

Efek Penambahan Sn pada Aktivitas Katalitik NiGa Tersangga Karbon Mesopori untuk Reaksi Karboksilasi Fenilasetilena dengan CO = Effect of Sn Addition on Catalytic Activity of NiGa Supported by Mesoporous Carbon for Carboxylation of Phenylacetylene with CO

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Abstrak

Reaksi karboksilasi fenilasetilena dengan CO dilakukan dengan menggunakan katalis Ni-Ga dan Ni-Ga termodifikasi timah (Sn) yang disangga oleh karbon mesopori (MC). MC disintesis dengan metode soft template dan dianalisis dengan TGA diperoleh kestabilan termal hingga 850 °C. Katalis dikarakterisasi dengan FTIR, XRD, Raman, SEM, TEM, dan BET. Hasil XRD menunjukkan puncak pada 24,47°; 32,73°; 43,83°; 50,96°; 74,32° yang merupakan difraksi NiGa dan partikel Sn(0) pada penyangga MC. Penambahan Sn diketahui tidak mengubah kestabilan katalis NiGa/MC yang dikonfirmasi melalui FTIR dan Raman. Hasil SEM dan TEM juga menunjukkan partikel Ni-Ga-Sn tersebar merata pada permukaan karbon mesopori. BET menunjukkan katalis termasuk dalam ukuran mesopori 2-50 nm. Uji aktivitas katalitik berdasarkan analisa HPLC menunjukkan hasil optimum diperoleh dengan menggunakan katalis NiGaSn_{0.5}/MC pada suhu 50°C selama 8 jam. Sedangkan berdasarkan LC-MS, diketahui terbentuk produk asam sinamat dan asam fenil propiolat dengan yield masing-masing 2,14% dan 3,04% dengan konversi fenilasetilena mencapai 93,06%.

.....The carboxylation reaction of phenylacetylene with CO₂ was carried out using Ni-Ga and Ni-Ga-modified tin catalysts supported by mesoporous carbon (MC). MC was synthesized using the soft template method and analyzed using TGA and obtained thermal stability up to 850 °C. To determine the modification effect of Sn addition, catalysts were synthesized with variations of Ni₅Ga₃/MC, Ni₅Ga₃Sn_{0.1}/MC, Ni₅Ga₃Sn_{0.3}/MC, Ni₅Ga₃Sn_{0.5}/MC, Ni₅Ga₃Sn_{0.7}/MC, Ni₅Ga₃Sn_{0.9}/MC. The catalysts were characterized by FTIR, XRD, Raman, SEM, TEM, and BET. XRD results show peaks at 24.47°; 32.73°; 43.83°; 50.96°; and 74.32° which is the diffraction of the Ni₅Ga₃ phase and Sn (0) particles on the MC support. The addition of Sn metal is known not to change the stability of the Ni₅Ga₃/MC catalyst which was confirmed through FTIR and Raman spectra. SEM and TEM results also show that Ni-Ga-Sn particles are evenly distributed on the mesoporous carbon surface with a spherical shape. BET-SAA shows the pore diameter size of the materials Ni₅Ga₃/MC, Ni₅Ga₃Sn_{0.1}/MC, Ni₅Ga₃Sn_{0.3}/MC, Ni₅Ga₃Sn_{0.5}/MC, Ni₅Ga₃Sn_{0.7}/MC, Ni₅Ga₃Sn_{0.9}/MC respectively, are 6.24 nm; 6.22nm; 7.22nm; 6.24 nm, 7.22 nm, and 10.46 nm which are included in the mesopore size of 2-50 nm. The catalytic activity test was carried out through the carboxylation reaction of phenylacetylene with CO₂ using variations of catalyst, time and temperature. HPLC analysis showed that optimum results were obtained using the Ni₅Ga₃Sn_{0.5}/MC catalyst at a temperature of 50°C for 8 hours. Meanwhile, based on LC-MS, it is known that cinnamic acid and phenyl propiolic acid products were formed with yields of 2.14% and 3.04% respectively with 93.06% phenylacetylene conversion.