Growth and Passive Q-Switching Application of Cr:TiTe₃O₈ Crystal

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Abstract: A Cr⁴⁺-doped TiTe₃O₈ crystal with dimensions up to 21 mm × 21 mm × 11 mm was grown successfully by the top-seeded solution growth method. A high-resolution X-ray diffraction experiment showed that the full width at half-maximum of the rocking curve was 41.90'. Energy-dispersive spectrometry (EDS) and X-ray fluorescence spectroscopy (XRF) proved that Cr⁴⁺ ions have been doped into the TiTe₃O₈ crystal with molar percentage of 4.9%. The Cr:TiTe₃O₈ crystal exhibited an absorption range from 995 to 1565 nm, which is suitable for passive Q-switch at 1064 nm. Then, a passive Q-switching solid-state laser operating at 1064 nm was realized using a Cr:TiTe₃O₈ crystal as the saturable absorber. The maximum laser pulse energy is calculated to be 0.7 µJ, and the maximum peak power reaches 0.7 W.

Keywords: Cr:TiTe₃O₈; passive Q-switching; top seed solution method

1. Introduction

Solid-state lasers have been widely used in the scientific research, medical treatment and military fields [1–5]. In order to obtain high energy outputs, the time domain modulation in laser operations is a kind of laser technology with much importance. Different from the acousto-optic and electro-optic Q-switching, passive Q-switching with saturable absorbers is a simple and economical method. Doping transition metal ions such as Cr⁴⁺, V³⁺ or Co²⁺ into crystals is the common method to obtain saturable absorbers [6–9]. The saturable absorber properties are mostly determined by doping transitions. Currently, the most widely used saturable absorber is Cr⁴⁺:YAG crystal due to the good optical and thermal properties of YAG [10] crystal. It is well-known that there are valence differences between the Cr⁴⁺ and Y³⁺ ions [11]. Therefore, the excellent optical crystals with positive ions exhibiting M⁴⁺ could be the ideal host crystals for the saturable absorbers.

TiTe₃O₈ belongs to the space group of Ia-3 in cubic system with a = 10.9585(5) Å [12,13]. The bulk crystal was firstly grown as a novel acousto-optic crystal in our group. The thermal conductivity coefficient of the TiTe₃O₈ was determined to be 3.3 Wm⁻¹K⁻¹ at room temperature [12], which indicates the crystal exhibits good thermal property and is suitable for optical applications. We consider the TiTe₃O₈ crystal as a suitable material for Cr⁴⁺ doping due to the valence balance and appropriate ion radii.

In this work, the Cr⁴⁺:TiTe₃O₈ crystal with dimensions up to 21 mm × 21 mm × 11 mm was successfully grown by the top-seeded solution growth method. The molar percentage of Cr⁴⁺ in TiTe₃O₈ crystal was determined to be 4.9%. The Cr:TiTe₃O₈ crystal exhibits a wide absorption band range from 995 to 1565 nm due to the doping of the Cr⁴⁺ ions. Additionally, the Cr:TiTe₃O₈ crystal was successfully used as a saturable absorber in the laser operating at 1064 nm.
2. Materials and Methods

2.1. Synthesis and Growth of Cr:TiTe$_3$O$_8$ Single Crystal

The polycrystalline powders of Cr:TiTe$_3$O$_8$ were prepared using the traditional high-temperature solid-state reaction method. The TiO$_2$ (Alfa Aesar, 99.99%), TeO$_2$ (Alfa Aesar, 99.99%) and CrO$_2$ (Aladdin, 99.0%) with mole ratio of 0.95:3:0.05 were thoroughly ground together using an agate pestle and mortar to ensure homogeneity, and they were pressed into a column. Then, the column was heated from room temperature to 700 °C with a rate of 10 °C/min in air. After holding at 700 °C, the column was quenched to room temperature directly. It was ground, pressed and sintered two more times as described above.

The single crystal Cr:TiTe$_3$O$_8$ was grown by the flux method because TiTe$_3$O$_8$ is an incongruently melting compound. The flux consists of TeO$_2$ and Li$_2$CO$_3$ with molar ration of 1:5. Then, a mixture of the Cr:TiTe$_3$O$_8$ polycrystalline powder and flux was placed in a dedicated temperature-controlled furnace. The furnace was heated up to 950 °C slowly and held for 3 days to ensure the mixture melted into a uniform and stable solution. A platinum wire was inserted into the surface of the solution. The temperature was decreased from 950 °C to 750 °C at a rate of 1 °C per hour. The saturation temperature was determined by observing whether the crystal was dissolved or grown in the solution. A crystal seed was cut from the millimeter-sized crystals which were grown by spontaneous crystallization on the platinum wire. Then, the selected crystal seed was tied to a platinum wire and put into the flux at the temperature of 4 °C above saturation point. The solution was cooled to the saturation temperature in two hours and then cooled at a rate of 0.5–0.8 °C per day. A single crystal was obtained after one month. When the growth process was finished, the crystal was pulled out from the solution and cooled to room temperature for three days.

2.2. Powder X-ray Diffraction (PXRD)

A multifunctional X-ray powder diffractometer D8 ADVANCE machine made by Bruker AXS was used in the experiment. The PXRD pattern was taken at room temperature by the Cu-K$\alpha$ radiation over an angle range (2$\theta$) from 10° to 90° with a scanning step size of 0.02°.

2.3. Laue Back-Reflection Measurement

The orientation and structure of the crystal can be obtained by analyzing the Laue X-ray back-diffraction image. A Laue X-ray backscattering diffractometer developed by Multiwire Laboratories Company was used in the experiment. The test was carried out on a Cr:TiTe$_3$O$_8$ wafer along the (200) plane with a size of 5 mm × 5 mm × 1 mm, which is mechanically polished on both sides.

2.4. High-Resolution X-ray Diffraction

It was carried out on a Bruker-AXS D8 DISCOVER diffractometer equipped by Cu-K$\alpha$1 radiation (\(\lambda = 1.54056\) Å) source. The accelerating voltage and tube current were 60 kV and 80 Ma, respectively. The step size and step time were 0.001° and 0.1 s, respectively.

2.5. Transmission and Absorption Spectroscopy

The optical transmission spectra of Cr:TiTe$_3$O$_8$ single crystal was collected on an Agilent Cary 5000 UV-vis-NIR spectrometer in the 200–2000 nm region. The spectra of the crystal in the 2000–8000 nm region were collected on a Nicolet NEXUS 670 FTIR spectrometer. A crystal wafer in size of 5 mm × 5 mm × 1 mm polished on both sides was used in the test.

2.6. Energy Dispersive Spectrometer (EDS)

According to the characteristics of different X-ray photon characteristic energy of each element, the component analysis was carried out by the energy spectrometer. The experiment was carried out on a 5 mm × 5 mm × 1 mm wafer, polished on both sides, employing a Hitachi S-4800 field emission scanning electron microscope.
2.6. Energy Dispersive Spectrometer (EDS)

According to the characteristics of different X-ray photon characteristic energy of Ti element, the component analysis was carried out by the energy spectrometer. The Cr:TiTe3O8 crystal exhibits a dark green color.

Figure 2. Cr:TiTe3O8 single crystal grown along [100] crystallographic axis.

3. Results and Discussion

3.1. Characterization of Polycrystalline Cr:TiTe3O8

After rapid cooling, there are dark green polycrystalline precipitates on the seed crystal rod. The polycrystal is ground into powder, and the XRD pattern of the powder is shown in Figure 1. It can be seen that each peak matches the peak at the corresponding position of the standard card (TiTe3O8 PDF 24-1348) [14] and no other impure peaks. XRD pattern indicates that there is no significant structure change in the crystal by Cr doping.

![Figure 1](image1)

Figure 1. (a) Polycrystals obtained by spontaneous nucleation; (b) XRD patterns of Cr:TiTe3O8 and the pure TiTe3O8.

3.2. Crystal Growth

The Cr:TiTe3O8 crystal was grown using flux method by slowly temperature dropping for a month. A piece of directional crystal seed cut along the [100] axis was used. Finally, a single crystal 21 mm \( \times \) 21 mm \( \times \) 1 mm in size was obtained, as shown in Figure 2. It has a square shape from top to bottom. The Cr:TiTe3O8 crystal exhibits a dark green color.

![Figure 2](image2)

Figure 2. Cr:TiTe3O8 single crystal grown along [100] crystallographic axis.

3.3. Laue Photograph and HRXRD

The single crystal orientation was determined by the back-beam Laue method. Four points on a 5 mm \( \times \) 5 mm \( \times \) 1 mm wafer were randomly selected, and the related Laue diffraction images are shown in Figure 3a. Each diffraction spot was clear and symmetrical without splitting and had good axial symmetry. The X-ray rocking curve of Cr:TiTe3O8 wafer (200) diffraction plane is shown in Figure 3b. It can be seen that the FWHM (full
The single crystal orientation was determined by the back beam Laue method. Four images are shown in Figure 3a. Each diffraction spot was clear and symmetrical without splitting and had good axial symmetry. The X-ray rocking curve images are shown in Figure 3b. It can be seen that the FWHM (full width at half maximum) is as low as 41.90 arcsec, and the peak is symmetrical and sharp. Both the Laue photograph and X-ray rocking curve measurements indicate that the crystal is of good quality.

![Laue Photograph and X-ray rocking curve](image)

**Figure 3.** (a) Laue photographs of Cr:TiTe$_3$O$_8$ single crystal; (b) X-ray rocking curve of the Cr:TiTe$_3$O$_8$ single crystal.

### 3.4. Optical Properties

The transmittance spectrum of the Cr:TiTe$_3$O$_8$ with a thickness of 1 mm (Figure 4a) at room temperature is shown in Figure 4b. It can be seen that the transmittance is more than 70% in the range from 1 to 5 μm. The UV-Vis spectra are shown in Figure 4c, which exhibit an absorption peak at 687 nm and an absorption region from 995 to 1565 nm. Magnified absorption spectra at about 1 μm are shown in Figure 4d. The spectra are in good agreement with the characteristic absorptions of Cr$^{4+}$ ions [15].

![Optical Properties](image)

**Figure 4.** (a) A Cr:TiTe$_3$O$_8$ crystal wafer with dimensions of 5 mm × 5 mm × 1 mm; (b) transmission spectra of the Cr:TiTe$_3$O$_8$ single crystal perpendicular to the (100) plane; (c) the absorption spectrum of the Cr:TiTe$_3$O$_8$ crystal; (d) the magnified absorption spectrum at 1 μm.

### 3.5. EDS and XRF

The chemical composition of Cr:TiTe$_3$O$_8$ crystal was determined using EDS technique, and the recorded spectrum is shown in Figure 5. The elemental mapping proves the
presence of Te, O, Ti and Cr elements in the crystal. EDS test can only be used for semi-quantitative analysis. In order to quantitatively analyze the concentration of Cr element in the crystal, the XRF test was carried out. The result is shown in Table 1. The molar percentage of Cr element is determined to be 4.9%.

![Figure 5. EDS image of Cr:TiTe$_3$O$_8$ crystal.](image)

Table 1. The content of each element in the crystal tested by XRF.

| No. | Element | wt.% | mol% |
|-----|---------|------|------|
| 1   | Ti      | 12.9 | 13.81|
| 2   | Cr      | 0.689| 4.9  |
| 3   | Te      | 86.4 | 34.74|

3.6. Passive Q Switch Test

In this section, the Cr:TiTe$_3$O$_8$ single crystal was tried to use as a saturable absorber in the laser operating. The schematic diagram of the experimental setup is shown in Figure 6. A Cr:TiTe$_3$O$_8$ crystal in size of 4 mm × 4 mm × 1 mm and a Nd:YVO$_4$ crystal in size of 3 mm × 3 mm × 10 mm were inserted into a 30 mm long resonant cavity. The Nd:YVO$_4$ crystal is coated with AR films working at 808 and 1064 nm. The cavity consists of an input-coupling mirror M$_1$ and an output-coupling mirror M$_2$ with the curvature radii of 100 mm and 200 mm, respectively. The mirror M$_1$ exhibits a high transmission of 808 nm (AR, R < 0.2%) and a high reflectivity at 1064 nm (HR, R > 99.8%). The mirror M$_2$ is coated with HR at 808 nm and partially reflective film at 1064 nm (R = 60%). The cavity is pumped by a fiber-coupled 808 nm CW-diode-pumped laser. Both the Nd:YVO$_4$ and Cr:TiTe$_3$O$_8$ crystals are wrapped with indium foil and mounted in water-cooled copper blocks and cooled to 15 °C. The output energy is measured by a PM100 digital power meter. The temporal behaviors of the pulse are recorded by a digital oscilloscope (DPO4104B-L, Tektronix).

![Figure 6. Schematic diagram of experimental setup.](image)

In our experiments, the Q-switching pulses were observed, which means that the Cr:TiTe$_3$O$_8$ has realized the saturable absorber property. The performances of the Cr:TiTe$_3$O$_8$ Q-switching laser are shown in Figure 7. The curve of output power changing with pump
power is shown in Figure 7a. The maximum output power can reach 50 mW with the pump power 1.194 W. Then, the repetition frequency and pulse width are determined to be 67 kHz and 1 μs. The maximum laser pulse energy is calculated to be 0.7 μJ, and the maximum peak power reaches 0.7 W. The laser pulse trains are displayed in Figure 7b. The pulses are equally spaced and there are no impurity peaks. Figure 7c is a typical monopulse with a pulse width of 1.667 μs.

Figure 7. (a) The average output powers versus pump powers; (b) the passive Q-switching laser pulse train; (c) a typical pulse shape of 1.667 μs width.

4. Conclusions

In this work, the Cr:TiTe$_3$O$_8$ crystal, which can be used as a saturable absorber, was successfully grown by top-seeded solution growth method. The molar percentage of Cr$^{4+}$ ions in TiTe$_3$O$_8$ crystal is determined to be 4.9%. Cr:TiTe$_3$O$_8$ crystal shows the absorption range from 995 to 1565 nm. A passive Q-switching solid-state laser with Cr:TiTe$_3$O$_8$ crystal as the saturable absorber operating at 1064 nm was realized.

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References

1. Waynant, R.W.; Ilev, I.K.; Gannot, I. Mid–infrared laser applications in medicine and biology. Philos. Trans. R. Soc. A Math. Phys. Eng. Sci. 2001, 359, 635–644. [CrossRef]

2. Stupl, J.; Neuneck, G. Assessment of Long Range Laser Weapon Engagements: The Case of the Airborne Laser. Sci. Glob. Secur. 2010, 18, 1–60. [CrossRef]

3. Tunc, B.; Gulsoy, M. The Comparison of Thermal Effects of a 1940-nm Tm:fiber Laser and 980-nm Diode Laser on Cortical Tissue: Stereotaxic Laser Brain Surgery. Lasers Surg. Med. 2020, 52, 235–246. [CrossRef] [PubMed]

4. Zhao, J.L.; Wu, R.; Wang, L.; Wu, Y. The Application of 3D Laser Scanning Technology in Ancient Architecture Protection. Appl. Mech. Mater. 2013, 353–356, 3476–3479. [CrossRef]

5. Shepelev, A.E.; Antipov, A.A.; Putilov, A.G.; Osipov, A.V.; Arakelian, S.M. Methods for the solid-state lasers generation modes control for the material laser processing efficiency improving. J. Phys. Conf. Ser. 2021, 1822, 012015. [CrossRef]

6. Lin, H.-Y.; Sun, D.; Copner, N.; Zhu, W. Nd:GYSGG laser at 1331.6 nm passively Q-switched by a Co:MgAl2O4 crystal. Opt. Mater. 2017, 69, 250–253. [CrossRef]

7. Hecht, H.; Burshtein, Z.; Katzir, A.; Noach, S.; Sokol, M.; Frumker, E.; Galun, E.; Ishaya, A.A. Passive Q-switching of a Tm:YLF laser with a Co2+ doped silver halide saturable absorber. Opt. Mater. 2017, 64, 64–69. [CrossRef]

8. Lin, H.; Liu, H.; Li, Y.; Sun, D. Passively Q-switched Nd:GGG laser at 1110.2 nm by Cr:YAG. Laser Phys. 2018, 28, 125001. [CrossRef]

9. Song, T.; Li, P.; Chen, X.; Ma, B.; Dun, Y. Passively Q-switched Nd:GYSGG laser operating at 1.3 μm with V:YAG as saturable absorber. Optik 2016, 127, 10621–10625. [CrossRef]

10. Lefever, R.A.; Torpy, J.W.; Chase, A.B. Growth of Single Crystals of Yttrium Aluminum Garnet from Lead Oxide—Lead Fluoride Melts. J. Appl. Phys. 1961, 32, 962–963. [CrossRef]

11. Shannon, R.D. Revised effective ionic radii and systematic studies of interatomic distances in halides and chalcogenides. Acta Crystallogr. Sect. A 1976, 32, 751–767. [CrossRef]

12. Lu, W.; Gao, Z.; Wu, Q.; Tian, X.; Sun, Y.; Liu, Y.; Tao, X. Tailored fabrication of a prospective acousto–optic crystal TiTe3O8 endowed with high performance. J. Mater. Chem. C 2018, 6, 2443–2451. [CrossRef]

13. Meunier, G.; Galy, J. Sur une déformation inédite du réseau de type fluorine. Structure cristalline des phases MTe3O8 (M = Ti, Sn, Hf, Zr). Acta Crystallogr. 1971, B27, 602–608. [CrossRef]

14. Jiao, X.; Zhong, C.; Zhang, S.; Liu, X.; Li, B. Microwave dielectric properties of BaO–TiO2–TeO2 ternary system. J. Mater. Sci. 2010, 45, 3331–3335. [CrossRef]

15. Li, C.; Xu, J.; Liu, W.; Zheng, D.; Zhang, S.; Zhang, Y.; Lin, H.; Liu, L.; Liu, J.; Zeng, F. Synthesis and characterization of Cr4+-doped Ca3GeO4 tunable crystal. J. Alloys Compd. 2015, 636, 211–215. [CrossRef]