Recent progress of laser performance in Nd-doped yttrium lanthanum oxide transparent ceramics

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 $Nd_{0.02}Y_{1.78}La_{0.2}O_3$ transparent ceramic is fabricated by conventional ceramic sintering process. The transparent ceramic has excellent spectroscopic properties, with the absorption cross section of 1.53×10^{-20} cm² and broad full width at half maximum (FWHM) of about 8 nm at 806 nm, the emission cross section of 3.57×10^{-20} cm² at 1 078 nm, and the decay lifetime of 232 μ s. For $Nd_{0.02}Y_{1.78}La_{0.2}O_3$ ceramic laser, a maximum continuous wave (CW) laser output power of 1.38 W with a slope efficiency of 22.9% is obtained at 1 079 nm under an 808 nm diode pump.

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The cubic sesquioxide Re_2O_3 (Re=Y, Sc, and Lu) with excellent thermal conductivities (12.5-16.5 W/mK)isotropic construction are excellent laser materials. However, it is difficult to grow large-size high-quality Re_2O_3 single crystals because of their high melting temperature of $2430 \sim 2480$ °C. Now it is easy to fabricate high quality Re_2O_3 ceramics at a relatively low temperature, which is about 700 °C lower than their melting temperature by a nanocrystalline and ceramics sintering technology. The first laser oscillation in Nd-doped Re_2O_3 ceramics materials was reported in $1973^{[1]}$, the laser material is Nd^{3+} doped Y_2O_3 -Th O_2 (NDY) solid-solution transparent ceramics. Since 2000, the optical quality of Re_2O_3 transparent ceramic has improved greatly and is good enough to obtain laser oscillations. In 2001, 1.5 at.-% $Nd:Y_2O_3$ transparent ceramics fabricated by nanocrystalline and vacuum sintering technology realized laser output under LD pumping, a maximum output power of 160 mW with a slope efficiency of 32% was obtained^[2]. Next year, 0.15 at.-% Nd:Lu₂O₃ transparent ceramics fabricated by the same technology realized laser output under LD pumping, output power of only 10 mW with a slope efficiency of 13.6% was obtained^[3]. This is the first report of Nd:Lu₂O₃ ceramic laser. In 2009, another 1 at.-% Nd:Y₂O₃ ceramics fabricated by a new technology including laser synthesis of nanopowders and the magnetic pulsed compaction got laser oscillation, the highest output power near 100 mW with a slope efficiency of 15% under LD pumping has been obtained^[4]. Almost at the same time, 1.5 at.-% Nd:Y_{1.8}La_{0.2}O₃ transparent ceramic fabricated by solid-state reaction method realized laser output, a maximum output power of 62 mW was obtained under diode pumping^[5].

In this letter, $Nd_{0.02}Y_{1.78}La_{0.2}O_3$ transparent ceramic was fabricated by a solid-state reaction method, the spectroscopic properties and laser performance were investigated. $Y_{2-2x}La_{2x}O_3$, a solid solution of Y_2O_3 and La_2O_3 , has lower sintering temperature than Y_2O_3 ceramics, and what is more, it has cubic glass-like structure, which make it possible to have broad absorption bandwidth.

High purity Y_2O_3 , La_2O_3 and Nd_2O_3 nanopowders were used as starting materials (All purchased from Rare-Chem. Hi-Tech Co., Ltd, China). The composition of $Nd_{0.02}Y_{1.78}La_{0.2}O_3$ nanopowders was mixed with ZrO_2 balls in anhydrous alcohol for $5\sim10$ h, calcined at $1\,100\sim1\,200^\circ$ for $5\sim10$ h. Finally the disks with 15 mm in diameter and $6\sim8$ mm in thickness were isostatically pressed at 200 MPa and sintered at 1 650~1 700° for $40\sim50$ h in H₂ atmosphere.

The sintered specimen was double polished for spectral analysis and laser performance. The absorption spectrum was measured with a spectrophotometer using Xe light as pump source (Model V-570, JASCO, Japan). The fluorescence spectra excited by 808 nm LD was measured with a fluorescence spectrum analyzer (Fluorolog-3, Jobin Yvon Spex, France). All the spectroscopic analyses were made at room temperature.

The ceramic employed in continuous wave (CW) laser experiment has a dimension of $5\times3\times3$ (mm). Both faces of the ceramic are finely polished and antireflection coated at 808 nm (R < 1.8%) and from 1 040 to 1 090 nm (R < 0.2%). A plano-concave cavity was used. The pump source is a commercial fiber-coupled diode laser (LIMO GmbH, Germany) emitting at 808 nm with a core diameter of 200 μ m (NA=0.22) which was focused into the crystal with a refocus module with a magnification of 1.

Figure 1 shows the transmittance of Nd_{0.02}Y_{1.78}La_{0.2}O₃ transparent ceramic. The in-line transmittance around wavelength of 1 100 nm is over 80%. The room-temperature absorption spectrum is presented in Fig. 2. The strongest absorption peak is located on 580 nm and the absorption band from 780 to 850 nm is suited for LD pumping. The effective bandwidth ($\Delta \lambda_{\rm eff}$) is used to calculate the full-width at half-maximum (FWHM) due

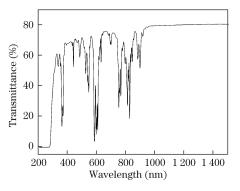


Fig. 1. Transmittance of $\rm Nd_{0.02}Y_{1.78}La_{0.2}O_3$ transparent ceramic.

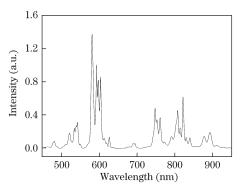


Fig. 2. Room-temperature absorption spectrum of $Nd_{0.02}Y_{1.78}La_{0.2}O_3$ ceramic.

to the unsymmetry of absorption peaks.

$$\Delta \lambda_{\rm eff} = \int \frac{I(\lambda)}{I_{\rm p}} \mathrm{d}\lambda,\tag{1}$$

where $I(\lambda)$ is the absorption intensity, and I_p is the intensity at the absorption peaks. The calculated FWHM of Nd_{0.02}Y_{1.78}La_{0.2}O₃ ceramic at 806 nm is about 8 nm, which is two times that of Nd:Y₂O₃ ceramic $(4 \text{ nm})^{[5]}$. Since Y_{1.8}La_{0.2}O₃ transparent ceramic has a disorder glasslike structure, the absorption spectrum is broaden imhomgeneously. With broad absorption bands, Nd_{0.02}Y_{1.78}La_{0.2}O₃ ceramic is lower dependent on temperature control of diode pumping and is advantageous to be a kind of material for miniaturization laser devices application.

Based on the absorption spectrum, the absorption cross section $\sigma_{\rm abs}$ of Nd³⁺ at wavelength λ can be calculated as

$$\sigma_{\rm abs} = \frac{2.303 \lg(I_0/I)}{L \cdot N},\tag{2}$$

where $\lg(I_0/I)$ is the optical density, N is the number of active ions in unit volume, and L is the thickness of specimen. The absorption cross section $\sigma_{\rm abs}$ at wavelengths of 806 and 820 nm have been calculated as 1.53×10^{-20} and 1.96×10^{-20} cm², respectively.

Figure 3 shows the emission spectrum, and the strongest peak is at 1 078 nm, corresponding to the ${}^{4}F_{3/2} \rightarrow {}^{4}I_{11/2}$ transition of Nd³⁺, with the FWHM of 5 nm, which is wider than those of Nd:Y₂O₃ (2.6 nm)^[6]. With broader absorption and emission bandwidths

 $\rm Nd_{0.02}Y_{1.78}La_{0.2}O_3$ can be a very promising laser material that is favourable to achieve the miniaturized LD pumping apparatus, widely tunable and mode-locked pulse laser.

The decay lifetime $(\tau_{\rm s})$ at 1 078-nm wavelength is 232 μ s. Based on the Judd-Ofelt theory, the calculated radiation lifetime of the Nd_{0.02}Y_{1.78}La_{0.2}O₃ ceramic is 477 μ s. The stimulated cross sections $\sigma_{\rm em}$ of the ceramic is determined by

$$\sigma_{\rm em}(\lambda) = \frac{1}{8\pi n^2 c} \frac{1}{\tau_{\rm rad}} \frac{\lambda^5 I(\lambda)}{\int \lambda I(\lambda) d\lambda},$$
 (3)

where the $I(\lambda)$ is the intensity of emission spectrum, n is the refractive index of material, and c is the light velocity.

The stimulated emission cross section calculated is 3.57×10^{-20} cm² at 1 078 nm. The values is smaller than the reported values of 7.63×10^{-20} and 6.35×10^{-20} cm² at 1 074.6- and 1 078.6-nm wavelengths in 1.5 at.-% Nd:Y₂O₃ ceramic, respectively^[6].

Figure 4 shows the output power of $Nd_{0.02}Y_{1.78}La_{0.2}O_3$ ceramic laser versus the absorbed power. The laser had a threshold of 1.0 W. With a pump power of 7.2 W, a maximum output power of 1.38 W at wavelength of 1 079 nm was obtained. The slope efficiency was measured to be 22.9%. This is, to our best knowledge, the first report of Watt-level CW laser output power in Nd-doped Re_2O_3 laser materials.

 $Nd_{0.02}Y_{1.78}La_{0.2}O_3$ transparent ceramic has broader absorption and emission bandwidth than that of $Nd:Y_2O_3$, so it is a promising high power and ultrafast pulse solid-state laser material.

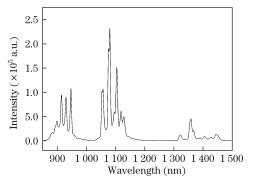


Fig. 3. Room-temperature emission spectrum of $\rm Nd_{0.02}Y_{1.78}\hdotsLa_{0.2}O_3$ ceramic.

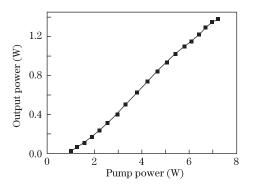


Fig. 4. The output power versus the absorbed power of $\rm Nd_{0.02}Y_{1.78}La_{0.2}O_3$ ceramic laser.

Nd_{0.02}Y_{1.78}La_{0.2}O₃ transparent ceramics has broader absorption and emission spectra due to its glass-like structure. The FWHM of absorption peak at LD pumping wavelength of 806 nm is about 8 nm and the absorption cross sections is 1.53×10^{-20} cm². The strongest emission peak was centered at 1078 nm with the stimulated emission cross section of 3.57×10^{-20} cm², FWHM of 5 nm and the decay lifetime of 232 μ s. For Nd_{0.02}Y_{1.78}La_{0.2}O₃ ceramic laser, a maximum output power of 1.38 W with a slope efficiency of 22.9% was obtained at 1 079 nm under an 808-nm diode pump.

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References

- C. Greskovich and J. P. Chernoch, J. Appl. Phys. 44, 4599(1973).
- J. Lu, J. Lu, T. Murai, K. Takaichi, T. Uematsu, K. Ueda, H. Yagi, T. Yanagitani, and A. A. Kaminskii, Jpn. J. Appl. Phys. 40, L1277(2001).
- J. Lu, K. Takaichi, T. Uematsu, A. Shirakawa, M. Musha, K. Ueda, H. Yagi, T. Yanagitani, and A. A. Kaminskii, Appl. Phys. Let. 81, 4324(2002).
- S. N. Bagayev, V. V. Osipov, M. G. Ivanov, V. I. Solomonov, V. V. Platonov, A. N. Orlov, A. V. Rasuleva, and S. M. Vatnik, Opt. Mater. **31**, 740(2009).
- Q. Yang, S. Lu, B. Zhang, H. Zhang, J. Zhou, Z. Yuan, Y. Qi, and Q. Lou, Opt. Mater. 33, 692(2010).
- G. A. Kumar, J. Lu, A. A. Kaminskii, and K. Ueda, J. Quantum. Electro. 42, 643 (2006).