

Supporting information of

Direct thermo catalytic transformation of pine wood into low oxygenated bio-fuel

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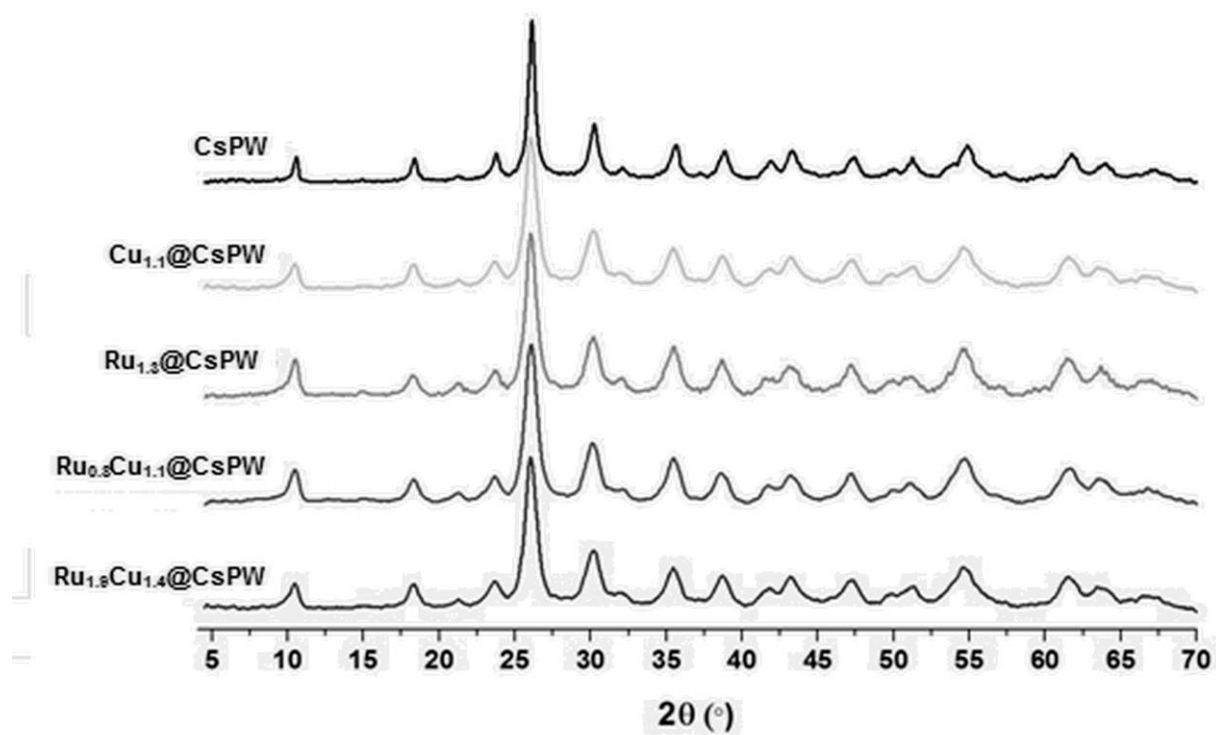


Fig. S1 X-ray diffractograms of the different samples.

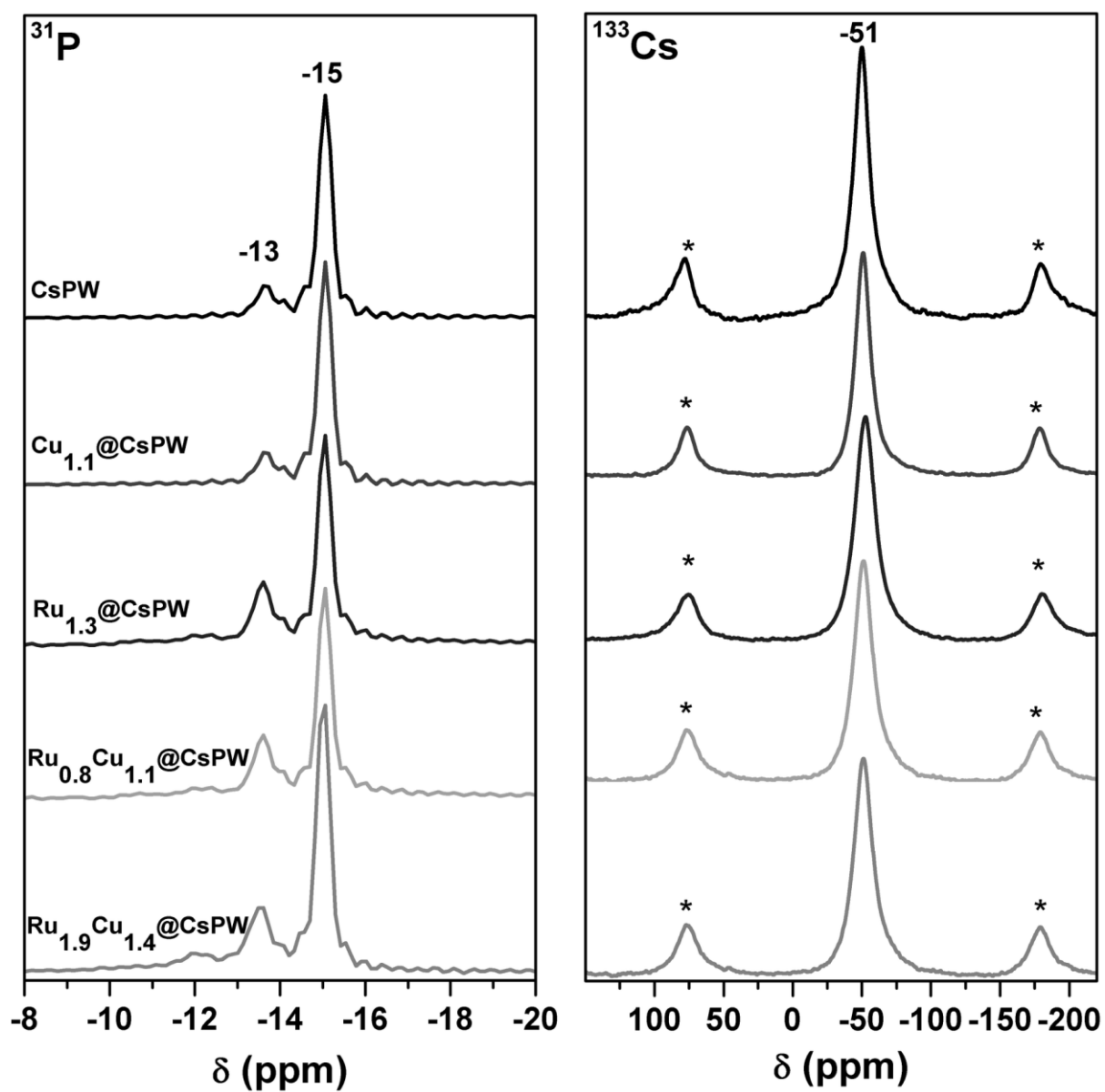


Fig. S2 ^{31}P and ^{133}Cs solid state MAS NMR of the samples after thermal treatment at 350 °C under oxygen and hydrogen.

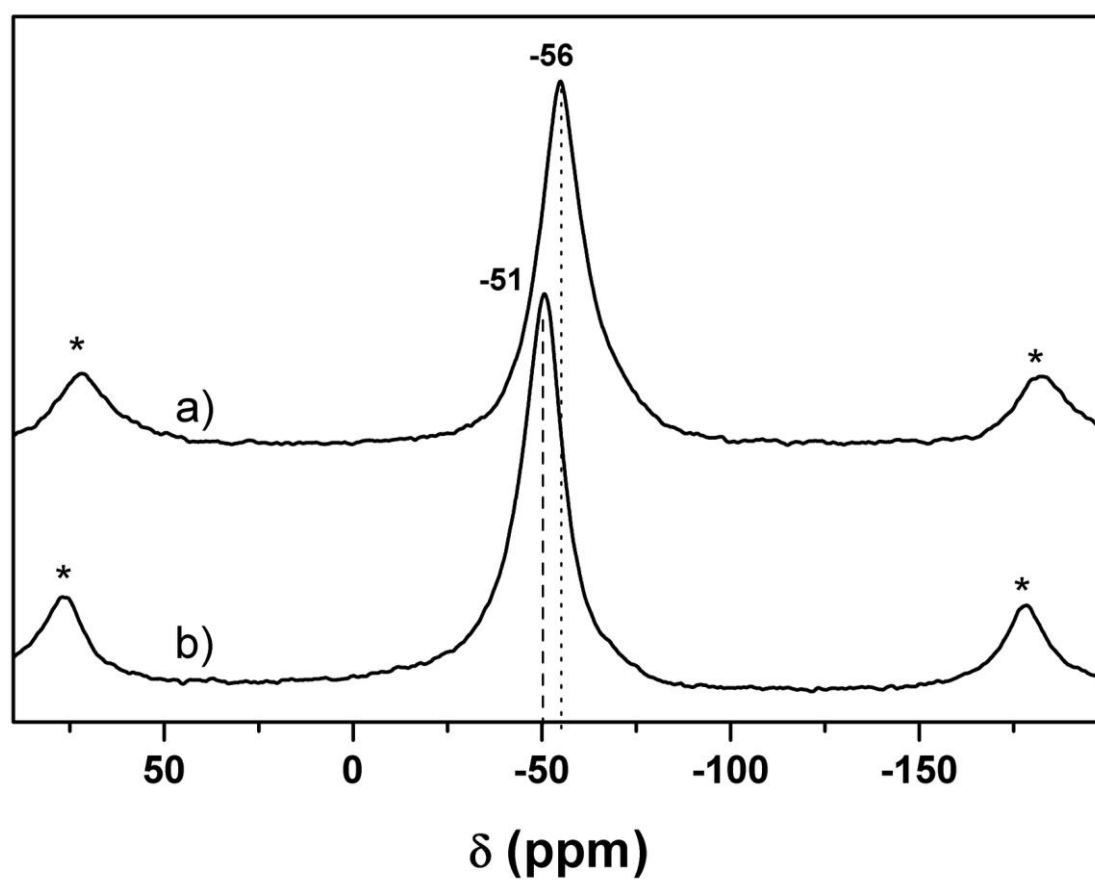


Fig .S3 ^{133}Cs solid state MAS NMR of cesium exchange heteropolyanion support dried at room temperature (a) and dried at 200 °C (b).

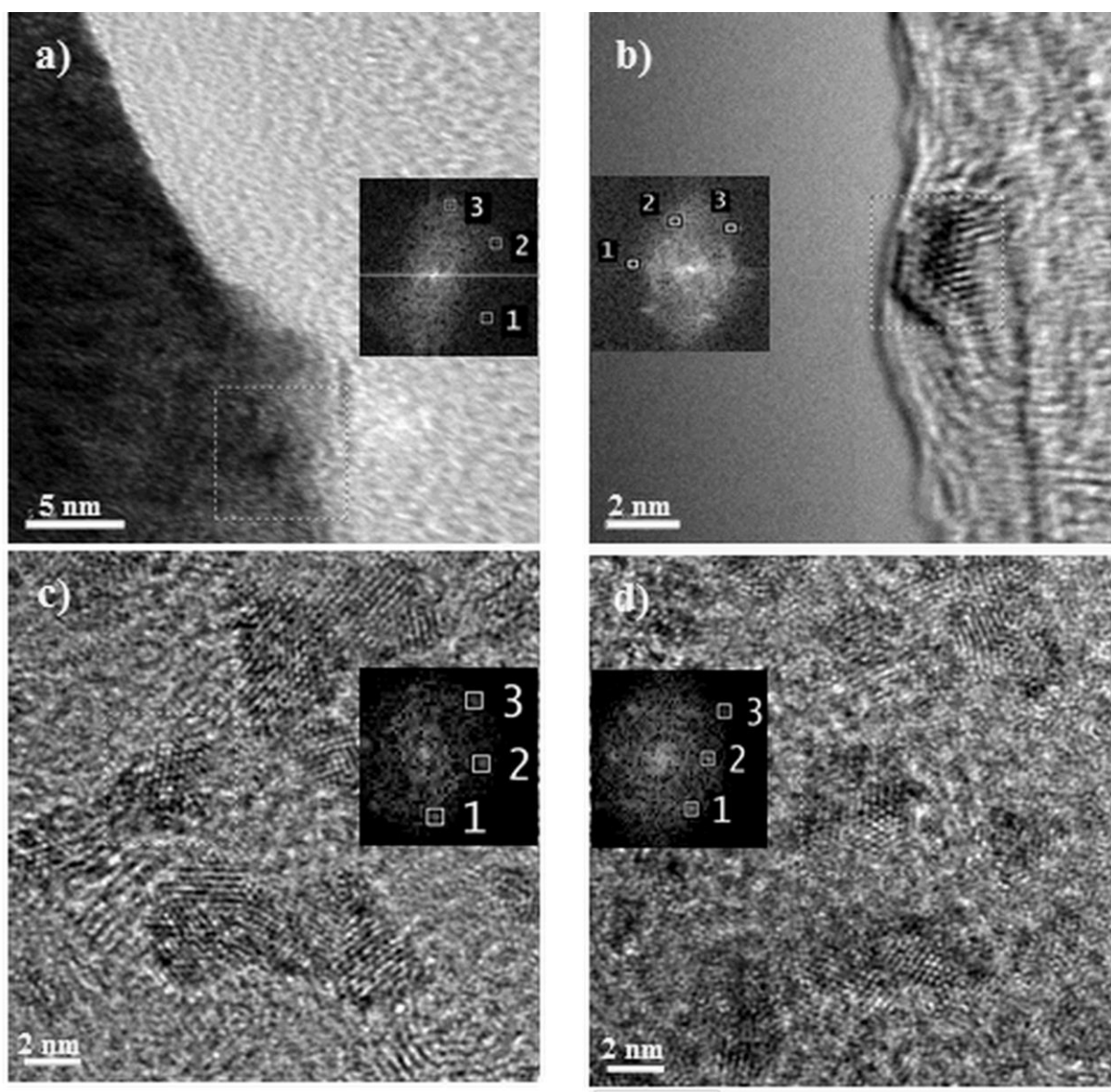


Fig. S4 High resolution transmission electron microscopy images of $\text{Ru}_{1.9}\text{Cu}_{1.4}@\text{CsPW}$ catalyst showing the different structures a) hexagonal $\text{Ru}(0)$, b) tetragonal RuO_2 , c) cubic RuO_4 and d) monoclinic CuO . Inset representative of fast fourier transformation power spectra of the particles.

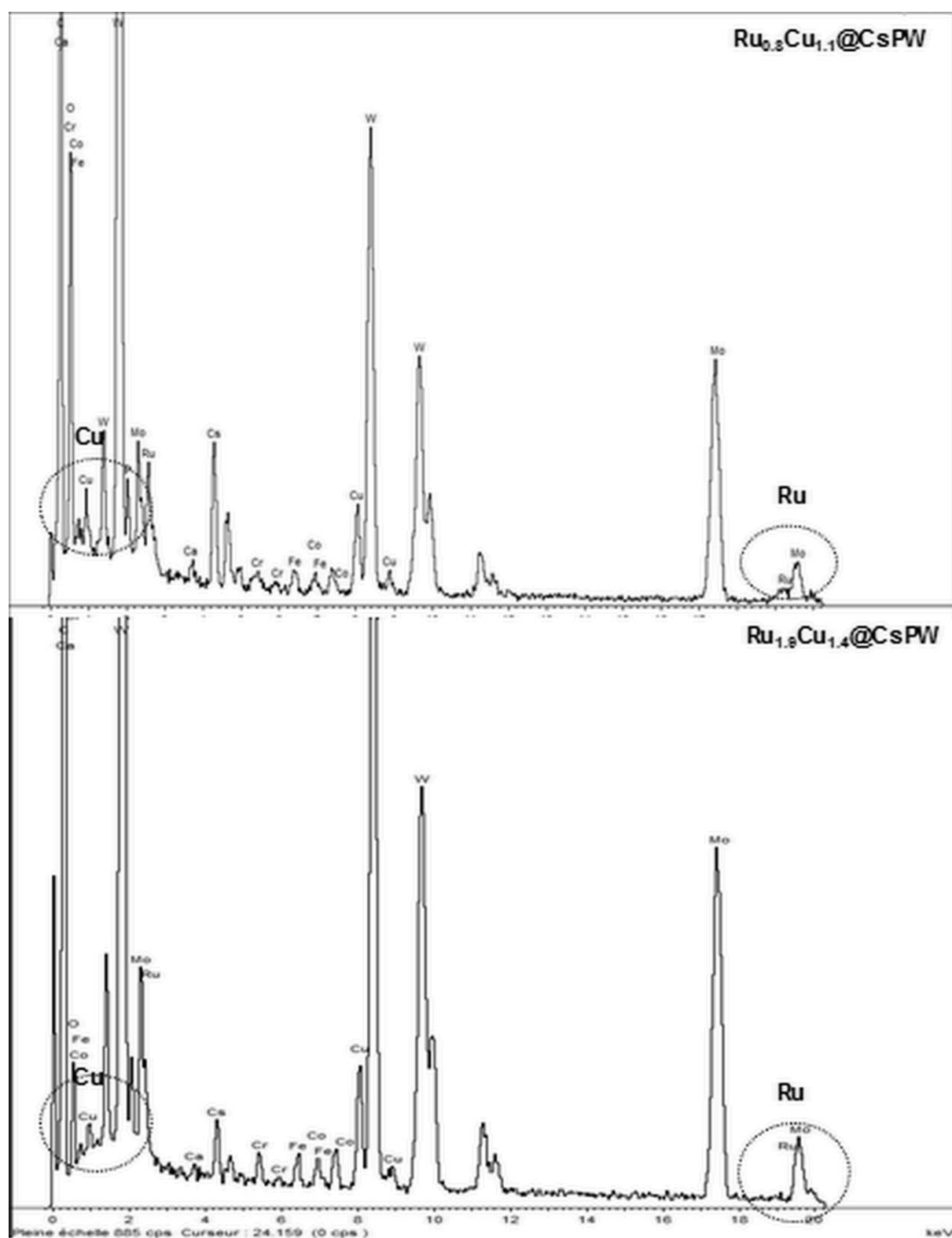


Fig. S5 Energy-dispersive X-ray spectroscopy of two catalysts $\text{Ru}_{1.9}\text{Cu}_{1.4}@CsPW$ and $\text{Ru}_{0.3}\text{Cu}_{1.1}@CsPW$, showing the presence of copper and ruthenium.

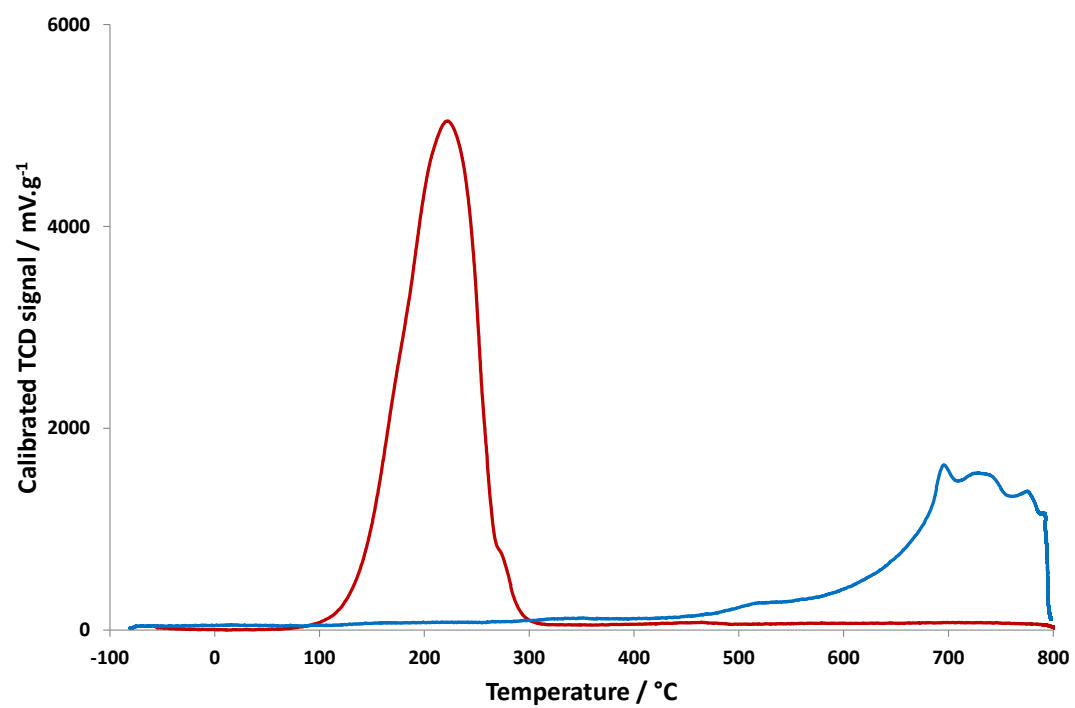


Fig. S6: s-TPR profile of Cu10wt%@Al₂O₃ (red) and Cu1wt%@CsWP (blue) catalysts.

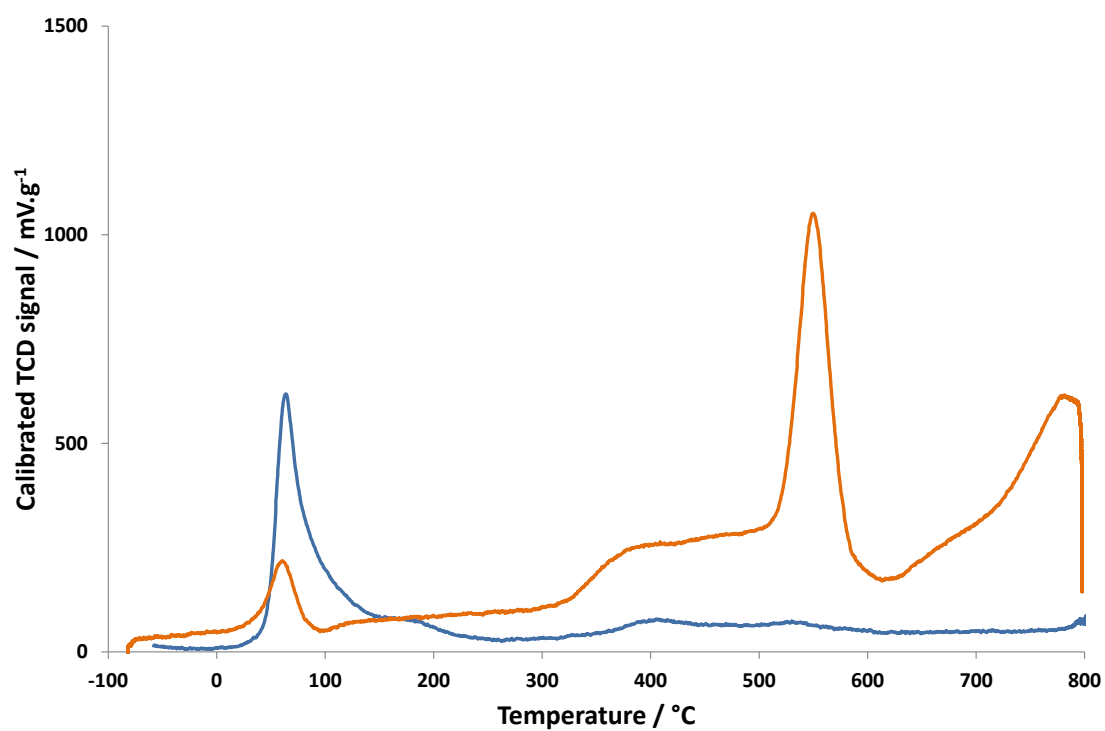


Fig. S7: s-TPR profile of Ru1wt%@CsWP (orange) and Ru2wt%@ Al₂O₃ (blue) catalysts.

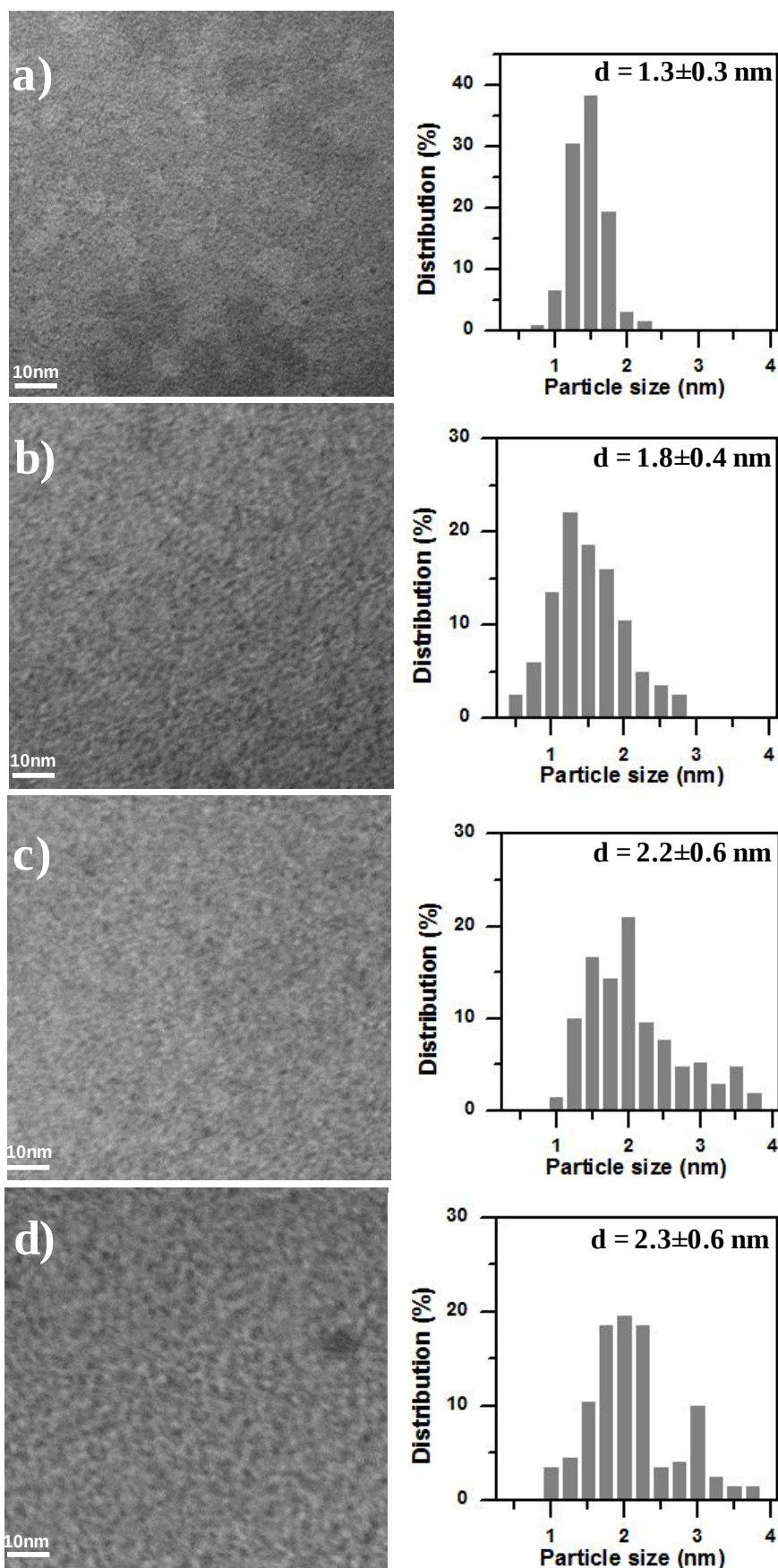


Fig. S8 Transmission electron microscopy of a) as prepared $\text{Ru}_{1.9}\text{Cu}_{1.4}@\text{CSPW}$ catalyst, b) after the first catalytic cycle, c) after second catalytic cycle and d) after fourth catalytic cycle.