



1 Supporting Information

Pt Nanocluster Co-Catalysts for Photocatalytic Water Splitting

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Figure S1. Attenuated total reflection Fourier transform infrared (ATR-FTIR) spectra showing
absorbance of H₂O (navy series) and D₂O (purple series) placed onto the ATR crystal which shows
that the D₂O is pure.





Figure S2. ATR-FTIR spectra of glass microfibre discs loading with increasing mass/area P25-Ptnc photocatalyst showing bare (0 mg/cm), low (~1.5 mg/cm), medium (~3 mg/cm) and high (~5 mg/cm) loading. From the disappearance of the Si-O stretch at 1000 cm⁻¹, the surface is coated for the medium and high loadings and the low loading has a very small fraction of uncoated surface.



Figure S3. Repeated photocatalysis experiments for a Pt-nc sample (1 wt%) showing the production of D₂ (purple, m/z = 4), O₂ (grey, m/z = 32) and CO₂ (brown, m/z = 44) for repeated experiments. In the first experiment, CO₂ is produced and O₂ is consumed. In subsequent experiments CO₂ and O₂ are not produced or consumed. Investigation is continuing to determine why O₂ is not produced. Previous experiments have shown a delay of 10 h for O₂ production in gas phase D2O splitting which was attributed to O₂ capture by the semiconductor.¹.



31Figure S4. Pt mass loading % for each sample determined by EDS. >10 individual areas were32investigated by EDS for each sample.



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Figure S5. Comparison of D₂ production rates with photocatalyst (P25-Ptnc-2wt%) loading. The data
 are presented with (upper figure) and without (lower figure) dividing by photocatalyst mass.



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Figure S6. Digital photographs of P25-Pt-nc-1wt% photocatalytic disc (a) before and (b) after 3
 repeated UV photocatalytic D₂ production experiments.

39 References

40 1. Kawai, T; Tadayoshi Sakata, T. Photocatalytic decomposition of gaseous water over TiO2 and
 41 TiO2–RuO2 surfaces. *Chem. Phys. Lett.* 1980, 72,87-89. 10.1016/0009-2614(80)80247-8



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