

CO2 UV Absorption and LIF

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Shock Heated Carbon Dioxide UV Absorption

Absorption of UV light by carbon dioxide in high temperature & high pressure conditions is studied in our shock tube facility. Spectrally resolved UV absorption cross-sections between 190 and 320nm are measured in shock heated CO₂ between 880 and 3050K and H₂O between 1230 and 2860K. Absorption spectra were acquired with a kinetic spectrograph, thereby enabling comparisons with time-dependent chemical kinetic modeling of post-shock thermal decomposition and chemical reactions. CO₂ absorption cross-sections were calculated as function of temperature and wavelength.

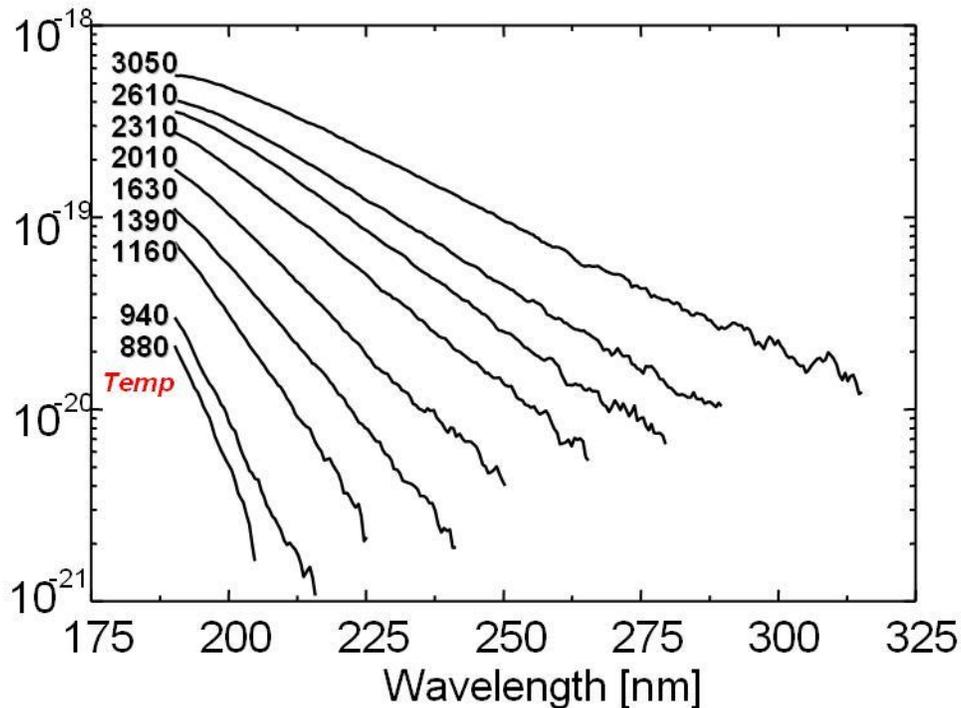


Figure 1: Absorption cross-section of CO₂ versus temperature and wavelength

Carbon Dioxide UV LIF in High-Pressure Flames

Laser-induced fluorescence (LIF) of carbon dioxide was investigated with excitation between 215 and 255nm with spectrally resolved detection in 5-40 bar premixed CH₄/O₂/Ar flames for different air/fuel ratios. Carbon dioxide LIF signal consists of a broad (200-450nm) continuum with a faint superimposed structure. Evidence of this signal of being carbon dioxide is from the fact that signal variations with excitation wavelength, equivalence ratio and flame temperature all correlate with CO₂ absorption cross-sections. The signal is linear with pressure and laser fluence within the investigated ranges.

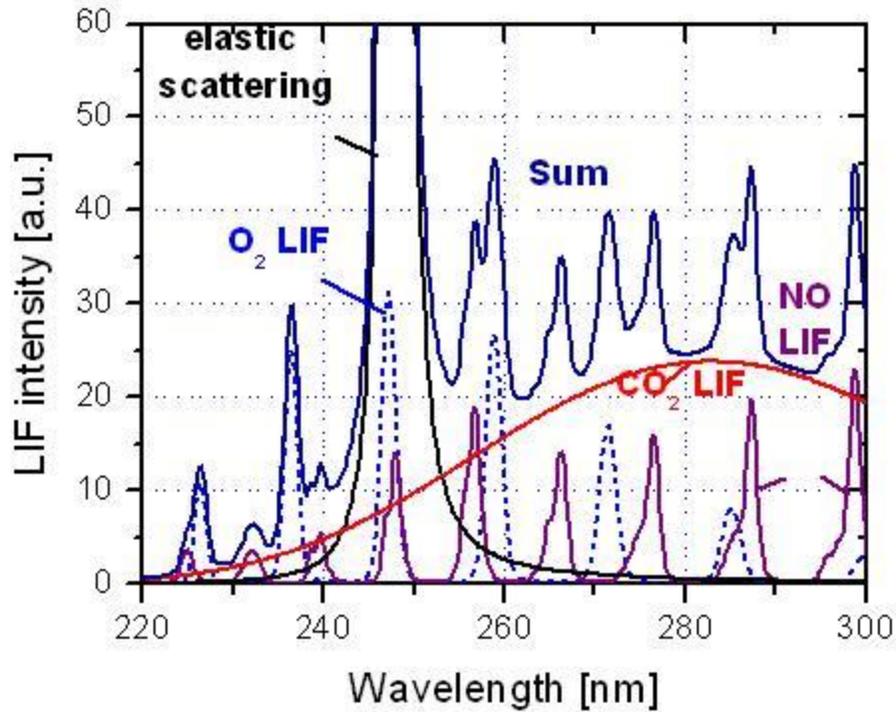


Figure 2: CO₂ with 248 nm laser excitation

PLIF Imaging with CO₂ UV LIF

Since carbon dioxide UV-LIF signal increases linearly with pressure, it offers possibilities for diagnostics in isothermal high-pressure mixing processes or for temperature field imaging in high-pressure systems with constant CO₂ concentrations. This offers a unique advantage to other signal tracers which are adversely affected by pressure due to collisional quenching of the LIF signal. UV planar laser-induced fluorescence (PLIF) images of hot carbon dioxide (CO₂) were obtained in a laminar flame (CH₄/air) at high pressure (20 bar) with excitation wavelength at 239.34 nm and 242.14 nm. Excitation wavelengths are chosen to minimize the contribution of nitric oxide and molecular oxygen LIF signals. Spectrally resolved single point measurements are used for correction of remaining oxygen LIF interference. The continuum LIF signal from electronically excited CO₂ is detected in a broad (280–400 nm) emission region. The UV PLIF of hot CO₂ has the potential for application to a wide variety of diagnostic needs in high-pressure flames, combustors, and engines.

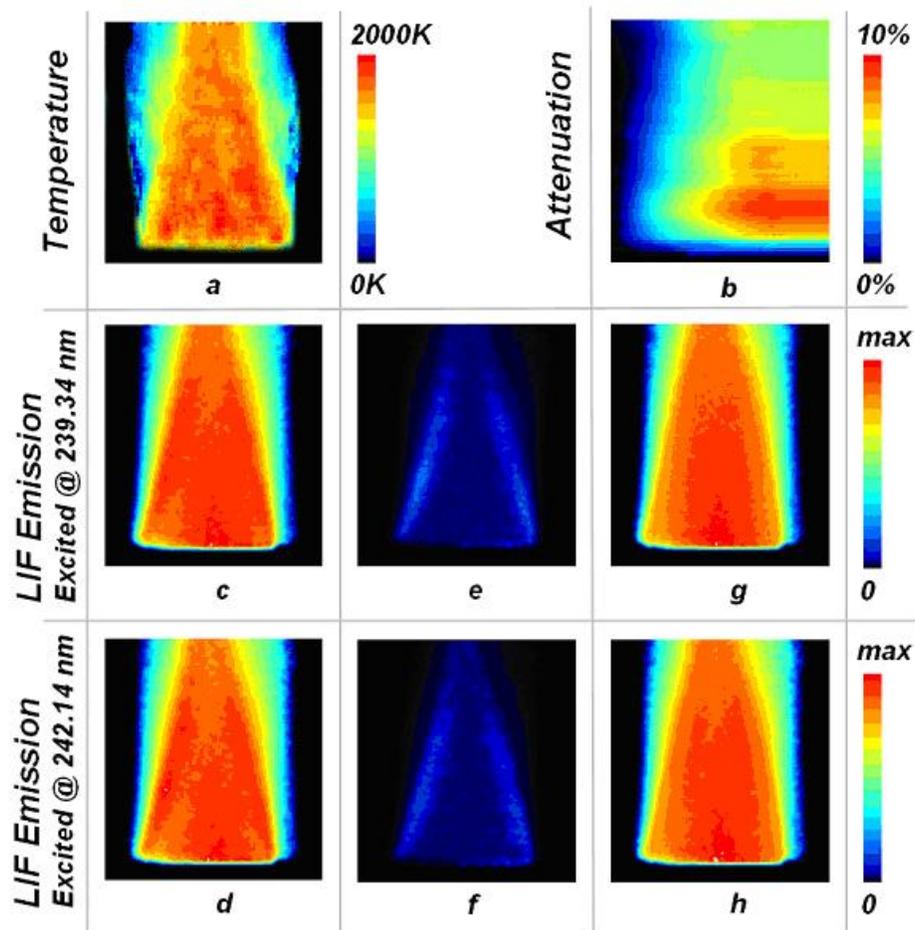


Figure 3: Results of imaging analysis of CO₂ UV-PLIF. **a** temperature field obtained from NO-LIF thermometry. **b** total attenuation due to hot CO₂ and H₂O (239.34 nm excitation); **c** and **d** CO₂-LIF and O₂-LIF combined image for 239.34 nm and 242.14 nm excitation, respectively; **e** and **f** O₂-LIF contribution for 239.34 nm and 242.14 nm excitation, respectively; **g** and **h** final CO₂-LIF image with O₂-LIF correction for 239.34 nm and 242.14 nm excitation, respectively. Images c-h share same scale.

References

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