

**Supplement information for Nanoplasmonic sensing to study CO and oxygen adsorption
and CO oxidation on size-selected Pt₁₀ clusters.**

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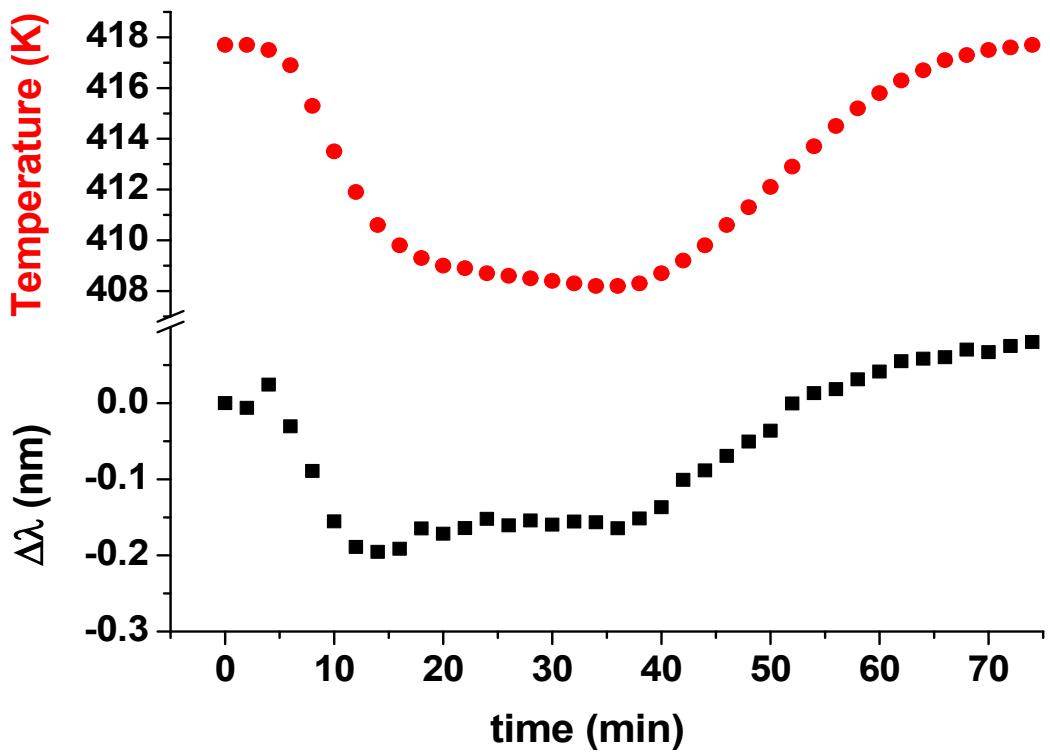


Fig. S1 Top: variation of the sample temperature (T) versus time. Bottom: Experimental LSPR shift $\Delta\lambda$ versus time of Au disks ($h = 17.5$ nm, $p = 300$ nm, $d = 140$ nm) covered with a SiO₂ layer (6 nm) deposited on an ITO film (21 nm) supported on a HQ-float glass window versus time. $\Delta\lambda$ is defined with respect to a reference which is the plasmonic sample at $T = 418$ K under high vacuum ($P \sim 10^{-6}$ Pa).

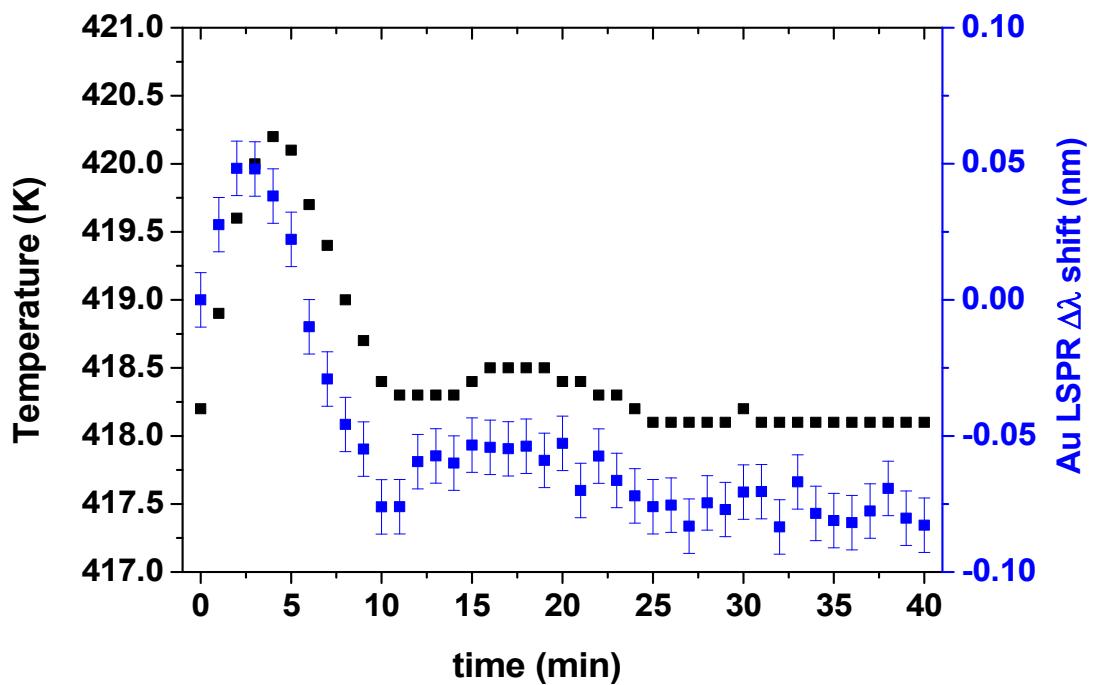


Fig. S2 Experimental Au LSPR shift and sample temperature versus time at $T = 418$ K and $P_{CO} = 2.3$ Pa for Pt_{10} clusters deposited on Au disks ($h = 17.5$ nm, $p = 300$ nm, $d = 140$ nm) covered with a SiO_2 layer (6 nm) and deposited onto an ITO-coated film (21 nm) glass window. The LSPR shift is defined with respect to a reference which is the plasmonic sample before CO adsorption. The accuracy of the wavelength measurements in the LSPR response is 0.01 nm (error bars).

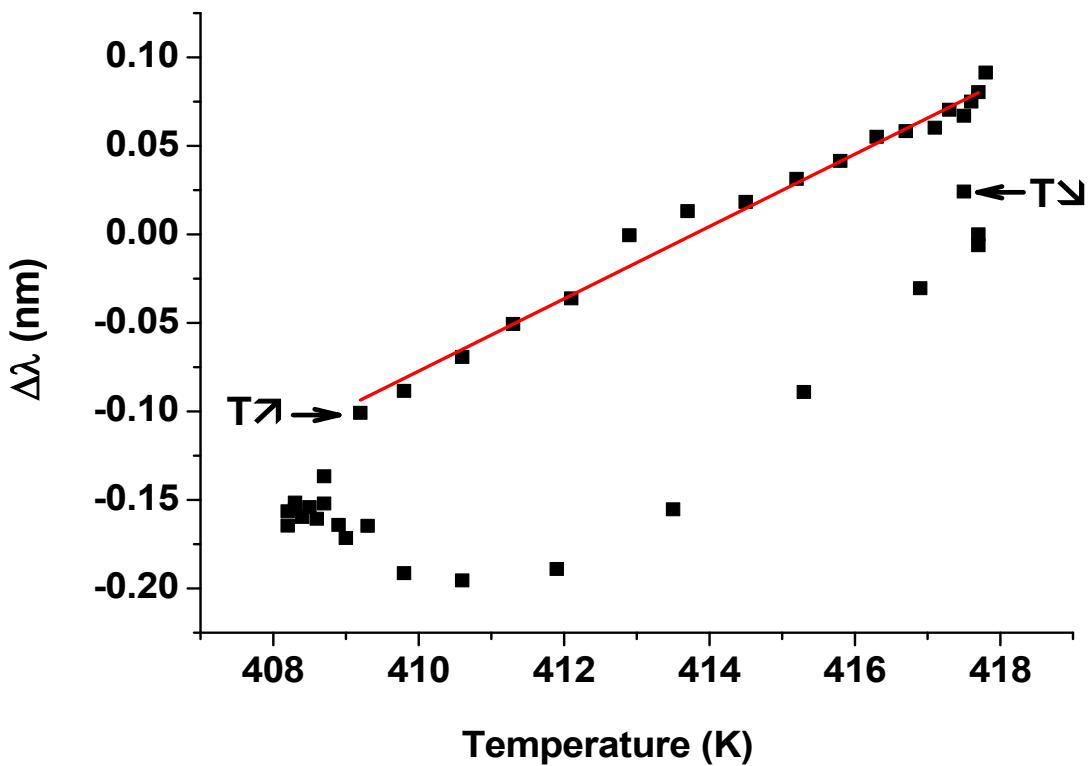


Fig. S3 Experimental LSPR shift $\Delta\lambda$ versus sample temperature (T) of Au disks ($h = 17.5$ nm, $p = 300$ nm, $d = 140$ nm) covered with a SiO₂ layer (6 nm) deposited on an ITO film (21 nm) supported on a HQ-float glass window. $\Delta\lambda$ is defined with respect to a reference which is the plasmonic sample at $T = 418$ K under high vacuum ($P \sim 10^{-6}$ Pa). There is a linear (fitted) part corresponding to the increase in T and a non-linear part corresponding to the decrease in T .