

Supplementary Materials

Using Surface Phonons as a Guide for Optimizing SERS and Light-Driven Processes †

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† This article is dedicated to Prof. Giuseppe Zerbi in recognition of his outstanding scientific contributions to Spectroscopy.

Figure S1. Raman spectra of T-horex@Au acquired with a 0.5 N.A. objective as a function of the power attenuation.

Figure S2. Temporal stability of the 700 cm⁻¹ mode.

Figure S3. Background-subtracted Raman spectra of Z-scan experiments and evolution of the 700 cm⁻¹ mode.

Figure S4. Far-field Vis reflectance of T-horex and T-horex@Au.

Figure S5. Raman spectrum of Au (8 nm) on glass slide substrates.

Using surface phonons as a guide for optimizing SERS and light-driven processes

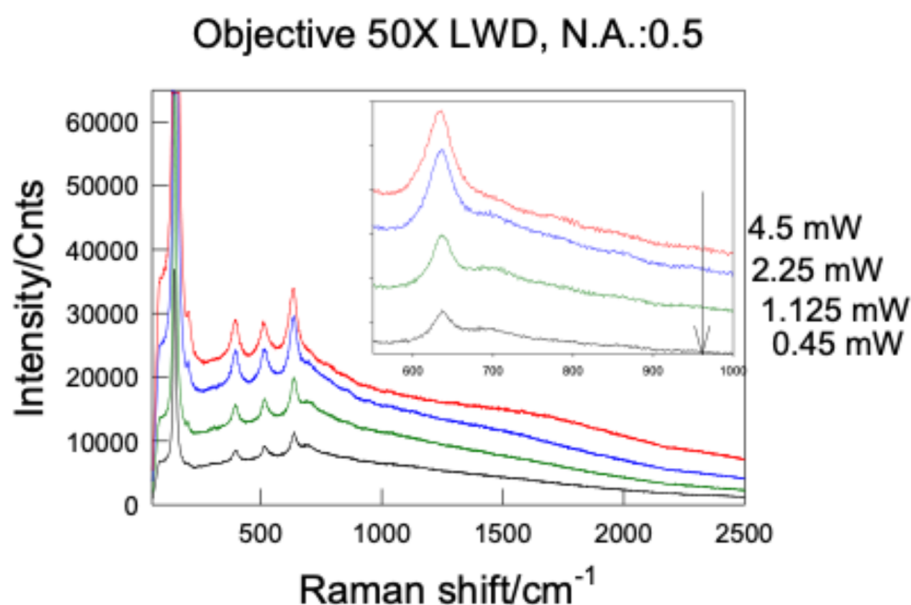


Figure S1. Raman spectra of T-horex@Au acquired with a 0.5 N.A. objective as a function of the power.



Using surface phonons as a guide for optimizing SERS and light-driven processes

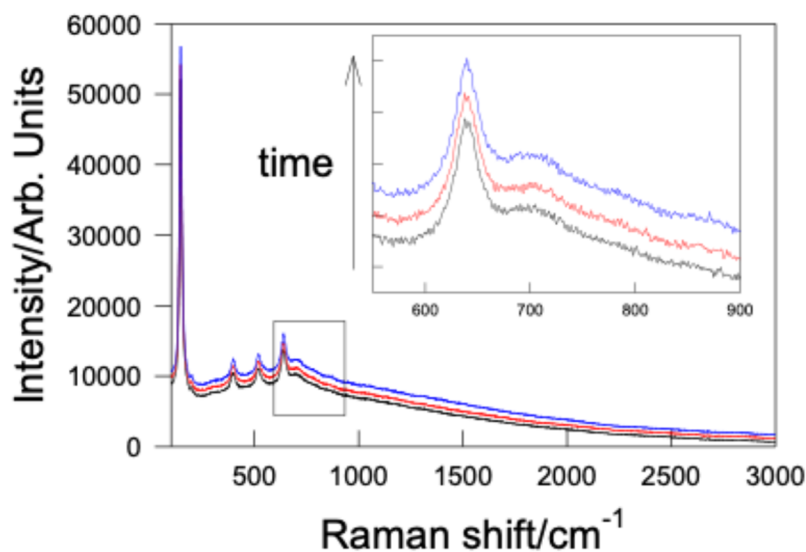


Figure S2. Temporal stability of the 700 cm^{-1} mode. The spectra have been stacked for clarity. The spectra were collected under the same conditions (power: 0.45 mW , acquisition time: 3 s) at $t = 0$ and after continuous irradiation for 5 and 10 min .

Using surface phonons as a guide for optimizing SERS and light-driven processes

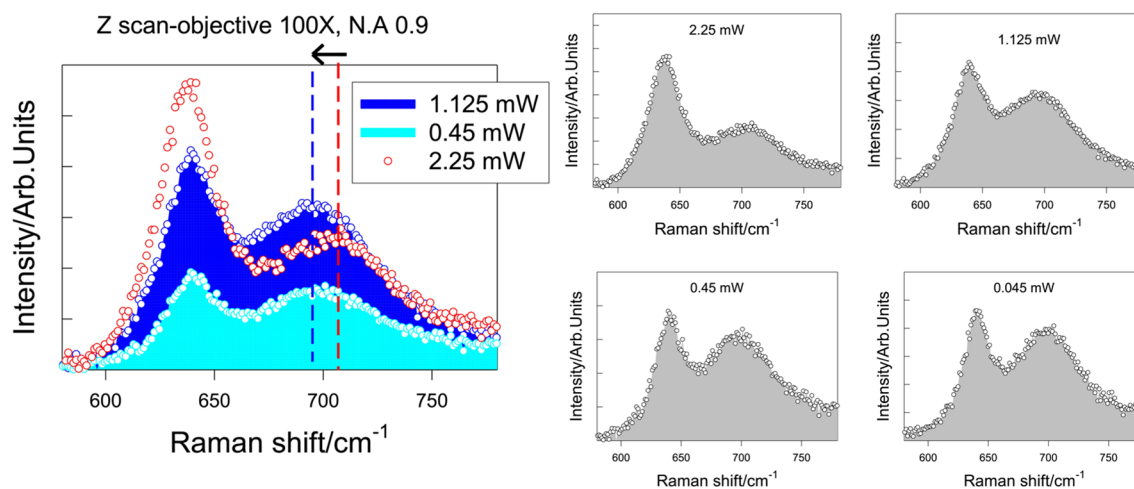


Figure S3. Background-subtracted Raman spectra of T-horex@Au acquired with a 0.9 N.A. objective as a function of the power (see Figure 3 in the main text). The phononic mode has been fitted with Pseudo-Voigt curves, the g -factor for 2.25 and 1.125 mW is 0.425 and 0.572 , respectively, whereas it is 0 in the other cases, which indicates a progressive increase of the band symmetry. The maximum of the center of the mode is also shifted by -5 cm^{-1} on passing from 2.25 to 1.125 mW .

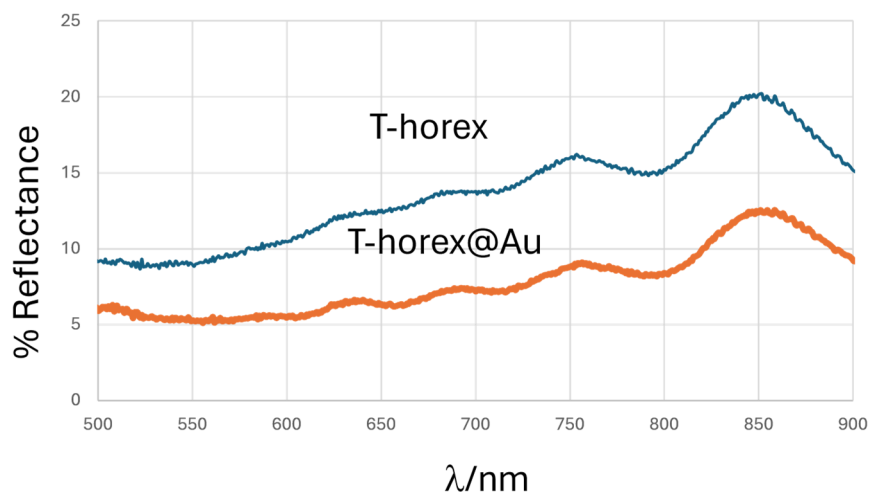


Figure S4. Far-field Vis reflectance of T-horex and T-horex@Au.

Using surface phonons as a guide for optimizing SERS and light-driven processes

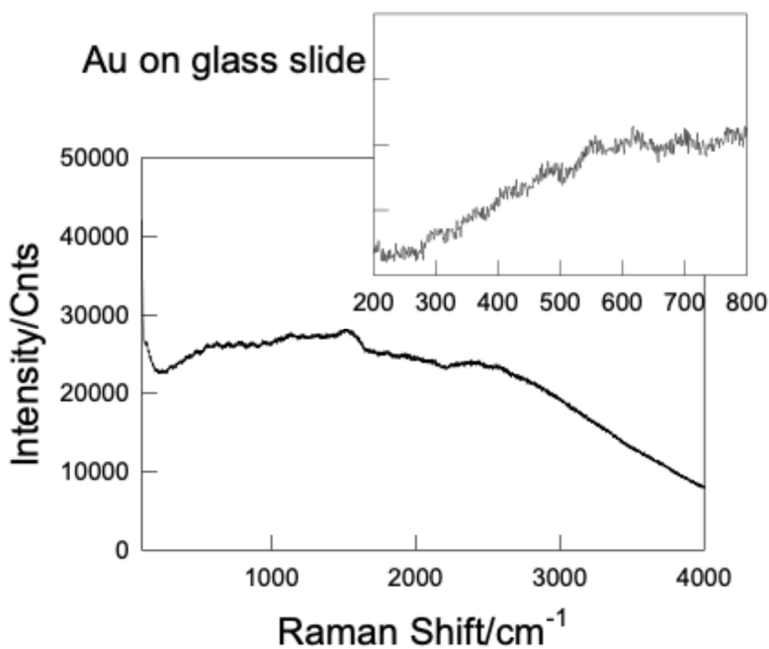


Figure S5. Raman spectrum of Au (8 nm) on glass slide substrates, acquired under the same conditions of Raman experiments show in Figure 5 of main text.