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- Convenient Preparations of 4-Methyl- and 4,6-Dimethyl-10,11-dihydro-5*H*-dibenzo[*a,d*]cycloheptene using the t-Butyl Function as a Positional Protective Group Masashi Tashiro, Takashi Furusawa, Akihiko Tsuge, Shuntaro Mataka, and Takehiko Yamato
 - N.B. The numbers in parentheses, prefaced by M, indicate the first frame occupied by the *full-text version* of the paper in J. Chem. Research (M). Where no such number is given, the paper as published in J. Chem. Research (S) is complete in itself, and there is no extra material in Part M.

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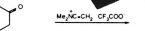
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A, is converted to the Mannich reagent, N,N-acetate, an excellent reagent for the α -dimethyl-ill. Soc. Chim. Fr., 2707 (1970). Compare dimethyl-31, p.423:



Deprotonation by lithium diisopropylamide at low temperature gives the unstable azomethine ylide, which undergoes 1,3-dipolar addition even with unactivated alkenes, to give pyrrolidines: *J.Chem.Soc.,Chem.Commun.*, 31 (1983):



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Conjugate addition of cis-alkenylzirconium reagents, from the hydrozirconation of alkynes, to Michael acceptors, with retention of configuration: *J.Am.Chem.Soc.*, **102**, 1333 (1980).

Coupling of Grignard reagents to give biaryls: *J.Org.Chem.*, **41**, 2252 (1976). Coupling of Grignard reagents with silyl enol ethers of both aldehydes and ketones, to give alkenes. In contrast to dichlorobis(triphenylphosphine)nickel, 0369, p.335, this reagent gives the thermodynamically more stable alkene: *Tetrahedron Lett.*, 3915 (1980):

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