Supplementary Materials: SERS-Based Flavonoid Detection Using Ethylenediamine-β-Cyclodextrin as a Capturing Ligand

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Figure S1. Schematic illustration of the synthesis of Et-β-CD@Ag@SiO2 NPs.



Figure S2. 1D NMR spectra of Et- β -CD. (a) ¹H NMR spectrum; (b) ¹³C NMR spectrum. The NMR experiments were carried out on a Brucker Avance spectrometer at 600 MHz in a D₂O solution at 25 °C.



Figure S3. UV-Vis absorbance of SiO₂@Ag@Et-β-CD NPs reacted with Nar, Hes, Lut, and Que.



Figure S4. TEM images of Ag@SiO₂ NPs in EtOH Ag@SiO₂ NPs measured using an energy-filtering transmission electron microscope (EF-TEM, LIBRA 120, Carl Zeiss, Oberkochen, Germany). The accelerating voltage was 120 kV.



Raman Shift (nm)

Figure S5. Surface-enhanced Raman scattering (SERS) spectra. (a) SERS spectra of organic molecu-le s after mixing with Et- β -CD@Ag@SiO₂ NPs (i. no target, ii. ethylene glycol, iii. β -estradiol, iv. Isop-ro pyl alcohol, v. naphthalene, vi. toluene and vii. Lut); (b) SERS spectra of (i) SiO₂@Ag NPs, (ii) SiO₂@ Ag NPs reacted with Aniline, (iii) SiO₂@Ag@Et- β -CD reacted with Aniline, and (iv) SiO₂@Ag@Et- β --CD NPs reacted with mixing solution of Lut and aniline.



Figure S6. Surface-enhanced Raman scattering (SERS) spectra and normalized SERS intensity graph. SERS spectra of SiO₂@ Ag NPs mixed with Lut at 1×10^{-2} M to 1×10^{-7} M.