THE USE OF LASER EXCITED RO-VIBRATIONAL SPECTRA OF OH RADICALS AS A PROBE OF A GAS PHASE COMPOSITION

<u>A. Nikiforov¹</u>, L. Li¹, N. Britun², R. Snyders^{2,3}, Ch. Leys¹

¹ Department of Applied Physics, Research Unit Plasma Technology, Ghent University, Jozef Plateaustraat 22, Ghent B-9000, Belgium, E-mail:Anton.nikiforov@ugent.be ² CIRMAP, Université de Mons, 20 Place du Parc, B-7000 Mons, Belgium ³ Materia Nova Research Center, Parc Initialis, B-7000 Mons, Belgium

Atmospheric pressure plasmas characterized by high concentration of radicals and low gas temperatures are found to be suitable for many applications [1]. Often the gas composition in the discharge is unknown because of the air diffusion in to the stream of the feed gas. Correspondingly, methods to probe the gas composition in the discharge zone with high spatial resolution are strongly desirable. In the present work the high resolved laser-induced fluorescence (LIF) ro-vibrtational spectroscopy is used to probe the gas composition through the excitation of OH radicals generated in the discharge region and measurement of the fluorescence signal decay time accompanied by rotational/vibrational energy transfer. Two types of discharge are investigated: glow discharge above water surface [2] and so called Ar "plasma jet" working in ambient air [3].

Ground $X^2\Pi$ state of OH radicals was excited by the tunable dye laser to the vibrational level v=1 of $A^2\Sigma^+$ state. The effect of N₂, O₂ and H₂O addition as OH quenchers on LIF signal intensity and decay time was measured. From LIF spectra it is found that at 20-40 ns after the laser excitation the intensity of the vibrational band (1,1) drastically decreases because of collisions with N₂, whereas much stronger vibrational band (0,0) starts to dominate in all the spectra. Time decay of the LIF signal at band (0,0) reveals exponential decay with time constant of 39 ns in "plasma jet" and even shorted decay in the glow discharge. The radiative decay time of OH $A^{2}\Sigma^{+}$ (v'=1, v'=0) to ground state is about 0.6x10⁻⁵ s and hence experimentally observed much shorter decay time is related to collisional quenching of OH by N₂, O₂, H₂O. The high resolution LIF rotational spectra of OH ($A^2\Sigma^+$ -X² Π) in the plasma at 0 ns, 50 ns and 150 ns after the laser pulse is used in order to investigate the rotational energy transfer processes. Based on experimental results the model of LIF ro-vibrational transfer has been developed in order to calculate mole fractions of N₂, O₂ and H₂O in the plasmas with high spatial resolution. The method is used to determine the gas composition in the "plasma jet" working in ambient air with a spatial resolution of better than 1 mm and to calculate the water vapor in the center of glow discharge above water where as high as 10% of H₂O was detected which is explained by non-equilibrium evaporation and sputtering of water surface.

The authors thank the financial supports of the China Scholarship Council (CSC), Belgium government research support FWO (project 1.5.005.13N), one of the authors N. Britun is a postdoc researcher of the FNRS Belgium.

References

- [1] E. Stoels, I.E. Kieft, R. E.J. Sladek, L. van den Bedem, E.P. van der Laan, M. Steinbuch Plasma Sources Sci. Technol. **15**, 169 (2006).
- [2] A. Nikiforov, L. Li, Q. Xiong, X. Lu, C.Leys. J. of Phys. D Applied Physics. 45(12), 125201, (2012).
- [3] L. Li, A.Yu. Nikiforov, Q. Xiong, N. Britun, R. Snyders, X.P. Lu and C. Leys Physics of plasmas. 20(9), 093502 (2013).