

# Fluorescence Anisotropy Decays of 4-Cyano-N,N-dimethylaniline in Propylene Glycol Studied by Frequency-Domain Fluorometry

A. Kawski, G. Piszczek, I. Gryczyński<sup>a</sup>, and Z. Gryczyński<sup>a</sup>

Luminescence Research Group, Institute of Experimental Physics, University of Gdańsk,  
ul. Wita Stwosza 57, 80-952 Gdańsk, Poland

<sup>a</sup> Center for Luminescence Spectroscopy, Department of Biological Chemistry, University of Maryland,  
660 West Redwood Street, Baltimore MD 21201, USA

Z. Naturforsch. **53a**, 711–716 (1998); received May 13, 1998

*Dedicated to Professor J. R. Lakowicz on the occasion of his 50th birthday.*

The fluorescence lifetimes of 4-cyano-N,N-dimethylaniline (CDMA), measured in propylene glycol at 293 K using frequency-domain fluorometry, in the short emission (SE) and long emission (LE) bands are 20 ps and 1.65 ns, respectively. The higher emission anisotropies in the SE band compared to that in the LE band are due to weaker rotational depolarization of fluorescence. Emission anisotropy decays imply that the initial limiting emission anisotropy,  $r(0)$ , is the same in the SE and LE bands and amounts to 0.28. This reflects the fact that the directions of transition moments in these bands are parallel. In the case of CDMA in propylene and ethylene glycol at temperatures from 293 to 343 K, viscosity more strongly affects the relaxation to the TICT state than does the change in polarity of the solvents used.

Reprint requests to Prof. Dr. Alfons Kawski.