

Characterization of CO₂ splitting in atmospheric nanosecond pulsed microwave plasma with a highly time resolved OES

Sergey Soldatov¹, Guido Link¹, Lucas Silberer¹, Clara Marie Schmedt³, John Jelonnek^{1,2}, Roland Dittmeyer³, Alexander Navarrete³

¹Institute for Pulsed Power and Microwave Technology (IHM), Karlsruhe Institute of Technology (KIT), Hermann-von-Helmholtz-Platz 1, 76344, Eggenstein-Leopoldshafen, Germany.

²Institute of Microwaves and Electronics (IHE). Karlsruhe Institute of Technology (KIT), Kaiserstraße 12, 76131, Karlsruhe, Germany.

³Institute for Micro Process Engineering (IMVT), Karlsruhe Institute of Technology (KIT), Hermann-von-Helmholtz-Platz 1, 76344, Eggenstein-Leopoldshafen, Germany

CO₂ conversion to carbon monoxide and oxygen is considered more efficient in non-equilibrium plasmas as compared with thermal ones [1]. To achieve a non-equilibrium state in CO₂ plasma, the microwave pulsing is one of promising approaches. It is based on the interruption the energy supply before the VT relaxation starts to deteriorate the population of vibrational states. At atmospheric pressure, however, the VT relaxation time reduces significantly as compared with vacuum conditions [2] and to sustain a non-equilibrium, the nanosecond pulse times are necessary. Systematically investigations on the conversion rate and efficiency in atmospheric plasmas sustained with nanosecond microwave pulses are performed in KIT since 2018 [3]. To understand the phenomena taking place at a pulse time scale, we examined the atmospheric plasma by ultrafast optical emission spectroscopy (OES). It is enabled with the synchronizing the ICCD camera gate and microwave pulse by means of microwave detector and delay generator. The sketch of experiment and example of spectra recorded within pulse time of 2500 ns in atmospheric microwave plasma are presented in Figure 1. To interpret molecular emission bands in terms of temperature, the recorded spectra are fitted with synthetic spectra by utilizing the software MassiveOES [4]. It enables to resolve the rotational and vibrational temperatures of CO and C₂ species within a period of microwave power modulation.

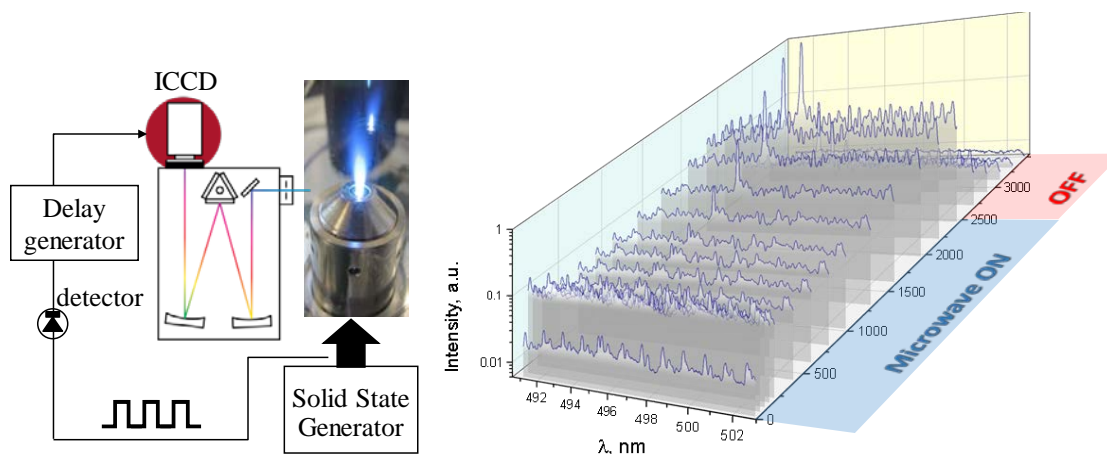


Figure 1: Sketch of time resolved OES system (left). Example of spectral characterization of CO₂ plasma emission for 2500 ns pulse time and 1 bar pressure (right).

References:

- [1] Fridman, A. Plasma Chemistry; Cambridge University Press: New York, USA, 2008.
- [2] Vermeiren, V.; Bogaerts, A. *The Journal of Physical Chemistry C* **2019**, *123*, 17650
- [3] Soldatov, S.; Navarrete, A.; Jelonnek, J.; Link, G.; Schmedt, C.; Dittmeyer, R.; 24th International Symposium on Plasma Chemistry (ISPC24 2019), Naples, Italy, 9.–14. Juni 2019: 2019
- [4] Voráč, J.; Synek, P.; Potočnáková, L.; Hnilica, J.; Kudrle, V. *Plasma Sources Science and Technology* **2017**, *26*, 025010