

Supporting Information

Cobalt-bilayer Catalysts Decorated Ta₃N₅ Nanorod Array as Integrated Electrodes for Photoelectrochemical Water Oxidation

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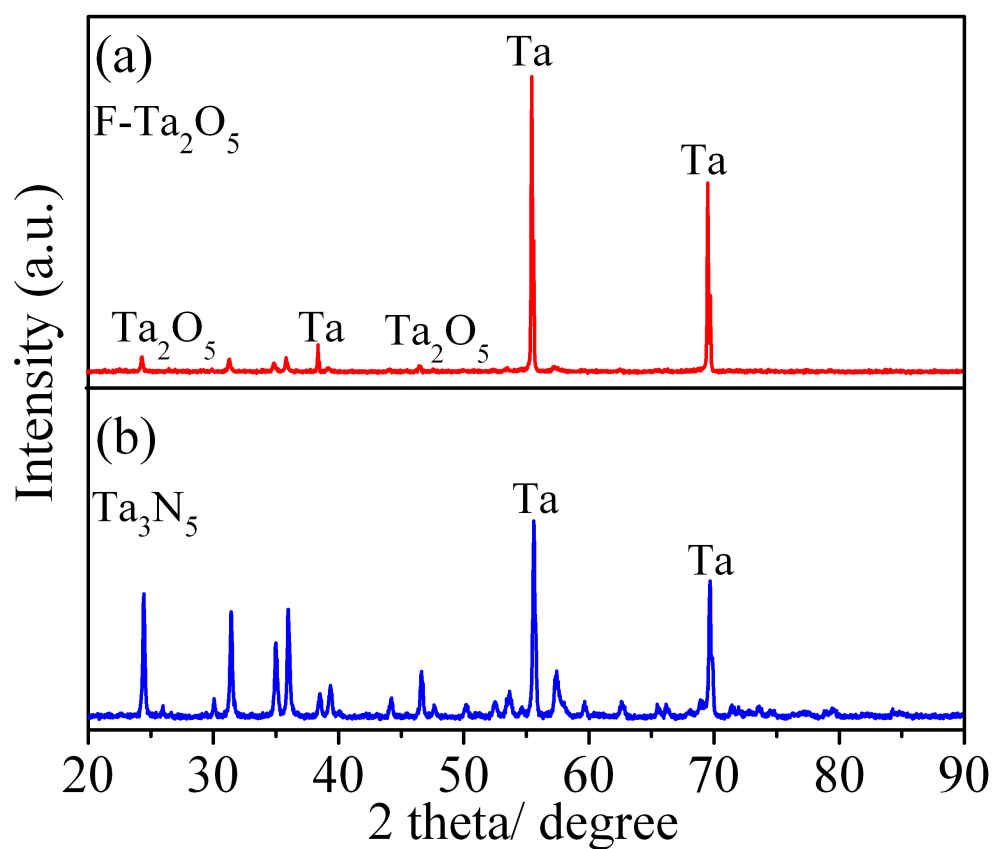


Figure S1. XRD patterns of (a) F-Ta₂O₅ nanorod arrays and (b) Ta₃N₅ nanorod arrays on the Ta substrates.

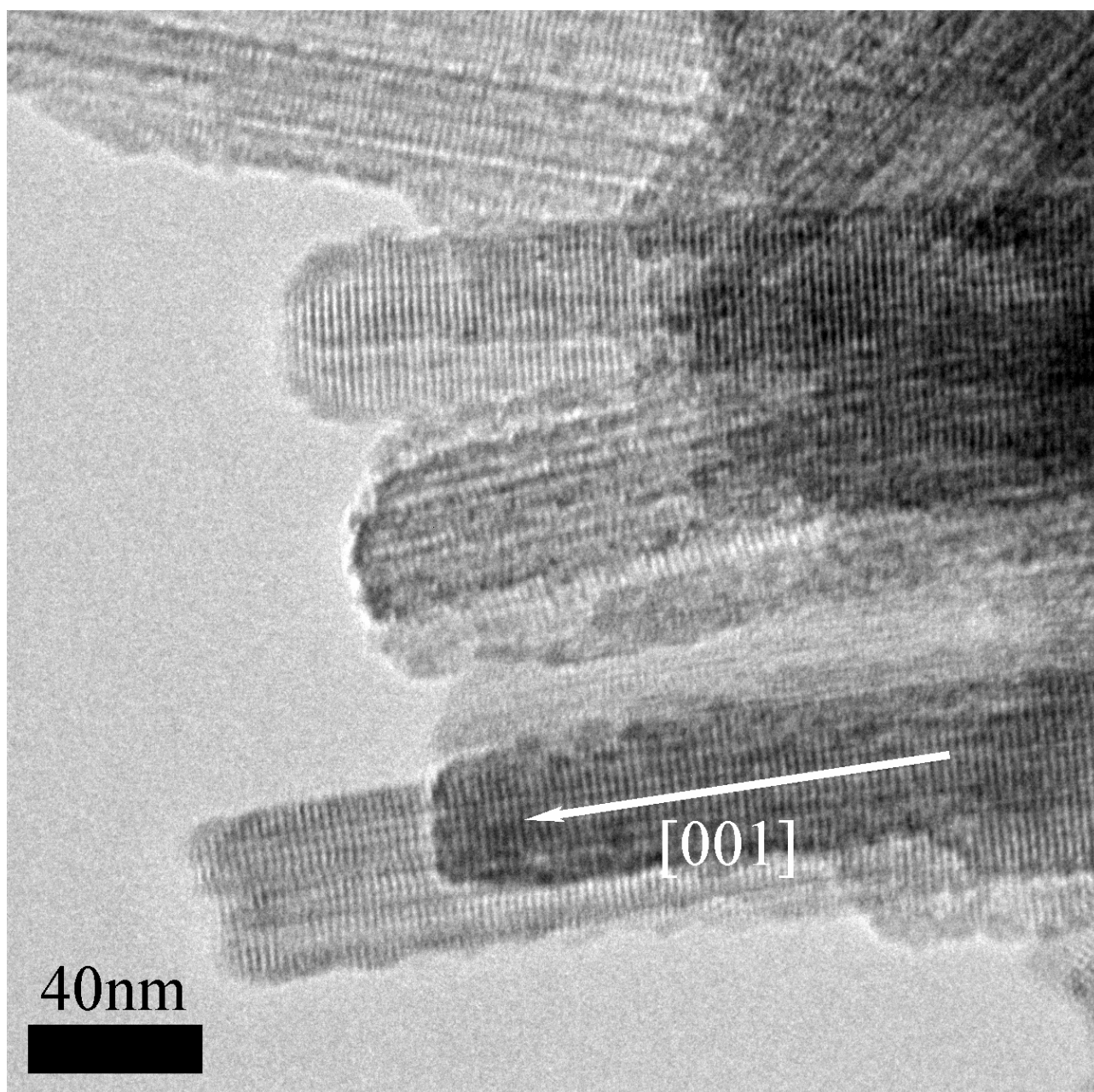


Figure S2. High resolution TEM image recorded from a single F-Ta₂O₅ nanorod with preferential growth along the [001] direction.

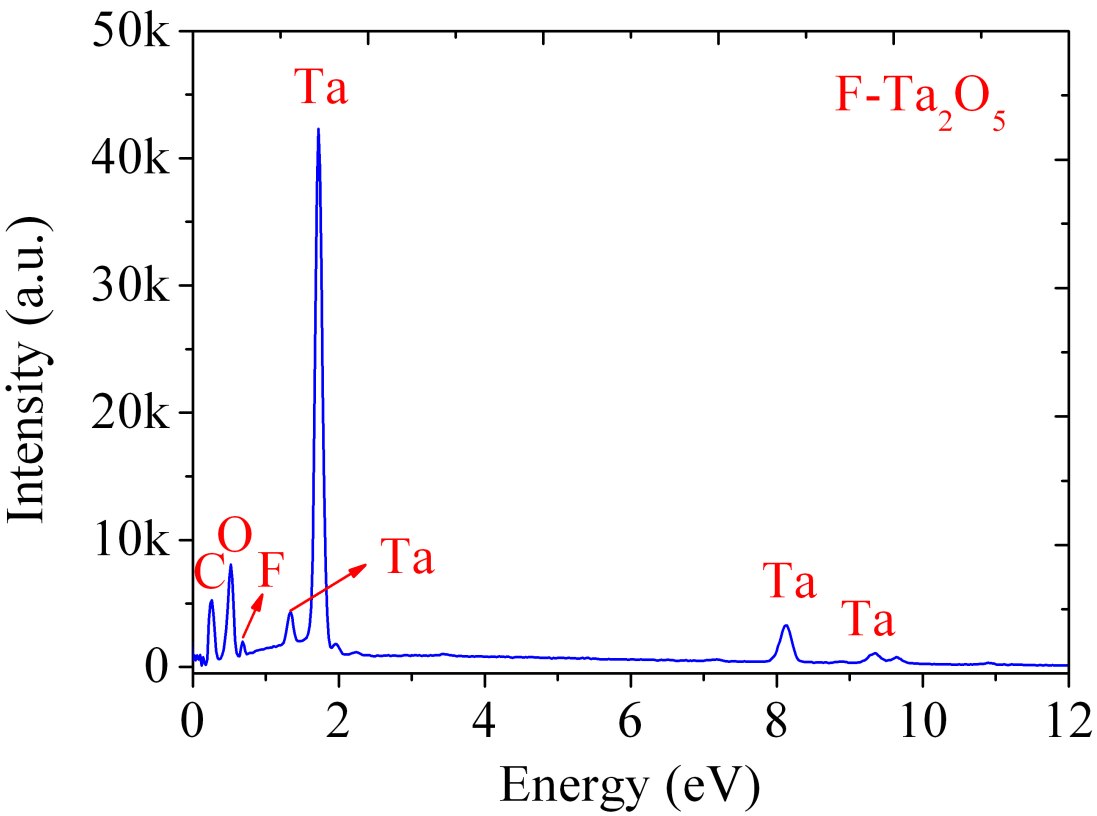


Figure S3. Energy-dispersive X-ray spectroscopy of the elements Ta, O and F in the F-Ta₂O₅ nanorod samples.

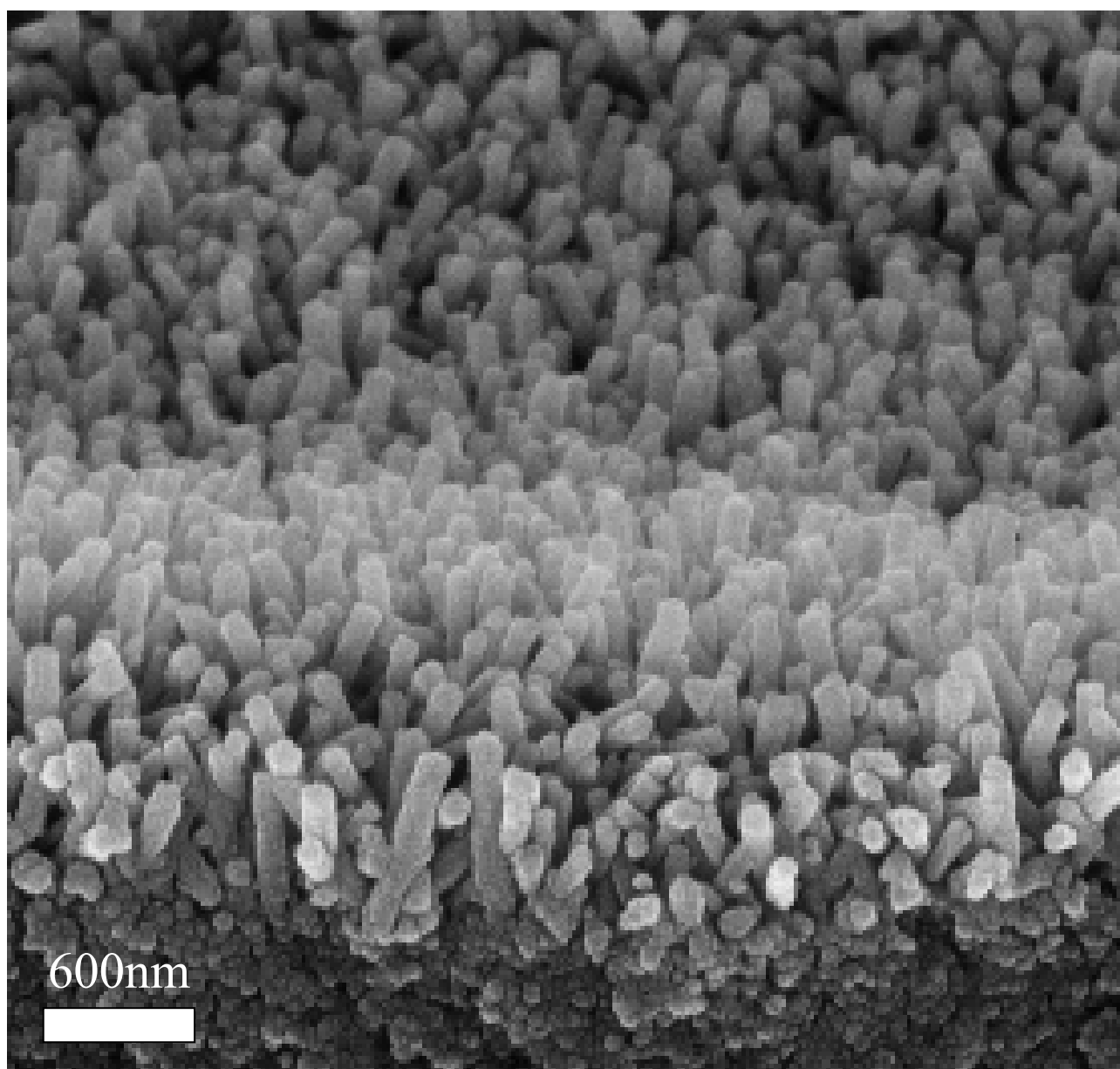


Figure S4. Typical cross-sectional SEM image of the Ta₃N₅ nanorod array.

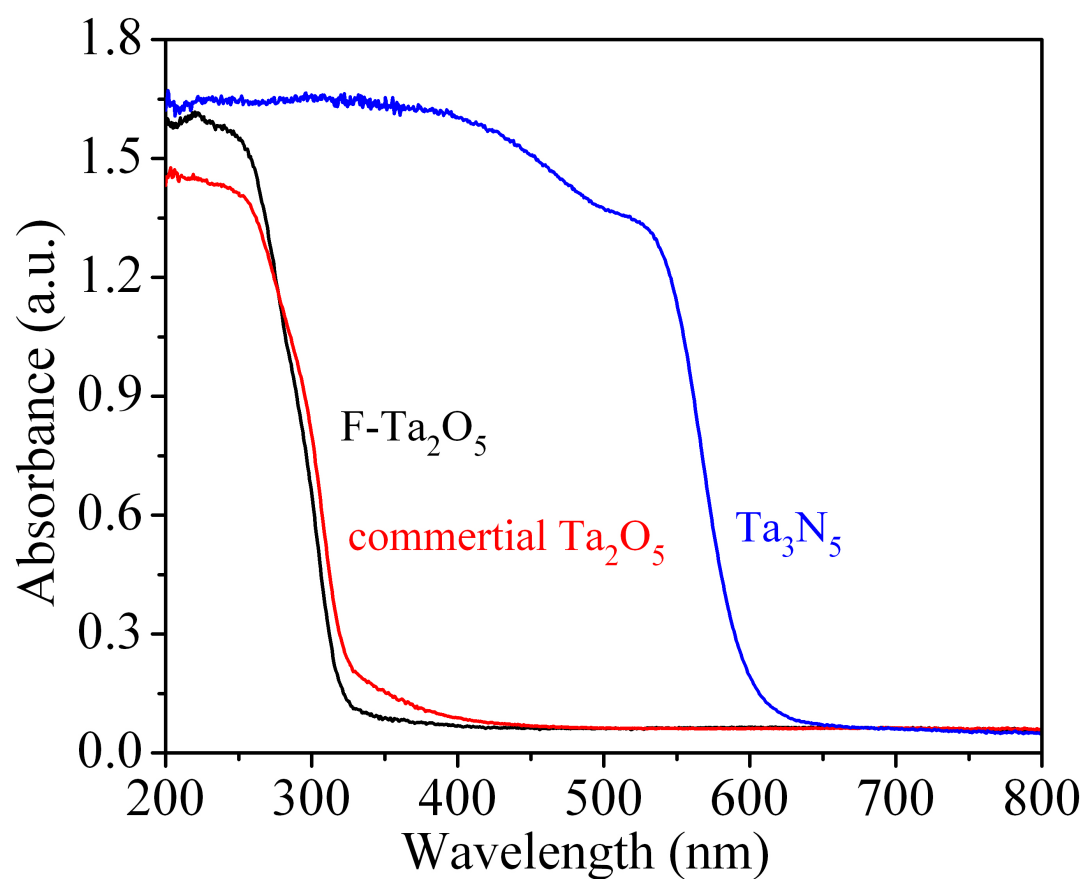


Figure S5. UV-visible absorption spectrum of commercial Ta₂O₅, F-Ta₂O₅ nanorods, and Ta₃N₅ nanorods samples.

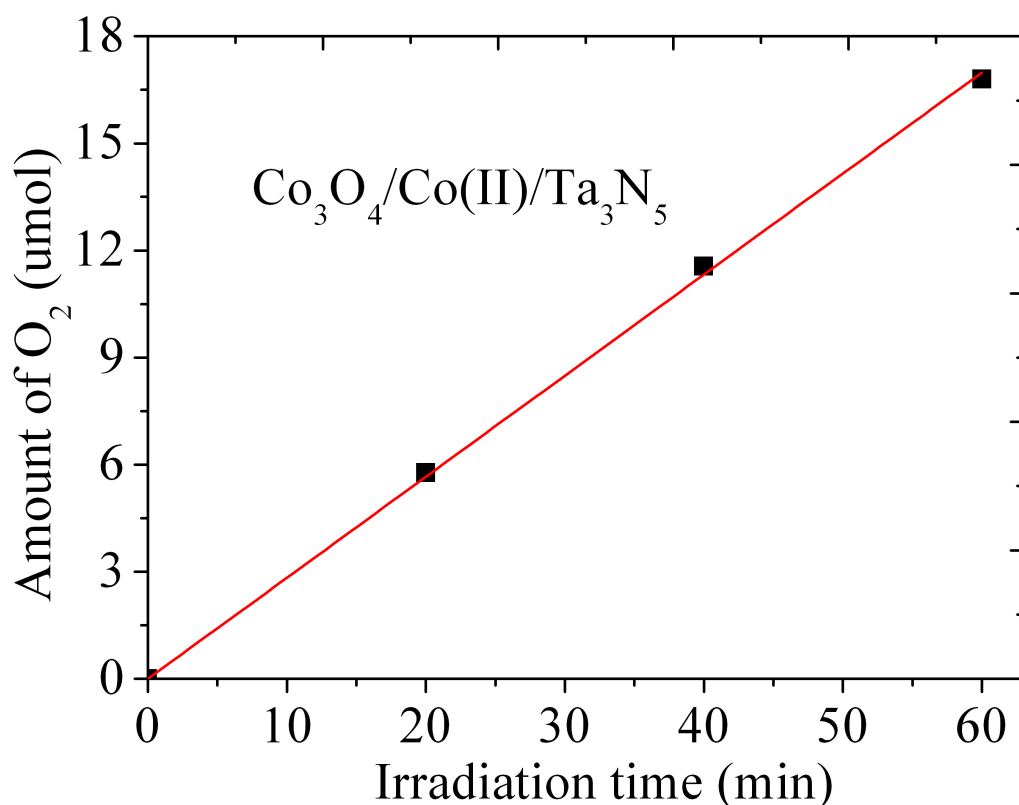


Figure S6. The O₂ production of Co₃O₄/Co(II)/Ta₃N₅ nanorods measured in 1.0 M NaOH solution (pH=13.6) in the three electrode system at an applied potential of 1.23 V vs. RHE AM 1.5G simulated sunlight at 100 mW/cm². The oxygen evolution by photoelectrochemical water splitting was conducted in the airtight reactor connected to a closed gas circulation system. The 300W Xe lamp (MAX-302, Asahi Spectra) under AM 1.5G illumination with a density of 100 mW cm⁻² (Newport) was used to irradiate only visible light. The amount of oxygen was determined by a gas chromatography (GC-3240) equipped with TCD (molecular sieve 5 Å column, Ar carrier gas).

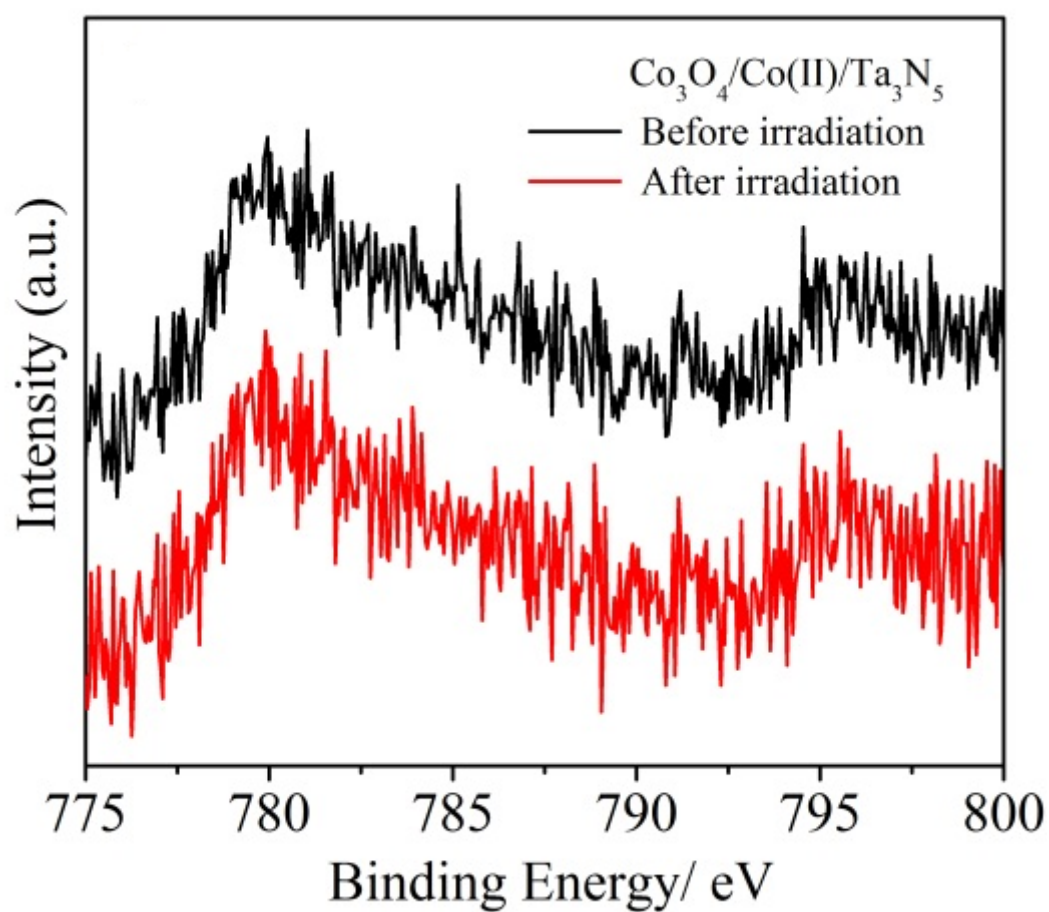


Figure S7. XPS spectra of Co 2p for Co₃O₄/Co(II)/Ta₃N₅ nanorods.