Supporting Information for

Arrangement effect of di- μ -oxo dimanganese catalyst and $\text{Ru}(\text{bpy})_3^{2+}$ photoexcitation centers adsorbed in mica on visible-light-derived water oxidation

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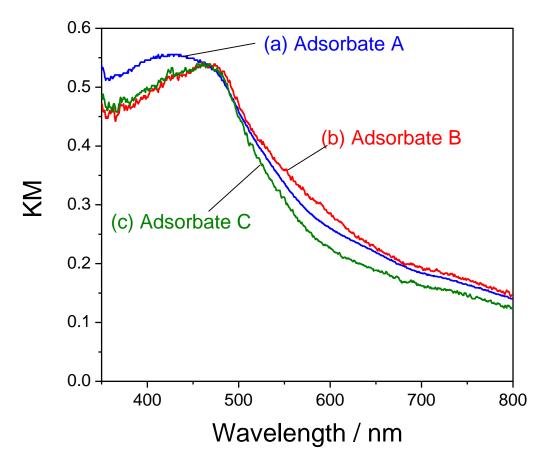


Figure S1. UV-vis DR spectra of the mica adsorbates (20 mg). (a) Adsorbate A (mica / **1** / Ru(bpy)₃²⁺) ($w_{\text{Mn}} = 164 \, \mu \text{mol g}^{-1}, \, w_{\text{Ru}} = 30 \, \mu \text{mol g}^{-1}$), (b) Adsorbate B (mica / Ru(bpy)₃²⁺ / **1**) ($w_{\text{Mn}} = 164 \, \mu \text{mol g}^{-1}, \, w_{\text{Ru}} = 100 \, \mu \text{mol g}^{-1}$), and (c) Adsorbate C (mica / (**1** + Ru(bpy)₃²⁺)) ($w_{\text{Mn}} = 164 \, \mu \text{mol g}^{-1}, \, w_{\text{Ru}} = 75 \, \mu \text{mol g}^{-1}$)

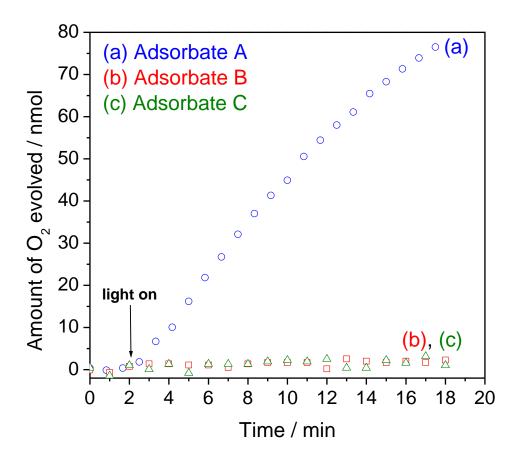


Figure S2. Time courses of the amount of O₂ evolved in photochemical water oxidation in an aqueous suspension of 0.2 M acetate buffer (2.0 ml, pH = 6.2) and 15 mM S₂O₈²⁻ containing (a) Adsorbate A (mica / **1** / Ru(bpy)₃²⁺) (w_{Mn} = 164 μmol g⁻¹, w_{Ru} = 30 μmol g⁻¹, 20 mg mica), (b) Adsorbate B (mica / Ru(bpy)₃²⁺ / **1**) (w_{Mn} = 164 μmol g⁻¹, w_{Ru} = 100 μmol g⁻¹, 20 mg mica), and (c) Adsorbate C (mica / (**1** + Ru(bpy)₃²⁺)) (w_{Mn} = 164 μmol g⁻¹, w_{Ru} = 75 μmol g⁻¹, 20 mg mica). A 150 W halogen lamp with a UV-cut filter, (λ > 420 nm, 127 mW cm⁻²) was used as a light source.