

**N₂ ROTATIONAL TEMPERATURE ANALYSIS BY
LASER INDUCED FLUORESCENCE FOLLOWING
RESONANT ENHANCED MULTIPHOTON
IONIZATION ***

Steven F. Adams, Jared A. Miles, Dustin M. Fisher, and
Amber L. Hensley
*Air Force Research Laboratory,
Wright-Patterson AFB, Ohio 45433 USA*

It has been demonstrated that the rotational temperature of molecular nitrogen at atmospheric pressure can be determined by direct optical probing of the N₂ (X,v=0) ground state with subsequent analysis of the rotational state distribution. A tunable probe laser was scanned over resonant enhanced multi-photon ionization transitions initiating from various N₂ (X,v=0, J) states. At atmospheric pressure, the laser photo-ionization also induced nitrogen fluorescence bands. Analysis of the relative fluorescence as a function of laser wavelength produced a calculated N₂ (X,v=0, J) rotation state distribution and the assignment of a rotational temperature.

This laser-based temperature probe may be more accurate and applicable in more environments than optical emission spectroscopy (OES), which is commonly applied as a passive remote temperature measurement technique. OES most often involves the analysis of the N₂ (C-B) emission bands, or the second positive system, where the rotational temperature is typically determined from a contour analysis of the unresolved rotational structure. OES analysis can be acceptable, but requires that (i) the nitrogen gas is electronically excited so that optical emission is observable and (ii) the rotational distribution of the N₂ (C) state is a Boltzmann distribution and it has the same distribution as the N₂ (X,v=0) ground state. The rotational temperature measurement technique introduced in this work involves remote sensing with a laser probe and can be applied to a more broad variety of nitrogen systems in thermal equilibrium or non-equilibrium, such as hot gases, flames, or atmospheric plasmas.

1. S. F. Adams, C. A. DeJoseph, and J. M. Williamson, "Formation and electron-ion recombination of N₄⁺ following photoionization in near-atmospheric pressure N₂", *The Journal of Chemical Physics*, April 2009, pp. 1-9.
2. J. Y. Roncin, J. L. Subtil, and F. Launay, "The high-resolution vacuum ultraviolet emission spectrum of molecular nitrogen from 82.6 to 124.2 nm: Level energies of 10 excited singlet electronic states", *Journal of Molecular Spectroscopy*, April 1998, pp. 128-137.

* Work supported by Air Force Office of Scientific Research