

COHERENT ANTI-STOKES RAMAN SPECTROSCOPIC MEASUREMENT OF
AIR ENTRAINMENT IN ARGON PLASMA JETS

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ABSTRACT

The concentration and temperature of air entrained into an argon plasma jet has been measured using coherent anti-Stokes Raman spectroscopy (CARS). The flow field is characterized by a short region of well behaved laminar flow near the nozzle exit followed by an abrupt transition to turbulence. Once the transition to turbulence occurs, air is rapidly entrained into the jet core. The location of the transition region is thought to be driven by the rapid cooling of the jet and the resulting increase in Reynolds number.

INTRODUCTION

In the plasma spray fabrication of coatings the composition of the surroundings into which the plasma jet flows is known to exert considerable influence on the behavior of the jet and can alter the characteristics and quality of the coatings produced [1]. For example, while the plasma spraying of oxides is successfully conducted in oxygen containing environments, tungsten carbide-cobalt, when similarly sprayed thermally decomposes via oxidation of the carbides leading to inferior, carbon deficient coatings [2,3]. The presence of oxygen is due to entrainment of the surrounding atmosphere into the plasma flow.

The complexity of the spray process requires a more detailed understanding of chemical reactions at the particle surface, entrainment mechanisms, and the influence of the surrounding atmosphere on both the flow field and the behavior of reactive particles. As an initial step in understanding this process we have begun to investigate the entrainment of the surrounding atmosphere into plasma jets using coherent anti-Stokes Raman spectroscopy (CARS). This technique is applicable to the measurement of the concentration of any Raman active species, including O_2 , N_2 , and CO . In the work described here we have chosen N_2 as the species to be probed. The CARS spectra of N_2 is well known, and the dissociation and ionization characteristics result in non-depleted populations of neutral, molecular nitrogen at temperatures in excess of 6000 K. The initial results on an argon plasma, issuing into a stagnant atmospheric pressure air environment, are presented here.

MEASUREMENT TECHNIQUE

Non-intrusive optical diagnostic techniques such as CARS have become standard tools in high temperature flow-field research. CARS has the advantage of high conversion efficiency, a laser-like coherent signal beam for high collection efficiency, excellent fluorescence and luminosity discrimination, and high spatial and temporal resolution. The theory of CARS and its application as a combustion diagnostic tool has been detailed in several reviews [4-6]. A CARS signal, Figure 1, is generated when two laser beams at frequency ω_1 , (termed the pump beams) and one laser beam at frequency ω_2 (termed the Stokes beam) interact through the third-order nonlinear susceptibility of the medium $\chi^{(3)}$ to generate an oscillating polarization and thus coherent (laser like) radiation at frequency $\omega_3 = 2\omega_1 - \omega_2$. The intensity of the CARS signal, I_3 , is proportional to:

$$I_3 \propto I_1^2 I_2 (3\chi^{(3)})^2$$

where I_i is the intensity at frequency ω_i . This relation assumes that the required momentum phase matching is satisfied and that both beams have the same focal diameter. The third-order nonlinear susceptibility, $\chi^{(3)}$, which governs CARS can be expressed as:

$$\chi^{(3)} = \chi_n + \chi_r$$

where χ_n is the nonresonant contribution and χ_r is the Raman resonant contribution. The nonresonant susceptibility is proportional to the number density of the species present, and is generally a slowly varying

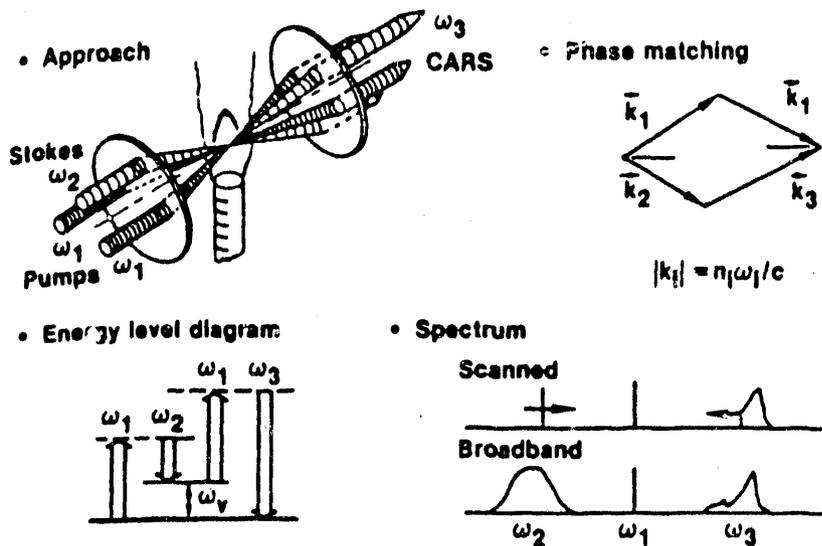


Figure 1. Coherent anti-Stokes Raman spectroscopy.

function of wavelength. The resonant contribution, far from electronic resonance, is given by:

$$\chi^{(3)} = (2c^4/h\nu^2) N \sum [\sigma \Delta(v,j) / (\omega_r - (\omega_1 - \omega_2) - iG_{v,j})]$$

where N is the number density of the species probed; $\Delta(v,j)$ is the population difference between the states involved in the Raman Transition; σ is the Raman scattering cross section; ω_r is the frequency of a Raman active rotational or vibrational state; and $G_{v,j}$ is the Raman half-width. When the frequency difference $(\omega_1 - \omega_2)$ is close to the frequency of a Raman resonance of a species, ω_r , the magnitude of the CARS radiation at ω_3 is resonantly enhanced, resulting in a signature unique to that molecular species.

CARS spectra can be acquired in two basic ways, Figure 1. The scanned approach uses a spectrally-narrow Stokes beam to generate one small segment of the CARS spectrum. The Stokes beam is then scanned across the Raman frequency range to generate a high-resolution CARS spectrum. This method requires considerable time and is limited to temporally stationary flow fields. The broadband approach employs a Stokes source that has a broad spectrum ($150-200 \text{ cm}^{-1}$). This allows the entire CARS spectrum to be generated simultaneously, on the time scale of the laser pulse. This method is preferred for most practical flow field work, where turbulence and temperature fluctuations are present and temporal fluctuation intensities are of interest. High spatial resolution is achieved by using a crossed beam geometry [6].

The CARS spectrometer consists of a frequency-doubled, Q-switched Nd:YAG laser. The second-harmonic green beam is used both as a pump beam for the dye laser which generates the Stokes beam, and as the CARS pump beam. The general beam geometry is shown in Figure 1. The focusing lenses are 100 mm focal length, resulting in an ellipsoidal measurement volume approximately $200 \mu\text{m}$ in diameter by 1.5 mm in length. In generating the spatial dependence of concentration and temperature, the plasma torch is moved relative to the stationary CARS measurement volume

Temperatures are derived from the spectral distribution of the CARS signal and concentration is derived from the strength of the signal. In the work performed here, a quick-fit technique [7], based on the full-width at half maximum of the spectral signal, has been used to derive temperatures from spectral data. The temperature determination is calibrated against a platinum, platinum-rhodium thermocouple, while the concentration measurement is calibrated by varying the concentration of air mixed into a cold argon jet. A correction to the measured concentration is derived by assuming vibrational equilibrium and calculating a Boltzmann correction factor using the measured temperature.

Alternately, spectral fitting procedures such as the CARSFT [8] computer code, or similar technique may be used to derive the same information from data acquired. The uncertainties in the concentration and temperature measurements are estimated to be 6 and 8 percent respectively.

RESULTS

The 1.27 cm. nozzle diameter plasma torch is operated at 20 volts and a current of either 250 or 500 amperes for a total power input of 5 or 10 kW. The torch is 26 and 30 percent efficient at 5 and 10 kW respectively. The argon gas flow rate is 0.59 SCMH, and the atmospheric pressure is 86 kPa. Shadowgraphs of the resulting jet at 5 and 10 kW appear in Figures 2 and 3. The flow field is characterized by a short region of well behaved laminar flow near the nozzle exit, followed by an abrupt transition to turbulence. The laminar region persists for a slightly longer distance for the higher-power case. The location of the transition region is thought to be driven by the rapid cooling of the jet and the resulting increase in Reynolds number. The estimated Reynolds number at the torch exit is approximately 100, based on the nozzle diameter. At the location of the transition to turbulence, the Reynolds number is on the order of 10,000. In the higher-power case a hotter and more viscous region maintains the Reynolds number at a lower value for a longer distance, delaying the onset of turbulence.

Contour plots of temperature and relative concentration appear in Figures 4 and 5 for the 5 and 10 kW cases respectively. The relative concentration is normalized to the number density of N_2 at ambient



Figure 3. Shadowgraph of plasma jet, 5 kW torch power.

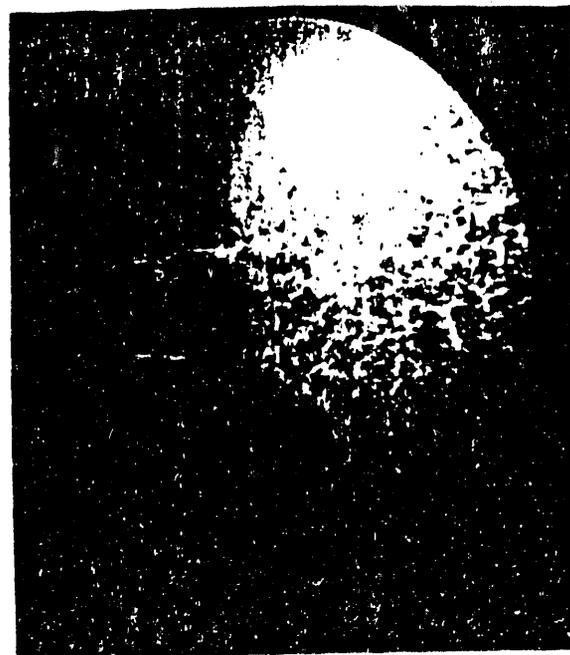


Figure 4. Shadowgraph of plasma jet, 10 kW torch power.

conditions; $N_0 \approx 1.7 \times 10^{19}$. The rapid entrainment of air into the core flow of the jet, coincident with the onset of turbulence and jet breakup, is evident in the concentration contours. This transition region is not as evident in the associated temperature fields. It is interesting to note that even though the onset of turbulence occurs earlier in the 5 kW case, a laminar, all-argon core appears to persist for approximately the same distance in both cases and initially the mixing rates are comparable. Further downstream the lower power 5kW case spreads more rapidly and entrains air at a greater rate.

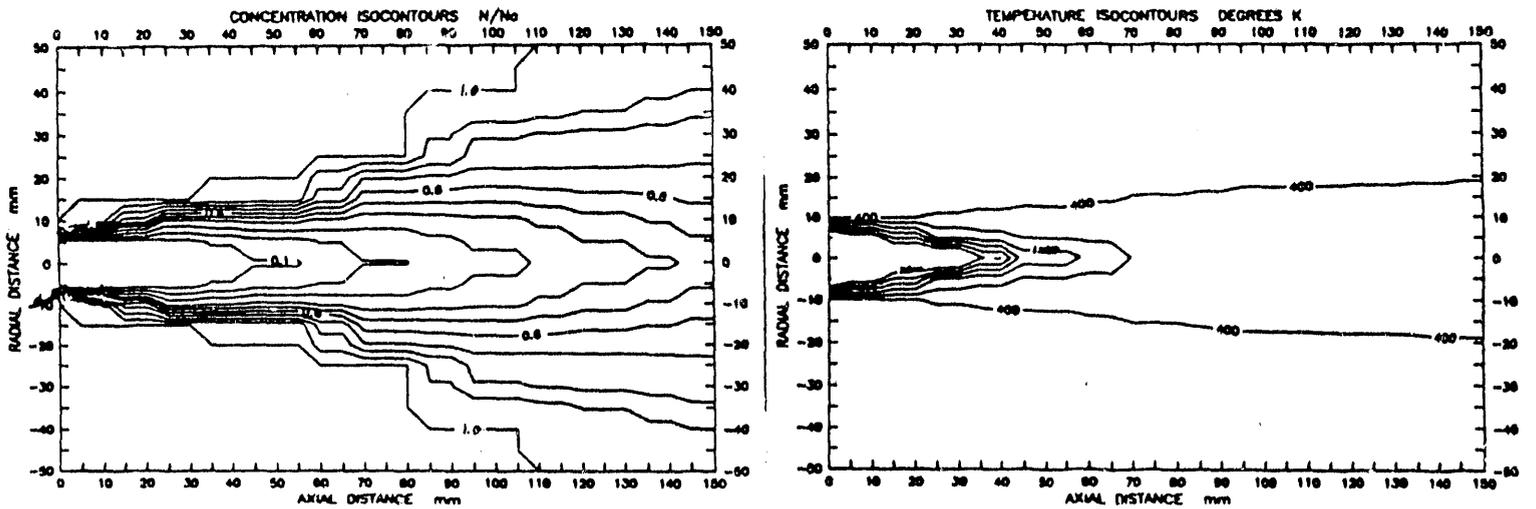


Figure 5. Contour plots of nitrogen concentration and temperature for 5 kW torch power.

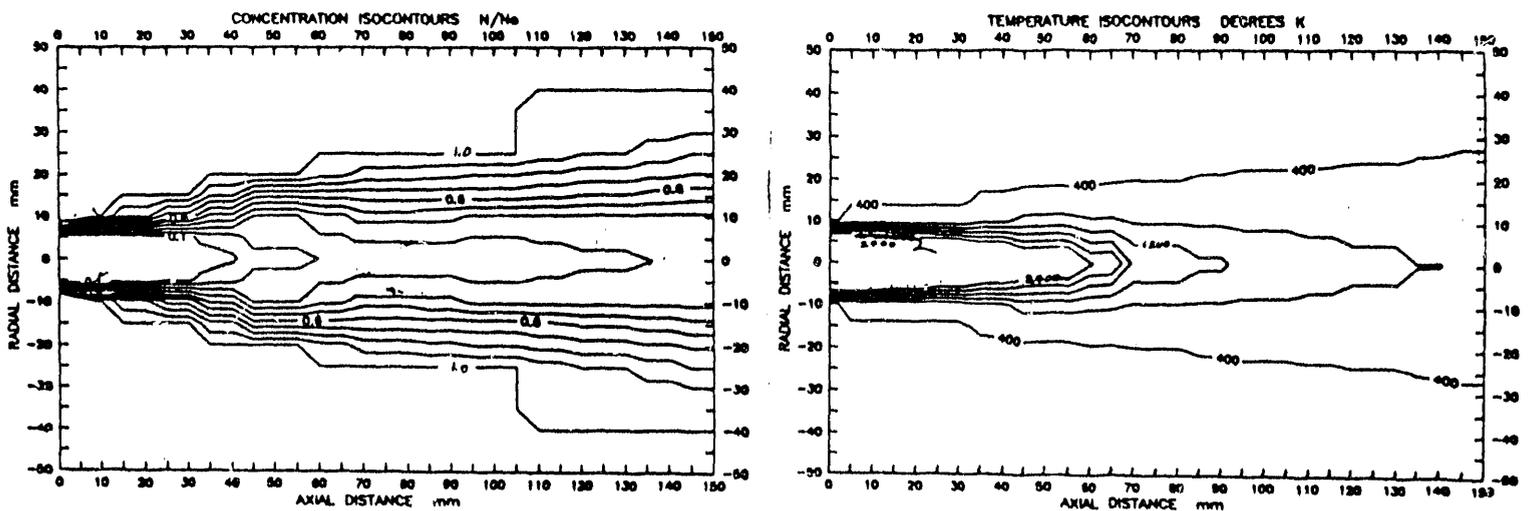


Figure 6. Contour plots of nitrogen concentration and temperature for 10 kW torch power.

CONCLUSIONS

The CARS technique has been successfully applied to the measurement of N_2 temperature and entrainment in argon plasma jets issuing into stagnant air environments. The entrainment process was found to be characterized by an initial laminar region followed by the rapid onset of turbulence and jet breakup. The location of the transition is dependent on the Reynolds number which is determined primarily by the temperature field. The spreading characteristics of the two cases studied were significantly different, as were the measured entrainment rates. The corresponding temperature fields were not as affected by the transition to turbulent flow.

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