

Imaging of dark spins using nitrogen-vacancy centers in diamond

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The negatively-charged nitrogen-vacancy center (NV) belongs to the most-studied paramagnetic defects in diamond as it shows coherence times up to milliseconds at room temperature¹ and enables optical polarization and coherent control of the electron spin². To optimize the sensitivity of NV-based quantum sensors, the spin environment in diamond has to be well controlled and understood.³ However, even in engineered diamond grown by chemical vapor deposition (CVD) dark spins like substitutional nitrogen (P1) and negatively charged nitrogen-vacancy-hydrogen (NVH) defects out-number the nitrogen vacancy defect by usually at least one order of magnitude.⁴ Therefore, for minimizing magnetic noise originating from dark spins, the nitrogen-to-NV conversion during CVD growth has to be maximized while keeping the density of P1 and NVH at a minimum.⁵ To develop such optimized CVD processes usually electron paramagnetic resonance (EPR) spectroscopy is the method of choice for quantifying dark spin densities with sub-ppm precision.⁴ In nm-thick epitaxial layers, however, the total number of paramagnetic spins becomes eventually too low for state-of-the-art EPR spectrometers.⁴ As a consequence, we exploit double-electron-electron-resonance spectroscopy with NV centers as local sensors⁶ (NV-DEER) to quantify P1 and NVH concentrations in epitaxially grown (100)-diamond layers with thicknesses of about 500 nm. Furthermore, by moving from a confocal to a wide-field microscope with a CCD camera, we even image the spatial distribution of P1 and NVH spins on the sub- μm scale. Comparing then the NV-fluorescence with the density maps for P1 and NVH, we get access to the correlation of nitrogen-related spins on the micron scale in diamond which yields valuable information on the defect formation in diamond.

References

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