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Charge-Sensitive Fluorescent Nanosensors Created from Nanodiamond

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We show that fluorescent nanodiamonds (FNDs) are among the few types of nanosensors that enable direct optical reading of noncovalent molecular events. The unique sensing mechanism is based on switching between the negatively charged and neutral states of NV centers which is induced by interaction of FND surface with charged molecules.

A nitrogen-vacancy (NV) center is a point lattice defect in a diamond consisting of a substitutional nitrogen atom and a vacancy at an adjacent lattice point. This color center is a near-infrared emitter providing a broad emission peak with a maximum at ~700 nm. Unlike other fluorophores, NV centers provide exceptionally stable fluorescence and do not photobleach or blink even after extreme continuous excitation used for example in super-resolution fluorescence microscopy. NV centers thus represent a promising alternative to small-molecule labels, fluorescent proteins, and quantum dots. Fluorescent nanodiamonds (FNDs) bearing NV centers have been used in biology primarily as non-toxic fluorescent labels in cell and animal bioimaging. Moreover, electron transitions between NV quantum states can be manipulated using microwave radiation at ~2.87 GHz which results in changes in its emission intensity that can be further modulated by external magnetic field. This behavior can be measured using optically detected magnetic resonance (ODMR), which has been used for construction of FND-based sensors for detection of electric or magnetic field, for NV tracking with precision of several nanometers as well as for chemosensing even in living cells.

Here, we provide evidence that FNDs containing NV centers are among the few types of nanosensors that enable direct optical reading of noncovalent binding events based on modulation of fluorescence. Moreover, their specificity to detect changes in charge goes significantly beyond the functionality offered by conventional fluorescent nanoprobes. In contrast to techniques that require special setups (such as ODMR), fluorescence reading can be performed routinely with a confocal microscope or spectrofluorimeter.

Although many highly sensitive and selective nanoparticle-based detection systems for biomolecular detection have been developed, they often include cytotoxic reagents and work only “in tube”. In contrast, only a limited number of signaling modes is available for intracellular and in vivo applications. These include mainly resonance energy transfer (Förster, bioluminescence, and chemiluminescence resonance energy transfer), charge transfer in semiconductor nanoparticles, modulation of localized surface plasmon resonance and surface-enhanced Raman scattering. Recently, it was reported that the emission state of NV centers in FNDs can be controlled by changing the ND surface chemistry at an atomic level. NV centers in NDs exist in two electronic states: negatively charged (NV-) and neutral (NV0). The NV- state emits in the red region with zero phonon line (ZPL) at 637 nm, which is followed by broad phonon replica side band luminescence with the highest intensity around 700 nm. The NV0 state emits orange luminescence with ZPL around 575 nm, also followed by similar broad side bands. The reversible switching between the NV- and NV0 states is enabled by shifting the position of the NV0 and NV- ground state levels with respect of the Fermi level at the ND surface. Thus, when the NV- ground state level is shifted below the Fermi level, an electron is lost from the NV center and vice versa. Depending on the surface termination, complementary changes in the occupancy of the NV- and NV0 energy states occur, which consequently affect the shape of fluorescence spectra.
We hypothesized that manipulations of the ND surface that affect the NV⁻ and NV⁰ states are not limited to changes in atomic surface composition, but also include interactions in aqueous solution. We therefore designed a set of fundamental experiments to test whether surface-charge-induced modulation of fluorescence can be observed for noncovalent surface chemical processes. Next, we explored an approach for direct two-color fluorescent monitoring of molecular binding events in proximity to the ND surface, which is an essential feature for construction of future ND-based fluorescent biosensors. For the experiments we used previously described FNDs of mean diameter ~49 nm and containing on average 1.7 NVs per particle (see Supporting Information and Figure S1 for experimental details and particle characterization).

We took advantage of the high level of polymer adsorption on NDs and created a well-controlled model system for binding of macromolecules to ND surfaces. By simply incubating NDs with various polymers, we achieved the desired surface changes without the need to construct sophisticated recognition architectures. Based on the previous findings that the NV state is more populated in oxidized, negatively charged NDs, we focused on ND surface charge reversal upon binding cationic polymers, which should lead to depletion of NV⁻ occupancy and subsequent decrease in its emission intensity.

To control the surface charge density after adsorption, we chose two types of flexible linear polymers: poly(allylamine) (1) and poly(diallyldimethylammonium chloride) (2; Scheme 1). While amino groups in 1 can participate in acid-basic equilibria (average pKₐ = 9.7) and reversibly switch between positively charged and neutral states, the charge of the quaternary ammonium salts is always positive and inert to pH changes.

First, we focused on confirming the electrostatic adsorption of polymers 1 and 2 onto the FND surface. We measured the pH dependence of the particles’ zeta potentials, which correspond to the electrical potentials near the nanoparticle surface at the “slipping plane” – the border between bulk water and the tightly adsorbed hydration sphere (Figure 1). The zeta potential of oxidized FNDs was negative over the entire pH range tested (Figure 2). The highly acidic FND surface resisted full protonation even at strongly acidic pH (pH ≤ 2). Upon addition of cationic polymers, the zeta potential reversed from ~40 mV to +40 mV, confirming successful binding of polymers on the FND surface. However, at higher pH, the zeta potential of the FND-1 complex began to drop due to gradual deprotonation of the surface-bound polymer. The FND-1 complex thus dynamically responds to changes in pH and allows us to manipulate charge density on the FND surface. The zeta potential of the FND-2 complex remained positive and roughly constant over the entire pH range.

For FND, FND-1, and FND-2, we measured the dependence of fluorescence spectra on pH (Figure 3A; for complete data, see Figure S2 in Supporting Information). The spectra show typical shapes with expected ZPLs of the NV⁰ (575 nm) and NV⁻ (638 nm) states accompanied by their characteristic red-shifted phonon replicas (see Figure S3 for full range spectrum of FND).
the FND surface clearly contributes to depletion of the NV$^-$ band and reduction of its phonon replicas. The situation is analogous for pH $\approx 2$, at which the polymers are similarly charged. After alkalinization to pH $\approx 11$, we observed a difference in the behavior of FND-1 and FND-2 complexes. While FND-2 remains positively charged, FND-1 loses its charge upon deprotonation (see also Figure 2), the charge density at the FND-1 interface drops, and the depletion of NV$^-$ emission is therefore weaker for FND-1.

For convenient and quantitative representation of these changes, we plotted the NV$^0$/NV$^-$ ZPL emission ratios versus pH for FND, FND-1, and FND-2 (Figure S2D in Supporting Information). Modulation of fluorescence induced by binding of positively charged polymers on FND surfaces results consistently in a dramatic increase in the NV$^0$/NV$^+$ ratio along the entire pH range. The expected decrease of this parameter for FND-1 in the strongly alkaline region (pH $\approx 11$) due to deprotonation was, however, not significant, and the polymers 1 and 2 cannot be distinguished on this base. We also performed analogous control experiments with negatively charged and neutral polymers, revealing that these have indeed an insignificant effect on FND fluorescence (Figure S4 in Supporting Information).

To demonstrate the potential of the sensing mechanism for molecular imaging, we performed a set of experiments on cover slide with deposited FND particles. To avoid the influence of refractive index on the fluorescence intensity, we covered the surface with a water droplet and measured always in aqueous environment. We recorded time-resolved fluorescence from 20x20 µm frame using two spectral regions corresponding to ZPLs of the NV$^0$ and NV$^-$. Then we added a small amount of 2-solution to the water droplet and repeated the measurement with the same particles. We processed the data using intensity and fluorescence lifetime thresholds. For imaging we chose calculated NV$^0$/NV$^-$ ZPL emission ratios (see Supporting information for details). A clear color shift caused by formation of FND-2 complex can be seen (compare Figure 3B, left and right). The shift towards higher NV$^0$/NV$^-$ ZPL emission ratios upon formation of FND-2 complex is apparent also from histograms in Figure 3C. As a control experiment we recorded similar images for FNDs interacting with a negatively charged polymer (Figure 5S in Supporting Information). Both the images and histograms remained in this case without a significant change.

Interestingly, the histograms reflect also heterogeneity in the particle brightness (related mainly to different contents of NV centers per particle) and sensitivity of the individual particles to the observed molecular interactions. Nevertheless, the spectral properties of the system combined with a proper data processing demonstrated to be robust enough for imaging and selective distinguishing the distinctive molecular events (such
as formation of FND complexes with cationic or anionic polymers).

The theoretical model we recently described\textsuperscript{20} can explain the observed results (the simplified schematics in Figure 1 can serves as a guideline). The NV\textsuperscript{0} and NV\textsuperscript{−} ground states with energy levels of 1.2 eV and 2.0 eV, respectively, are both below the Fermi level (E\textsubscript{F}) inside a FND crystal. However, near the surface, these levels are bent due to the presence of a surface potential. The negatively charged surface of an oxidized diamond with negative zeta potential (Figure 1A) is responsible for downward band bending. Upon adsorption of a positively charged polymer, the surface potential and zeta potential are reversed (Figure 1B), and the NV\textsuperscript{0} and NV\textsuperscript{−} levels are bent upward. Thus, the NV\textsuperscript{−} state switches to NV\textsuperscript{0}, resulting in corresponding change in fluorescence. Deprotonation of the FND-I complex at basic pH reduces its positive charge density, the upward bending effect is lowered, the NV\textsuperscript{−} center becomes occupied again, and its emission is partially restored (Figure 1C).

Our sensing mechanism based on surface-charge-induced modulation of fluorescence has several important features. First, it is fundamentally different from methods based on resonance energy transfer, which is a common strategy for fluorescent biosensor construction.\textsuperscript{19} Although NV centers allow FRET,\textsuperscript{30–32} its efficacy is inversely proportional to the sixth power of the distance between chromophores, and therefore only very small FNDs can be used. Because the proportion of NV\textsuperscript{−} centers (which roughly corresponds to the brightness) drastically decreases with the FND size,\textsuperscript{22} this issue is a major limitation for FND-based FRET sensors. In contrast, the sensitivity of an NV center to surface-charge-induced alterations decreases according to the square of its distance from the surface and can be effective for distances between the NV center and surface up to \textasciitilde20 nm.\textsuperscript{20} Based on this fact, one can expect that even bigger FNDs can be sensitive over the entire volume. This may represent a significant advantage over any type of FRET-based FND sensor (which requires bright sub-10-nm particles) and deserves further examination.

Second, a diamond sensor based on surface-charge-induced modulation of fluorescence will have an intrinsic ratiometric reading mode (i.e., one that includes both reference and measurement signals) because the information is contained in the shape of the emission spectrum and not only in the intensity. Merging the reference and responsive signal into one emitter is possible due to the unique non-photobleachable character of NV centers.

Third, the optical behavior of FNDs is qualitatively different from the majority of other fluorescent nanoparticles: their emission spectra do not show quantum confinement effects known for semiconductor nanomaterials such as quantum dots.\textsuperscript{4} Possible size polydispersity of FND particles\textsuperscript{33} therefore does not contribute to broadening of fluorescent lines and other spectral features caused by polydispersity at semiconductor fluorescent nanoparticles.\textsuperscript{34}

### Conclusions

In summary, we describe a new detection mechanism based on modulation of fluorescence induced by changes in the FND surface charge that occur upon noncovalent interaction with charged molecules. In a well-controlled model system, we optically detected gradual changes in charge density on the FND surface caused by deprotonation of a polymer containing amino groups. This sensing mechanism is inherently insensitive to particle size polydispersity and operates as an intrinsic ratiometric system containing a natural non-photobleachable reference. This finding opens new possibilities for construction of biocompatible and extremely photostable two-color fluorescent (bio)nanosensors with potential high-impact applications in nanobiological and medical imaging and chemical sensing. We envision applications in time-unlimited detection of molecular events on FND surfaces, selective recognition of molecules, and quantification of chemical and biochemical entities. We are currently working in these directions such as development of intracellular nanosensors for detection and quantification of prototypal charged biopolymers – nucleic acids.

### Notes and references

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