## **Supporting Information**

## Vanadium-based oxyhalide photocatalysts for visible-light-driven Zscheme water splitting: advancing conduction band engineering

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Figure S1. XRD patterns of PbVO<sub>3</sub>Cl synthesized by (a) solid-state reaction and (b) hydrothermal reaction.



Figure S2. XPS spectra of Pb 4f, V  $2p_{3/2}$ , Cl 2p and O 1s for the hydrothermally synthesized PbVO<sub>3</sub>Cl. The oxidation states were evaluated with reference to the NIST XPS Database.



Figure S3. XRD pattern of Pb<sub>5</sub>(VO<sub>4</sub>)<sub>3</sub>Cl synthesized by precipitation method.



Figure S4. Le Bail refinements using XRD patterns of  $Pb_{14}(VO_4)_2O_9Cl_4$  prepared at 550 °C and 600 °C.



Figure S5. SEM images of (a)  $PbVO_3Cl$ , (b)  $Pb_5(VO_4)_3Cl$ , and (c)  $Pb_{14}(VO_4)_2O_9Cl_4$ . The  $PbVO_3Cl$  sample was prepared by hydrothermal reaction, and the  $Pb_{14}(VO_4)_2O_9Cl_4$  sample was prepared by solid-state reaction at 600 °C.



Figure S6. Tauc plots of  $Pb_5(VO_4)_3Cl$ ,  $Pb_{14}(VO_4)_2O_9Cl_4$ , and  $PbVO_3Cl$ . Based on the band structure calculations (Figure S7), these materials were suggested to be indirect bandgap semiconductors; accordingly, the coefficient for indirect transition was applied in the calculations.



Figure S7. Band structures of Pb<sub>5</sub>(VO<sub>4</sub>)<sub>3</sub>Cl, Pb<sub>14</sub>(VO<sub>4</sub>)<sub>2</sub>O<sub>9</sub>Cl<sub>4</sub>, and PbVO<sub>3</sub>Cl.



Figure S8 (a) Photoelectron yield spectra of PbBiO<sub>2</sub>Cl and PbVO<sub>3</sub>Cl. The spectrum of PbBiO<sub>2</sub>Cl was cited from our previous work.<sup>[S1]</sup> (b) Band diagram estimated from the obtained ionization energies.

[S1] H. Suzuki, M. Higashi, O. Tomita, Y. Ishii, T. Yamamoto, D. Kato, T. Kotani, D. Ozaki, S. Nozawa, K. Nakashima, K. Fujita, A. Saeki, H. Kageyama, R. Abe, *Chem. Mater.* 2021, **33**, 9580-9587.



Figure S9. Band formation in solids from isolated atoms adopted and modified from Ref [S2].

[S2] P. A. Cox, *The Electronic Structure and Chemistry of Solids; Oxford Science Publications: Oxford*, 1986, 146.



Figure S10. Photocatalytic O<sub>2</sub> evolution using unmodified or (Fe,Ru)O<sub>x</sub>-loaded PbVO<sub>3</sub>Cl in aqueous Fe(NO<sub>3</sub>)<sub>3</sub> solution (5 mM, 250 mL, pH 2.4) under visible light irradiation ( $\lambda > 400$  nm).



Figure S11. Current–potential curves for electrodes composed of PbVO<sub>3</sub>Cl, Pb<sub>5</sub>(VO<sub>4</sub>)<sub>3</sub>Cl, and Pb<sub>14</sub>(VO<sub>4</sub>)<sub>2</sub>O<sub>9</sub>Cl<sub>4</sub> in a phosphate-buffered solution (0.1 M, pH 6.0) under chopped visible light from a 300-W Xe lamp with a cutoff filter (L-42).



Figure S12. XRD pattern of (Fe,Ru)O<sub>x</sub>-PbVO<sub>3</sub>Cl after photocatalytic O<sub>2</sub> evolution in aqueous Fe(NO<sub>3</sub>)<sub>3</sub> solution (5 mM, 250 mL, pH 2.4) under visible light irradiation ( $\lambda > 400$  nm), shown in Figure 6a.



Figure S13. Schematic of Z-scheme water splitting using a mixture of Ru/SrTiO<sub>3</sub>:Rh as a H<sub>2</sub>evolution photocatalyst and (Fe,Ru)O<sub>x</sub>/PbVO<sub>3</sub>Cl as an O<sub>2</sub>-evolution photocatalyst. The dotted arrows represent the electron flow.



Figure S14. Time courses of photocatalytic evolution of H<sub>2</sub> and O<sub>2</sub> using a mixture of Ru/SrTiO<sub>3</sub>:Rh and (Fe,Ru)O<sub>x</sub>/PbVO<sub>3</sub>Cl under visible light ( $\lambda > 400$  nm) in an aqueous Fe(NO<sub>3</sub>)<sub>3</sub> solution (5 mM, 250 mL, pH 2.4).



Figure S15. Z-scheme water splitting using unmodified PbVO<sub>3</sub>Cl as an OEP and Ru/SrTiO<sub>3</sub>:Rh as a HEP under visible light ( $\lambda > 400$  nm) in an aqueous Fe(ClO<sub>4</sub>)<sub>3</sub> solution (5 mM, 250 mL, pH 2.4).