

Formation of Quantum Emitters in Air-Suspended Carbon Nanotubes Using Vapor-Phase Reaction

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High quality carbon nanotubes such as an air-suspended will lead to improve single photon properties. Air-suspended SWCNTs are an ideal platform as a single photon emitter because they exhibit brighter photoluminescence than liquid-dispersed SWCNTs [1,2]. Such SWCNTs keep their length and pristine surface, which leads to suppress end quenching often observed in liquid-dispersed SWCNTs [3]. Introducing color centers, however, requires to disperse SWCNTs in solution [4–7]. This process results in contaminating tube surface and quenching photoluminescence due to an interaction between SWCNTs and surrounding environments [8]. Air-suspended SWCNTs are ideal as a host of color centers, but directly immersing air-suspended SWCNTs in water inevitably destroys the suspended structures due to the high surface tension of the solvent. To use the excellent optical properties of the air-suspended carbon nanotubes with color centers, an intelligent design of chemical reaction is required without compromising the air-suspended structure.

In this presentation, we propose a vapor-phase reaction in air-suspended SWCNTs, where adsorbing precursor vapor with a weak mechanical perturbation preserves the suspended structures [9]. We present the formation of color centers in air-suspended SWCNTs using the vapor-phase reaction and evaluate the photoluminescence properties of the functionalized SWCNTs. We use confocal spectroscopy to measure photoluminescence spectra and time-resolved photoluminescence, focusing at individual carbon nanotubes. We build a theoretical model considering the strain along the curvature of SWCNT and relate diameter-dependent photoluminescence properties to the chemical reactivity. We also describe the diameter dependence of trapping potential at the sp^3 defect by statistically analyzing emission energies. We estimate formed defect density from the change in photoluminescence intensity by the functionalization.

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