

Supporting Information

Solar water splitting over Rh_{0.5}Cr_{1.5}O₃-loaded AgTaO₃ of a valence-band-controlled metal oxide photocatalyst

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Table S1 Photocatalytic water splitting over $\text{Rh}_{0.5}\text{Cr}_{1.5}\text{O}_3(0.3 \text{ wt}\%)/\text{AgTaO}_3$ synthesized by a solid state reaction with various excess amount of Ag under various calcination conditions under UV irradiation.

Calcination condition		Excess Ag / mol%	Activity / $\mu\text{mol h}^{-1}$	
Temperature / K	Time / h		H ₂	O ₂
1173	10	0	130	166
1273	5	0	156	80
1273	10	0	171	87
1273	15	0	181	95
1273	15	3	149	78
1273	15	5	180	95
1273	15	7	151	76
1273	20	0	163	83
1373	10	0	120	60

Photocatalyst: 0.3 g, reactant solution: distilled water (120 mL), cell: top-irradiation cell with a Pyrex

window, light source: 300 W Xe-arc lamp ($\lambda > 300 \text{ nm}$).

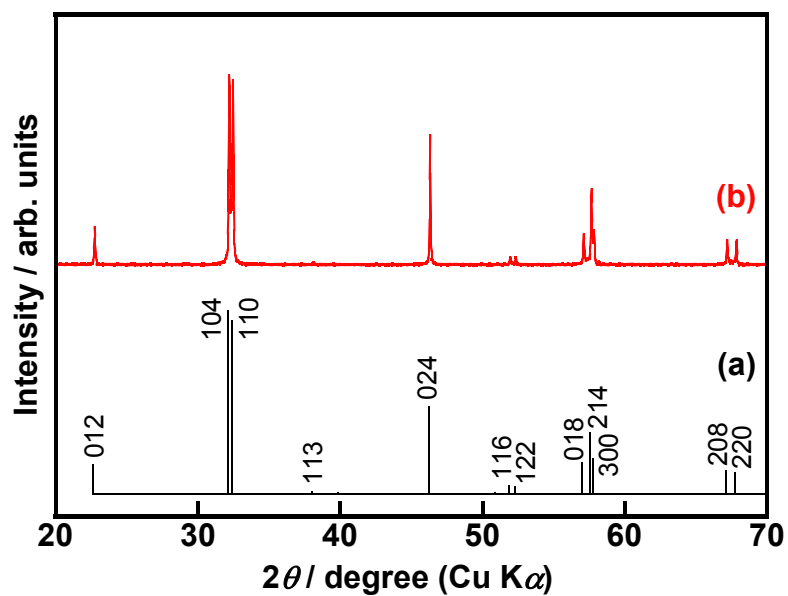


Fig. S1 (a) PDF of trigonal AgTaO₃ (1-72-1383). (b) XRD pattern of AgTaO₃ prepared by a solid state reaction at 1273 K for 15 h without excess Ag. Peaks due to K α_2 were removed.

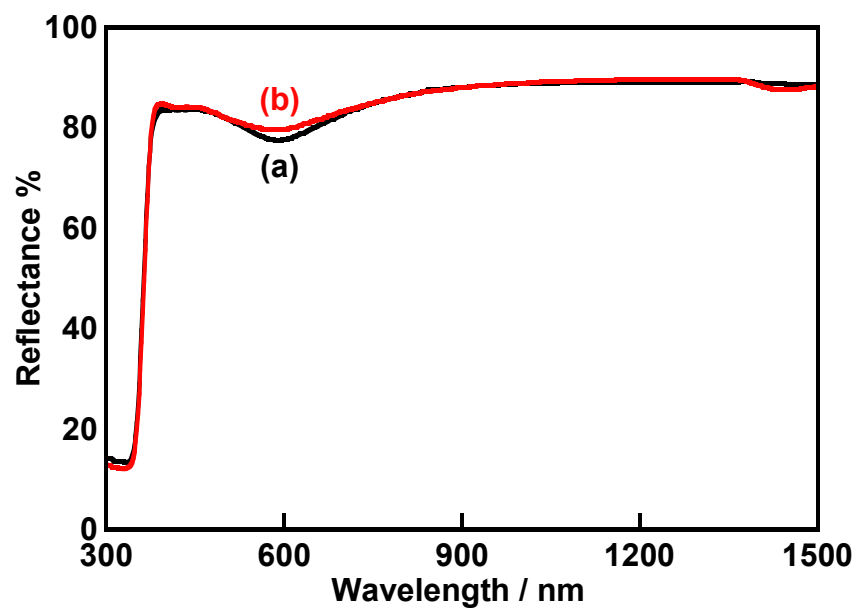


Fig. S2 A diffuse reflectance spectrum of AgTaO₃ (a) before and (b) after washing with an aqueous HNO₃ solution. AgTaO₃ was prepared by a solid state reaction at 1273 K for 15 h without excess Ag.

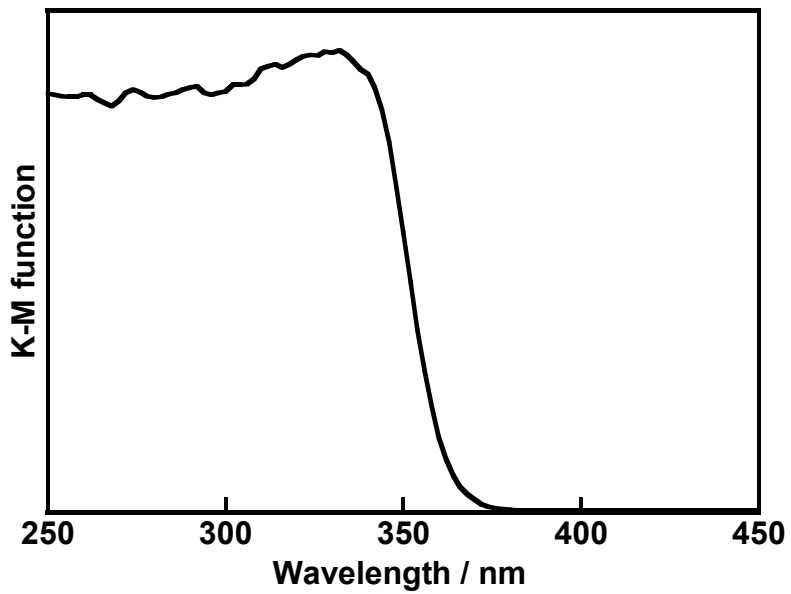


Fig. S3 A diffuse reflectance spectrum of AgTaO_3 prepared by a solid state reaction at 1273 K for 15 h without excess Ag.

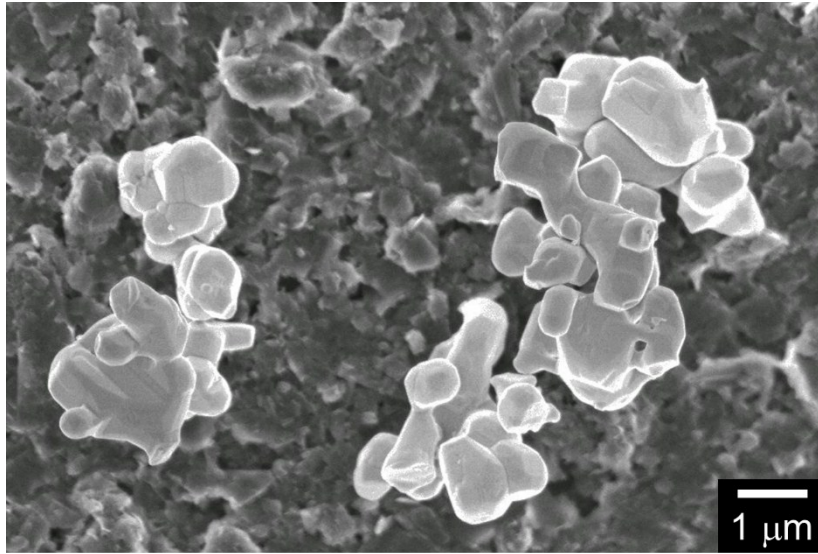


Fig. S4 Scanning electron microscopy image of AgTaO₃ prepared by a solid state reaction at 1273 K for 15 h without excess Ag.

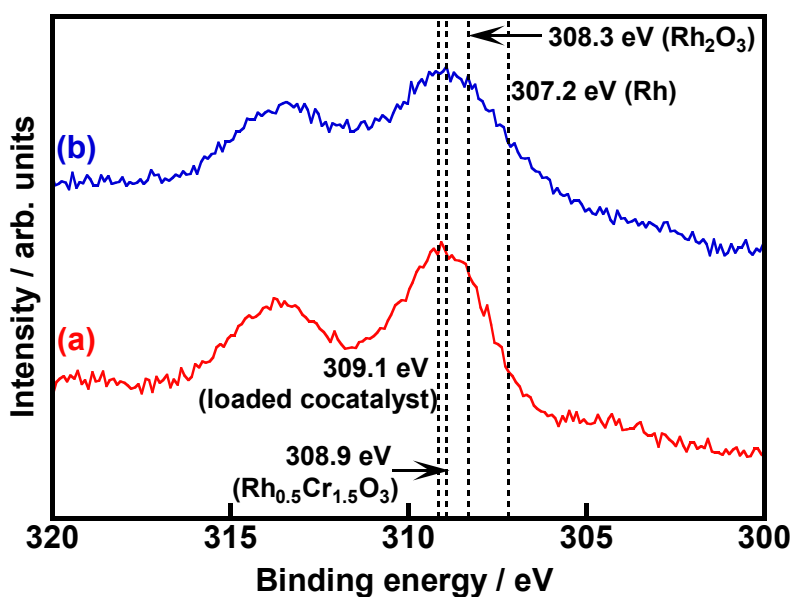


Fig. S5 X-ray photoelectron spectra of Rh 3d for Rh_{0.5}Cr_{1.5}O₃(0.2 wt%)/AgTaO₃ (a) before and (b) after photocatalytic water splitting under UV irradiation using a 300 W Xe-arc lamp. AgTaO₃ was synthesized by a solid state reaction at 1273 K for 15 h without excess Ag. Rh_{0.5}Cr_{1.5}O₃ was loaded by an impregnation method at 623 K for 1 h. Binding energies of all peaks were calibrated with C 1s (284.2 eV). All assigned binding energies for Rh, Rh₂O₃, Rh_{0.5}Cr_{1.5}O₃ and loaded cocatalyst were referred to a previous report.^{1,2)}

References

- 1) C. D. Wagner, W. M. Riggs, L. E. Davis, J. F. Moulder and G. E. Muilenberg, *Perkin-Elmer Corporation, Physical Electronics Division, Eden Prairie, Minn.*, 1979.
- 2) K. Maeda, K. Teramura, D. Lu, T. Takata, N. Saito, Y. Inoue and K. Domen, *J. Phys. Chem. B*, 2006, **110**, 13753–13758.