

Supporting Information

**Silica/Titania Sandwich-like Mesoporous Nanosheets Embedded
with Metal Nanoparticles Templated by Hyperbranched
Poly(ether amine) (hPEA)**

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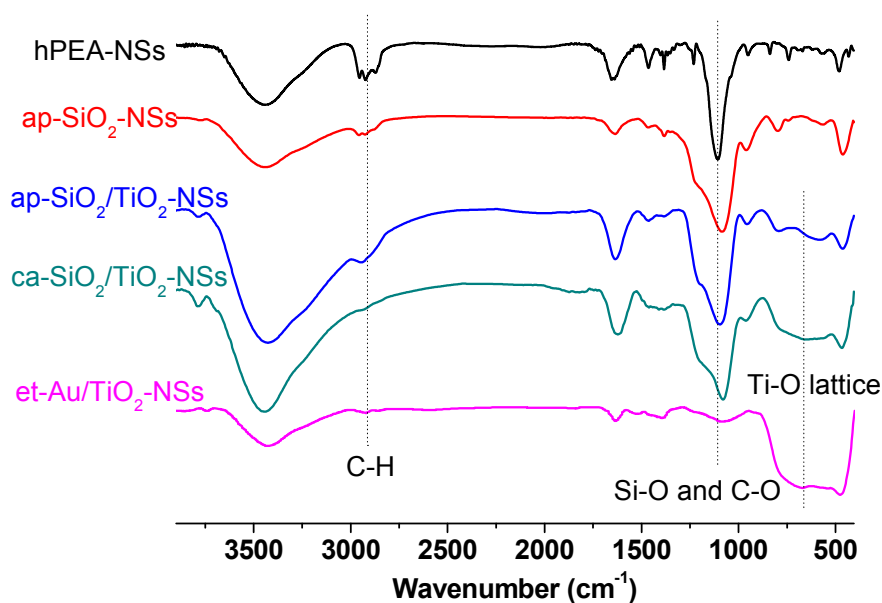


Figure S1. FT-IR spectra of hPEA-NSs, ap-SiO₂-NSs, ap-SiO₂/TiO₂-NSs, ca-SiO₂/TiO₂-NSs, and et-Au/TiO₂-NSs, respectively.

The FTIR spectra of SiO₂-NSs and SiO₂/TiO₂-NSs were also investigated (Figure S1). Compared with the template hPEA-NSs, the peak at 1098 cm⁻¹ assigned to Si-O stretching vibration was obviously strengthened in the FTIR spectrum of ap-SiO₂-NSs, indicating the SiO₂ has been deposited on hPEA-NSs. Meanwhile, the very broad peak at 450-800 cm⁻¹ assigned to Ti-O lattice vibration appeared, which

proved TiO₂ grains had grown on the nanosheets in ap-SiO₂/TiO₂-NSs. After calcinations, the C-H stretching vibration at 2941 cm⁻¹ nearly disappeared which indicated the hPEA-NSs template had been removed. The etching of the SiO₂ middle-layer was exhibited by the nearly disappearance of the Si-O stretching vibration at 1098 cm⁻¹.

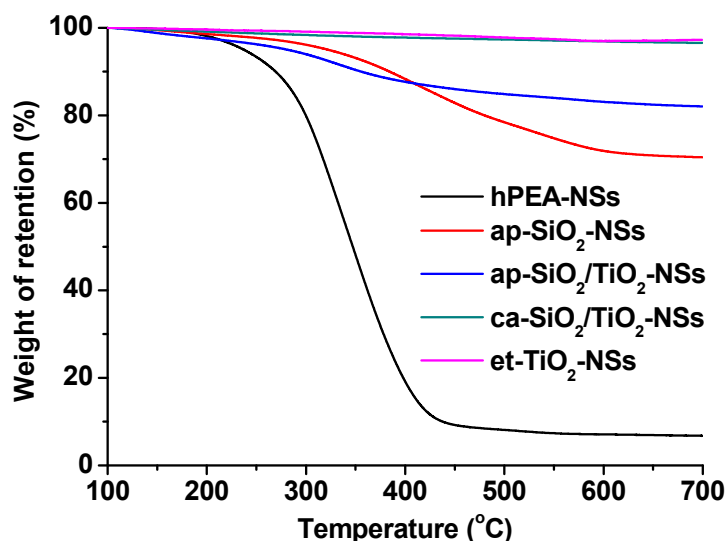


Figure S2. TGA spectra of hPEA-NSs, ap-SiO₂-NSs, ap-SiO₂/TiO₂-NSs, ca-SiO₂/TiO₂-NSs, and et-TiO₂-NSs, respectively.

As shown in Figure S2, the first weight loss below 150 °C was mostly due to water evaporation from the nanosheets, and the second weight loss occurring at 200-450 °C was attributed to the decomposition of hPEA-NSs templates. The weight of retention increased while SiO₂ and TiO₂ were deposited on hPEA-NSs, respectively, which indicated the existence of SiO₂ and TiO₂ in ap-SiO₂-NSs and ap-SiO₂/TiO₂-NSs, respectively. The samples after calcinations show only 3% weight loss at higher temperature, which implies the completely removing of the hPEA-NSs templates. The weight percentage of SiO₂ and TiO₂ can also be calculated from the TGA results.

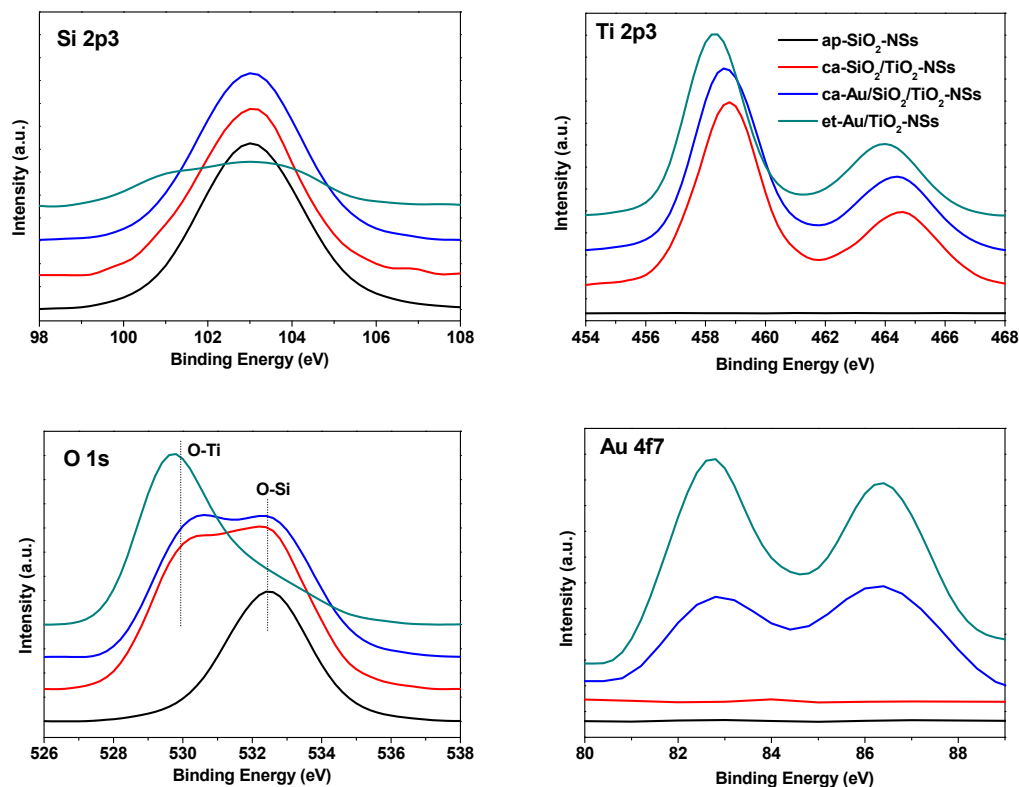


Figure S3. High-resolution XPS spectra of Si 2p₃, Ti 2p₃, O 1s, Au 4f₇ for ap-SiO₂-NSs, ca-SiO₂/TiO₂-NSs, ca-Au/SiO₂/TiO₂-NSs, and et-Au/TiO₂-NSs, respectively.

The chemical composition of the samples was also analyzed by XPS, and the XPS spectra were shown in Figure S3. There are two peaks at the binding energies of 464.4 and 458.6 eV for Ti 2p XPS spectra of SiO₂/TiO₂-NSs, which can be assigned to Ti 2p_{1/2} and Ti 2p_{3/2}, respectively. This result indicates the existence of TiO₂ in these SiO₂/TiO₂-NSs. There are two peaks at the binding energies of 529.8 and 532.4 eV for O 1s in XPS spectra of SiO₂/TiO₂-NSs, which can be assigned to O 1s for TiO₂ and SiO₂, respectively. The peak for O 1s of et-Au/TiO₂-NSs at 532.4 eV was weakened compared with SiO₂/TiO₂-NSs, which indicates the successful etching of SiO₂ in Au/SiO₂/TiO₂-NSs, and this result is consistent with the weakening of the peak at the binding energy of 103.0 eV for Si 2p₃ XPS spectrum of et-Au/TiO₂-NSs. Meanwhile, the binding energy of Ti 2p_{1/2} and Ti 2p_{3/2} for et-Au/TiO₂-NSs shift about 0.6 eV to the lower energy direction, and this phenomenon may be attributed to the heterostructure effect between the SiO₂ layer and TiO₂ layer in Au/SiO₂/TiO₂-NSs. Moreover, the embedding of AuNPs can be confirmed by the appearance of two peaks

at 82.8 and 86.4 eV, which correspond to Au 4f_{5/2} and Au 4f_{7/2}, respectively in XPS spectra of Au/SiO₂/TiO₂-NSs.