

O₂ dissociation in plasma and problem of O₂ cross sections set

J.P. Booth¹, O. Guaitella¹, A. Chatterjee¹, S. Zyryanov², D. Lopaev², D. Voloshin² and T. Rakhimova²

¹ *Laboratoire de Physique des Plasmas, CNRS, Ecole Polytechnique, UPMC Univ Paris 06, Univ Paris-Sud*

² *Skobeltsyn Institute of Nuclear Physics, Lomonosov Moscow State University, Russian Federation*

DC glow discharges in pure O₂ in a Pyrex tube were studied to determine dissociation rate constant over a wide range of E/N and thereby to probe O₂ dissociation cross section close to threshold. Electric field, E, was found from probe measurements while the gas density, N, from the gas temperature derived from the O₂(b¹Σ_g⁺) → O₂(X³Σ_g⁻) emission spectrum. O atom density (as well ratio O/N ratio) was measured by HR TALIF while O/N ratio was also determined by Ar actinometry. Time-resolved actinometry of partially-modulated discharges was used to probe the O loss rate. The O₂ dissociation rate constant was determined as a function of E/N, and compared to calculations from different O₂ cross section sets. This comparison allowed validation of a the self-consistent cross section set for O₂.

1. Introduction

Chemical activity of oxygen plasma is mainly provided by odd oxygen (O atoms and ozone). O₂ dissociation by electron impact is the main channel of O atoms production, therefore the O₂ dissociation cross section, together with electron energy distribution function define the chemical efficiency of oxygen plasmas. Nevertheless, the cross-section for electron impact dissociation of molecular oxygen is the subject of active debate, especially near threshold. The available experimental cross sections near the threshold predict a much lower dissociation rate than that observed. Furthermore, O₂ dissociation is an important electron energy loss channel, influencing electron transport. Cross section sets consistent with observed transport coefficients require an unrealistically high value of the O₂ dissociation cross section. This motivated a study of the dissociation rate constant to validate the near-threshold dissociation cross section and, indirectly, the self-consistent cross sections set for O₂.

2. Experiment

Time-resolved absolute density measurements are needed to probe O atoms kinetics. The measurements were carried out in DC glow discharge in pure O₂, generated in water-cooled Pyrex glass tube of 2 cm diameter and ~50 cm length. The discharge current was controlled by a large (68kΩ) ballast resistor, and could be modulated (5-15%) by shunting a smaller resistor installed between the discharge and ground. The electric field was found from probe measurements, and the gas density was calculated using the gas temperature deduced from the O₂(b¹Σ_g⁺) → O₂(X³Σ_g⁻) emission spectrum. The O atom density

and O/N ratio was measured by the HR TALIF method while the O/N ratio was also determined from actinometry using Ar atoms. Time-resolved actinometry in partially-modulated discharges was used to probe the loss rate of O atoms.

The O₂ dissociation rate constant as a function of E/N, calculated from the balance between O loss and production rates is shown in figure 1.

This research was conducted in the scope of the KaPPA International Associated Laboratory (LIA), performed within the LABEX Plas@par project, and received financial state aid managed by the Agence Nationale de la Recherche, as part of the programme "Investissements d'avenir" under the reference ANR-11-IDEX-0004-02. It was also supported by the Applied Materials University Research Partnership Program. Russian team was also supported by RFBR grant 16-52-16024.

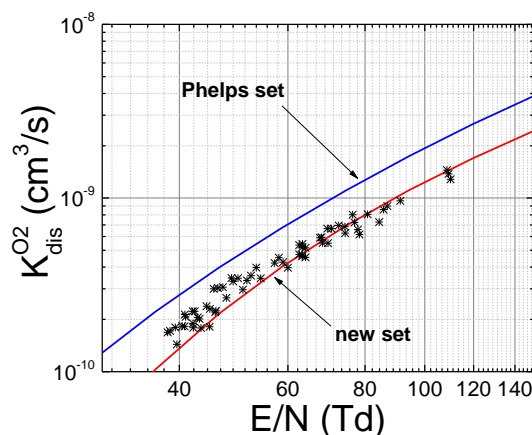


Figure 1. O₂ dissociation rate constant as a function of the reduced electric field E/N.