

deoxygenation¹



- The main cause of Fe⁰ deactivation is oxidation by
- At low pH, Pd adsorbs on
 - Fe_2O_3
- At high pH, Pd adsorbs on silica

Temperature Programmed Reduction (TPR) 5 nm Fe_2O_3

- Fe₂O₃ before reduction shows mostly $5nm \gamma - Fe_2O_3$
- Pd/Fe₂O₃ reduced at 475 °C shows large Fe⁰ peaks – avg. particle size is 40 nm

Conclusions

- Pd can be steered onto Fe₂O₃ using SEA
- These samples show enhanced reducibility
- Reactivity testing underway soon

water²



- Pd helps protect Fe⁰ sites from water oxidation^{2,3}
- Pd can be selectively steered with Strong Electrostatic



Pd adsorbed at high pH shows some enhanced

Future efforts to reduce the Fe particle size

References

- Hong, Y.; Wang, Y., Elucidation of reaction mechanism for m-cresol hydrodeoxygenation over Fe based catalysts: A kinetic study. *Catalysis Communications* **2017**, *100*, 43-47.
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- Hensley, A. J.; Hong, Y.; Zhang, R.; Zhang, H.; 3. Sun, J.; Wang, Y.; McEwen, J.-S., Enhanced Fe2O3 reducibility via surface modification with Pd: Characterizing the synergy within Pd/Fe catalysts for hydrodeoxygenation reactions. ACS Catalysis 2014, 4 (10), 3381-3392.

Adsorption (SEA)^{4,5}



reducibility – suggests H₂ spillover

Pd adsorbed at low pH

shows most reducibility – indicates an electronic effect

- Bulk Fe₂O₃ TPRs indicate the electronic effect has limited range, said to be 4
 - Angstroms³

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