

## Passive Q-switching of the Erbium:Glass Laser Using $\text{Er}^{3+}:\text{CaF}_2$

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### Abstract

The advantages of passive Q-switching with  $\text{Er}^{3+}:\text{CaF}_2$ , as well as a phenomenological model of the  $^4\text{I}_{13/2}$  fluorescence decay, are presented.

### Key Words

Rare earth doped materials, Fluorescence, Optical properties of materials, Infrared and far-infrared lasers.

### Introduction

Eyesafe Er:glass lasers (1.54  $\mu\text{m}$ ) are interesting and useful for a large number of applications such as communications, optical atmospheric studies, traffic enforcement, obstacle avoidance, and air defense [1-4]. Many applications require short 1.5  $\mu\text{m}$  pulses with high peak power which can be obtained by Q-switching the Er:glass laser. A saturable absorber Q-switch is the simplest and least expensive Q-switch option, also permitting a very compact resonator size.

Er:glass is a three-level laser where the 1.54  $\mu\text{m}$  transition,  $^4\text{I}_{13/2} \rightarrow ^4\text{I}_{15/2}$ , terminates in the ground-state. Consequently, many Er-doped materials may be used as saturable absorber Q-switches because of the significant overlap of the  $\text{Er}^{3+}$ :host absorption spectra with the Er:glass emission spectrum.

Er: $\text{CaF}_2$  possesses a broad, continuous absorption spectra nearly coincident with the emission spectrum of Er:glass (Fig. 1). The shorter effective relaxation lifetime of Er: $\text{CaF}_2$  avoids the free-running problems encountered with other Er-doped crystal Q-switches, such as  $\text{Er}^{3+}:\text{Ca}_2(\text{PO}_4)_3\text{F}$  (or Er:FAP) [5]. The Er: $\text{CaF}_2$  crystals used in these experiments, with 2.0 and 3.5% of Er, were obtained from Optovac, Inc.

Undoped  $\text{CaF}_2$  crystals have the cubic structure of fluorite with space group Fm3m [6]. The divalent cations ( $\text{Ca}^{2+}$ ) are at (0,0,0) with the fluorine ions at  $\pm(1/4,1/4,1/4)$  in an FCC lattice. The presence of  $\text{Er}^{3+}$ , which replaces the  $\text{Ca}^{2+}$  ion, distorts the otherwise cubic symmetry due to several possible charge compensation mechanisms, i.e. multiple sites with various crystalline field symmetries are possible. As a result, the optical spectra of  $\text{Er}^{3+}$  in  $\text{CaF}_2$  are complex and characterized by the presence of a large number of overlapping inhomogeneously broadened electronic and/or vibronic lines. The broad (full width at half maximum = 66 nm) absorption band of our (3.5%)Er: $\text{CaF}_2$  crystal with the maximum around 1.54  $\mu\text{m}$  is shown in Fig. 1, along with Er:glass (Kigre QE-7) fluorescence. The absorption spectrum for the (2%)Er: $\text{CaF}_2$  was the same, except the peak absorption coefficient was 2.7  $\text{cm}^{-1}$ , instead of 4.4  $\text{cm}^{-1}$ .

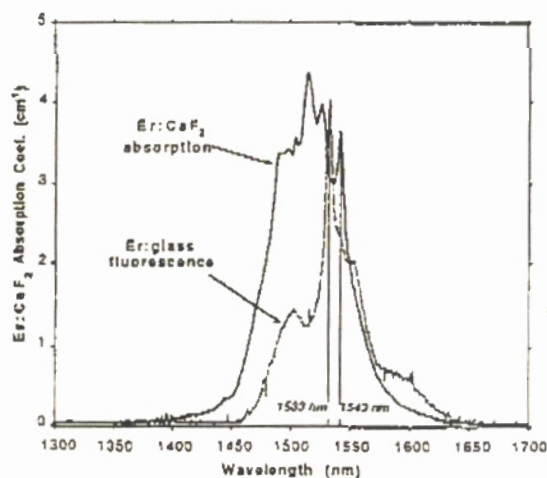


Figure 1. (3.5%)Er: $\text{CaF}_2$  absorption and Er:glass fluorescence.

## Spectroscopic Measurements

An Er:glass laser, passively Q-switched with a U:SrF<sub>2</sub> saturable absorber Q-switch [7], was used as the excitation source for the lifetime experiments. The laser output energy was 5 mJ and the pulsewidth was 138 ns, full width at half maximum (FWHM). This Q-switched laser had a wavelength of 1533 nm. The Er-crystals were placed near the focus of a +15 cm focal length lens, used to concentrate the laser beam into a 400 μm spot and produce an incident fluence of about 4 J/cm<sup>2</sup>.

The crystal fluorescence (Fig. 2) was collected using a fast (f/1.2) glass lens. The light signal was detected by a Ge photodiode (Judson, J16-series). A thin silicon crystal was placed in front of the Ge diode to block possible emissions associated with Er<sup>3+</sup> energy levels higher than <sup>4</sup>I<sub>13/2</sub>.

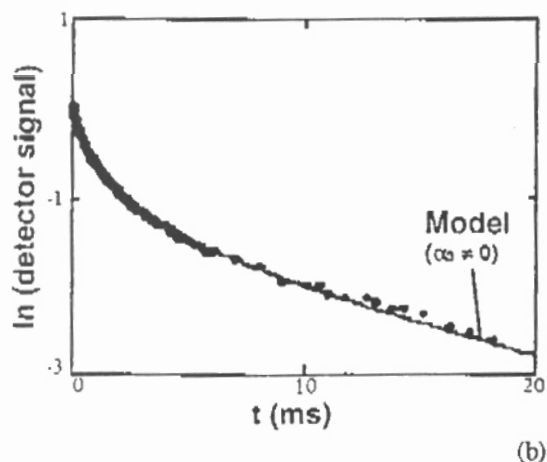
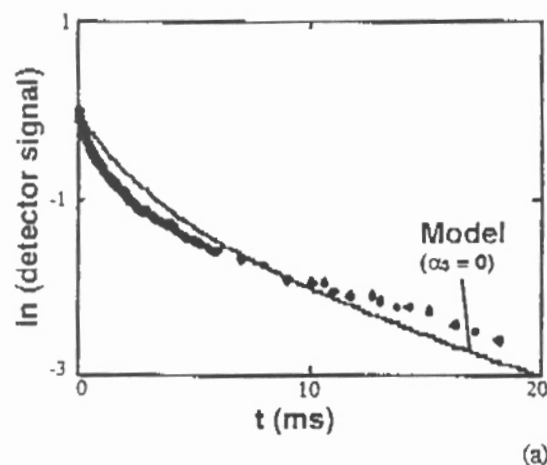


Figure 2. Semi-log plots of (3.5%)Er:CaF<sub>2</sub> fluorescence, modeled (a) without cubic term in Eq. (1), and (b) with the cubic term.

Multi-ion processes are strong in Er:CaF<sub>2</sub>, typically resulting in highly nonexponential <sup>4</sup>I<sub>13/2</sub> fluorescence decays, even with relatively low Er concentrations. Pair theory [8] proved inadequate to model the (3.5%)Er:CaF<sub>2</sub> fluorescence (Fig. 2a). Instead, we used the following rate equation for the Er<sup>3+</sup> <sup>4</sup>I<sub>13/2</sub> population density (*m*), which includes both quadratic and cubic terms:

$$\frac{dm}{dt} = -\alpha_1 m - \alpha_2 m^2 - \alpha_3 m^3 \quad (1)$$

Using the method of partial fractions, Eq. (1) can be put in the following closed transcendental form:

$$\mu \left( \frac{1+a\mu}{1+a} \right)^{\frac{a}{b-a}} \left( \frac{1+b\mu}{1+b} \right)^{\frac{b}{a-b}} = e^{-\alpha_1 t} \quad (2)$$

where  $\mu = m/m_0$ ,  $m_0$  is the initial ( $t = 0$ ) value of  $m$ , and  $a, b$  are defined by:

$$a = \frac{F_1 + [F_1^2 - 4F_2\alpha_1]^{0.5}}{2\alpha_1} \quad (3)$$

$$b = \frac{F_1 - [F_1^2 - 4F_2\alpha_1]^{0.5}}{2\alpha_1} \quad (4)$$

where  $F_1 = \alpha_2 m_0$ ,  $F_2 = \alpha_3 m_0^2$ . We numerically solved Eq. (2), and adjusted the parameters  $a, b$ , and  $\alpha_1$  to yield the best fit to the measured (3.5%)Er:CaF<sub>2</sub> fluorescence (Fig. 2b). The coefficients used in the model, are given in Table 1.

Table 1. Fluorescence model parameters.

Crystal	$m_0$	$\alpha_1$	$\alpha_2$	$\alpha_3$
(3.5%) Er:CaF <sub>2</sub>	$\sim 5 \times 10^{18}$ cm <sup>-3</sup>	67 s <sup>-1</sup>	$10^{-17}$ cm <sup>3</sup> /s	$4 \times 10^{-35}$ cm <sup>6</sup> /s

The absorption cross-section for Er:CaF<sub>2</sub> was obtained by bleaching the 2% crystal with a short pulse (14 ns, FWHM) of 1543 nm light from a Raman-shifted Nd:YAG laser, with a spectral linewidth of less than 1 nm. The measured Frantz-Nodvik saturation fluence ( $h\nu/\sigma_{eff}$ ) yielded an effective absorption cross section ( $\sigma_{eff}$ ) of  $1.43 \times 10^{-20}$  cm<sup>2</sup>.

Two Er:CaF<sub>2</sub> samples were used to perform the Q-switch experiments: a) 1.0 mm thick piece with 2.0% Er concentration, and b) a 1.1 mm thick piece with 3.5% Er concentration. Both switches were cut and polished with flat and parallel surfaces, and were used uncoated. Since the Er:CaF<sub>2</sub> absorption cross-section

value is comparable to that of glass, intracavity focusing was necessary to obtain Q-switching [5].

### Q-Switched Laser Results and Theory

The Q-switched resonator cavities, all of which possessed stable Gaussian modes, are shown in Fig. 3. The cavity shown in Fig. 3a was flat-flat and with mirror reflectivities of 100% and 88% (outcoupler), at 1533 nm. The length was 21 cm, and a +7.6 cm lens was used to focus the laser beam into the Q-switch. The output mirror in Fig. 3b had a 94% (1533 nm) reflectivity, with a 2.5 cm radius of curvature. The cavity length in this case was 17.5 cm, and a +5 cm intracavity lens was used. Fig. 3c resonator cavity had an internal 2x telescope. All the resonator internal components (except the Q-switches) were AR-coated at 1.54  $\mu\text{m}$ . The Q-switches were always positioned between the output mirror and the intracavity lens, close to the Gaussian beam waist.

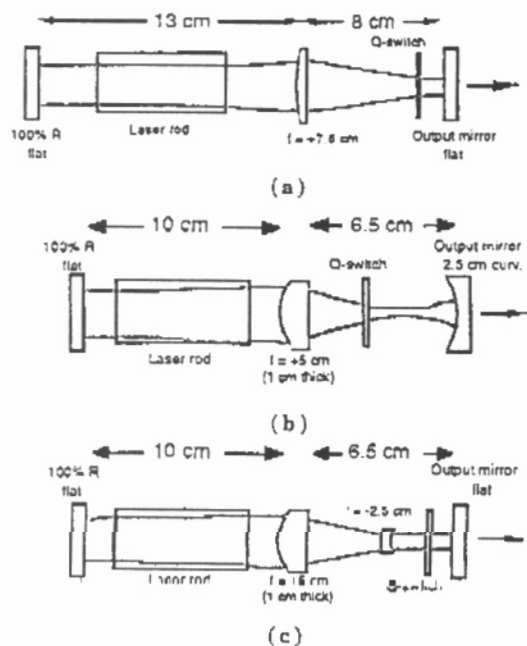


Figure 3. Er:glass resonator cavities.

The Er:CaF<sub>2</sub> Q-switch results were obtained using a QE-7S, 3 x 50 mm Er:glass rod, pumped by a Xenon flashlamp with a FWHM pulsewidth of approximately 600  $\mu\text{s}$ . The free-running laser had a threshold of about 14 J with an output slope efficiency of 0.51% for the cavity in Fig. 3b.

The results for both 2% and 3.5% Er:CaF<sub>2</sub> Q-switches are summarized in Table 2. The best results were obtained with the (2%)Er:CaF<sub>2</sub> switch, using the cavity in Fig. 3b. A typical output pulse obtained is

shown in Fig. 4, for this configuration. The Q-switched pulse was recorded using a fast InGaAs photodiode, and the Q-switched output energy was measured using a Scientech calorimeter.

A saturable absorber Q-switch rate equation model was applied to the (2%)Er:CaF<sub>2</sub> Q-switch. Using the parameters from Table 2 (losses due to Fresnel reflections from the uncoated surfaces were also included), and the measured value for the absorption cross-section ( $\sigma_{sp}$ ), the theoretical model predicted a pulsewidth of 18 ns and an output energy of 36.6 mJ. When the focusing parameter [5],  $A_s/A_a$ , was adjusted to 2.94 in the model, the simulated output pulse shape was identical to experiment (see Fig. 4). The corresponding theoretical output energy was 19.3 mJ.

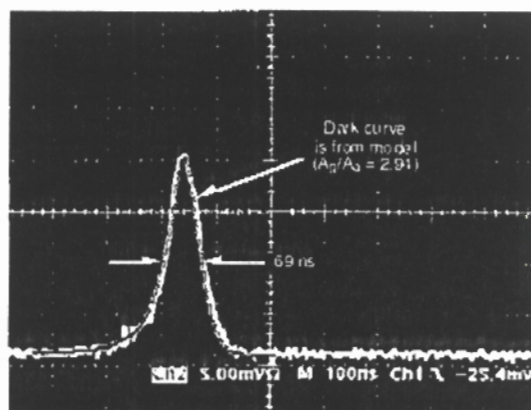


Figure 4. Q-switched pulse.

### Conclusion

In conclusion, we have demonstrated a new saturable absorber Q-switch material, Er<sup>3+</sup>:CaF<sub>2</sub>, for the 1.54  $\mu\text{m}$  Er:glass laser. We achieved reliable operation with output pulses of 11 mJ, 69 ns in a 17.5 cm laser cavity. The Er:CaF<sub>2</sub> did not exhibit any of the free-running problems previously encountered with Er:FAP, because of the rapid initial decay of its <sup>4</sup>I<sub>1,3/2</sub> level. The theoretical modeling for the Q-switched pulse shapes was in good agreement with experimental data. Further work is required to understand the physical processes involved with the fluorescence decay of the <sup>4</sup>I<sub>1,3/2</sub> level in this material.

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Table 2. Summary of Er:CaF<sub>2</sub> Q-switched laser experiments.

Q-switch: Thickness (mm)	Resonator cavity; Er:glass rod; Intracavity lens f.l.	Internal transmit. at 1533 nm	A <sub>1</sub> /A <sub>2</sub> (calc.)	Output mirror reflect. (%)	Output Energy (mJ)	Thresh. (J)	Free-run. output energy (mJ)	Q-sw. pulsewidth (ns)
(2.0%)Er: CaF <sub>2</sub> : 1.0	Fig. 3b; 3 x 50 mm QE- 7S; +5cm	0.80	12.9	94	11	46	162	69
(2.0%)Er: CaF <sub>2</sub> : 1.0	Fig. 3c; 3 x 50 mm QE- 7S; +5 & -2.5 cm	0.80	4.0	94	4	35	34	400
(3.5%)Er: CaF <sub>2</sub> : 1.1	Fig. 3a; 3 x 50 mm QE- 7S; +7.6cm	0.72	6.2	88	3.3	205	not measured	129