Monitoring Net CO₂ Dissociation Rates in the Effluent of Common Plasma Discharges with Optical Emission Spectroscopy

Andrew C. Herschberg, Nathan Bartlett, Jameson Crouse, Emily Greene, Jaime Robertson, and David N. Ruzic

Center for Plasma-Material Interactions · Department of Nuclear, Plasma and Radiological Engineering · University of Illinois at Urbana-Champaign, Urbana, IL 61801 USA

Carbon dioxide is an important gas for many plasma discharges, among these include carbon capture and chemical conversion technologies. Such plasma-based systems offer increased sustainability by reducing net carbon footprints and limiting waste from industrial processes. During plasma excitation, much of the CO₂ present in the inlet flow will be reduced into CO or other products. Therefore, the CO₂ dissociation fraction can be used as a metric for extent of reaction and to optimize process efficiency. Many methods can be employed for this purpose; in this work, an OES method of interest is compared against a standard QMS measurement. These metrologies are implemented into the exhaust gas from a flowing inductively coupled plasma containing CO₂ and N₂. The OES method employs a self-actinometry technique, comparing the line ratios from the CO Angstrom and N₂ second positive spectroscopic systems. This is implemented through a Gencoa OPTIX Remote Spectrometer for a more direct comparison to a differentially pumped SRS Residual Gas Analyzer. Overall both methods were comparable, measuring similar dissociation fractions under tested parameters, with a maximum dissociation of approximately 90%. Actinometric constants for the OES method were stable, deviating by as little as 2% across tested conditions. Implementation of the OES self-actinometric method will require calibration on system of interest, but showed to be more consistent with lower error than the QMS method.