

Characterization of Palladium (Pd) on Alumina Catalysts Prepared Using Liquid Carbon Dioxide

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Supported palladium (Pd) catalysts were prepared using an environmentally benign route. Palladium (II) hexafluoroacetylacetonate ($\text{Pd}(\text{hfac})_2$) dissolved in liquid carbon dioxide (L-CO_2) was used to deposit Pd nanoparticles onto low-surface-area α -alumina ($13.3 \text{ m}^2/\text{g}$) and high-surface area γ -alumina ($207.0 \text{ m}^2/\text{g}$) supports. $\text{Pd}(\text{hfac})_2$ dissolved in L-CO_2 at 6.9 MPa and 28.5 °C was impregnated into the supports by slowly venting gaseous CO_2 until L-CO_2 was completely evaporated. After depressurization, the impregnated $\text{Pd}(\text{hfac})_2$ was reduced in hydrogen at a relatively low temperature of 75 °C. The adsorption isotherm of $\text{Pd}(\text{hfac})_2$ on γ -alumina suggests a weak interaction between the organometallic compound and the support. The average particle size on the low-surface-area α -alumina increased from 13.1 ± 3.5 to 59.9 ± 11.3 nm and the metal dispersion, measured by pulsed CO chemisorption, decreased from 11% to 3% as the Pd loading increased from 0.15 to 1.54 wt%. When high-surface-area γ -alumina was used, Pd particle size increased from 3.1 ± 1.9 to 7.0 ± 5.9 nm and metal dispersion decreased from 56 to 5 % as Pd loading increased from 0.58 to 3.94 wt%.