Impurity intrusion in radio-frequency micro-plasma jets operated in ambient air

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Abstract
Space and time resolved concentrations of helium He (3S1) metastable atoms in an atmospheric pressure radio-frequency micro-plasma jet were measured using tunable diode laser absorption spectroscopy. Spatial profiles and lifetime measurements show significant influences of air entering the discharge from the front nozzle and of impurities originating from the gas supply system. Quenching of metastables was used to deduce quantitative concentrations of intruding impurities. The impurity profile along the jet axis was determined from optical emission spectroscopy as well as their dependence on the feed gas flow through the jet.

((Some figures in this article are in colour only in the electronic version)

1. Introduction

In recent years the research on atmospheric pressure micro-plasmas became a strong focus in plasma sciences, mostly due to their high potential for new plasma applications without the need for expensive vacuum equipment [1]. Among the large variety of micro-plasma sources, that make use of dc, pulsed dc and ac ranging from mains frequency to RF, atmospheric pressure plasma jets provide a simple design by featuring an α-mode RF discharge between two bare metallic electrodes.

Jet discharges are particularly well suited for applications in ambient atmosphere, such as biological, medical or a variety of surface treatment applications. Operating the discharge in direct contact with air, however, makes the plasma vulnerable to intrusions of nitrogen and oxygen molecules from the surrounding environment. This may reduce the operational space, induce instabilities and create unwanted reactive species. To study these influences in detail, excited species in the discharge provide an indirect measure for the amount of impurities in the discharge, since they react very sensitively to collisions with these species. In particular, metastable species carry large amounts of potential energy and are a source of ionization. In this context helium metastable species pose an appropriate indicator, since helium is a widely used carrier gas in atmospheric pressure micro-discharges. Compared with molecular gases, the excitation threshold of helium metastable atoms is high, and can be exceeded just by high energetic electrons in the tail of the EEDF. Thereby the metastable excitation process is highly sensitive to impurities in the discharge. Due to their long lifetime they collide more frequently with other particles (about 15 000 times per µs at atmospheric pressure), which makes them important for plasma chemistry processes and very sensitive to even a tiny amount of molecular gases in the discharge. In particular, nitrogen and oxygen molecules have some of the largest quenching cross sections for metastable helium atoms [2]. The metastable density in micro-discharges is several orders of magnitude lower than the density of the ground-state atoms. However, compared with most other species the density is significant and the electron collision excitation cross sections of some helium levels out of the metastable states exhibit values which are several orders of magnitude larger and have much lower thresholds than those for the ground state [3, 4]. Thus, metastables strongly contribute to the ionization.

A reliable technique for the systematic investigation of metastable species is tunable diode laser absorption spectroscopy (TDLAS), since it is not invasive and provides absolute population densities. We applied TDLAS to record the spectral profiles and the lifetime of the lowest helium metastable state, deducing absolute densities with high sensitivity and spatial resolution, revealing detailed information about the distribution of impurities in the discharge. In addition to TDLAS we applied optical emission spectroscopy (OES) to observe nitrogen emissions lines that provide direct information about the nitrogen content in the discharge.
2. The atmospheric pressure micro-plasma jet

The atmospheric pressure micro-plasma jet is a capacitively coupled, non-thermal glow-discharge at high pressures. The design concept of this discharge is based on the plasma jet introduced by Selwyn et al in 1998 [5] and advanced by Schulz-von der Gathen et al [6]. Feed gas flows between two closely spaced stainless-steel electrodes driven at 13.56 MHz radio-frequency in a parallel plate configuration (figure 1). Electrodes and plasma volume are enclosed by quartz windows, giving direct optical access to the discharge. The jet is operated in helium at various flow rates from 200 sccm to 5 slm. The electric field between the electrodes causes a breakdown in the gas and produces a plasma with electron temperatures and densities of about 1–2 eV and 10^{10} cm^{-3}, respectively [7, 8]. Atoms and molecules in the feed gas become excited, dissociated or ionized by electron impacts. Since the electrons are not in thermal equilibrium with the ions and neutrals, the gas temperature remains a few tens of K above room temperature [9]. The distance between the two windows is fixed to 1 mm while the electrode gap size is variable between 0.2 and 3 mm. In all here presented configurations the discharge operates as a typical α-mode rf glow discharge.

The whole jet setup is located in an airtight stainless-steel vessel, whose atmosphere can be defined accurately. Thus, the jet can either be operated in ambient air or in a pure helium environment, and a precise retracement of leak channels is possible.

3. Spectroscopic setup

The small dimensions of micro-discharges and their operation at atmospheric pressures are a challenge for optical diagnostics, since high sensitivity and high spatial resolution are required. For the absorption spectroscopic measurements a standard TDLAS setup was used. The absorption profile was recorded by scanning the laser frequency across the absorption line. Figure 2 shows a sketch of the experimental setup. The laser beam from the DL passed through two beam splitters. A part of the beam was guided to a Fabry–Perot interferometer (1 GHz free spectral range), a second part through a low pressure reference cell to perform the calibration of the laser frequency. The part of the beam transmitted through the first beam splitter was attenuated by neutral density filters with an optical density in the order of 3, and focused into the discharge with a beam power of less than 2 µW at 100 µm spot size, to avoid any saturation effects. After passing the discharge, the beam was guided through a set of apertures and filters to suppress the emission from the plasma by reducing the collection angle and blocking wavelengths different from the observed transitions. The transmitted beam intensity was measured either by a photodiode with on-chip transimpedance amplifier for accurate density measurements or with a biased high speed photodiode for lifetime measurements. For an effective measurement of the absorption signal across the jet axes, the discharge casing was mounted on a small movable stage featuring three electronically controlled stepping motors to adjust, with high precision, the discharge cell position in all spatial dimensions. This set up allows the positioning of the jet with an accuracy of about 5 µm and automated xz-mapping of the complete plasma volume. In the y-axis all measurements presented in this paper are line averaged.

OES measurements were carried out using a lens system to image the plasma emissions into a fibre, guiding the signal to a spectrometer with a spectral resolution of 0.75 nm. Emission profiles are averaged in y and z dimensions. The spatial resolution in x is 1 mm.

Since the absorption rate of the laser light by metastable atoms is very low, in the order of 10^{-3} after 1 mm absorption length, lock-in technique was used to measure the changes in signal intensity. Applying the lock-in technique requires the pulsing of the signal to be measured. This was realized by pulsing the RF-power coupled into the system, which consequently leads to a pulsing of the metastable density in the discharge. The pulse frequency was chosen as 4 kHz and a duty cycle of 50%.

Absolute metastable densities were derived from the transmittance I/I_0 and the Beer–Lambert law. Therefore, four signals have to be acquired:

- \( L(v) \) — Plasma and Laser on,
- \( L_0(v) \) — Plasma off, Laser on,
- \( P(v) \) — Plasma on, Laser off,
- \( B(v) \) — Plasma and Laser off (background),

to calculate the transmittance spectra and correlate them with the plasma properties by

\[
\frac{I(v)}{I_0(v)} = \frac{L(v) - P(v)}{L_0(v) - B(v)} = e^{-k(v)\ell},
\]  

where \( I(v) \) and \( I_0(v) \) are the intensities of transmitted radiation with and without the presence of absorbing species, \( k(v) \) is the absorption coefficient and \( \ell \) the path length through the absorbing medium [10]. The absorption coefficient is connected to the population density of metastable atoms by

\[
k(v) = \frac{1}{4\pi e_0} \frac{\pi (\varepsilon^2)}{cm_e} f_{ik} N_i F(v),
\]

where \( f_{ik} \) is the oscillator strength of the line, \( N_i \) the density of the lower level, and \( F(v) \) a normalized function (\( \int_0^\infty F(v) \cdot dv = 1 \)) representing the absorption lineshape. All other terms
have their usual definitions. The absolute metastable density can then be given by

\[ \int_{0}^{\infty} \ln \left( \frac{I_0(\nu)}{I(\nu)} \right) d\nu = S = \frac{e^2 f_{ikl}}{4\epsilon_0 m_e c} \cdot N_i, \]  

where \( S \) is the area under the absorption curve that provides the line-averaged density of the absorbing species.

4. Results and discussion

4.1. Metastable quenching by air

The data shown in figure 3 (upper graph) represent the metastable density in dependence on the feed gas flow. They have been taken in the centre of the discharge (2 cm from the exit nozzle) while the jet was running surrounded by ambient air. The density increases with the gas flow rate and shows a weak tendency to saturate at higher flows. This strong dependence cannot be explained by a fundamental change in the excitation mechanism, since neither the pressure nor the outer power coupling to the discharge is changing. To determine the origin of this mechanism in detail the decay rate of the density in the afterglow was measured under different conditions.

Figure 3 (bottom graph) indicates that the lifetime of the species increases also with the gas flow and, as with the density, shows a tendency to saturate for higher flows. Assuming infinite purity of the helium gas and proposing that the metastable lifetime is mainly determined by the three-body collision process:

\[ \text{He}^* + 2\text{He} \rightarrow \text{He}_2^* + \text{He}, \]  

the theoretical lifetime of \( \text{He}^* (^3S_1) \) in pure helium is expected to be

\[ \tau = (K_3 \cdot N_{\text{He}}^2)^{-1} = 5.8 \mu s, \]  

with \( K_3 = 2.5 \times 10^{-34} \text{ cm}^6 \text{ s}^{-1} \) the rate coefficient for three-body collisions and \( N_{\text{He}} = 2.68 \times 10^{19} \text{ cm}^{-3} \) the helium ground-state density [12, 13]. The loss of metastable atoms by two-body collisions as well as by diffusion can be omitted since the diffusion loss frequency is 2 orders of magnitude lower than the loss frequency by three-body collisions. Three situations have to be distinguished in this context. In the first setup (red triangles in figure 3) the airtight vessel around the jet was evacuated and filled with pure helium. Furthermore, the gas supply system, that is the stainless-steel tube connection from the gas bottle to the micro-plasma jet, was kept under a vacuum of 1 Pa for some time, to clean it from traces of water and air. As a result, for high flows of 5 slm, the theoretical lifetime for a clean helium feed gas can be reached, but the lifetime is dropping significantly for lower flows. Taking into account that the leakage rate of the gas supply system can be neglected, the metastable quenching can be attributed to tiny
amounts of water and air molecules desorbing from the surface and contaminating the feed gas in the order of a 2–40 ppm. This fraction of contamination decreases linearly with the feed gas flow.

In a second setup (pink triangles in figure 3) the airtight vessel was filled with ambient air. The measured lifetimes in this case are significantly below the theoretical value for a clean feed gas. Again the discrepancy can be attributed to the loss of metastable atoms by molecular excitation processes. Due to the low ionization thresholds these excitation processes are mostly Penning ionizations, in this case, with air introduced through the front nozzle:

\[
\begin{align*}
\text{He}^+ + \text{N}_2 & \rightarrow \text{He} + \text{N}_2^+ + e, \quad (6) \\
\text{He}^+ + \text{O}_2 & \rightarrow \text{He} + \text{O}_2^+ + e. \quad (7)
\end{align*}
\]

Assuming that the dominant impurity contribution is due to residual \( \text{N}_2 \) and \( \text{O}_2 \) molecules, the impurity level can be estimated by a simple model. Taking into account quenching collisions with nitrogen and oxygen molecules the lifetime is limited to

\[
\tau = [K_3 \cdot N_{\text{He}}^2 + K_{\text{N}_2} \cdot (N_{\text{N}_2, \text{Bottle}} + N_{\text{N}_2} (\Gamma))]^{-1}, \quad (8)
\]

\[
+ K_{\text{O}_2} \cdot [(N_{\text{O}_2, \text{Bottle}} + N_{\text{O}_2} (\Gamma))]^{-1}, \quad (9)
\]

with \( K_{\text{N}_2} = 7 \times 10^{-11} \text{ cm}^3 \text{ s}^{-1} \) and \( K_{\text{O}_2} = 2.5 \times 10^{-10} \text{ cm}^3 \text{ s}^{-1} \) the rate constants for Penning ionization processes with molecular nitrogen and oxygen and \( N_{\text{N}_2} (\Gamma) \) as well as \( N_{\text{O}_2} (\Gamma) \) the flow dependant nitrogen and oxygen densities \([13,14]\). \( N_{\text{N}_2, \text{Bottle}} \) and \( N_{\text{O}_2, \text{Bottle}} \) are the base contaminations of the feed gas, given by the purity of the helium used in the experiment. This base contamination sums up to about 2 ppm. Since we are assuming the intrusion of air into the system, the ratio between \( \text{N}_2 \) and \( \text{O}_2 \) is fixed to 3:7. Furthermore, we presume the intrusion air to vary anti-proportionally with the flow rate. The fitting parameter is the amount of air in the system and the fitting was done until the modelled data optimally reproduced the measured data. The result of the model is shown in figure 3 (blue line). It agrees well with the measured values over the whole flow rate range. We determine the level of air introduced through the nozzle to be in the order of a few dozen ppm (120 ppm for 1 slm flow, 25 ppm for 5 slm flow).

In a third setup (green triangles in figure 3) the microplasma jet as well as the gas supply system was exposed to ambient air. Repeating the flow-dependent lifetime measurements, the values show again a significant decrease. Nevertheless, the amount of air introduced into the system through the front nozzle must be the same as in the previous case. Additional metastable quenching can only be caused by traces of air and water desorbing from the surfaces of the gas supply system as well as from the metallic electrodes. This assumption is supported by mass spectrometry and OES measurements that show significant amount of water ions as well as nitrogen and oxygen emissions in the plasma and the effluent region. Matching these measurements with the previously described model (orange line) the observed quenching corresponds to a few hundred ppm of air molecules in the feed gas stream (500 ppm for 1 slm flow, 100 ppm for 5 slm flow).

For metastable density and lifetime measurements no error bars are shown, since statistical as well as systematic errors are negligible. For the presented data they are in the order of \( 10^9 \text{ cm}^{-3} \). Due to the use of the lock-in technique a sensitive measurement of low densities was possible with a high signal-to-noise ratio. A minor systematic error is caused by the unknown metastable profile in the y-axis, and the implications on the absorption length. But since the metastable production and loss processes take place on a very local scale at atmospheric pressure, the profile in the y-axis is expected to be homogeneous with the exception of a tiny volume in front of the glass windows that confines the plasma channel and acts as a sink for metastable atoms. Lifetime measurements were performed with a time resolution of less than 30 ns and revealed a perfectly exponential decay of the species. Thus, errors can be omitted in this case as well.

4.2. Longitudinal impurity profile

Figure 4 shows a 2D-map of the \( \text{He} (3S_1) \) metastable density in the discharge volume. Both, horizontal and vertical axes show the exact area between the electrodes. The map covers 2.000 reading points (40 vertical \( \times \) 50 horizontal) of the absorption signal in the plasma volume. A variety of effects determine the metastable distribution in the vertical and in the longitudinal axis. While the vertical profile reflects the RF-sheaths structure and is determined by the electron density and temperature distribution, the longitudinal profile is governed mainly by environmental factors \([11]\).

As discussed before, the impurities entering the jet through the front nozzle are quenching the metastable atoms. Since the jet is running permanently in contact with the ambient air the intrusion of nitrogen and oxygen into the plasma channel is significantly high, and decreases the metastable density especially in the first millimetres from the nozzle. This observation is supported by OES measurements showing the emission of various nitrogen lines. Figure 5 presents the longitudinal profile of the molecular nitrogen emissions at 357 nm, representing the second positive system. The measurements reveal strong nitrogen emissions in the first 10 mm from the front nozzle. The intrusion depth increases with decreasing gas velocities, ranging from about 10 to
100 ms\(^{-1}\). In particular, for small gas flows a back diffusion of nitrogen can be observed over the whole length of the jet. The back diffusion of air into the system is promoted by the geometry of the gas channel, whose shape is rectangular. In particular, in the corners of the channel the gas flow is likely not to be laminar and turbulences may occur.

OES measurements have been performed with a very limited spectral and spatial resolution, as described in the previous section. Therefore the expected errors are much larger than for TDLAS measurements. The signal-to-noise ratio strongly depends on the amount of nitrogen in the discharge. In the present case the errors for lower flows (around 0.5 slm) and therefore high nitrogen content are expected to be in the order of 10%, while for higher flows (around 5 slm) the errors are much larger, in the order of 50%.

5. Summary

It was shown that in atmospheric pressure micro-plasma jets operated in contact with ambient air the back diffusion of nitrogen and oxygen into the gas channel is a significant effect with strong impact on the energy transfer processes of the discharge. Furthermore, the inner surfaces of the gas supply system and of the jet discharge itself pose a source of impurities contaminating the feed gas. These influences and the likely consequences on the plasma chemistry have to be taken into account for any application where the use of such discharges in ambient atmosphere is envisaged.

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