For Submission to AVS 71 Charlotte, North Carolina, USA September 21-26, 2025 JON: 67-A1N3-0-5

## Spatiotemporal analysis of a submerged water plasma driven with nanosecond long voltage pulses

Michael J. Johnson<sup>1\*</sup>, David R. Boris<sup>1</sup>, Tzvetelina B. Petrova<sup>1</sup>, Mackenzie E. Meyer<sup>2</sup> and Scott G. Walton<sup>1</sup>

 <sup>1</sup> Plasma Physics Division at the U.S. Naval Research Laboratory, Washington, DC 20375
<sup>2</sup> NRC Postdoctoral Research Associate, U.S. Naval Research Laboratory, Washington, DC 20375 \*Corresponding Author Email: michael.j.johnson895.civ@us.navy.mil

## Abstract

Atmospheric pressure plasmas generate a distinct chemical and electrical environment ideal for treating water, making them attractive for applications in wound healing, chemical synthesis, nanomaterial fabrication, and water remediation. These plasmas can operate in a nonequilibrium regime when driven by short pulses of power, lasting tens to hundreds of nanoseconds, that energize electrons but are too short to significantly heat the surrounding gas. This study investigates the impact of pulse width on plasma-water interactions by applying 70-350 ns pulses to an argon plasma submerged in water. Plasma properties are analyzed using optical emission spectroscopy and electrical measurements. Results indicate that within the first 15 ns of the pulse, the plasma fully fills the gap between the electrodes. After this initial stage, the plasma expands to occupy the entire inter-electrode space for the remainder of the pulse, forming an arc-like plasma where current flow is regulated by the power supply. Essentially, pulse width determines how long the plasma remains in this high-current state. Optical emission spectroscopy revealed that argon dominates the emission immediately after plasma formation, but over time, hydrogen emission becomes more prominent as the plasma dissociates water molecules. This results in higher power consumption at longer pulse widths due to increased energy transfer to the water. Spatial emission profiles show uniform hydrogen emission across the reactor, whereas argon emission weakens near the positive electrode. Significant broadening of emission lines was observed during the pulse, with Stark broadening of hydrogen lines used to estimate electron density. Measurements indicate that a substantial electron density persists for several microseconds after the pulse, likely due to residual voltage on the electrodes during power supply neutralization. At the longest tested pulse width (350 ns), the post-pulse current lasted nearly 10 µs, highlighting not only the influence of pulse width on plasma dynamics but also the importance of other system parameters in determining plasma lifetime.

This work was partially supported by the U.S. Naval Research Laboratory Base Program.