

Oxidation mechanism of tetrachloroplatinate(II) by hydrogen peroxide in hydrochloric acid solution

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Electronic Supplementary Information (ESI)

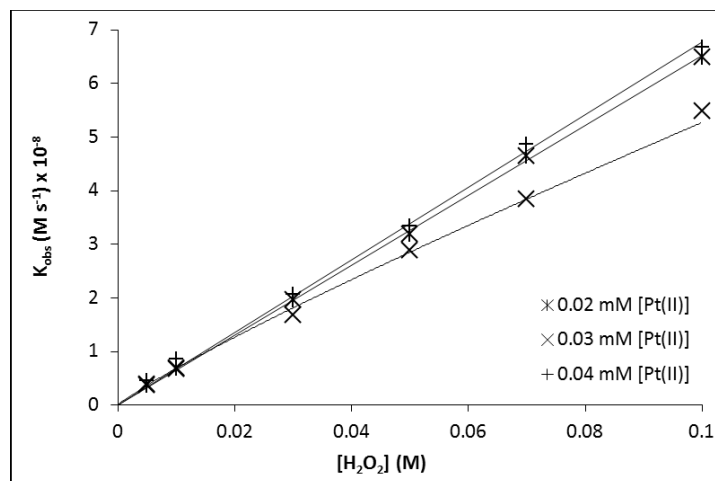


Figure S1A. Dependence of k_{obs}^0 on the concentration of H_2O_2 for the oxidation of 0.02, 0.03 and 0.04 mM $[\text{PtCl}_4]^{2-}$.

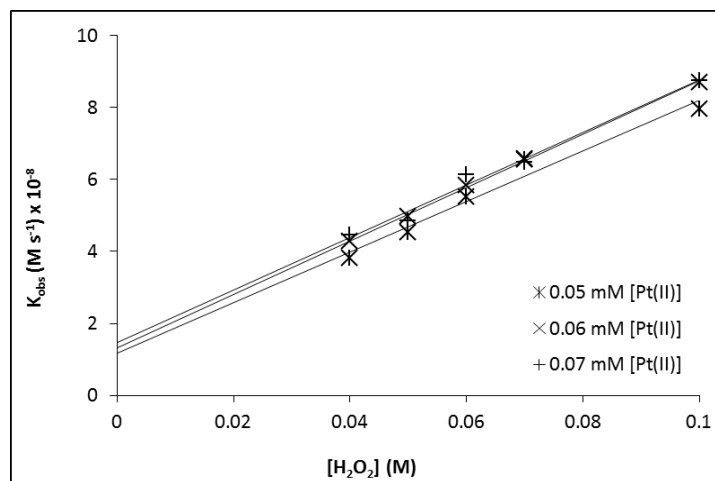


Figure S1B. Dependence of k_{obs}^0 on the concentration of H_2O_2 for the oxidation of 0.05, 0.06 and 0.07 mM $[\text{PtCl}_4]^{2-}$.

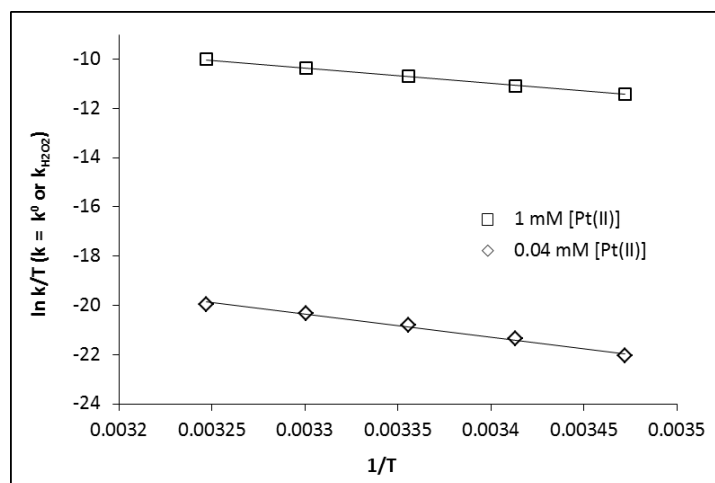


Figure S2. Eyring plots for the oxidation of 0.04 mM $[\text{PtCl}_4]^{2-}$ by 100 mM H_2O_2 , and of 1 mM $[\text{PtCl}_4]^{2-}$ by 300 mM H_2O_2 .

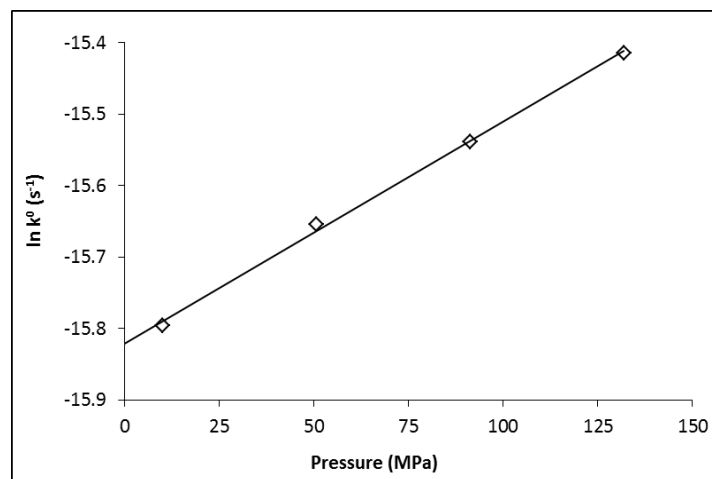


Figure S3. Plot of $\ln k^0$ (s^{-1}) as a function of pressure to calculate the activation volume for the zero-order mechanism.

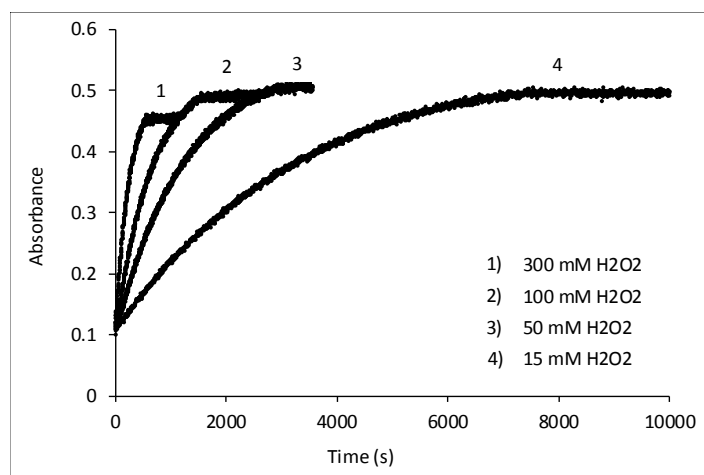


Figure S4. Absorbance vs. time plots at 353 nm and 35 °C for the oxidation of 1 mM $[\text{PtCl}_4]^{2-}$ by various concentrations of H_2O_2 , $[\text{H}^+]$, $[\text{Cl}^-] = 1 \text{ M}$.

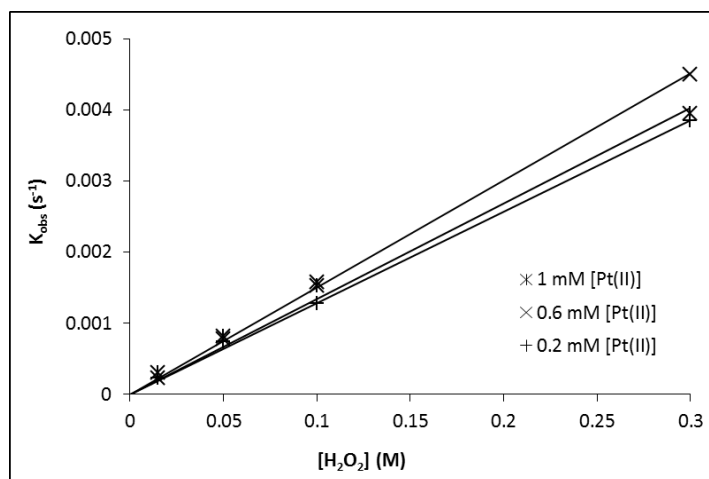


Figure S5. The dependence of k_{obs} on the H_2O_2 concentration for oxidation of 1, 0.6, and 0.2 mM $[\text{PtCl}_4]^{2-}$.

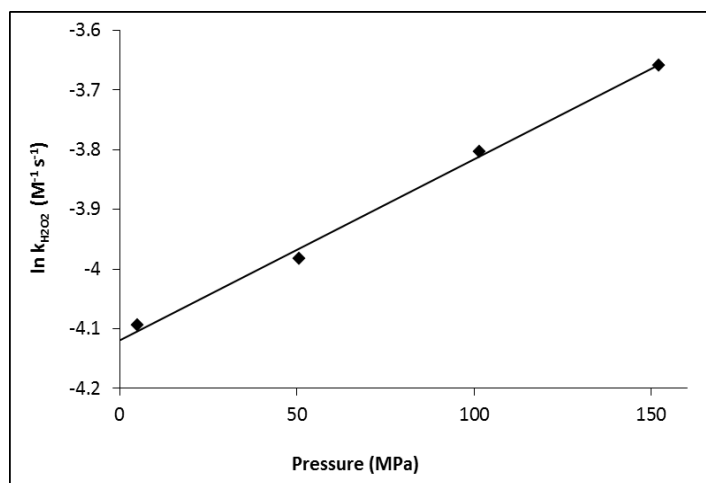


Figure S6. Plot of $\ln k_{\text{H}_2\text{O}_2}$ as a function of pressure (MPa) to calculate the activation volume for the pseudo-first order mechanism.

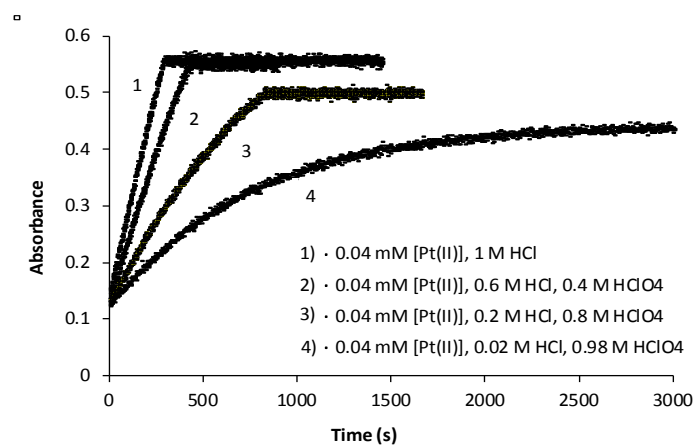


Figure S7. Absorbance vs. time plots at 353 nm and 35 °C for the oxidation of 0.04 mM $[\text{PtCl}_4]^{2-}$ as a function of chloride concentration. Concentrations of $[\text{PtCl}_4]^{2-}$ was kept constant at 0.04 mM, $[\text{H}_2\text{O}_2]$ at 80 mM and ionic strength 1 M.

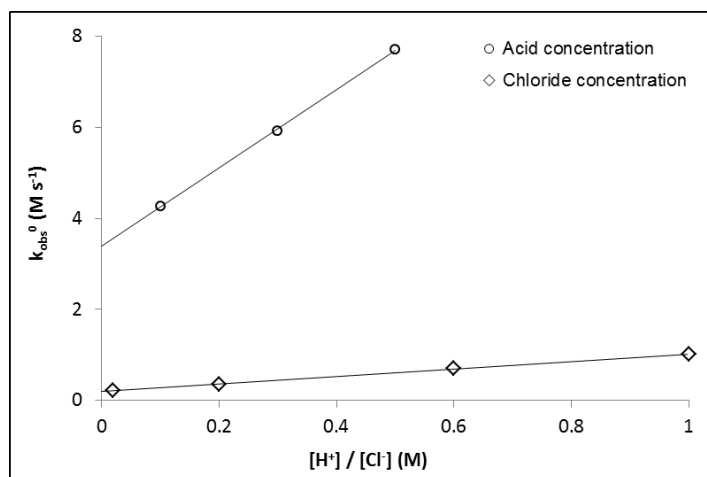


Figure S8. Plots of k_{obs}^0 (M s^{-1}) as a function of acid and chloride concentration. $[\text{PtCl}_4]^{2-} = 0.04$ mM; $[\text{H}_2\text{O}_2] = 80$ mM and ionic strength = 1 M.

