Plasmon enhanced Raman scattering effect for an atom near a carbon nanotube

I.V. Bondarev
Math & Physics Department, North Carolina Central University, 1801 Fayetteville Str, Durham, NC 27707, USA

Quantum electrodynamics theory of the resonance Raman scattering is developed for an atom in a close proximity to a carbon nanotube. The theory describes both weak and strong atomic coupling to nanotube plasmon near fields, and predicts a dramatic enhancement of the Raman intensity in the strong coupling regime. This resonance scattering is a manifestation of the surface enhanced Raman scattering effect, and can be used in designing efficient nanotube based optical sensing substrates for single atom detection, precision spontaneous emission control, and manipulation.

Surface-enhanced Raman spectroscopy (SERS) has received much of attention recently due to a very broad range of its applications ranging from optics and plasmonics to biochemistry and medicine \cite{1,2}. High scattering intensities within narrow spectral bands reduce the probability for spectral overlapping to allow for better recognition of multiple markers, making SERS one of the most efficient optical sensing techniques. With the development of advanced nanomaterials, various SERS substrates are demonstrated \cite{3,4}. However, there is still a need for the substrates of improved sensitivity and signal reproducibility, which require clear understanding of the underlying scattering mechanisms to be developed.

In this Letter, the quantum electrodynamics (QED) theory of the resonance Raman scattering is presented for an atom near a carbon nanotube (CN). Nanotubes offer precise tunability of their electromagnetic properties by simply varying their diameters and chiralities. CNs of different diameters and chiralities feature similar electronic band structure peculiarities, yet shifted in frequency relative to one another \cite{5,6}. This yields similar optical properties over a broad range of excitation frequencies both in the far- and in the near-field zone, originating from exciton and plasmon excitations, respectively. Excitons and plasmons are different in their physical nature, but originate from the same circumferentially quantized electronic transitions. Due to the circumferential quantization of the longitudinal electron motion, real axial (along the CN axis) optical conductivities of single wall CNs consist of series of peaks \(E_{11}, E_{22}, ...,\) representing the 1st, 2nd, etc. excitons, respectively [see Fig. 1 (a)]. Imaginary conductivities are linked with the real ones of the neighboring elementary cells of the CN. When excited, their associated (plasmon-induced) quasi-static fields can be strong enough to result in the enhanced Raman scattering effect by atomic type species (extrinsic atoms, ions, molecules, or semiconductor quantum dots) in the CN vicinity. This work derives and analyzes the differential cross-section for such scattering.

In absence of an external electromagnetic (EM) radiation, an atom, modeled here by a two-level system (TLS) positioned at the point \(r_A\) near an infinitely long single wall CN, interacts with the quantum EM field of the CN via an electric dipole transition \(\Delta_{\omega}\) of frequency \(\omega_A\) with the quantization axis being the CN symmetry axis [Fig. 1(b), inset]. Transverse dipole orientations can be neglected due to the transverse depolarization effect in individual CNs. The Hamiltonian of the coupled CN–TLS system is given by (see Refs. \cite{15,16})

\[
\hat{H} = \hat{H}_F + \hat{H}_A + \hat{H}_{AF} \\
= \int_0^\infty d\omega \hbar \omega \int d\mathbf{R} \hat{f}^\dagger (\mathbf{R}, \omega) \hat{f}(\mathbf{R}, \omega) + \frac{\hbar \omega_A}{2} \hat{\sigma}_Z \\
+ \int_0^\infty d\omega \int d\mathbf{R} \left[ g^{(+)}(\mathbf{r}_A, \mathbf{R}, \omega) \hat{\sigma}^\dagger + g^{(-)}(\mathbf{r}_A, \mathbf{R}, \omega) \hat{\sigma} \right] \hat{f}(\mathbf{R}, \omega) + \text{h.c.}
\]

with the three terms representing the (medium-assisted) quantum EM field of the CN, the TLS, and their interaction, respectively. Here, \(\hat{f}(\mathbf{R}, \omega)\) and \(\hat{f}^\dagger (\mathbf{R}, \omega)\) are the scalar bosonic field operators defined on the surface of the CN \([\mathbf{R} = (R_{cn}, \varphi, z)]\) is the radius-vector of a point on the CN surface] to represent the CN field subsystem. Pauli operators, \(\hat{\sigma}_Z = |u\rangle\langle u| - |l\rangle\langle l|, \hat{\sigma} = |l\rangle\langle u|\) and \(\hat{\sigma}^\dagger = |u\rangle\langle l|\), describe the TLS and its (dipole) transitions between the two states, upper \(|u\rangle\) and lower \(|l\rangle\), with the transition frequency \(\omega_A\) modified by the diamagnetic \(\mathbf{A}^2\)-term (vector potential) to result in the new renormalized transition frequency \(\tilde{\omega}_A = \omega_A[1 - 2/(\hbar \omega_A)^2] \int_0^\infty d\omega \int d\mathbf{R} |g^{(+)}(\mathbf{r}_A, \mathbf{R}, \omega)|^2\). The matrix elements of the CN field interaction with the TLS are of the form \(g^{(\pm)} = g^{(\pm)}(\mathbf{r}_A, \mathbf{R}, \omega) = -i(4\omega_A/c^2)\sqrt{\pi \hbar \omega} \text{Re} \sigma_{zz}(\omega) d_z \vert \delta(\mathbf{r}_A, \mathbf{R}, \omega)\rangle\). The \(\delta(\mathbf{r}_A, \mathbf{R}, \omega)\) is the zz-component of the transverse (longitudinal) Green tensor (with respect to the first variable) of the CN assisted quantum field, \(\sigma_{zz}(\omega)\) is the CN

\*E-mail: ibondarev@nccu.edu
surface axial conductivity [Fig. 1(a)]. Functions $g^{\perp(1)}$ have the property as follows. \( \int dR |g^{\perp(1)}(\mathbf{r}_A, R, \omega)|^2 = (\hbar^2/2\pi)(\omega_A/\omega)^2 \Gamma_0(\omega)\xi^{\perp(1)}(\mathbf{r}_A, \omega) \), where $\xi^{\perp(1)}(\mathbf{r}_A, \omega) = \text{Im}^{\perp(1)}(\mathbf{r}_A, \omega)$. The transverse (longitudinal) photonic density of states (DOS) as seen from the TLS location relative to vacuum, and $\Gamma_0(\omega) = 3\pi\omega^2\delta^2 \text{Im} G^{\|}_A(\omega)/3\hbar c^2$ is the TLS spontaneous decay rate in vacuum with $\text{Im} G^{\|}_A(\omega) = \omega/6\pi\hbar$ being the vacuum imaginary Green tensor $z\cdot z$-component.

Hamiltonian (11) involves only two standard approximations, the electric dipole and two-level approximation [15, 16], while conveniently representing the coupled TLS-CN system in terms of the relative distance dependent DOS functions $\xi^{\perp(1)}(\mathbf{r}_A, \omega)$. For short TLS-CN separation distances EM retardation effects play no role, and so one has $\xi^{\perp} = \xi^{\parallel} = \xi(r_A, \omega)$ for the DOS functions (16). Figure 1(b) shows the calculations of $\xi(r_A = R_{CN} + 2b, x) = \rho_\omega(2\pi\gamma_0$ being the dimensionless energy, with $b = 1.42$ Å and $\gamma_0 = 2.7$ eV representing the C-C interatomic distance and overlap integral, respectively, for the three semiconducting CNs of increasing diameter. We see the sharp single-peak resonances originating from the interband plasmons of the respective CNs [cf. Fig. 1(a) and Fig. 1(b)]. These are responsible for the CN–TLS coupling in the near-field. In the linear coupling regime, quite generally, the coupled CN–TLS system can be represented as a four-level system with the eigenvectors of the Hamiltonian (11) being as follows

\[
\begin{align*}
|0\rangle &= |l\rangle|\{0\}\rangle, \\
|1\rangle &= C^{(1)}_{ul}|u\rangle|\{0\}\rangle + \int_0^\infty d\omega \int dR C^{(1)}_{l}(R, \omega)|l\rangle|\{1(R, \omega)\}\rangle, \\
|2\rangle &= C^{(2)}_{ul}|u\rangle|\{0\}\rangle + \int_0^\infty d\omega \int dR C^{(2)}_{l}(R, \omega)|l\rangle|\{1(R, \omega)\}\rangle, \\
|3\rangle &= |u\rangle|\{1(R, \omega)\}\rangle.
\end{align*}
\]

Here, $|\{0\}\rangle$ and $|\{1(R, \omega)\}\rangle$ are, respectively, the vacuum and single-quantum excited states of the CN field subsystem, and $C^{(1,2)}_{ul}$ are the mixing coefficients of the spontaneous decay type transition $|u\rangle|\{0\}\rangle \to |l\rangle|\{1(R, \omega)\}\rangle$ in the coupled CN–TLS system [16, 17], while the similar mixing of the $|l\rangle|\{0\}\rangle$ and $|u\rangle|\{1(R, \omega)\}\rangle$ states, known to be responsible for the long-range dispersive van der Waals interaction [15, 16], is neglected for simplicity.

To proceed to the Raman scattering cross-section calculations, it is necessary to fully determine scatterer’s eigenvectors in Eq. (2) together with their respective energy eigenvalues. Approximating the sharp DOS resonance [Fig. 1(b)] at the plasma frequency $\omega_p$ by the Lorentzian function of half-width-at-half-maximum $\Delta_0$ of the form $\xi(r_A, \omega) \approx \xi(r_A, \omega_p)\Delta_0^2/(\omega - \omega_p)^2 + \Delta_0^2$ allows one to obtain the analytical solution to the eigenvalue problem that is valid both in the weak and in the strong CN–TLS coupling regime [13]. Namely, applying the Hamiltonian (11) to the four-vector set (2), one obtains the (dimensionless) eigen energies as follows

\[
\begin{align*}
\varepsilon_0 &= -\frac{\tilde{x}_A}{2} + \varepsilon_{1,2} = \frac{1}{2}(x_p \mp \sqrt{\delta^2 + X^2} - i\Delta x_p), \\
\varepsilon_3 &= \frac{\tilde{x}_A}{2} + x_p - i\Delta x_p,
\end{align*}
\]

where $X = (\hbar/2\gamma_0)\sqrt{2\Delta_0\Gamma_0(\omega_p)(1 + \omega^2/\omega_p^2)} \xi(r_A, \omega_p)$, $\delta = \tilde{x}_A - x_p$, $(\tilde{x}_A, x_p, \Delta x_p) = (\hbar(\omega_A, \omega_p, (\Delta_0)/2\gamma_0$, and $\Delta x_p$ is added to phenomenologically account for the finite half-width of the plasmon resonance, as seen in Fig. 1(a), which is assumed to be much broader than the excited atomic level natural half-width dropped here on this account for simplicity. For the mixing coefficients, one has

\[
|C^{(1,2)}_{ul}|^2 = \frac{1}{2} \left( 1 + \frac{1 \mp \sqrt{1 + X^2/\delta^2}}{1 + X^2/\delta^2} \right).
\]

FIG. 1: (Color online) (a) Fragment of the energy dependence of the dimensionless (normalized by $\bar{\epsilon}/2\pi\hbar$) axial surface conductivities $\sigma_{zz}$ for the semiconducting (6,4), (10,0) and (11,0) CNs of increasing diameter. Peaks of Re $\sigma_{zz}$ represent excitation modes ($E_{11}$, $E_{22}$,...); peaks of Re $\sigma_{zz}$ are inter-band plasmons ($P_{11}$,...). (b) Photonic DOS functions for the CNs in (a) with the TLS placed at $r_A = R_{CN} + 2b$ from the CN symmetry axis (inset). Dimensionless energy is [Energy]/$2\gamma_0$. Conductivities are obtained using the (k, p)-scheme of Ref. 3. DOS functions are calculated as described in Refs. 15, 16, 18.
The entire scattering process includes three sequential steps [19, 20]. They are: (a) excitation of the system by an incident photon of the frequency \( \omega \) with the unit polarization vector \( \mathbf{e}_i \), described by the interaction matrix element \( \langle \mathbf{n}|\hat{H}_R(\omega_i)|0\rangle = (-i/c)\sqrt{2\pi\omega_i}d_z\cos\theta_i \) \((n = 1, 2)\) with \( \cos\theta_i = \mathbf{e}_i \cdot \mathbf{e}_z \); (b) plasmon emission (or absorption) on the CN surface, described by the matrix element \( \langle 1|\hat{H}_{AF}^{(c)}|2\rangle \) (or \( \langle 2|\hat{H}_{AF}^{(a)}|1\rangle \)) with \( \hat{H}_{AF}^{(c)} \) and \( \hat{H}_{AF}^{(a)} \) being the emission term (\( \sim \hat{j}^f \)) and the absorption term (\( \sim \hat{j} \)), respectively, of the interaction Hamiltonian \( \hat{H}_{AF} \) in Eq. (1); (c) de-excitation of the CN–TLS system by the scattered (Raman) photon emission of the frequency \( \omega_s \) with the unit polarization vector \( \mathbf{e}_s \), described by the interaction matrix element \( \langle \mathbf{n}|\hat{H}_R(\omega_i)|0\rangle^* \). There are four Feynman diagrams, shown in Fig. 2, bottom, to contribute to this process. They are two for the plasmon emission (bottom left) and two for the plasmon absorption (bottom right), to represent two indistinguishable ways for the emission and absorption to occur. Two types of the emission (absorption) diagrams should be summed up and squared, followed by adding the emission and absorption contributions together [19, 20]. This results in the differential Raman scattering cross-section of the form

\[
\frac{d\sigma}{d\Omega} = \frac{(2\gamma_0)^2|d_z|^4}{\hbar^2 c^2} \cos^2\theta \cos^2\phi P(x_i, x_s)
\]

with the dimensionless (angle-free) scattering probability

\[
P(x_i, x_s) = \frac{x_i x_s^4 X^8}{2^6(\delta^2 + X^2)^2(\delta^2 + \Delta x_p^2)} \times \frac{1}{\left| (x_i - x_p - \delta_+ / 2)^2 + \Delta x_p^2 \right|} \left( \frac{x_s - x_p - \delta_- / 2}{2} + \Delta x_p^2 \right)
\]

\[
+ \frac{1}{\left| (x_i - x_p - \delta_- / 2)^2 + \Delta x_p^2 \right|} \left( \frac{x_s - x_p - \delta_+ / 2}{2} + \Delta x_p^2 \right)
\]

where \( x_{i,s} = \hbar \omega_{i,s} / 2\gamma_0, \delta_{\pm} = \delta \pm \sqrt{\delta^2 + X^2} \), and two terms...
stand for the contributions from the plasmon emission and absorption, respectively.

Each term in Eq. (7) has the product of two resonance energy denominators that include $x_i$ and $x_s$, incident (incoming) and scattered (outgoing) photon energies. This is what makes this Raman scattering resonant. For fixed $x_i$, only one term contributes, resulting in either Stokes scattering with $x_i < x_s$ and a plasmon created in the CN, or in anti-Stokes scattering with $x_i > x_s$ and a plasmon absorbed from the CN. The absolute value of the Raman shift is $\sqrt{\delta^2 + X^2}$, yielding a quantity $\sim X$ in resonance, where $\delta \approx 0$ whereby $X^2/\delta^2 \gg 1$, and that $\sim \delta$ out of resonance with $X^2/\delta^2 \ll 1$. In the latter case, the pre-factor in Eq. (7) tends to zero as $X^4/\delta^4$, totally ruling out the probability of the scattering process. When in resonance, on the other hand, the maximum of $P(x_i, x_s)$ goes as $(X^4/\Delta x_p^4)/(X^2 + \Delta x_p^2)$, being strongly suppressed in the weak TLS-plasmon coupling regime where $X^2/\Delta x_p^2 \ll 1$, and being dramatically enhanced in the opposite case where $X^2/\Delta x_p^2 \gg 1$ so that the TLS-plasmon coupling is strong. The scattering enhancement factor is about square of that for the resonance absorption by atomically doped CNs [2], the way it should be for scattering as a two-step process of absorption followed by emission viewed as “reversed absorption”.

Figure 3 shows an example of the numerical calculations for the scattering probability $P(x_i, x_s)$ as given by Eq. (7) for $x_p = 0.35$ [4] plasmon in Fig. (1 (a)] with $\Delta x_p$ and $X$ varying independently [columns (b) and (a)] within a typical (conservative) range of parameters [21], to see the plasmon decoherence effect and the role of DOS resonance variation as the TLS–CN separation distance changes. As discussed above, Raman scattering is seen to be very sensitive to the strong TLS-plasmon coupling, blowing up by a factor over $10^3$ for $X/\Delta x_p \sim 10$ and totally vanishing when $X/\Delta x_p \sim 1$. Raising $X$ increase both the Raman shift and the intensity, while greater $\Delta x_p$ quench the intensity with no Raman shift change.

In summary, the QED theory of the resonance Raman scattering is developed for a two-level dipole emitter coupled to a weakly-dispersive interband plasmon resonance of a carbon nanotube. Such scattering is a manifestation of the general SERS effect, in which the enhancement is due to plasmon-induced near-fields that affect the TLS in close proximity to the CN surface. Raman cross-section obtained covers both weak and strong TLS-plasmon coupling, and shows a dramatic increase in the strong coupling regime. The effect may be used to detect individual atomic type objects trapped near CNs. More advanced applications, which require further theoretical development, may include highly efficient CN based SERS substrates for single molecule/atom/ion detection, precision spontaneous emission control, and optical manipulation.

This work is supported by DOE (de-sc0007117). I.V.B. acknowledges hospitality of Munich Advanced Photonics Center at TU-Minuch, Germany, where this work was started, as well as fruitful discussions with its staff members, Prof. W.Domcke and Dr. M.Gelin.

[1] K.Kneipp, M.Moscovits, and H.Kneipp, Surface-enhanced Raman scattering: Physics and applications (Springer-Verlag, Berlin, 2006).
[2] R.Zhang, Y.Zhang, Z.C.Dong, S.Jiang, C.Zhang, L.G.Chen, L.Zhang, Y.Liao, J.Aizpurua, Y.Luo, J.L.Yang, and J.G.Hou, Nature 498, 82 (2013).
[3] M.Peng, H.Xu, and M.Shao, Appl. Phys. Lett. 104, 193103 (2014).
[4] D.M.Aandrade, H.S.Vieira, M.M.Oliveira, A.P.Santos, L.Yin, R.Saito, M.A.Pimenta, C.Fantini, and C.A.Furtado, Carbon 56, 235 (2013).
[5] Q.Hao, S.M.Morton, B.Wang, Y.Zhao, L.Jensen, and T.J.Huang, Appl. Phys. Lett. 102, 011102 (2013).
[6] R.Lv, Q.Li, A.R.Botello-Mendez, T.Hayashi, B.Wang, A.Berkdemir, Q.Hao, A.L.Elias, R.Cruz-Silva, H.R Gutierrez, Y.A.Kim, H.Muramatsu, J.Zhu, M.Endo, H.Terrones, J.-C.Charlie, M.Pan, and M.Terrones, Scientific Reports 2, 586 (2012).
[7] D.Z.Lin, Y.P.Chen, P.J.Huang, J.Y.Chu, J.T.Yeh, and J.-K.Wang, Optics Express 19, 4337 (2011).
[8] R.Saito, G.Dresselhaus, and M.S.Dresselhaus, Science of Fullerens and Carbon Nanotubes (Imperial College Press, London, 1998).
[9] T.Ando, J. Phys. Soc. Jpn. 74, 777 (2005).
[10] T.Hertel and I.V.Bondarev (eds.), Photophysics of Carbon Nanotubes and Nanotube Composites (Special Issue), Chem. Phys. 413, 1 (2013).
[11] I.V.Bondarev, L.M.Woods, and K.Tatur, Phys. Rev. B 80, 085407 (2009).
[12] I.V.Bondarev, Phys. Rev. B 85, 035448 (2012).
[13] I.V.Bondarev and T.Antonijevic, Phys. Stat. Sol. C 9, 1250 (2012).
[14] T.Pichler, M.Knupfer, M.S.Golden, J.Fink, A.Rinzler, and R.E.Smalley, Phys. Rev. Lett. 80, 4729 (1998).
[15] I.V.Bondarev and Ph.Lambin, Phys. Rev. B 72, 035451 (2005).
[16] I.V.Bondarev and Ph.Lambin, Near-field electrodynamics of atomically doped carbon nanotubes, in: Trends in Nanotubes Research (Nova Science, New York, 2006). Chapter 6, p. 139.
[17] I.V.Bondarev and Ph.Lambin, Phys. Rev. B 70, 035407 (2004).
[18] Actual DOS resonance frequencies are slightly red shifted relative to their respective plasmon resonance frequencies [cf. Figs. 1 (a) and (b)]. The shifts are within plasmon resonance widths though, and so are neglected, thereby reducing the number of relevant theory parameters here.
[19] V.B.Berestetskii, E.M.Lifshitz, and L.P.Pitaevskii, Quantum Electrodynamics (Pergamon, Oxford, 1982).
[20] P.Y.Yu and M.Cardona, Fundamentals of Semiconductors, 4th edn. (Springer-Verlag, Berlin 2010).
[21] I.V.Bondarev and B.Vlahovic, Phys. Rev. B 74, 073401 (2006).