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SYNTHESIS AND COORDINATION CHEMISTRY OF *N*-(DIPHENYLPHOSPHINO) DIALLYLAMINE

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SYNTHESIS AND COORDINATION CHEMISTRY OF N-(DIPHENYLPHOSPHINO) DIALLYLAMINE

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N-(diphenylphosphino)diallylamine (1) is a potential tridentate ligand containing both phosphorus and olefinic donor sites. Its preparation is straightforward and involves the addition of chlorodiphenylphosphine to a 2-fold excess of diallylamine in toluene.¹

A study of the coordination chemistry of 1 has shown some interesting results. Reactions of 1 with $[PtCl_2(cod)]$ and $[PdCl_2(cod)]$ (cod = 1,5-cyclooctadiene) has furnished a variety of complexation products. Analysis of these products by multi-nuclear NMR, FAB mass spectra, and elemental analysis has provided suggested structures of the complexes formed. Single crystal x-ray crystallographic analysis of the 1:1 $[PdCl_2(cod)]$:1 product shows that the ligand adopts a bidentate P/η^2 allyl coordination mode resulting in the formation of a five-membered $PdPNC_2$ metallacycle.

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