

STUDY OF 1.8 NM Pt NANOPARTICLES ANCHORED ON DIFFERENT AMORPHOUS SILICA SUPPORTS IN ETHANOL DECOMPOSITION REACTION

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Abstract

1.8 nm Pt nanoparticles with narrow size distribution were anchored on mostly identical, amorphous silica supports (SBA-15 [1], MCF-17 [2], Silica Foam [3]) and were tested in ethanol decomposition reactions at < 573 K. The reaction on the Pt/SF (0.117 molecules·site⁻¹·s⁻¹) was ~2 times faster compared to Pt/MCF-17 (0.055 molecules·site⁻¹·s⁻¹) and Pt/SBA-15 (0.063 molecules·site⁻¹·s⁻¹) at 573 K. In the case of Pt/SBA-15, selectivity towards acetaldehyde was ~4

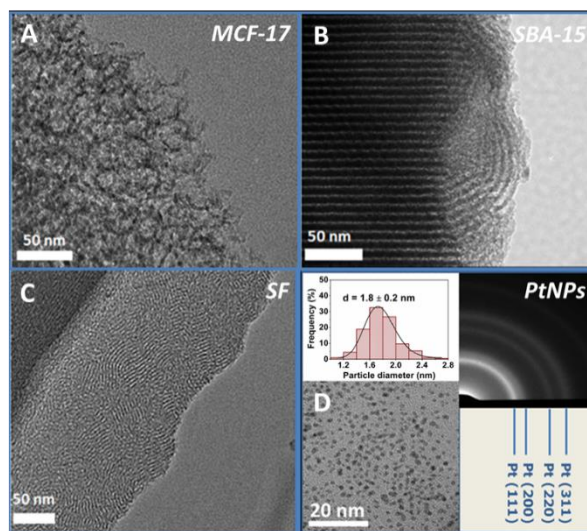


Fig. 1. Typical TEM images of MCF-17 (A), SBA-15 (B) and SF (C) silica supports as well as 1.8±0.2 nm metallic Pt nanoparticles with narrow size distribution (D).

times higher (68%) compared to the Pt/MCF-17 (18%) and Pt/SF (16%) catalysts. In the case of Pt/MCF-17 and Pt/SF, the methane to acetaldehyde ratio was 0.27 and 0.24, respectively, while it was ~ 10 times higher (1.97) for Pt/SBA-15 catalyst. The ethene selectivity was ~2 times higher

in the case of Pt/MCF-17 (0.99%) and Pt/SF (0.93%) compared to Pt/SBA-15 (0.41%). Pt/MCF-17 and Pt/SBA-15 produces ~ 50% more hydrogen (~27%) compared to Pt/SF catalyst (21 %). Small Angle X-ray Scattering (SAXS) and Transmission Electron Microscopy (TEM) studies showed striking differences in the porosity, pore- and mesostructure, sintering and Pt-SiO₂ interface altering effect of the silica supports as well as the Pt nanoparticles decorated catalysts which may have significant effect on the catalytic activity.

References

- [1] D. Y. Zhao, Q. S. Huo, J. L. Feng, B. F. Chmelka, G. D. Stucky *J. Am. Chem. Soc.* 120 (1998) 6024.
- [2] P. Schmidt-Winkel, W. W. Lukens, P. D. Yang, D. I. Margolese, J. S. Lettow, J. Y. Ying, G. D. Stucky *Chem. Mater.* 12 (2000) 686.
- [3] S. A. Bagshaw, *Chem. Comm.* (1999) 767-768.