

Polymerization of phenylacetylene catalysed by RhTp(cod) and RhBp(cod) in ionic liquids: effect of alcohols and of tetraammonium halides.

Autorzy

Anna M. Trzeciak

Józef J. Ziólkowski

Rok wydania

2004

Czasopismo

Applied Organometallic
Chemistry

Numer woluminu

18

Strony

124-129

DOI

10.1002/aoc.589

Kolekcja

Naukowa

Język

Angielski

Typ publikacji

Artykuł

Streszczenie

RhTp(cod) (**1**) and RhBp(cod) (**2**), almost inactive in CH₂Cl₂, became good catalysts of phenylacetylene polymerization in ionic liquids ([bmim]Cl, [bmim]BF₄: *bmim* = 1-butyl-3-methylimidazolium, [mokt]BF₄: *mokt* = 1-methyl-3-oktylimidazolium, [bumepy]BF₄: 1-butyl-4-methylpyridinium) and in CH₂Cl₂ in the presence of tetraammonium halides ([R₄N]X, R = Bu, Et; X = Cl, Br). The highest yields of polyphenylacetylene with catalyst **1** were obtained in [bmim]Cl at 65°C (64% after 2 h) and in [mokt]BF₄ at 20°C (56% after 24 h). In alcohols (CH₃OH, (CH₃)₂CHOH, (CH₃)₃COH) as solvents, up to 100% of the polymer was produced. When a mixture of an ionic liquid and CH₃OH was used as the reaction medium, the polymer yield was similar to the yield achieved in an ionic liquid only, but the molecular weight increased remarkably. Tetraammonium salts, [R₄N]X, are co-catalysts for **1**, and the yield of the polymer increased in the order [Et₄N]Br < [Bu₄N]Br < [Et₄N]Cl < [Bu₄N]Cl. Polymers with molecular weights from 6900 to 38 800 Da were obtained with catalyst **2** in [R₄N]Br or [R₄N]Cl, whereas in ionic liquids ([bmim]Cl, [bmim]BF₄) the corresponding molecular weights were higher, from 51 300 to 60 300 Da.

Adres publiczny

<https://doi.org/10.1002/aoc.589>

Strona internetowa wydawcy

onlinelibrary.wiley.com