

# Surface optimization of nanodiamonds using non-thermal plasma

**Michal Gulka<sup>1,\*</sup>, Priyadharshini Balasubramanian<sup>2</sup>, Ekaterina Shagieva<sup>3</sup>, Jakub Copak<sup>1,4</sup>, Josef Khun<sup>5</sup>, Vladimir Scholtz<sup>5</sup>, Fedor Jelezko<sup>2</sup>, Stepan Stehlik<sup>3</sup>, Petr Cigler<sup>1</sup>**

<sup>1</sup>Institute of Organic Chemistry and Biochemistry, of the CAS, Flemingovo nam. 2, Prague, Czechia

<sup>2</sup>Institute of Quantum Optics, Ulm University, Ulm 89081, Germany

<sup>3</sup>Institute of Physics of the Czech Academy of Sciences, Cukrovarnická 10, 162 00 Prague 6, Czechia

<sup>4</sup>Department of Physical and Macromolecular Chemistry, Faculty of Science, Charles University in Prague, Czechia

<sup>5</sup>Department of Physics and Measurements, University of Chemistry and Technology in Prague, Czechia

\*[gulka.michal@gmail.com](mailto:gulka.michal@gmail.com)

Possibility of local sensitive monitoring of intracellular processes using quantum sensing with nitrogen-vacancy (NV) centers in nanodiamonds (NDs), would greatly advance cell biology and would enable numerous medical applications. Although such probes have been studied extensively [1], they are still limited in sensitivity. Whereas the NV center in bulk diamond holds record room-temperature electron spin coherence time among solid-state qubits, NVs in NDs exhibit much poorer relaxation times. As suggested in [2], prolongation of the NVs relaxation time should be possible by ND surface optimization creating mixed H/O termination.

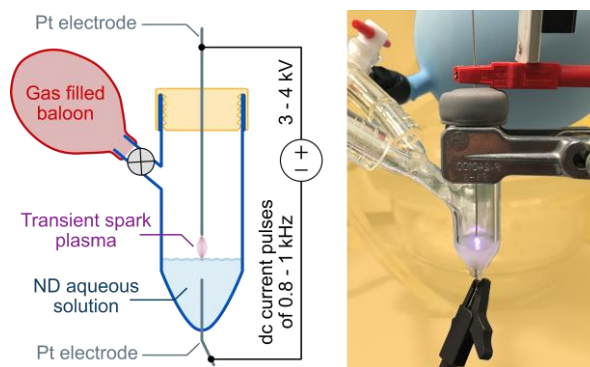


Fig. 1. Non-thermal plasma system for ND modifications

Here we employ a non-thermal plasma (NTP) [3], with the aim of prolongation of the NV electron spin relaxation times (Figure 1). We have used various nanodiamond samples (HPHT/detonation, H-/O-terminated surface) and ambient gases (air, O<sub>2</sub>, He, and H<sub>2</sub>). The NTP modification induces changes in C=O bonds, restructuring of C-H bonds, and differences in OH termination. Using hydrogen as the ambient gas results in the biggest changes in FTIR spectra. For oxidized HPHT NDs the zeta potential dropped from an initial ~ -40 mV to ~ -50 mV. Both T<sub>1</sub> and T<sub>2</sub> NV electron relaxation times improved significantly after the modification.

## Acknowledgement

This project has received funding from the European Union's Horizon 2020 research and innovation program. The work was further supported by the Czech Science Foundation project No. 23-04876S, the Ministry of Education, Youth, and Sports of the Czech Republic (project No. CZ.02.01.01/00/22\_008/0004558, co-funded by the European Union), the European Union project C-QuENS (Grant No. 101135359), the Czech Academy of Sciences – Strategy AV21 – Research program VP29 (to P.C.), TACR project Nr. TH90010001 (QuantERA – ERA-NET Cofund, project EXTRASENS), and Horizon project No 101086142 (FLORIN).

## References

- [1] T. Zhang et al., ACS Sens. **6**, 2077–2107 (2021).
- [2] M. Kaviani et al., Nano Lett. **14**, 4772–4777 (2014).
- [3] J. Khun et al., Plasma Sources Sci. Technol. **27**, 065002 (2018).