Highlighting excitonic optical properties of bundled carbon nanotubes to tailor novel nanomaterials-based devices

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The original excitonic optical properties of carbon nanotubes (CNT) [1,2] confer great potential on these 1D-nanomaterials for direct telecommunications applications around 1,55µm. Nowadays, all-optical regeneration of telecom signal, in high-bit-rate optical fibre transmission systems, need still more efficient optical devices, such as saturable absorbers (SA) with ultrafast optical response time and great ratio between ON/OFF states. Excitonic optical properties of nanomaterials- based SA, such as quantum wells (QW) nanostructures (2Dnanomaterials), have already demonstrated huge potential for such signal regeneration [3], in comparison to bulk materials. However, fabrication of these ultrafast 2D-nanostructures-based SA need complex and expensive growth technique, and QW-doping [4] or QW-irradiation [5] techniques have been developed to reduce typical nanosecond QW nanostructures optical response time down to subpicosecond range, required for high-bit-rate signal regeneration. Thus, ultrafast absorption dynamics and large 1D-excitonic nonlinearities of bundled CNT [6] make them intrinsically and potentially as prime candidate to replace QW-based SA. In this way, we have deposited CNT on glass, Si or photolithographied-InP substrates, from nitrogen-brushed NMP-dispersion containing CNT. We also extend the well-controlled microtechnology techniques developed for III-V semiconductor nanostructures-based devices (quantum wells, wires and dots) to CNT-based SA realization. Quality of technological process used for CNT deposition is analyzed by Scanning Electron Microscopy (SEM). Vibrational properties of CNT-based SA are characterized by Raman spectroscopy using 785nm-pump laser diode, and will be presented too. This invasive and useful optical experiment reveals vibrational modes and typical features of CNT [7]: radial breathing mode (RBM), defect-mode D and tangential modes G. The position of RBM mode provides first useful information, as it depends inversely to CNT-diameter [8]. This estimation of CNT-diameters provides first optical Van-Hove transition energy of semiconductor CNT, thanks to Kataura plot, which can be exploited for telecom applications near 0,8eV (1,55µm). Furthermore, the Raman spectra and SEM pictures show a relatively good reproductibility of our CNT-based SA, from fabrication (HiPCO from CNI *Inc.*) to deposit technological process.

Dynamics of nonlinear optical properties of CNT-based SA are highlighted by femtosecond-pulses cross-polarized degenerate pump-probe experiment. Figure 1 shows temporal evolution of normalized differential transmissions ($\Delta T/T_0$) of CNT-based and doped-QW-based SA for comparison (see ref [4] for details on doped-QW SA). A significant reduction of monoexponential response time by

using CNT as efficient and relatively low-cost nanomaterials instead of doped-QW for SA applications is demonstrated (divided by ten factor), keeping an encouraging good contrast ratio for CNT-based SA. Studies on power and wavelength dependence of these excitonic nonlinear optical properties in telecom wavelength range (ultrafast response time and contrast ratio) are in progress.

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References:

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Figure:

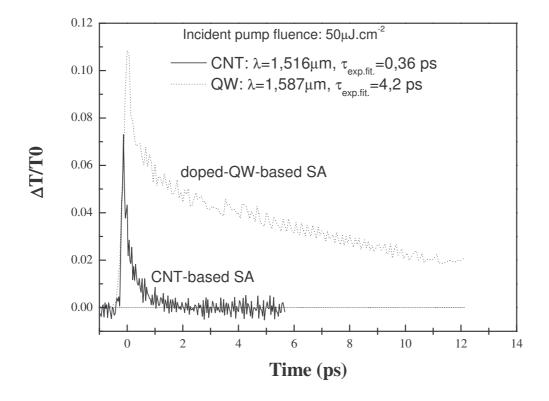


Figure 1: Ultrafast dynamics of CNT-based and doped-QW-based-SA.

Degenerate pump and probe wavelengths, in resonance with first semiconductor nanomaterials optical transition, and fitted monoexponential response times are mentioned. Dashed line at zero level is a guide for the eye.