

# Ozone recovery in the presence of CO

A.P. Torbin, A.A. Pershin, V.N. Azyazov

Samara University. 443086, Samara, Moskovskoye shosse, 34.

E-mail: anchizh93@gmail.com, torbinalex@gmail.com, azyazov@rambler.ru

The temporal profiles of ozone number densities after pulsed UV laser photolysis in a gas mixture O<sub>2</sub>-O<sub>3</sub>-Ar-CO obtained by time-resolved absorption spectroscopy was presented. The experimental results demonstrated the dominance of the stabilization channel over the reactive one for the reaction of O<sub>3</sub>(v) with CO. The rate constants for the processes O<sub>3</sub>(v) + CO → O<sub>3</sub> + CO was obtained to be  $(1.5 \pm 0.2) \times 10^{-13}$  cm<sup>3</sup>/s using kinetics modeling of experimental data.

There are some experimental data showing the high reactive activity of vibrationally excited ozone O<sub>3</sub>(v) [1–4] which, in the presence of O atoms and O<sub>2</sub> molecule excess (fuel–air mixtures, upper atmosphere layers, oxygen-containing plasma, and others), are efficiently generated in the recombination process O + O<sub>2</sub> + M ↔ O<sub>3</sub>(v) + M, (1), where M is the third body.

The reaction of vibrationally excited ozone with carbon monoxide at moderate temperatures can proceed in two possible channels: reaction channel O<sub>3</sub>(v) + CO → CO<sub>2</sub> + O<sub>2</sub>, (2),

stabilization channel O<sub>3</sub>(v) + CO → O<sub>3</sub> + CO. (3)

In this work, we present the time dependences of the ozone concentration in the region after photolysis as a function of the carbon monoxide content in the O<sub>2</sub>-O<sub>3</sub>-Ar-CO initial mixture, measured using pulsed laser equipment and time-resolved emission spectroscopy.

Figure 2 shows the typical time dependences of the ozone concentration after laser photolysis of mixture O<sub>2</sub>-O<sub>3</sub>-Ar-CO at a wavelength of 266 nm for specific pulse energy E = 70 mJ/cm<sup>2</sup>, total gas pressure P<sub>tot</sub> = 720 Torr, oxygen pressure P<sub>O<sub>2</sub></sub> = 180 Torr, gas mixture temperature T = 300 K, and initial ozone pressure P<sub>O<sub>3</sub></sub> = 0.85 Torr.

Kinetic modeling of processes in the photolysis cell during experiments was performed. The best agreement with experimental data was achieved at the process (3) rate constant of  $(1.5 \pm 0.2) 10^{-13}$  cm<sup>3</sup>/s.

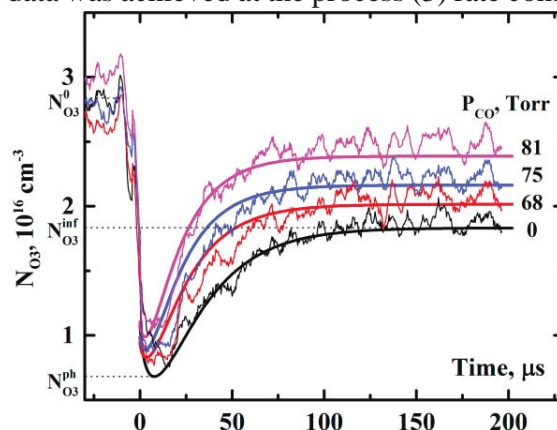


Figure 1 – Time profiles of the O<sub>3</sub> concentration at E = 70 mJ/cm<sup>2</sup>, P<sub>tot</sub> = 720 Torr, P<sub>O<sub>2</sub></sub> = 180 Torr, T = 300 K, and various CO pressures. Smooth curves are time profiles calculated by kinetic modeling.

## References

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