

Supporting Information

Cr³⁺ luminescence based ratiometric optical laser power meter

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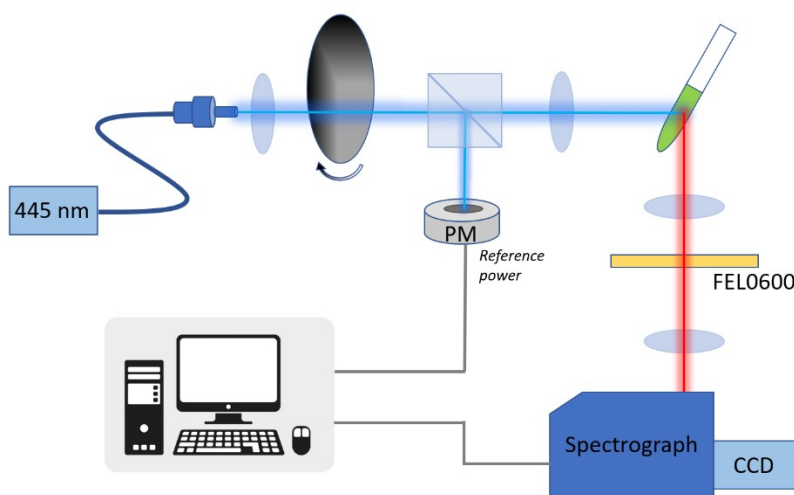


Figure S1. Schematic representation of the experimental setup used for emission spectra measurement as a function of excitation density.

Crystal field strength parameter Dq/B can be determined as follows:

$$Dq = \frac{E(4A_2 \rightarrow 4T_2)}{10} \quad (S1)$$

$$x = \frac{E(4A_2 \rightarrow 4T_1) - E(4A_2 \rightarrow 4T_2)}{Dq} \quad (\text{S2})$$

$$\frac{Dq}{B} = \frac{15(x - 8)}{(x^2 - 10x)} \quad (\text{S3})$$

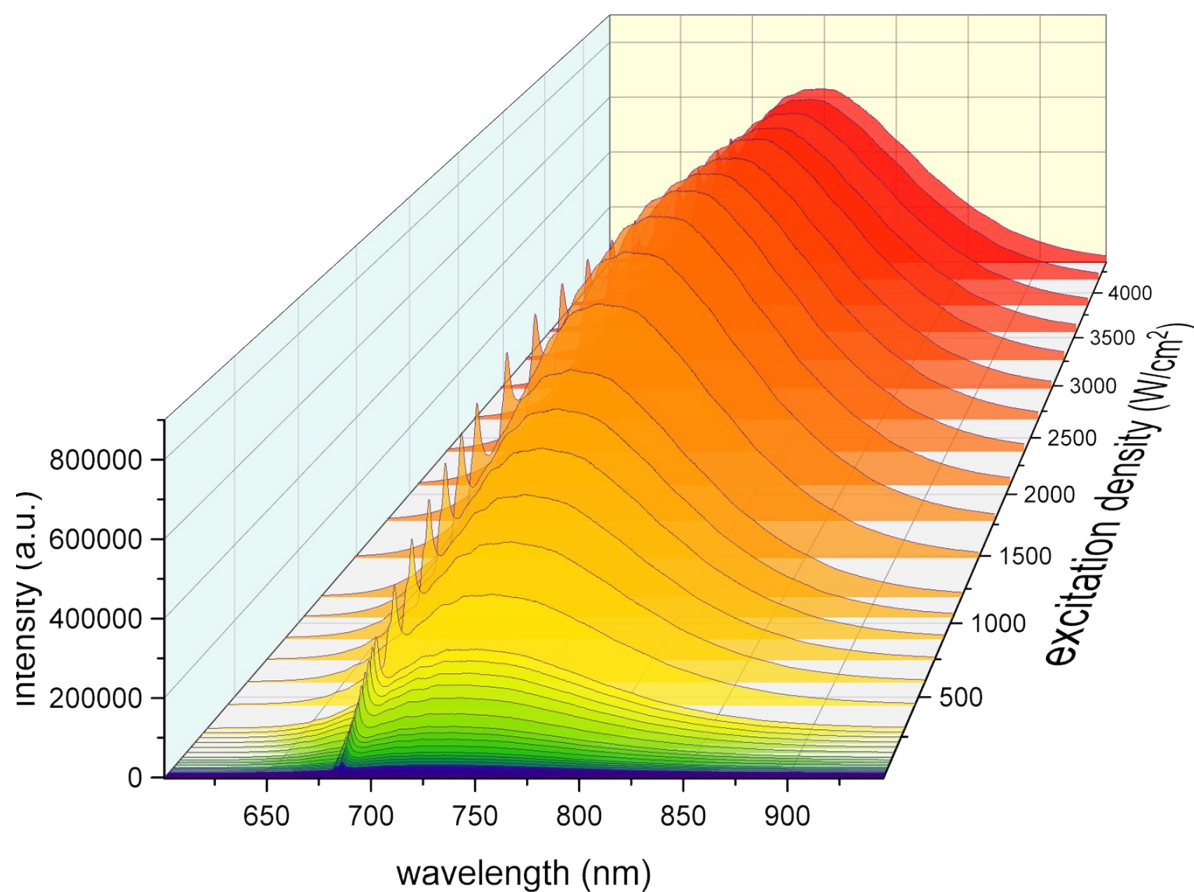


Figure S2. Emission spectra of GdAl₃(BO₃)₄:5%Cr³⁺ measured as a function of excitation density ($\lambda_{\text{exc}}=445$ nm).

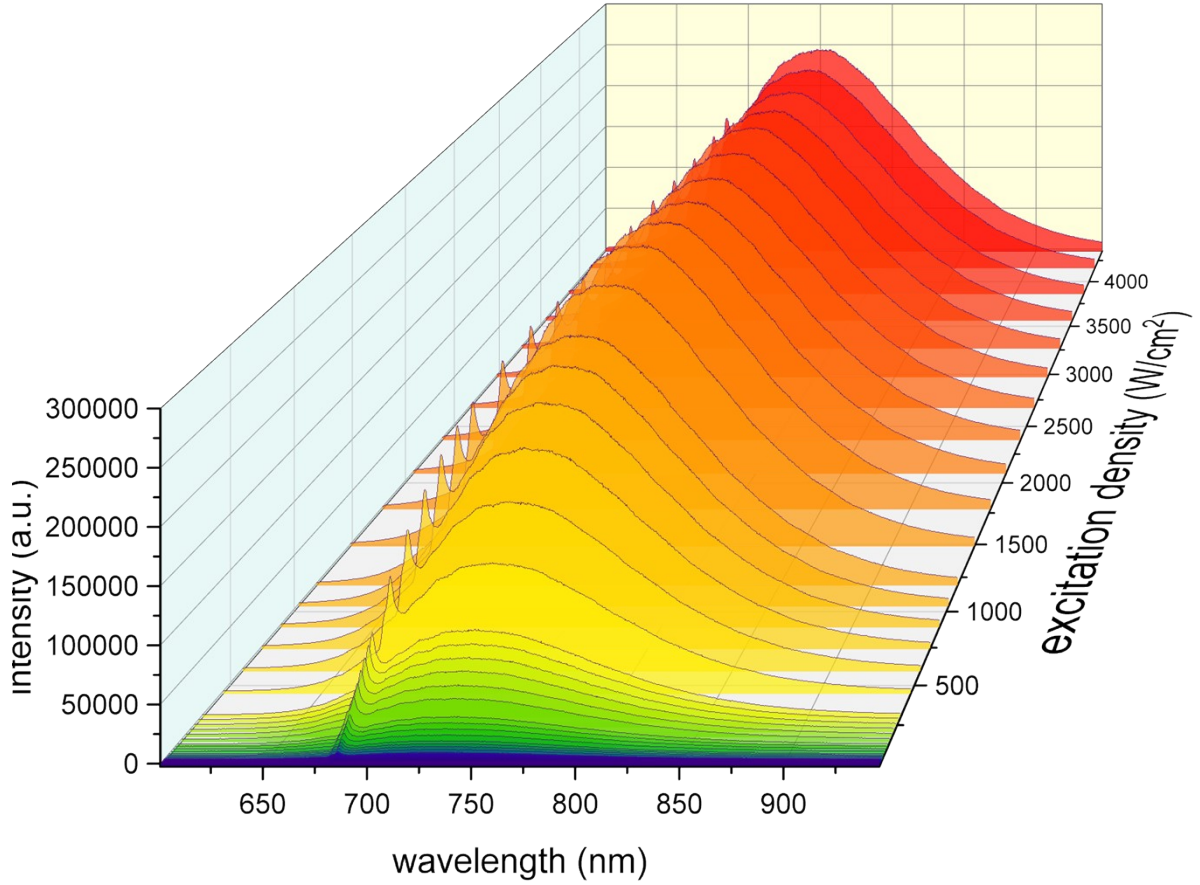


Figure S3. Emission spectra of $\text{GdAl}_3(\text{BO}_3)_4:10\%\text{Cr}^{3+}$ measured as a function of excitation density ($\lambda_{\text{exc}}=445$ nm).

The nonradiative transition probability can be determined as follows:

$$W_{nr} = 1/\tau_0 - 1/\tau_{exp} \quad (\text{S4})$$

where τ_0 and τ_{exp} represent the radiative and experimental values of lifetime.

In this particular case, we assumed that τ achieved for $0.01\%\text{Cr}^{3+}$ can be used as a $\tau_0=220 \mu\text{s}$.

The following values of τ_{avr} were achieved $\tau_{avr}=109 \mu\text{s}$, $\tau_{avr}=65 \mu\text{s}$ and $\tau_{avr}=26 \mu\text{s}$ for $1\%\text{Cr}^{3+}$, $5\%\text{Cr}^{3+}$ and $10\%\text{Cr}^{3+}$, respectively.

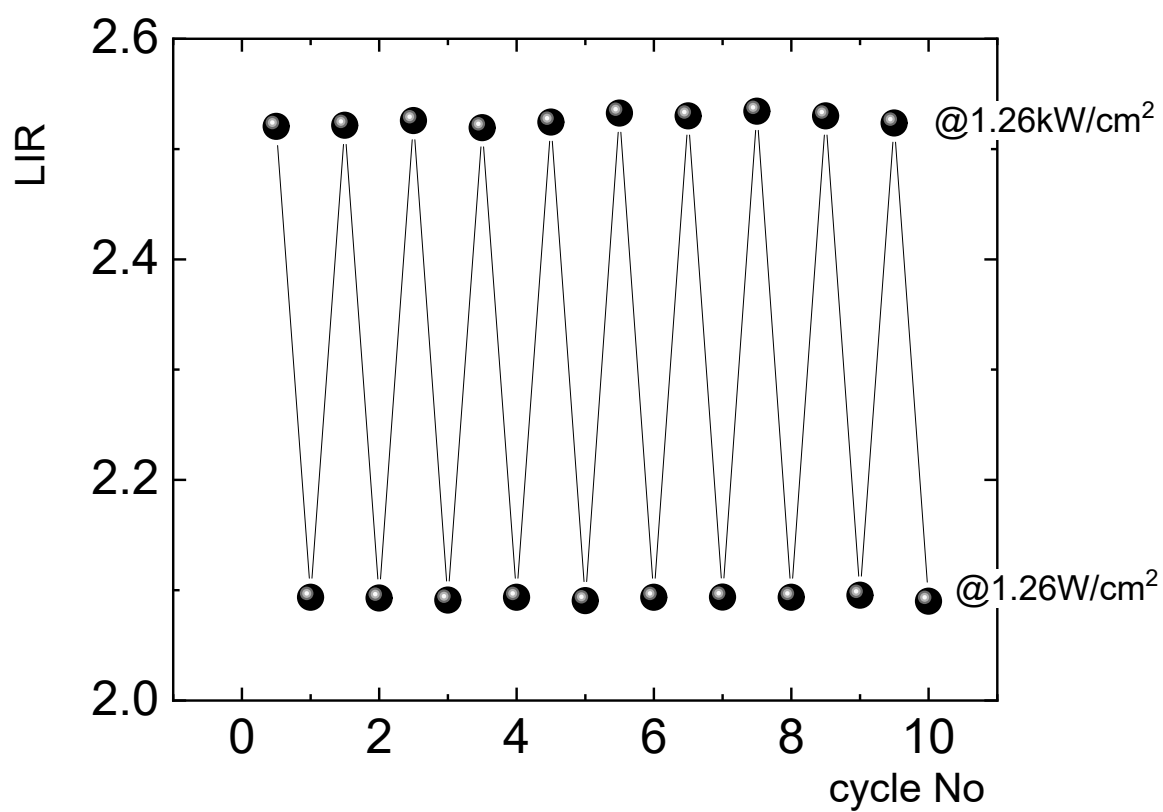


Figure S4. The measurement of LIR of $\text{GdAl}_3(\text{BO}_3)_4:1\%\text{Cr}^{3+}$ measured in a function of low and high excitation densities cycles.

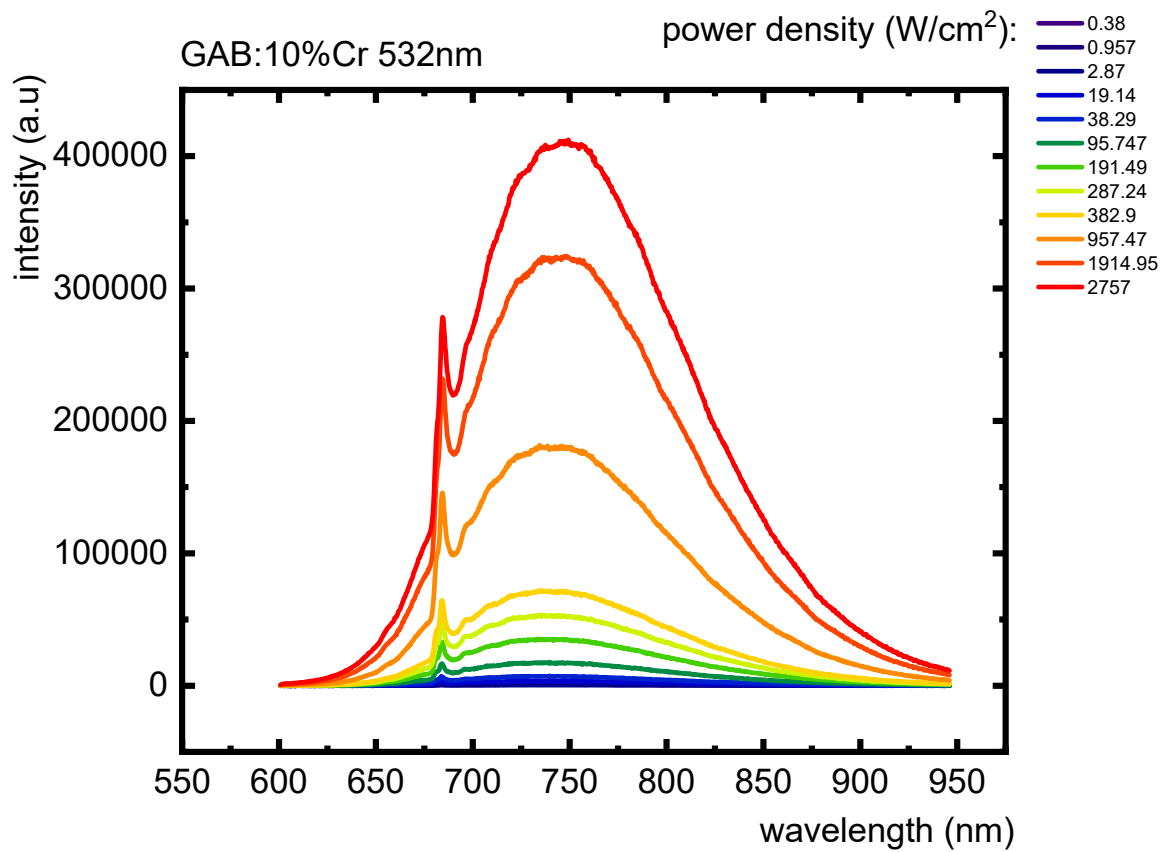


Figure S5. Emission spectra of $\text{GdAl}_3(\text{BO}_3)_4:1\%\text{Cr}^{3+}$ upon $\lambda_{\text{exc}}=532$ nm of different excitation densities.