

Supplemental Document: “Plasmon Resonant Enhancement of Dye Sensitized Solar Cells”

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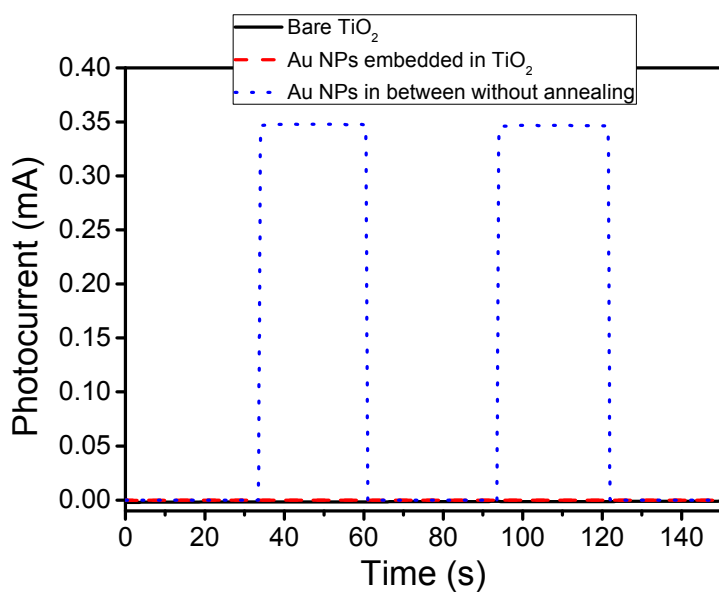


Figure S1. Short-circuit photocurrents of DSSCs with bare TiO₂, Au nanoparticles embedded in TiO₂ without dye molecules, and 5 nm Au thin film without the second annealing deposited between the TiO₂ layer and the dye monolayer as working electrodes.

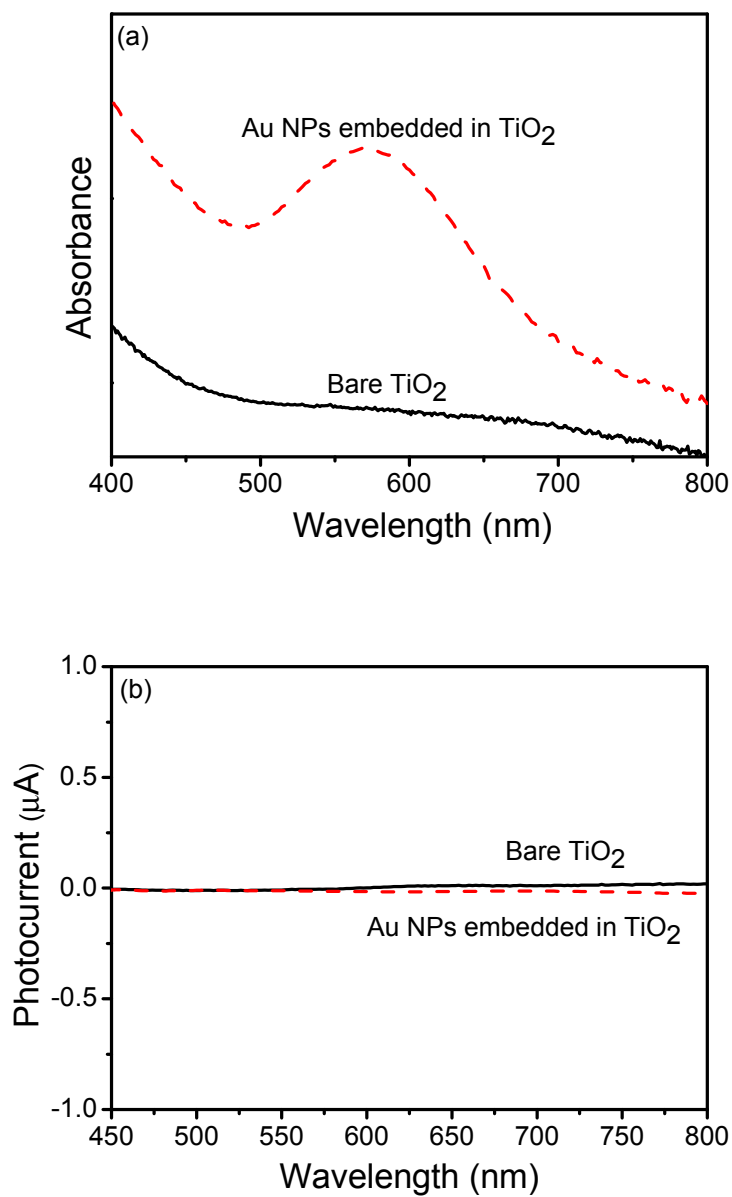


Figure S2. (a) Absorption spectra of bare TiO₂ and Au nanoparticles embedded in TiO₂ without dye molecules. (b) Photocurrent spectra of photovoltaic cells with bare TiO₂ and Au nanoparticles embedded in TiO₂ without dye molecules as working electrodes.

The power conversion efficiency η of the solar cells is determined by

$$\eta(\%) = \frac{V_{oc} I_{sc} FF}{P_{in} S} \times 100 \quad (1)$$

where V_{oc} is the open-circuit photovoltage, I_{sc} is the short-circuit photocurrent, and $P_{in} S$ is the incident laser power times the working electrode area (60 mW). The fill factor FF is given by

$$FF = \frac{V_m I_m}{V_{oc} I_{sc}} \quad (2)$$

where V_m and I_m are the voltage and the current at the maximum output power point, respectively.

Table S1 Comparison of photovoltaic device performances of Au nanoparticle/dye/TiO₂ configuration #2 with and without the second annealing.

Working Electrode	V _{oc} (V)	I _{sc} (mA)	FF(%)	η (%)
Without annealing	0.48	0.34	52	0.14
With annealing	0.73	0.86	58	0.60