

On how to turn quantum dynamical phase transitions into plasmonic applications.

Raúl A. Bustos Marún,^(1,2) Axel D. Dente,⁽¹⁾ Eduardo A. Coronado,⁽²⁾ and Horacio M. Pastawski. ⁽¹⁾

⁽¹⁾IFEG, Facultad de Matemática Astronomía y Física, Universidad Nacional de Córdoba, Córdoba, Argentina

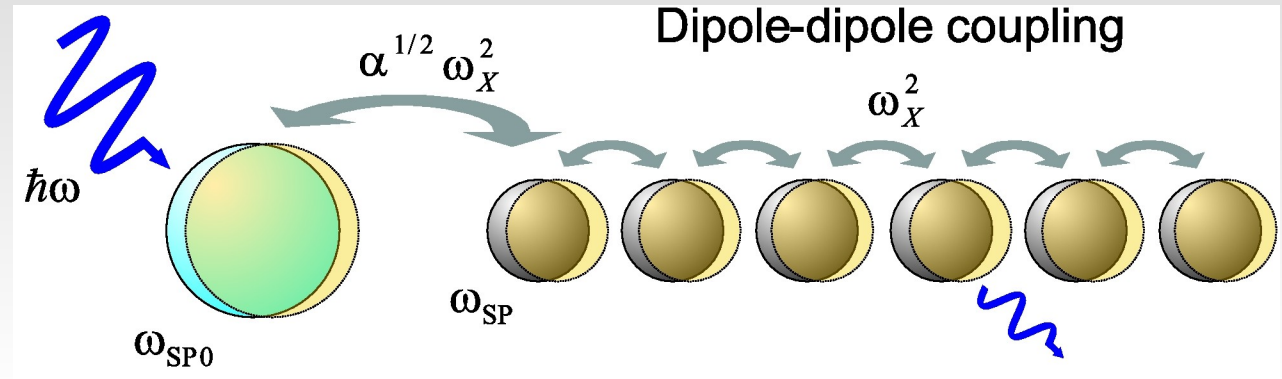
⁽²⁾INFIQC, Facultad de Ciencias Químicas, Universidad Nacional de Córdoba, Córdoba, Argentina

On how to turn quantum dynamical phase transitions into plasmonic applications.

- **Injection of excitations into waveguides.**
(Localized-delocalized transition and role of virtual states)
- **Using dynamical phase transition in plasmonics.**
(Nano-rulers, dielectric constant sensors, and deformation sensors)
- **Plasmonic synchronization.**
(New possibilities at the nanoscale)

Injection of excitations into waveguides.

The system



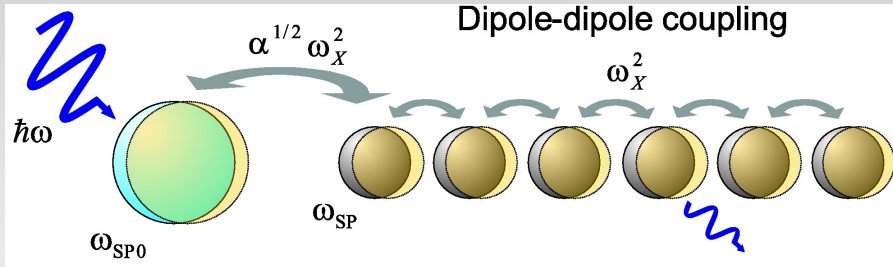
Coupled dipole approximation

$$[\omega_{SPi}^2 - \omega^2 - i\eta_i\omega] \vec{P}_i = V_i \omega_{Pi}^2 \epsilon_0 \left[\vec{E}_i^{(ext)} + \sum_{j \neq i}^N \vec{E}_{j,i}(\vec{P}_j, \vec{d}_{j,i}, \vec{k}) \right]$$

Near field approximation

$$\vec{E}_{j,i}(\vec{P}_j, \vec{d}_{j,i}, \vec{k})_{kd \rightarrow 0} \approx \frac{\vec{P}_j - 3\hat{d}_{j,i}(\vec{P}_j \cdot \hat{d}_{j,i})}{4\pi\epsilon_0 n^2 d_{j,i}^3} = \frac{\gamma^{T/L}}{4\pi\epsilon_0 n^2 d_{j,i}^3} \vec{P}_j$$

Response function



$$|P_m|^2 = |\chi_{m,0}|^2 |E_0^{(ext)}|^2$$

$$\chi_{m,0}(\omega) = \chi_{0,0}(\omega) \alpha^{1/2} e^{-m(\kappa + ik)}$$

$$\vec{P} = [\omega^2 \mathbf{I} - \mathbf{M}]^{-1} \mathbf{R} \vec{E} = \chi \vec{E}$$

$$\mathbf{M} = \begin{bmatrix} \omega_{SP0}^2 - i\eta\omega & \omega_{x0-1}^2 & 0 & 0 \\ \omega_{x1-0}^2 & \omega_{SP}^2 - i\eta\omega & \omega_x^2 & 0 \\ 0 & \omega_x^2 & \omega_{SP}^2 - i\eta\omega & \omega_x^2 \\ 0 & 0 & \omega_x^2 & \ddots \end{bmatrix}$$

$$\chi_{0,0}(\omega) = \frac{V_0 \omega_{P0}^2 \epsilon_0}{[\omega^2 - \tilde{\omega}_{SP0}^2] - \alpha \Pi(\omega)}$$

$$\alpha^{1/2} = \frac{\omega_{x1-0}^2}{\omega_x^2}$$

$$\kappa + ik = \ln(\omega_x^2 / \Pi)$$

$$\omega_{xi-j}^2 = \frac{\gamma^{T,L} \omega_{Pi}^2}{3n^2} \left(\frac{r_i}{d_{i,j}} \right)^3$$

$$\Pi = \frac{(\omega^2 - \tilde{\omega}_{SP}^2)}{2} - \text{sgn}(\omega^2 - \omega_{SP}^2) \sqrt{\left(\frac{\omega^2 - \tilde{\omega}_{SP}^2}{2} \right)^2 - \omega_x^4}$$

Poles of the response function

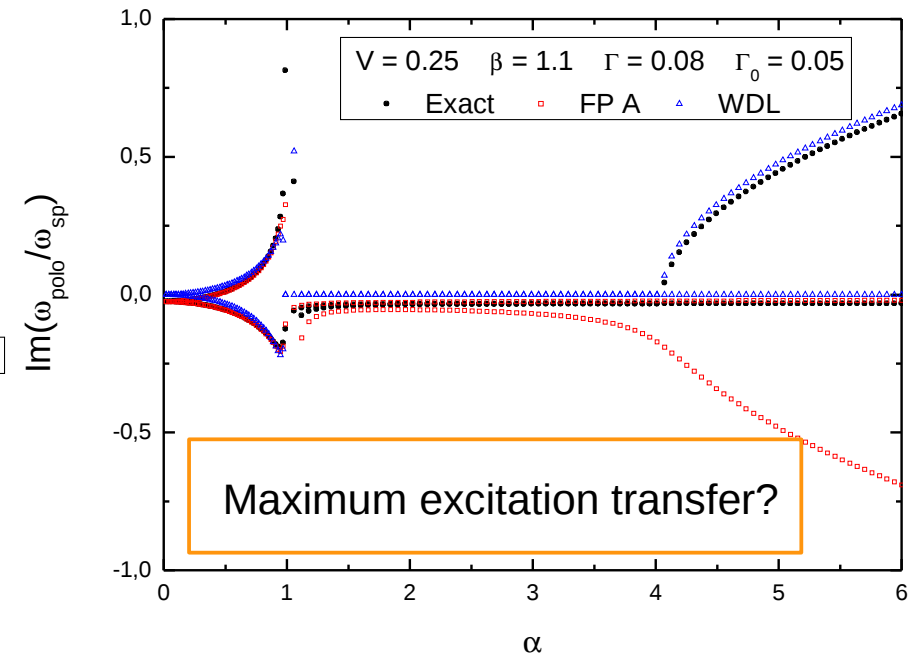
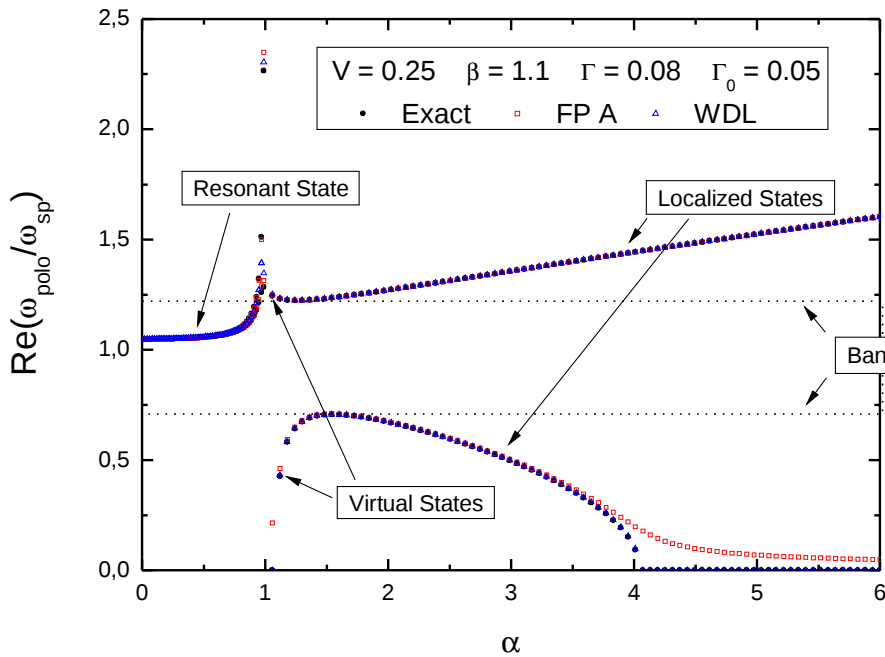
$$M = \begin{bmatrix} \omega_{SP0}^2 - i\eta\omega & \omega_{x0-1}^2 & 0 & 0 \\ \omega_{x1-0}^2 & \omega_{SP}^2 - i\eta\omega & \omega_x^2 & 0 \\ 0 & \omega_x^2 & \omega_{SP}^2 - i\eta\omega & \omega_x^2 \\ 0 & 0 & \omega_x^2 & \ddots \end{bmatrix}$$

$$X_{0,0}(\omega) = \frac{V_0 \omega_{P0}^2 \epsilon_0}{[\omega^2 - \tilde{\omega}_{SP0}^2] - \alpha \Pi(\omega)}$$

$$\Pi = \frac{(\omega^2 - \tilde{\omega}_{SP}^2) - \text{sgn}(\omega^2 - \omega_{SP}^2) \sqrt{\left(\frac{\omega^2 - \tilde{\omega}_{SP}^2}{2}\right)^2 - \omega_x^4}}{2}$$

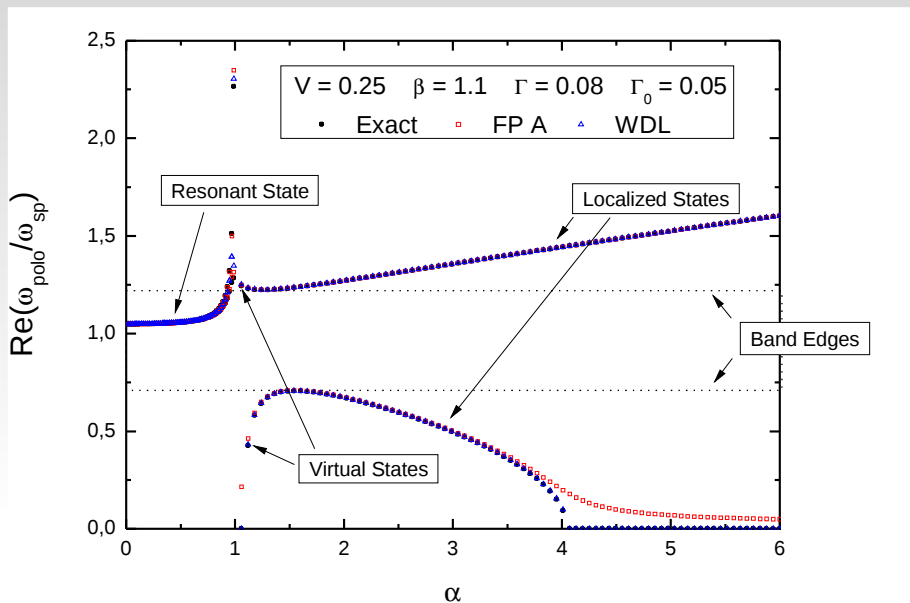
$$\tilde{\omega}_{pole}^2 = \frac{[\beta - \alpha(\beta+1)/2]}{(1-\alpha)} \pm \frac{\alpha}{2(1-\alpha)} \sqrt{(1-\beta)^2 - 4V^2(1-\alpha)}$$

$$\alpha = \frac{\omega_{x0-1}^2}{\omega_x^2} \quad \beta = \frac{\tilde{\omega}_{SP0}^2}{\tilde{\omega}_{SP}^2} \quad V = \frac{\omega_x^2}{\tilde{\omega}_{SP}^2}$$



Real and imaginary part of the poles of the response function as function of the relative coupling between nanoparticle "0" and the chain (α).

What is a virtual state?



They appear as solutions of the pole equation

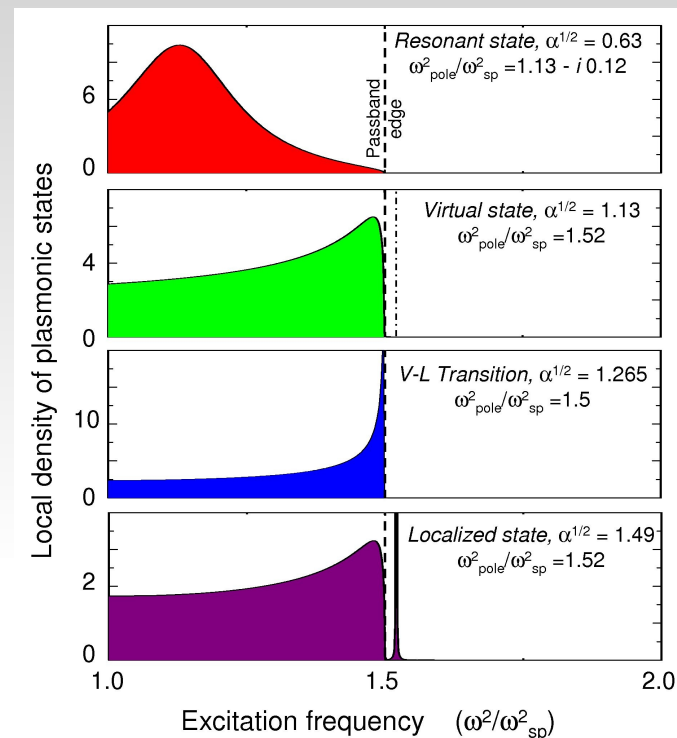
$$\tilde{\omega}_{pole}^2 = \frac{[\beta - \alpha(\beta + 1)/2]}{(1 - \alpha)} \pm \frac{\alpha}{2(1 - \alpha)} \sqrt{(1 - \beta)^2 - 4V^2(1 - \alpha)}$$

Pole of a non-physical response function

$$\chi'_{0,0}(\omega) = \frac{V_0 \omega_{p0}^2 \epsilon_0}{[\omega^2 - \tilde{\omega}_{sp0}^2] - \alpha \Pi'}$$

Non-physical self-energy

$$\Pi'(\omega) = \Delta(\omega) + i\Gamma(\omega)$$



Why care?

$$\Im(\chi_{0,0}) \approx \frac{-V_0 \omega_{p0}^2 \epsilon_0 \alpha \Gamma}{[\omega^2 - \tilde{\omega}_{sp0}^2 - \alpha \Delta]^2 + \alpha^2 \Gamma^2} \quad \Im(\chi'_{0,0}) \approx \frac{V_0 \omega_{p0}^2 \epsilon_0 \alpha \Gamma}{[\omega^2 - \tilde{\omega}_{sp0}^2 - \alpha \Delta]^2 + \alpha^2 \Gamma^2}$$

Within the passband, the poles of the LDOS are the same!!!

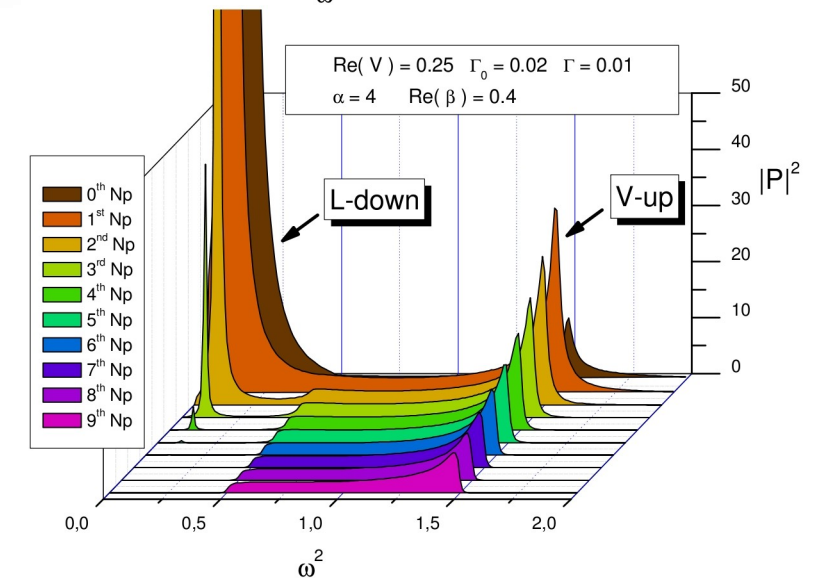
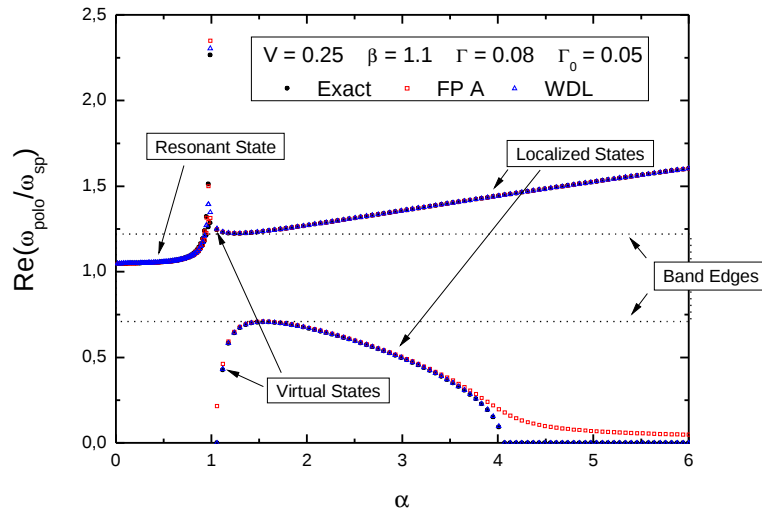
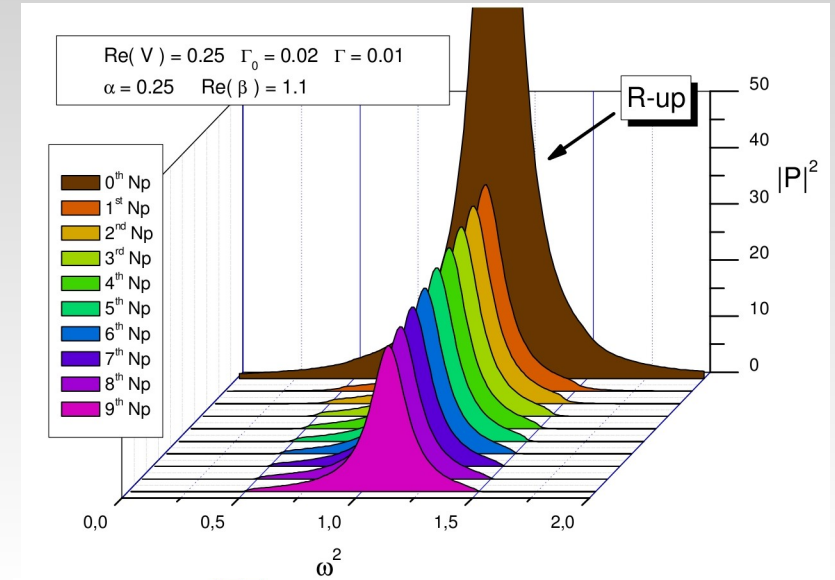
They describe the localized-delocalized transition.

Propagation of the excitation

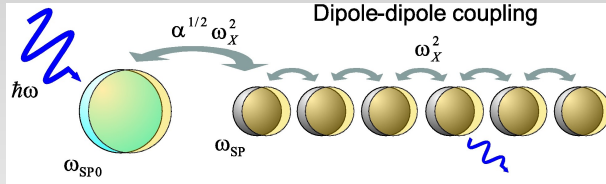
$$|P_m|^2 = |\chi_{m,0}|^2 |E_0^{(ext)}|^2$$

$$\chi_{m,0}(\omega) = \chi_{0,0}(\omega) \alpha^{1/2} e^{-m(\kappa + ik)}$$

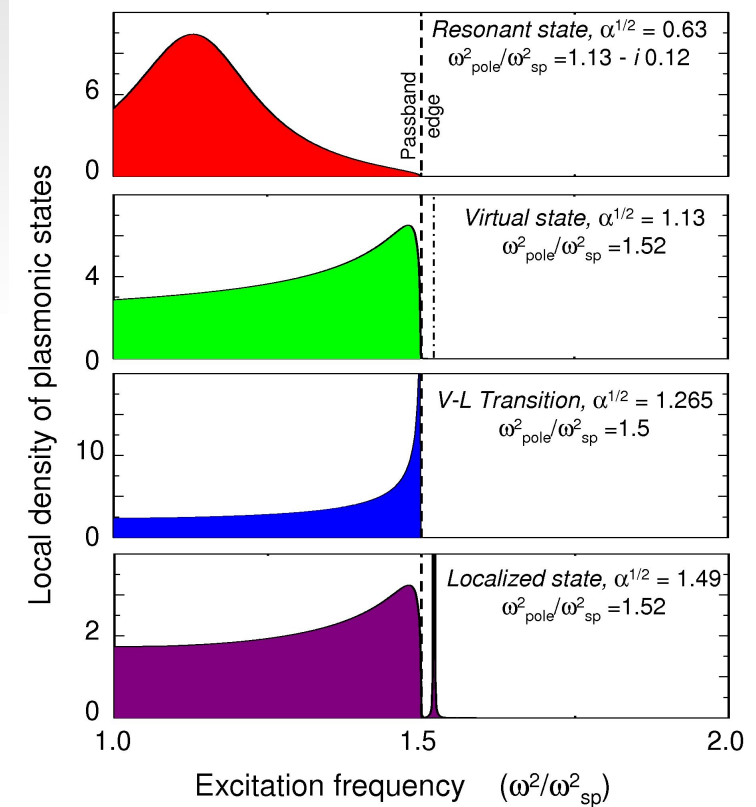
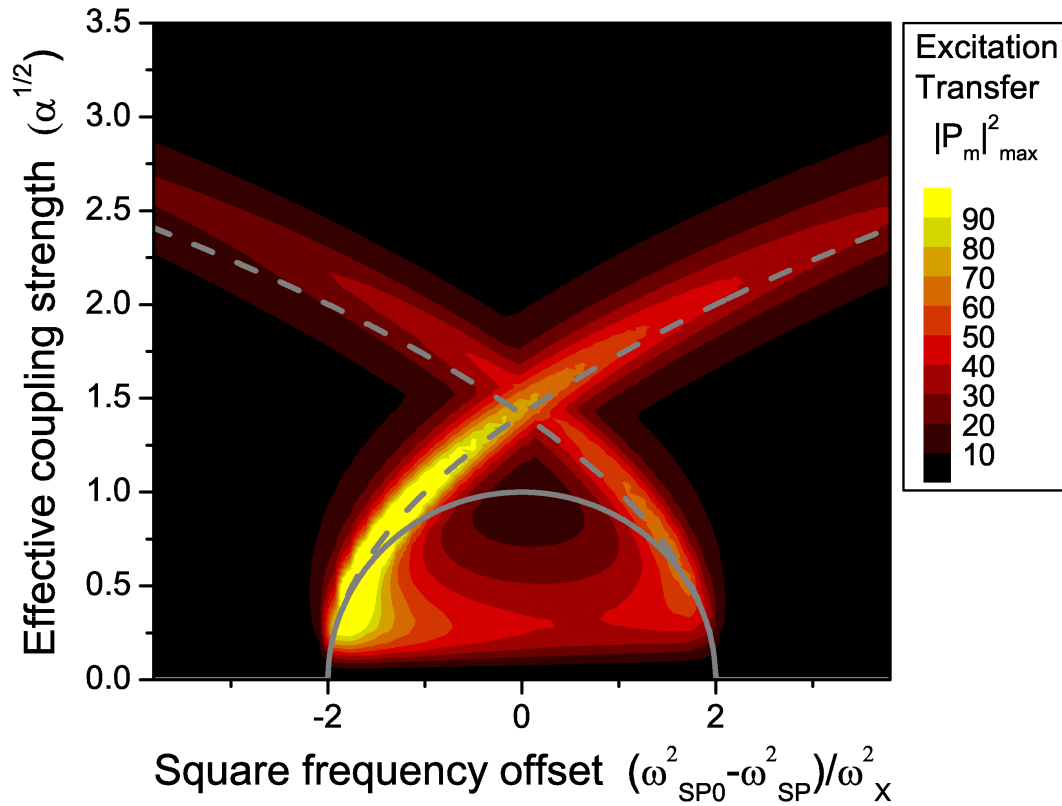
$$\kappa + ik = \ln(\omega_x^2 / \Pi)$$



Phase diagram



At the virtual-localized transition there is an accumulation of states at the passband edge.



Dynamical phase transitions
control excitation transfer!!!

On how to turn quantum dynamical phase transitions into plasmonic applications.

- **Injection of excitations into waveguides.**
(Localized-delocalized transition and role of virtual states)
- **Using dynamical phase transition in plasmonics.**
(Nano-rulers, dielectric constant sensors, and deformation sensors)
- **Plasmonic synchronization.**
(New possibilities at the nanoscale)

Using DPTs in a concrete plasmonic example.

Nps modeled as Dipoles

=> small Nps => size and shape corrections to damping

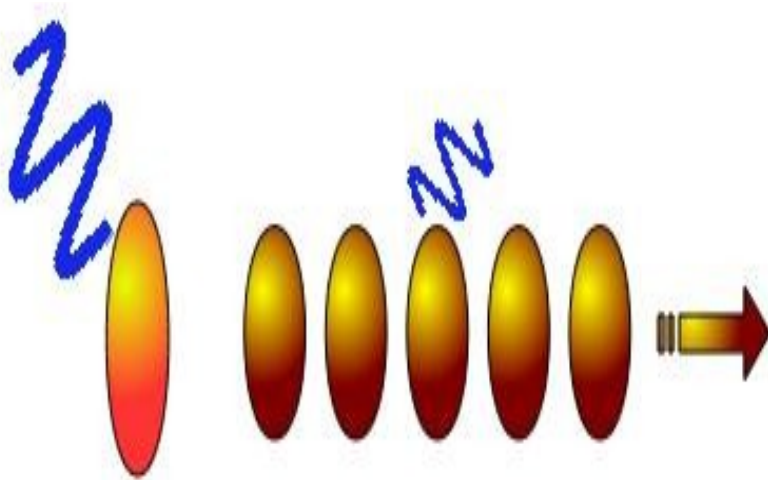
Are stronger damping compensated by stronger coupling?
Are “needed parameters” within actual limits?

Dipolar interaction

=> distance/radius not too small
...but...

Near field approximation

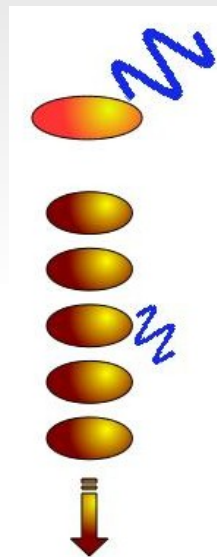
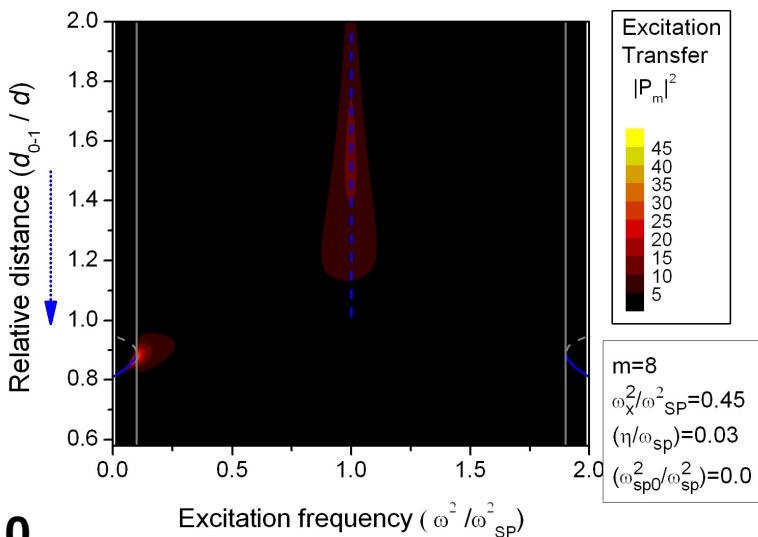
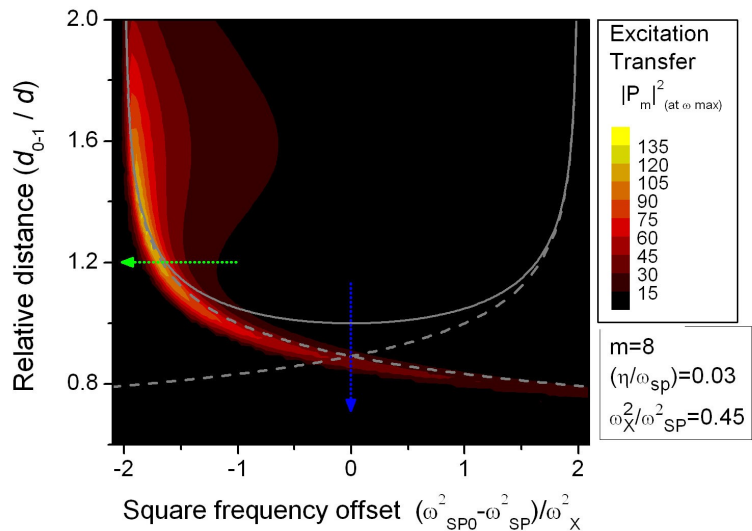
=> Small separation



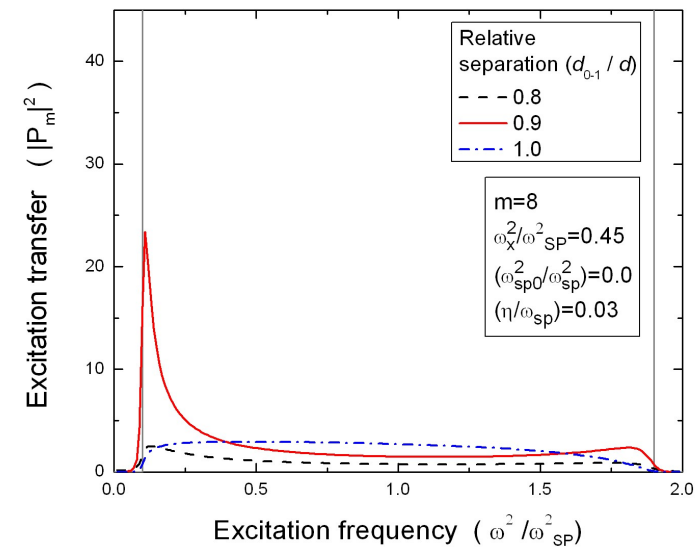
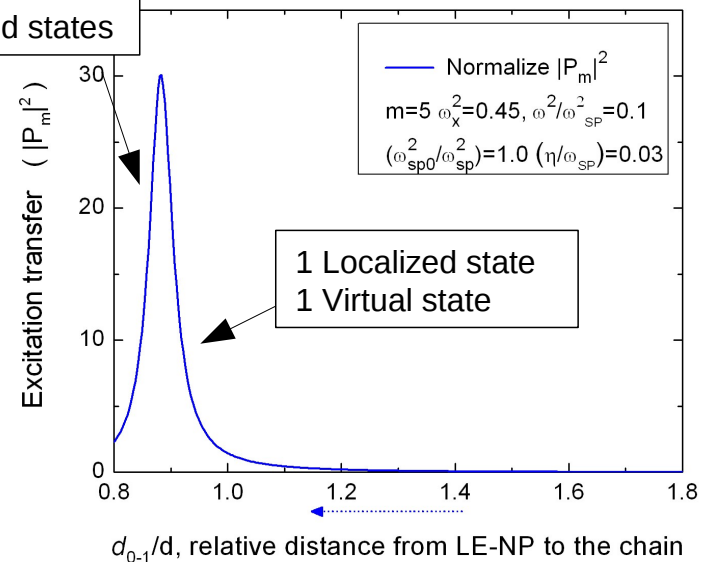
YES!!
Ag Spheroidal NPs
Dimensions: a=3nm, b=c=10.5nm
Separation d=9nm

$$\frac{\omega_x^2}{\omega_{SP}^2} = 0.45$$
$$\frac{\eta}{\omega_{SP}} = 0.03$$

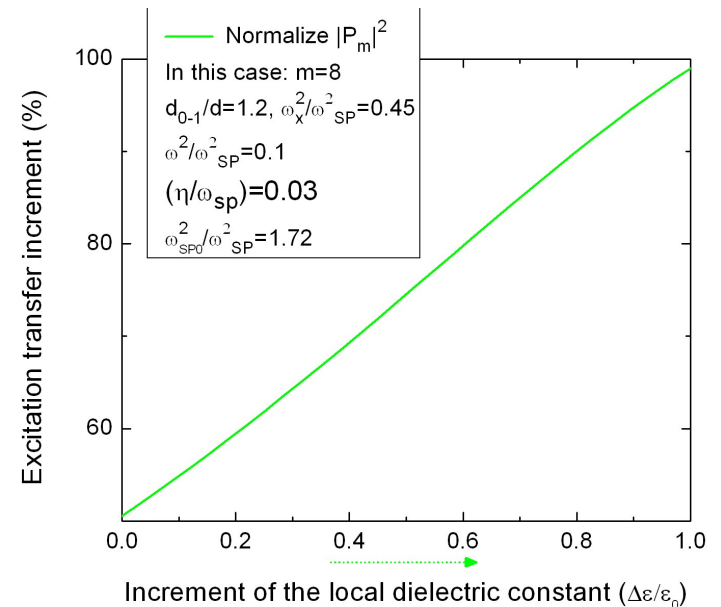
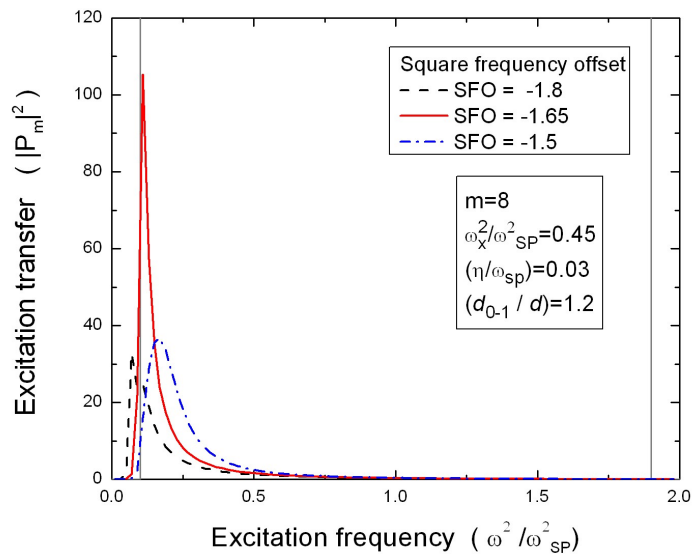
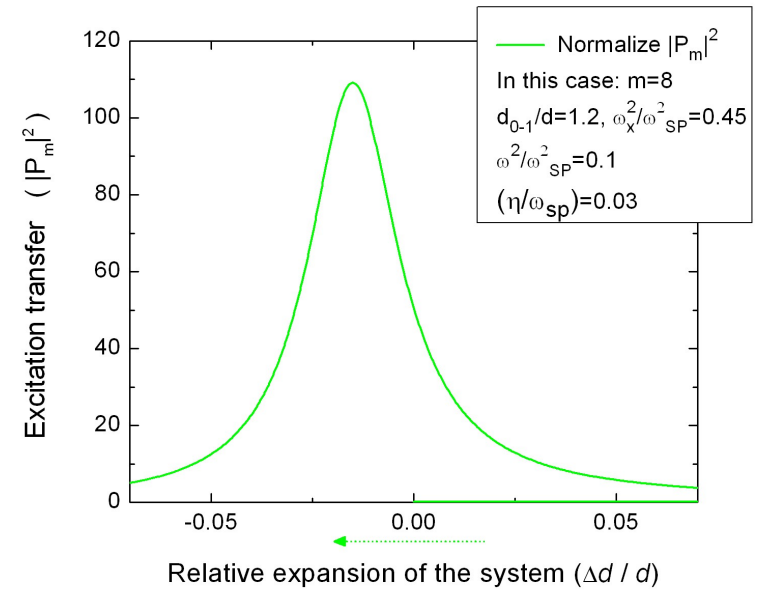
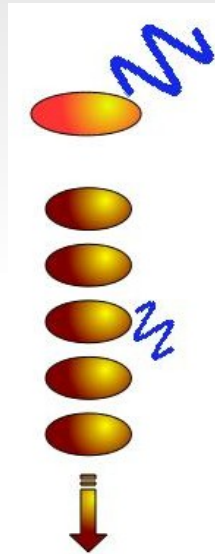
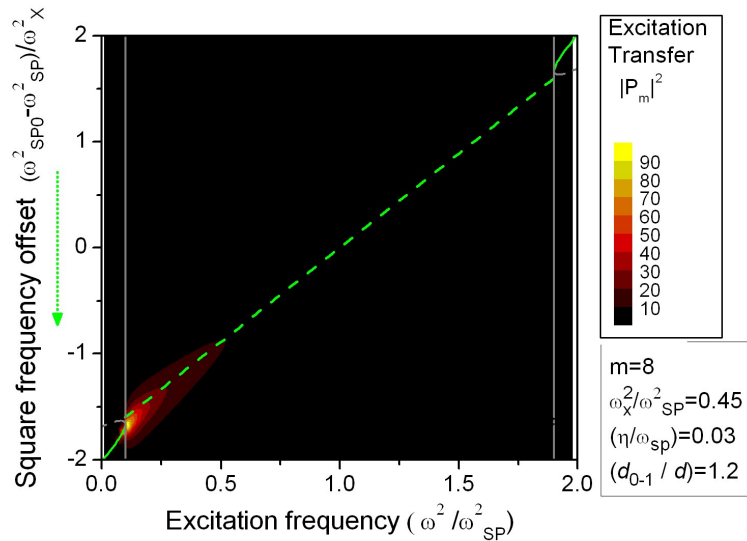
Using DPT for a Nano-ruler



2 Localized states



Using DPT to measure local dielectric constant and material expansion.



Conclusions:

Injection of excitations into waveguides.

- Contrary to common wisdom, the largest excitation transfer **does not** occur when the poles of the response function present the **biggest imaginary part** (resonant state) but when a virtual state is transformed into a localized state.
- Excitation transfer is controlled by dynamical phase transitions.
- Generality of the model => conclusions applicable to great number of physical situations.

Using dynamical phase transition in plasmonic.

- We show that even under realistic conditions, DPTs still provide new tools for plasmonics.
- Three examples were analyzed nano-rulers, dielectric constant sensors, and deformation sensors.

Plasmonic synchronization.

- Preliminary results shows that it is possible to build synchronized plasmonic circuits. Possible applications.... Initialization of plasmonic circuits?....

Thanks you very much.

References:

[1] Propagation in nanoparticle chain:

a) R. Baer, K. Lopata and D. Neuhausera, *Properties of phase-coherent energy shuttling on the nanoscale*, J. Chem Phys. 126, 014705 (2007). **b)** A. Alú and N. Engheta, *Theory of linear chain of metamaterial/plasmonic particles as subdiffraction optical nanotransmission lines*, Phys. Rev. B 74, 205436 (2006). **c)** S. Zou and G. Schatz, *Metal nanoparticles array waveguides: Proposed structures for subwavelength devices*, Phys. Rev. B. **74**, 125111 (2006). **d)** V. Markel and A. Sarychev, *Propagation of surface plasmons in ordered and disordered chains of metal nanospheres*, Phys. Rev. B **75**, 085426 (2007). **e)** D. S. Citrin, *Coherent excitation transport in metal-nanoparticles chain*, Nano Lett. 4, 1562 (2004). **f)** [4] M. Quinten, A. Leitner, J. R. Krenn, and F. R. Aussenegg, *Electromagnetic energy transport via linear chains of silver nanoparticles*, Opt. Lett. 23,1331 (1998).

[2] Localized Excitation:

a) M. Stockman, S. Faleev, and D. Bergman, *Coherent control of femtosecond energy localization in nanosystems*, Phys. Rev. Lett. **88**, 067402 (2002). **b)** X. Li and M.I. Stockman, *Highly efficient spatiotemporal coherent control in nanoplasmonics on a nanometer-femtosecond scale by time reversal*, Phys. Rev. B 77, 195109 (2008). **c)** S. Maier, P. Kik, H. Atwater, S. Meltzer, E. Harel, B. Koel, and A. Requicha, *Local detection of electromagnetic energy transport below the diffraction limit in metal nanoparticle plasmon waveguides*, Nature Materials **2**, 229 (2003). **d)** G. Colas des Francs and C. Girard, *Theory of near field optical imaging with a single fluorescent molecule used as a point-like detector*, Chem. Phys. **282**, 277 (2002). **e)** [2] A. V. Malyshev, V. A. Malyshev, and J. Knoester, *Frequency-Controlled Localization of Optical Signals in Graded Plasmonic Chains*, Nano Lett. 8, 2369 (2008).

[3] The model:

a) M. L. Brongersma, J. W. Hartman, and H. A. Atwater, *Electromagnetic energy transfer and switching in nanoparticle chain arrays below the diffraction limit*, Phys. Rev. B 62, R16 356 (2000). **b)** J. V. Hernandez, L. D. Noordam, and F. Robicieux , *Asymmetric response in a line of optically driven metallic nanospheres*, J. Phys. Chem. B 108, 15808 (2005).

[4] Virtual states and Green function formalism:

A. Dente, R. Bustos-Marún, and H. M. Pastawski, *Dynamical regimes of a quantum SWAP gate beyond the Fermi golden rule*, Phys. Rev. A **78**, 062116 (2008). **R. A. Bustos-Marun, E. A. Coronado, and H. M. Pastawski., Phys. Rev. B 82, 035434 (2010).**