

Multiband infrared luminescence of Er^{3+} - Ho^{3+} - Nd^{3+} / Tm^{3+} -codoped telluride glasses

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Abstract This paper reports the simultaneous emissions around 1.53, 1.80, 2.10, 2.70 and 3.00 μm in Er^{3+} - Ho^{3+} - Nd^{3+} / Tm^{3+} -codoped telluride glasses upon excitation of a conventional 808 nm laser diode. Both emission bands of 1.53 and 2.70 μm were assigned to the transitions of $^4\text{I}_{13/2}$ - $^4\text{I}_{15/2}$, $^4\text{I}_{11/2}$ - $^4\text{I}_{13/2}$ of Er^{3+} ions, respectively, the emission near 1.80 μm was assigned to the transition $^4\text{F}_4$ - $^4\text{H}_6$ of Tm^{3+} ions, and the emissions at 2.10 and 3.00 μm arose from the transitions of $^5\text{I}_7$ - $^5\text{I}_8$, $^5\text{I}_6$ - $^5\text{I}_7$ of Ho^{3+} ions. The materials are promising for ultra-broad band amplified spontaneous emission optical sources at near and middle infrared region.

Keywords multiband luminescence, Er^{3+} - Ho^{3+} - Nd^{3+} / Tm^{3+} , telluride glass

1 Introduction

Presently, near- and mid-infrared lasers operating in the wavelength region around 1.5–3.0 μm are attracting interest of researcher in fields of optical communication, medical surgery, eye-safe laser radar, remote sensing, atmosphere pollution monitoring, and so on. New materials, which are used to produce these lasers, are being searched. The glass, crystal and rare-earth ions should be considered because the hosts for dopants are cost-effective and easy to produce in large scale, and rare-earth materials are abundant in natural source. Rare-earth doped fluoride, fluoro-phosphate glasses [1–3], telluride and fluoro-telluride glasses [4–11], chalcogenide glasses [12,13] and other hosts [14–18] as well as fiber devices [19–23] for mid-infrared luminescence have been reported, and nearly all the literatures concentrated on the single emission at 2.70 μm or other wavelengths.

In our present work, we simultaneously observe the multiband emissions around 1.53, 2.10, 2.70 and 3.00 μm in Er^{3+} - Ho^{3+} - Nd^{3+} -codoped telluride glasses, and the multiband emissions around 1.53, 1.80, 3.00 μm in Er^{3+} - Ho^{3+} - Tm^{3+} -codoped telluride glasses with the excitation of 808 nm pump laser diode.

2 Fabrication and experiment

The Er^{3+} - Ho^{3+} - Nd^{3+} -codoped system has the following compositions (in mol %): 70 TeO_2 -20 ZnO -9.0 CaO -0.6 Er_2O_3 -0.1 Ho_2O_3 -0.3 Nd_2O_3 (EHN), and the Er^{3+} - Ho^{3+} - Tm^{3+} -codoped system (EHT) has the same compositions as EHN, but Nd_2O_3 is replaced by Tm_2O_3 .

The samples were prepared with conventional melting method. Before melting powders in a Si-C electric resistance furnace, the compositions were weighted and mixed. The mixtures were melted and homogenized at 800°C–900°C in an Al_2O_3 crucible in the furnace. The melting was poured onto a preheated brass mold and annealed for 3 hours around 100°C in a muffle furnace, and then cooled inside the furnace by turning the power supply off. The samples were cut and polished to form a plate shape with the size 10 mm \times 10 mm \times 1 mm for emission spectra measurements.

The near- and mid-infrared (mid-IR) emission spectra of the samples were measured with excitation of an 808 nm laser diode at room temperature by a Traix 320 type spectrometer (Jobin-Yvon Co., France) with resolution of 1 nm and detectable wavelength range 1.0–3.0 μm .

3 Results and discussion

The near- and mid-IR photoluminescence spectrum of the EHN is shown in Fig. 1. As can be seen, its peaks are at 1.53, 2.10, 2.70 and 3.00 μm , which correspond to the

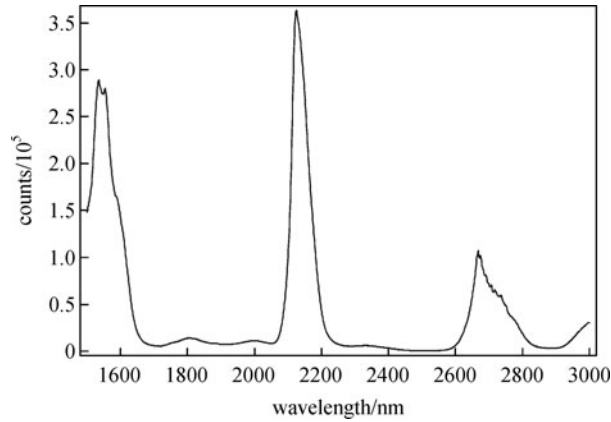


Fig. 1 Multiband emissions around 1.53, 2.10, 2.70 and 3.00 μm in Er³⁺-Ho³⁺-Nd³⁺-codoped (EHN) telluride glasses upon excitation of a conventional 808 nm laser diode

transitions Er³⁺: $^4I_{13/2} \rightarrow ^4I_{15/2}$, Ho³⁺: $^5I_7 \rightarrow ^5I_8$, Er³⁺: $^4I_{11/2} \rightarrow ^4I_{13/2}$ and Ho³⁺: $^5I_6 \rightarrow ^5I_7$, respectively. The photoluminescence spectrum of the EHT is shown in Fig. 2, its peaks are at 1.53, 1.80, 3.00 μm , which correspond to the transitions Er³⁺: $^4I_{13/2} \rightarrow ^4I_{15/2}$, Tm³⁺: $^4F_4 \rightarrow ^4H_6$ and Ho³⁺: $^5I_6 \rightarrow ^5I_7$, respectively.

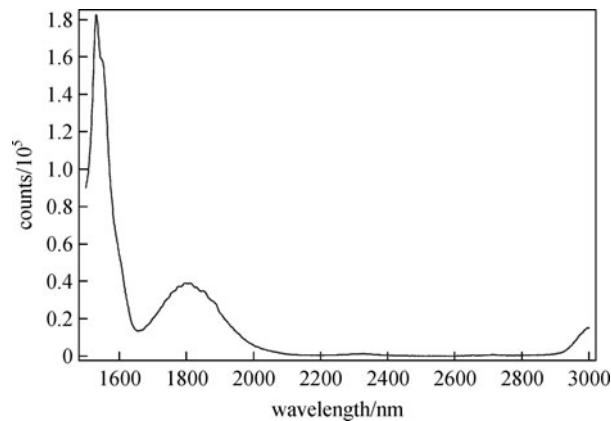


Fig. 2 Multiband emissions around 1.53, 1.80, 3.00 μm in Er³⁺-Ho³⁺-Tm³⁺-codoped (EHT) telluride glasses upon excitation of a conventional 808 nm laser diode

The processes of electron transitions and energy transfers in EHN and EHT systems are schematically shown in Figs. 3 and 4.

In EHN system, under the excitation of 808 nm laser diode, the electrons at the ground states $^4I_{9/2}$ of Nd³⁺, $^4I_{15/2}$ of Er³⁺ and 5I_8 of Ho³⁺ are excited to the states $^4F_{3/2}$ of Nd³⁺ and $^4I_{9/2}$ of Er³⁺ and 5I_5 of Ho³⁺, respectively. Due to smaller energy level difference between $^4F_{3/2}$ of Nd³⁺ and $^4I_{9/2}$ of Er³⁺, $^4F_{3/2}$ of Nd³⁺ and 5I_5 of Ho³⁺, the energy at the excited state ($^4F_{3/2}$) of Nd³⁺ is transferred (ET1, ET2) to $^4I_{9/2}$ of Er³⁺ and 5I_5 of Ho³⁺. For Er³⁺ ions, the electrons at the excited state relax into $^4I_{11/2}$, then further transit into

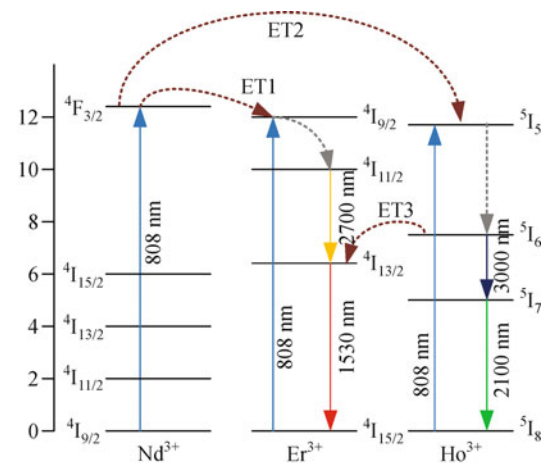


Fig. 3 Processes of electronic transitions and energy transfers in 70TeO₂-20ZnO-9.0CaO-0.6Er₂O₃-0.1Ho₂O₃-0.3Nd₂O₃ systems (EHN) upon excitation of a conventional 808 nm laser diode

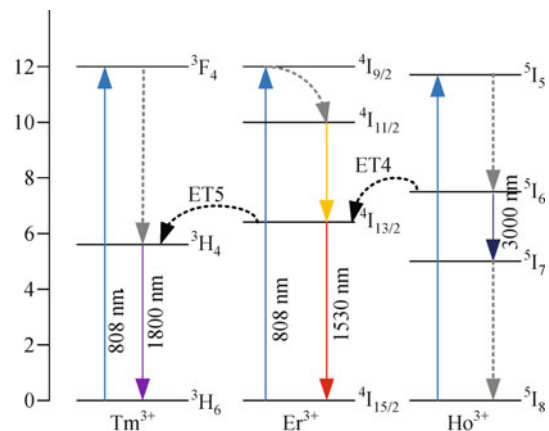


Fig. 4 Processes of electronic transitions and energy transfers in 70TeO₂-20ZnO-9.0CaO-0.6Er₂O₃-0.1Ho₂O₃-0.3Tm₂O₃ systems (EHT) upon excitation of a conventional 808 nm laser diode

$^4I_{13/2}$, $^4I_{15/2}$, with the emissions around 2.70, 1.53 μm . For Ho³⁺ ions, the electrons at the excited state relax into 5I_6 , then further transit into 5I_7 , 5I_8 , with the emissions around 3.00, 2.10 μm .

In EHT system, under the same excitation as the system above, the electrons at the ground states 3H_6 of Tm³⁺, $^4I_{15/2}$ of Er³⁺ and 5I_8 of Ho³⁺ are excited to the states 3F_4 of Tm³⁺ and $^4I_{9/2}$ of Er³⁺ and 5I_5 of Ho³⁺. Due to smaller energy level difference between 5I_6 of Ho³⁺ and $^4I_{13/2}$ of Er³⁺, $^4I_{13/2}$ of Er³⁺ and 3H_4 of Tm³⁺, a fraction of energy at 5I_5 of Ho³⁺ is transferred (ET3) to $^4I_{13/2}$ of Er³⁺, and a fraction of energy at $^4I_{13/2}$ of Er³⁺ is transferred (ET4) to 3H_4 of Tm³⁺. For Tm³⁺ ions, meanwhile, the electrons at the excited state relax into 3H_4 , then further transit into the ground state 4H_6 , with emissions around 1.80 μm . For Ho³⁺ ions, the electrons at the excited state relax into 5I_6 , then further transit into 5I_7 with emissions around 3.00 μm . For Er³⁺

ions, the electrons at the excited state relax into ${}^4I_{11/2}$, ${}^4I_{13/2}$, then transit into ${}^4I_{15/2}$, with emissions around 1.53 μm .

Additionally, it is shown in Figs. 1 and 2 that the emission bands at 2.10 and 2.70 μm in EHN system disappear in EHT system. It can be explained that in the latter, due to smaller energy level difference between 5I_6 of Ho^{3+} and ${}^4I_{13/2}$ of Er^{3+} , ${}^4I_{13/2}$ of Er^{3+} and 3H_4 of Tm^{3+} , a fraction of energy at 5I_6 of Ho^{3+} is transferred into the ${}^4I_{13/2}$ of Er^{3+} , and a fraction of whose energy is transferred into 3H_4 of Tm^{3+} . Consequently, the electron population inversions between the 5I_7 and 5I_8 of Ho^{3+} and between the ${}^4I_{11/2}$ and ${}^4I_{13/2}$ of Er^{3+} are so weak that the emissions caused by the transitions are too weak to be detected.

4 Conclusions

In conclusion, we presented Er^{3+} - Ho^{3+} - Nd^{3+} - and Er^{3+} - Ho^{3+} - Tm^{3+} -codoped telluride glasses. With excitation of a conventional 808 nm laser diode, the multiband emissions around 1.53, 1.80, 2.10, 2.70 and 3.00 μm were observed. The emission bands 1.53 and 2.70 μm were assigned to the transitions of ${}^4I_{13/2}$ - ${}^4I_{15/2}$, ${}^4I_{11/2}$ - ${}^4I_{13/2}$ of Er^{3+} ions, respectively, and 1.80 μm was assigned to the transition 4F_4 - 4H_6 of Tm^{3+} ions, and the emissions at 2.10 and 3.00 μm arose from the transitions of 5I_7 - 5I_8 , 5I_6 - 5I_7 of Ho^{3+} ions. The materials will be potential in use of ultra broad band amplified spontaneous emission optical sources at near- and mid-IR region.

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