HMOS Sensing of Oxidizing Species in an Atmospheric Plasma Jet

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ABSTRACT: A novel implementation of heated metal-oxide semi-conductor gas sensing (HMOS) is utilized to map the oxidizing species concentration in the effluent of a low-temperature plasma jet. Optical breadboard, posts and feet accessories are repurposed to position a small HMOS sensor accurately at coordinates in the plane of effluent emanating from the KinPEN MED® plasma jet, a commercial atmospheric plasma system. Operating with an argon carrier gas, oxidizing species concentrations of 240–11 ppb can extend 500 mm from the 1-mm-diameter nozzle encompassing a radius of 20 mm. For jet operation with compressed air, larger oxidizing species concentrations up to 1100 ppb were found surrounding the jet. This diagnostic approach shows potential for continuous and spatially resolved monitoring of collective concentrations of oxidizing/reducing reactive species (O₃, NOₓ, H₂O₂), which is of particular interest in characterizing treatment areas, informing dosage, and enabling control systems in emerging clinical and industrial application environments.

KEY WORDS: plasma medicine, plasma diagnostics, cold atmospheric pressure plasma, low temperature atmospheric pressure plasma jet

I. INTRODUCTION

Plasma-generated oxidizing species such as Ozone have been utilised for sterilization and antibacterial properties since the 19th century.¹ This utility has expanded recently to a wider family of plasma-generated species with promising bioactive capabilities. These species are often collectively referred to as reactive oxygen nitrogen species (RONS); they result from the higher enthalpy products of atmospheric gases (e.g., O, N, O₃, NO) precipitated via plasma generation. The field of “plasma biomedicine” has recently garnered considerable research interest,²–⁶ including potential for novel cancer therapies.⁷,⁸ Many prospective clinical treatments aim to manipulate the reduction-oxidation (REDOX) biochemistry of cells⁹ by administering RONS formed in a low-temperature plasma jet at atmospheric pressure. Gas flows are typically employed to deliver RONS to nearby target surfaces.¹⁰–¹²

Implementation of optical absorption spectroscopy diagnostics to atmospheric plasma jets¹³ has generated a fundamental understanding of key reactive species; however, detection typically requires relatively “bulky” laser and spectrometry systems. Similarly, mass spectrometry systems¹⁴ at atmospheric pressures have a large footprint incorporating several differential pumping stages and a quadrupole arm. These systems are
comparatively large in relation to the jet size (~ cm) and thus present significant practical challenges for deployment in application scenarios. The continuing miniaturization of heated metal-oxide semiconductor (HMOS) technology offers the potential for adaptable and low-cost diagnostics for future low-temperature plasma application scenarios. In situ HMOS sensors can potentially complement laboratory-based optical and particle diagnostics, enabling real-time monitoring and control in application environments.

HMOS technology along with electrochemical gas sensing is cross-sensitive to oxidant/reductant species present at the sensor. The auditing of reactive species prior to sensor deployment using species selective laboratory-based methods (e.g., optical adsorption techniques, mass spectrometry) is therefore a critical precursor and has been successfully employed in numerous fundamental studies. HMOS technology allows combined oxidizing/reducing reactive species density to be measured and could provide a practical (in terms of cost and size) monitoring and/or control solution in many future application scenarios.

HMOS gas sensors exploit the relative change in resistance of a heated thin metal-oxide layer (SnO2, WO3, In2O3, ZNO) in the presence of an oxidizing/reducing gas such as ozone. Adsorption on an n-type semi-conductor material such as SnO2 leads to a relative decrease in charge carriers (electrons) and a relative decrease in conductivity (increased resistance), which can be monitored via a DC circuitry. The range, reliability, and sensitivity of a particular thin-film oxide for gas sensing is dependent on the particular surface microstructure. The sophistication and reliability of HMOS gas sensors is an area of growing research interest driven in part by the need for long-term monitoring of air quality. This report aims to explore, in part, application of this technology to an atmospheric plasma jet system.

The NuWave MO35 sensor is a commercially available sensor based on HMOS technology that is employed here to map the effluent of a biomedical plasma jet. This sensor includes temperature-control circuitry to obtain accurate resistance measurements. The kINPen-MED is a commercially available plasma jet designed for use in medical applications such as wound healing and sterilization. This low-temperature corona source consists of a pin and plate geometry enclosed in a ceramic shower head for plasma generation. The pin electrode is biased at a MHz frequency (1 MHz) with voltage amplitudes of the order of 2–6 kV pulsed at a kHz frequency presumably to limit gas heating. In this report, the device is operated with compressed air and bottled argon (99.999% purity).

Optical absorption techniques in the ultraviolet and infrared bands have reported O3 and NOx species in close proximity to a plasma jet nozzle (~20 mm). In this report, we explore the spatial variation in these oxidizing species concentration in the far regions (25–500 mm) for this jet. RONS species (O3, NOx, O, N) share a competitive chemistry that varies with position inside and outside the plasma region. Lower-enthalpy RONS (O3, NOx, H2O2) are generally longer lived and most likely to dominant outside the immediate region of the nozzle.
II. METHODS

The plasma jet (kINPen-MED) and HMOS sensor (NuWave MO35) were laid out on a honeycomb structured optical breadboard as shown in Fig. 1. The kINPen-MED was aligned horizontally using an x–y positioning stage. The sensor head (diameter ~8 mm) is aligned with the jet nozzle (diameter ~1 mm) at fixed positions on the breadboard using the optical post and feet accessories.

This novel setup allows spatially resolved measurement of long-lived reactive species in a plasma jet plume using a small-area HMOS sensor. An oxidizing species map is formed by positioning the sensor head at the numerous bore holes on the optical breadboard (25 mm separation). Finer readings are then taken by offsetting the plasma jet position relative to the breadboard bore holes within 5–20 mm and repeating the measurement. Data are output in parts per billion (ppb) increments each second with a maximum concentration detectable of 2000 ppb. Steady-state readings during jet operation were reached after ~1 minute. Sensor data are taken as an average over ~60 seconds of steady-state data capture. The error is calculated from the standard deviation of this data.

III. RESULTS AND DISCUSSION

HMOS sensor measurements for operation of the kINPen-MED with an Argon carrier gas using a 5-s-lpm gas inlet flow are shown in Figs. 2 and 3. Background (ambient)
values of ~11 ppb were recorded. This value is typically constituent of ambient O$_3$ and NO$_2$ concentrations EPA16.

Figure 2 shows a relatively diffuse oxidizing species distribution around the plasma jet considering the nozzle diameter of 1 mm. Peak oxidizing species values of 238 ppb occur in the vicinity of the nozzle (x = 25 mm) and encompass a radius of ~5 mm. Downstream oxidizing species values of ~30 ppb, well above the background density, occur at x = 500 mm and encompass a radius of ~25 mm surrounding the nozzle. Axial symmetry is clearly observable about the central axis here. Given the circular nozzle opening the axial symmetry shown in Figs. 2 and 3 can be expanded to characterize the oxidizing species distribution within the 3-D plume (radial symmetry).

The spatial variation of oxidizing species concentration in Fig. 2 displays an (approximately) semicircular pattern with an expanding radius diffusing from the nozzle. Oxidizing species measurements along the central axis of the jet plume (y=0; see Fig. 2) are shown in Fig. 4. The oxidizing species concentration drops from 238 ± 19.8 ppb to 128 ± 2.2 ppb between x = 25 and x = 100 mm away from the nozzle. At 200 mm, oxidizing species values of 72 ± 2.8 ppb are recorded, and at 525 mm this drops to 27 ± 2 ppb. This trend displays an exponential rate of concentration decrease over this interval.

Figure 3 shows the HMOS sensor measurements for operation with air as the carrier gas (5 slpm). Air is captured from the surrounding environment via a compressor (Clarke WizAir compressor) and fed to the KINPen-MED plasma jet system. Figure 3 shows a
FIG. 3: Oxidizing species density (ppb) in the horizontal plane of plasma jet plume (25–500 mm): Air (5 slpm inlet).

FIG. 4: Oxidizing species density (ppb) along the central axis (y = 0 (see Fig. 2)): Argon (5 slpm inlet).
diffuse oxidizing species pattern similar to argon operation shown in Fig. 2; however, a significantly larger species concentration is measured here. In Fig. 5, oxidizing species density along the horizontal axis ($y = 0$) is given. At $x = 25$ mm, the oxidizing species concentration is $1088 \pm 187$ ppb which is approximately 5 times that of the equivalent argon value (Fig. 4). A similar exponential decay with increasing distance from the nozzle to argon operation is found for air. Oxidizing species values at $x = 125$ mm, 225 mm, and 525 mm were $378 \pm 26.7$, $129 \pm 3.8$, and $64 \pm 2.4$, respectively (Fig. 3).

**IV. CONCLUSION**

An exploratory application of heated metal-oxide sensor (HMOS) technology for reactive species detection in an atmospheric pressure plasma jet is reported. Spatial mapping of the oxidizing species concentration in the plume of the kINPen-MED plasma jet was carried out. A relatively diffuse oxidizing species distribution around the plasma plume for both argon and air carrier gases is found. Oxidizing species concentration decays exponentially along the central axis of the plasma plume, with concentrations exceeding background levels at over 500 mm downstream and within a 25-mm lateral radius from the device nozzle. The peak oxidizing species density recorded for argon operation ranged from 27 to 238 ppb along the central axis of the plume. Two-dimensional oxidizing species values of 240–211 ppb extend 500 mm from the nozzle, encompassing a radius of 20 mm. Operation with compressed air led to significantly larger oxidizing species values ranging from 64 to 1088 ppb along the central axis. Two-dimensional
oxidizing species values of 1100–1115 ppb extended 500 mm from the nozzle within a radius of 20 mm. Axial symmetry in the oxidizing species distribution occurred under both operating conditions. Given the circular nozzle opening design of the jet, radial symmetry can be asserted, and measurements can be extended to characterise the oxidizing species distribution within the 3-D plume.

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REFERENCES

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