Supporting Information

Surface Grafting of Ru(II) Diazonium-Based Sensitizers on Metal Oxides Enhances Alkaline Stability for Solar Energy Conversion

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Table of contents:

- Figure S1. TEM images of MOx nanoparticles
- Figure S2. Extinction coefficient of the ruthenium sensitizers
- Figure S3. non-normalized UV-vis spectra of MOx-bound sensitizers
- Figure S4. XPS spectra of $Ru-N_2^+$ grafted on SnO₂, ZnO and ZrO₂.
- **Figure S5.** Full spectra of Ru- N_2^+ on TiO₂ under 24 hr illumination

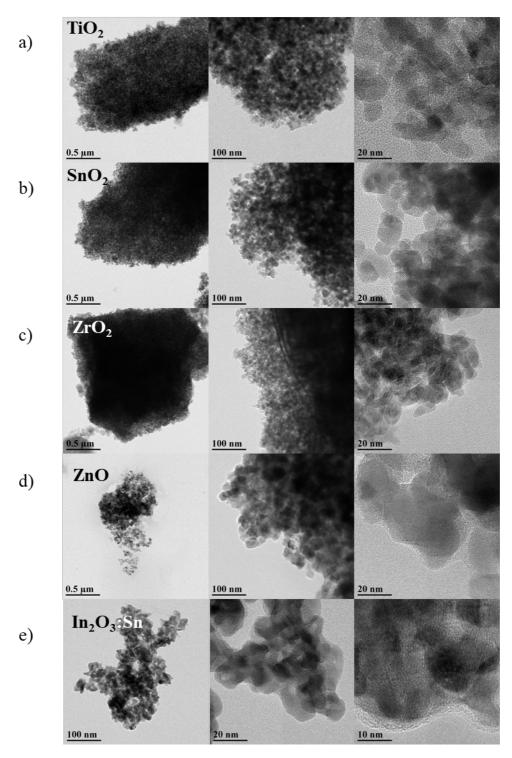


Figure S1: TEM images of MOx nanoparticles scraped from thin films. Types of MOx are separated as a) TiO_2 , b) SnO_2 , c) ZrO_2 , d) ZnO, and e) In_2O_3 :Sn.

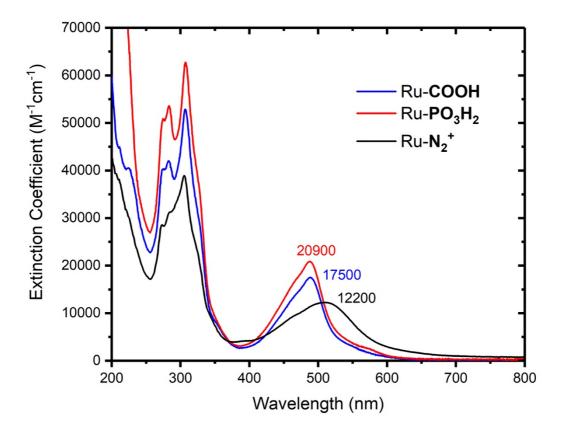


Figure S2: Extinction coefficients spectra of Ru-N_2^+ (black), Ru-PO_3H_2 (red), and Ru-COOH (blue) dissolved in CH₃CN.

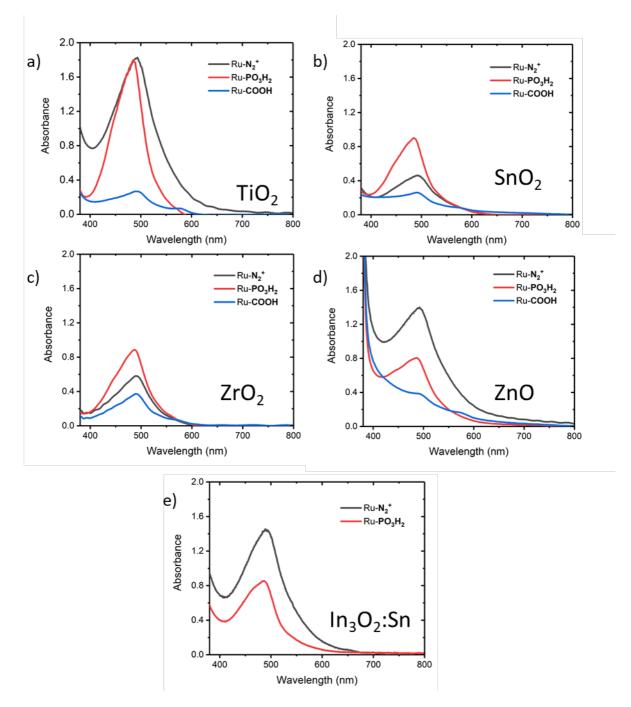


Figure S3: UV-Vis spectra of all sensitizers bound on MOx surfaces showing relative absorbance intensity. Ru-N_2^+ (black), $\text{Ru-PO}_3\text{H}_2$ (red), and Ru-COOH (blue) are shown for a) TiO₂, b) SnO₂, c) ZrO₂, d) ZnO, and e) In₂O₃:Sn. Ru-N₂⁺ grafted films were made by electrografting at held potentials given in Table 1 using a 2 mM Ru-N₂⁺ solution in 100 mM TBAClO₄ CH₃CN for 30 min. Ru-PO₃H₂ and Ru-COOH were adsorbed by soaking films in concentrated solutions in CH₃CN.

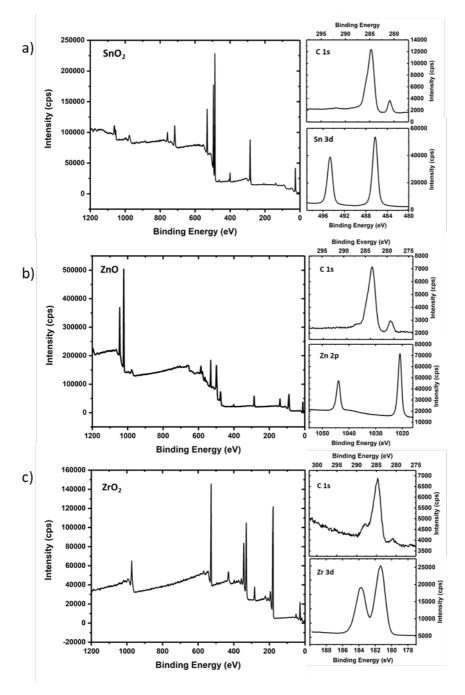


Figure S4: XPS spectra of a) SnO_2 , b) ZnO, and c) ZrO₂ films sensitized with Ru-N₂⁺. High definition spectral regions verify the presence of the sensitizer through the presence of a Ru3d peak (C1s) and show no detectable change in oxidation state of the metal.

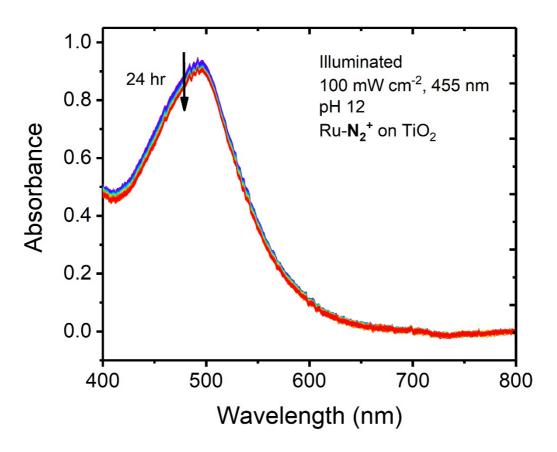


Figure S5: UV-visible spectra of Ru-N_2^+ on TiO₂ in pH 12 aqueous solution as it is illuminated with 455 nm, 100 mW cm⁻² light over 24 hrs. No spectral shift is observed, and little change in absorbance occurs.