupied states in the virtual electron (VE) process have been calculated for the VE process dominantly contributing to the overall SHG effects. As shown in Fig. 3a, the SHG-weighted electron density of the occupied states is mainly concentrated on O$^2$–. In the unoccupied states (Fig. 3b), the orbitals on both O$^2$– and I$^5$– were the major source of SHG-weighted electron densities. It means that the IO$_3$ groups have made the dominant contribution to the whole SHG effects. This is also confirmed by the real-space atom-cutting technique (see Table S5†), where the contribution of IO$_3$ groups (−0.83 pm V$^{-1}$) to the SHG coefficient $d_{31}$ was very large, while the YO$_6$F$_2$ polyhedra made a negligible negative contribution (0.07 pm V$^{-1}$). Besides, the birefringence was calculated to be 0.041 at a wavelength of 1064 nm, which is large enough for satisfy the phase-matching condition at 1064 nm. And the atom-cutting result also exhibited the dominant contribution of IO$_3$ groups to the birefringence.

A series of RE (RE = Pr, Nd, Dy, Ho, Er, Tm, and Yb) doped YIF crystals were synthesized in order to confirm its great potential for laser application. The synthesis process is quite easy, which should be attributed to the high compatibility of the spring-shaped helix structure with doped ions with different ionic radii and concentrations. Fig. 4 shows the fluorescence spectra of RE doped YIF crystals at room temperature. The detailed spectral properties are shown in Table S6,† including the main absorption peak and width, the typical emission peak and width, and the fluorescence lifetime. As demonstrated in Fig. 4b and c, the full width at half maximum (FWHM) of absorption and emission peaks for Dy:YIF was much wider than that of the traditional laser crystal Y$_3$Al$_5$O$_{12}$ (YAG) and nearly the same as that of the newly developed Na$_2$Gd$_4$(MoO$_4$)$_7$ (NGM) crystal. And the Pr:YIF sample exhibited a very broad absorption and emission peak in comparison with commercial YLiF$_4$ (YLF) and YAIO$_3$ (YAP) crystals. Besides, the fluorescence lifetime for both Dy and Pr doped YIF were comparable to that of the aforementioned crystals. Therefore, both Dy and Pr doped YIF are very promising laser gain media in the visible spectrum of 1064 nm.

Fig. 2  The powder SHG measurements of YIF and KDP at 1064 nm.

Fig. 3  The SHG-weighted densities for (a) occupied and (b) unoccupied electronic states in YIF.
range. As for the Nd:YIF sample (Fig. 4d), its fluorescence properties far exceeded those of the well-known traditional self-frequency-doubling crystals YAB, YCOB and GdCOB, including much broader absorption and emission peaks as well as longer lifetime.28 As plotted in Fig. 4e, the fluorescence properties of Yb:YIF were nearly the same as those of Yb:YAB. Besides, the NLO effect of YIF was close to those of the abovementioned crystals (1.1–1.3 pm V−1).29 Therefore, it can be inferred that both Nd and Yb doped crystals would be new remarkable SFD crystals in the near-infrared region. With regard to the samples doped with Er, Tm and Ho, one can see the significant feature of broadening absorption and emission peaks, which should be attributed to the varied and flexible coordination patterns of rare earth ions in RE:YIF. In brief, from the viewpoint of fluorescence, YIF is an excellent laser SFD host crystal particularly in the field of the Dy and Pr activated visible region as well as Nd and Yb activated near-infrared region.

Conclusions

In conclusion, via the introduction of F−, we successfully obtained the first trivalent rare-earth iodate fluoride nonlinear optical (NLO) crystal Y(IO3)2F (YIF). It featured unusual helical chains constructed from trans-YO4F2 polyhedra and IO3 groups, which are favorable for polar arrangement of the IO3 groups. Apart from the suitable balance of a wide transparency range of 0.26–10.0 μm, high laser damage threshold (LDT) of 39.6 × AgGaS2, and moderate second harmonic generation (SHG) effect of 2 × KDP, YIF showed great potential as a laser self-frequency doubling (SFD) host crystal. Benefiting from the spring-shaped helix structure, a series of doped RE:YIF (RE = Pr, Nd, Dy, Ho, Er, Tm, and Yb) crystals were easily synthesized, which possessed wide absorption and emission peaks as well as long lifetime, especially in the visible and near-infrared regions. Particularly, compared with those famous laser and SFD crystals including YAG, NGM, YLF, and YAP as well as YAB, YCOB, and GdCOB, the Dy, Pr, Nd and Yb doped YIF all exhibited more remarkable and comparable fluorescence properties. Hence, it can be concluded that YIF is a promising NLO and SFD host multifunctional crystal. Besides, constructing helical chains should be an effective strategy for the design of inorganic polar materials.

Conflicts of interest

There are no conflicts to declare.

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