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1-aryl-3-butanones. Particularly Well Suited For This Protocol Are Anilines Ortho-substituted With Electron-withdrawing Groups (alkoxycarbonyl, Cyano, Acetyl, Trifluoromethyl, Halogens). Apr 2th, 2024

Organic Syntheses, Coll. Vol. 6, P.190 (1988); Vol. 55, P ...

2. Thionyl Chloride Was Obtained From Anachemia Chemicals Ltd., Fisher Scientific Company (reagent Grade), Or Matheson, Coleman And Bell. The First Two Were Slightly Yellow, And The Latter Was Colorless; However, The Yields Of Final Product Were Identical With Each Brand. The Excess Thionyl Chloride Serves Mar 2th, 2024

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¹H, ³¹P{¹H}, And ¹³C{¹H} NMR Spectroscopy And Elemen-tal Analyses. All Characterization Data Are Consistent With The Expected Structures And Demonstrate The High Purity Of The Ligands. The ¹H And ¹³C{¹H} NMR

Resonances Were Partially Assigned By Using 2D Heteronuclear Multiple-bond Correlatio Feb 2th, 2024

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Stereoselective Completion Of The Synthesis H N N OMe H N N OH H OMe 65% 1. HF, MeCN 2. i. MsCl, Pyridine, CH₂Cl₂ ii. MeCN, Reflux i. NaH, DMSO ii. O₂ 78% Quinine 19 Steps, 0.96 % Yield N MeO N OTBDPS MeO H H NaBH₄, MeOH, THF Jan 3th, 2024

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An Intermediate (route B), While Bromonium Ion-initiated Cyclization Gives A Brominated Monocyclofarnesol Pyrophosphate As The Intermediate (route A, Scheme 1). Even If It Were Possible To Check The Chemical Viability Of One Of The Path-ways (a Or B) Of Scheme 1, The Sense In Which The Biological Transformation Mar 3th, 2024

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