Bulk Nd³⁺-doped tellurite glass laser at 1.37 μm

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Abstract We have demonstrated, for the first time to our knowledge, lasing at 1.37 µm in a tellurite-based glass host doped with 0.5 mol.% neodymium: Nd³⁺:(0.8)TeO₂–(0.2)WO₃. The gain-switched laser could be operated with 59 µJ threshold pulse energy as well as 5.5% slope efficiency. As high as 6 µJ-pulses with a duration of 1.74 µs were obtained. The pulse repetition rate was 1 kHz. The emission cross section from the threshold analysis turned out to be 1.57×10^{-20} cm² at 1370 nm by taking into account excited-state absorption from ⁴F_{3/2} to ⁴G_{7/2} energy level. Furthermore, the ratio of excited-state absorption to the emission cross section was found out to be 0.78 by using the slope efficiency value.

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1 Introduction

There is a growing demand for coherent light sources operating in the 1.3–1.4 μ m window for biomedical imaging applications [1, 2]. In comparison with more widely used sources at 800 nm and 1 μ m, use of higher wavelength lasers significantly decreases the losses due to Rayleigh scattering and enables deeper tissue penetration. Furthermore, staying below 1.4 μ m avoids water absorption losses. As potential laser hosts, glasses offer several advantages over single crystals, including far lower cost, simpler preparation techniques, and the possibility of power scaling when drawn

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Koç University, Rumelifeneri, Sariyer, 34450 Istanbul, Turkey e-mail: asennar@ku.edu.tr into fiber. Due to these advantages, there have been many studies aimed at the development of glass lasers doped with rare-earth ions. In the particular case of neodymium (Nd³⁺)doped systems, lasing has been reported in various glass matrices including fluorides [3, 4], chalcogenides [5], aluminosilicates [6], germinates [7], and tellurites [8–11]. Among these, tellurite-based glasses have attractive properties including a wide transparency range (0.35-5.0 µm), a high refractive index, lower non-radiative decay rates than silicates, phosphates or germanates, and resistance to corrosion [12–14]. In addition, tellurite-based glasses are considered to be strong candidates as laser host materials because they have the highest emission cross section among the Nd^{3+} doped oxide glasses [12]. To date, lasing could be obtained from a bulk tellurite glass from the ${}^4F_{3/2} \rightarrow {}^4I_{11/2}$ transition of the Nd^{3+} ion [8–11] which corresponds to the wavelength of around 1064 nm. However, the lasing operation has not been achieved so far for the ${}^{4}F_{3/2} \rightarrow {}^{4}I_{13/2}$ transition, which generates radiation in the 1.34–1.37 um spectral region.

In this letter, we report for the first time to our knowledge, lasing action at 1370 nm from ${}^{4}F_{3/2} \rightarrow {}^{4}I_{13/2}$ transition of Nd³⁺ ion in a bulk tellurite glass host where the emission cross section is lower than that for the ${}^{4}F_{3/2} \rightarrow {}^{4}I_{13/2}$ transition. The glass sample had a composition of (0.8)TeO₂-(0.2)WO₃ with 0.5 mol.% Nd₂O₃. The laser could be operated in gain-switched regime at 1 kHz with as low as 59 µJ of threshold pulse energy. In addition, the slope efficiency was determined to be 5.5%. As a pump source, we used a pulsed Ti:sapphire laser at the wavelength of 805 nm. From the analysis of threshold data, we determined the emission cross section to be 1.57×10^{-20} cm² at the wavelength of 1370 nm. Lasing could not be obtained over the broad emission band, believed to be due to excited-state absorption. By using the lasing efficiency data, we further esti-



mated the excited-state absorption cross section σ_{ESA} to be 1.22×10^{-20} cm². The ratio of the excited-state and stimulated emission cross sections (0.78) is consistent with the reported value obtained for SiO₂-based Nd:glass fiber operating at 1.36 µm [15].

2 Experimental

The tellurite glass sample was prepared with the composition of Nd^{3+} : (0.8)TeO₂-(0.2)WO₃ by melting powder mixtures of Nd₂O₃ (99.9% purity), TeO₂ (99.999% purity), and WO₃ (99% purity). The batch containing 0.5 mole of Nd_2O_3 to 100 moles of (0.8)TeO₂-(0.2)WO₃ was mixed and melted in a platinum crucible at 800°C for 1 hour in an electrically heated furnace in ambient air atmosphere. Then, the glass melt was rapidly quenched in a preheated stainless steal mold at the temperature around 150°C. To release the residual thermal stresses formed in the glass during the quenching process, the sample was annealed below the glass transition temperature at 250°C. Sample surfaces were then polished to a thickness of 4.1 mm. Using the measured density of 5.82 gm/cm³, the Nd³⁺ ion concentration was determined to be 2.02×10^{20} cm⁻³. The single-pass absorption at the pump wavelength of 805 nm was 92%.

Figure 1 shows a schematic of the standard astigmatically compensated x-cavity laser setup. The cavity consisted of two curved highly reflecting mirrors with radius of curvature 5 cm (M1 and M2), a flat end high reflector (M3), and a flat output coupler with transmission of 2.5% at 1370 nm (OC). The highly reflecting mirrors have reflectivity above 99.9% in the range of 1150-1450 nm and transitivity above 90% at 1065 nm, which prevents laser operation at that wavelength. As a pump source, we used a home-made, tunable, pulsed Ti:sapphire laser at 805 nm with a pulse repetition rate of 1 kHz. The output of the pump laser was focused inside the glass sample by using a lens with a focal length of 5 cm. The pump beam waist was measured to be 27 µm with the knife-edge technique. The high reflector and output coupler arm lengths were 31 and 32 cm, respectively, giving an estimated beam waist of 21 µm near the middle of the stability range. The time-dependent fluorescence signal and pulse traces of glass and pump lasers were measured with a digital oscilloscope and a Ge PIN detector (response time of 3.5 ns). Emission spectrum and wavelength measurements were performed with a 1/2-m Czerny-Turner-type monochromator and a PbS detector coupled with a lock-in amplifier.

3 Results and discussion

The power efficiency curve of the Nd³⁺:tellurite glass laser operating at 1370 nm is shown in Fig. 2. The slope efficiency of the laser was determined to be 5.5% with respect to the incident pump pulse energy. The laser could be operated with as low as 59 µJ of incident threshold pulse energy and we obtained as high as 6 µJ of output pulse energy. To determine the passive loss of the resonator, the threshold pump power of the laser was also measured by replacing the output coupler with a flat high reflecting mirror. The corresponding threshold energy decreased to 32 µJ. By comparing the threshold pulse energy for two different output coupling levels, we determined the round trip passive loss of the cavity (L) to be 2.9%. Here, we assumed that the threshold pump energy is directly proportional to (L + T)where T is the output coupling level. Furthermore, the optimum output coupling level can be found by using the formula, $T_{\text{opt}} = L(\sqrt{r} - 1)$, where r is the ratio of maximum available pump pulse energy to the threshold pulse energy $(r = E_p/E_{\rm th})$ for zero output coupling (32 µJ). In our case, the optimum output coupler level turned out to be about



Fig. 2 Energy efficiency curve of Nd^{3+} :tellurite glass laser. The slope efficiency is around 5.5%. The transmission of the output coupler (OC) is 2.5% at 1370 nm

3.8% at the maximum available pump energy of 169-µJ. The corresponding estimated output energy for optimum output coupling becomes 6.5 µJ, which is close to experimentally obtained pulse energy with the 2.5% output coupler.

Figure 3 shows the time-dependent fluorescence decay curve for the ${}^4F_{3/2} \rightarrow {}^4I_{13/2}$ transition. From the decay data, the fluorescence lifetime was determined to be 114 µs. In our previous study about 1065 nm laser action of the same host,



Fig. 3 Fluorescence decay curves for the Nd³⁺:tellurite glass sample at 1.37 µm. The fluorescence lifetime is around 114 µs

Fig. 4 Temporal profiles of the laser pulses at a pump energy of 110 and 170 µJ

Fig. 5 Emission spectrum of

the Nd³⁺:tellurite glass sample

laser was operating at 1.37 µm

the fluorescence lifetime for the ${}^4F_{3/2} \rightarrow {}^4I_{11/2}$ transition was measured to be 142 μ s [11] which is close to what we had for the ${}^{4}F_{3/2} \rightarrow {}^{4}I_{13/2}$ transition, as expected. Furthermore, Fig. 4 shows the temporal profile of the output laser pulses at different pump energies. As can be seen, the output pulse width drops from 3.37 to 1.74 us as the pump pulse energy increases from 110 to 170 µJ. In this pumping level range, the pump pulse width remains almost constant around 116 ns. In Fig. 4, the lower peaks preceding the laser pulses were the pump laser pulses which could not be completely eliminated by the filter. The delay between the pump and laser pulses depended on the pumping level as expected and decreased from 7.4 to 5.6 µs as the incident pump energy was increased from 110 to 170 µJ.

Figure 5 shows the emission spectrum of the glass sample and laser emission. As can be seen from the spectrum, there are two peaks around 1065 and 1342 nm, which correspond to the transitions ${}^4F_{3/2} \rightarrow {}^4I_{11/2}$ and ${}^4F_{3/2} \rightarrow {}^4I_{13/2}$, respectively. From the spectrum measurements, one would expect to get lasing at the peak wavelength of 1342 nm. However, previous studies show that there is a competing excited-state absorption (ESA) at the peak emission wavelength [15–17]. As can be seen from the energy level diagram of Nd^{3+}





Fig. 6 Schematic of the partial energy level diagram for the Nd^{3+} ion, showing the laser and excited-state absorption transitions

ion (Fig. 6), there are two excited-state absorption transitions ${}^{4}F_{3/2} \rightarrow {}^{2}G_{9/2}$ and ${}^{4}F_{3/2} \rightarrow {}^{4}G_{7/2}$ which correspond to the wavelengths around 1 µm and 1.34 µm, respectively. The transition ${}^{4}F_{3/2} \rightarrow {}^{2}G_{9/2}$ does not introduce any significant loss at 1065 nm laser operation, even though it is close to the lasing wavelength [4]. However, ESA due to ${}^{4}F_{3/2} \rightarrow {}^{4}G_{7/2}$ transition introduces extra loss near the peak of the 1342-nm emission and shifts the laser wavelength to 1.37 µm. To check whether the ambient humidity has any effect on the emission wavelength, we further purged the whole resonator with pure N₂ gas and lowered the relative humidity level from 50 to 12%. No observable change was recorded in the lasing wavelength or the output power of the laser.

The slope efficiency of the laser was used to estimate the ratio of ESA cross section (σ_{ESA}) to the emission cross section (σ_L), ($f_L = \sigma_{\text{ESA}}/\sigma_L$). By assuming that the pump and laser beams are nearly matched inside the gain medium, f_L can be estimated from [18],

$$\eta = \frac{T}{(T+L)} \frac{\lambda_p}{\lambda_L} \eta_a (1 - f_L).$$
(1)

Here, η is the slope efficiency of the laser, η_a is the absorption at the pump wavelength, and λ_p and λ_L are the wavelengths of the pump and the laser beams, respectively. For our case, we estimated f_L to be 0.78, which is in very good agreement with the reported value for Nd:SiO₂ glass fiber operating at 1.36 µm (0.78) [15]. In addition, we determined the emission cross section (σ_L) at 1.37 µm from the threshold pulse energy value by taking ESA into account. Under the approximation that the pump pulse width is shorter than

the fluorescence lifetime, the emission cross section can be determined by using the formula [18],

$$\sigma_L = \frac{\pi h \nu_p (w_L^2 + w_p^2) (T+L)}{4\eta_a E_{\text{th}} (1 - f_L)},$$
(2)

where $E_{\rm th}$ is the threshold pulse energy, v_p is pump photon frequency, h is the Planck constant, w_L and w_p are the laser and pump spot sizes inside the gain medium, respectively. Since the spotsize function varies within the gain medium due to diffractive spreading, we approximated w_L and w_p in (2) by their root-mean-squared (rms) values, which turned out to $w_L = 24 \ \mu\text{m}$ and $w_p = 35 \ \mu\text{m}$, respectively. If w(z) is the spot size function of one of the beams, the corresponding rms value $w_{\rm rms}$ is calculated by using

$$w_{\rm rms} = \sqrt{\frac{1}{L_0} \int_0^{L_0} w^2(z) \, dz}.$$
(3)

Here L_0 is the length of the glass. With $E_{\rm th} = 32 \,\mu\text{J}$, T = 0 (for the high reflector), L = 0.029, $\eta_a = 0.92$, the emission cross section, σ_L , was determined to be $1.57 \times 10^{-20} \,\text{cm}^2$ at 1.37 μm . Moreover, by using the calculated f_L value, the excited-state absorption cross section was estimated to be $1.22 \times 10^{-20} \,\text{cm}^2$ at the wavelength of 1.37 μm .

4 Conclusions

In conclusion, we have reported, for the first time to our knowledge, lasing action at 1370 nm in a bulk tellurite glass from the energy transition ${}^{4}F_{3/2} \rightarrow {}^{4}I_{13/2}$ of the Nd³⁺ ion. The laser was operated in gain-switched regime at 1 kHz with a threshold pulse energy of 59 μ J. The slope efficiency was further determined to be 5.5%. As high as 6 µJ-pulses with a duration of 1.74 µs were obtained. From the threshold analysis, the emission cross section turned out to be 1.57×10^{-20} cm² in the presence of excited-state absorption. In addition, the excited-state absorption cross section, $\sigma_{\rm ESA}$ was estimated to be 1.22×10^{-20} cm² by using the slope efficiency analysis. The ratio of excited-state to stimulated emission cross section is consistent with the value reported in the literature [15]. The performance of the Nd^{3+} doped tellurite glass laser was limited by excited-state absorption. However, the tellurite glass host is still promising when doped with other ions such as Pr^{3+} , which has optical transitions in the same spectral region and is less susceptible to excited-state absorption. Hence, it should be possible to develop efficient near infrared laser systems based on tellurite glass host.

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References

- B.E. Bouma, G.J. Tearney, I.P. Bilinsky, B. Golubovic, J.G. Fujimoto, Opt. Lett. 21, 1839–1841 (1996)
- G.J. Tearney, M.E. Brezinski, B.E. Bouma, S.A. Boppart, C. Pitris, J.F. Southern, J.G. Fujimoto, Science 276, 2037–2039 (1997)
- R.R. Petrin, M.L. Kliewer, J.T. Beasley, R.C. Powell, I.D. Aggarwal, R.C. Ginther, IEEE J. Quantum Electron. 27, 1031–1038 (1991)
- A.S.S. De Camargo, C. Jacinto, T. Catunda, L.A.O. Nunes, Appl. Phys. B 83, 565–569 (2006)
- T. Schweizer, D.W. Hewak, D.N. Payne, T. Jensen, G. Huber, Electron. Lett. 32, 666–667 (1996)
- D.F. De Sousa, L.A.O. Nunes, J.H. Rohling, M.L. Baesso, Appl. Phys. B 77, 59–63 (2003)
- J. Fernandez, I. Iparraguirre, R. Balda, J. Azkargorta, M. Voda, J.M. Fernandez-Navarro, Opt. Mater. 25, 185–191 (2004)
- J.C. Michel, D. Morin, F. Auzel, Rev. Phys. Appl. 13, 859–866 (1978)
- 9. N. Lei, B. Xu, Z.H. Jiang, Opt. Commun. 127, 263–265 (1996)

- I. Iparraguirre, J. Azkargorta, J.M. Fernandez-Navarro, M. Al-Saleh, J. Fernandez, R. Balda, J. Non-Cryst. Solids 353, 990–992 (2007)
- H. Kalaycioglu, H. Cankaya, G. Ozen, L. Ovecoglu, A. Sennaroglu, Opt. Commun. 281, 6056–6060 (2008)
- M.J. Weber, J.D. Myers, D.H. Blackburn, J. Appl. Phys. 52, 2944– 2949 (1981)
- 13. J.S. Wang, E.M. Vogel, E. Snitzer, Opt. Mater. 3, 187-203 (1994)
- B. Richards, Y. Tsang, D. Binks, J. Lousteau, A. Jha, Opt. Lett. 33, 402–404 (2008)
- F. Hakimi, H. Po, R. Tumminelli, B.C. Mccollum, L. Zenteno, N.M. Cho, E. Snitzer, Opt. Lett. 14, 1060–1061 (1989)
- M. Brierley, S. Carter, P. France, J.E. Pedersen, Electron. Lett. 26, 329–330 (1990)
- W.J. Miniscalco, L.J. Andrews, B.A. Thompson, R.S. Quimby, L.J.B. Vacha, M.G. Drexhage, Electron. Lett. 24, 28–29 (1988)
- A. Sennaroglu, C.R. Pollock, H. Nathel, J. Opt. Soc. Am. B 12, 930–937 (1995)