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Stereoselective and catalyzed halogenation reactions

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Contents

Announcement: Tetrahedron Symposia-in-Print Preface

pp 7145–7147 p 7149

ARTICLES

Highly *Z*/*E* **stereoselective approach to** β**-iodo aza Morita–Baylis–Hillman adducts** Cody Timmons, Adiseshu Kattuboina, Soham Banerjee and Guigen Li* pp 7151–7154



A multicomponent reaction between sulfonyl-protected imines, magnesium iodide, and acetylenic esters or ketones is described. The resulting β -iodo aza Morita–Baylis–Hillman adducts were obtained in good yields (68–84%) and excellent Z/E stereoselectivities (17:1–20:1) for 11 examples.

Planned and unplanned halogenations in route to selected oroidin alkaloids Shaohui Wang, Anja S. Dilley, Karine G. Poullennec and Daniel Romo* pp 7155-7161



Elemental fluorine. Part 19: Electrophilic fluorination of hexyl derivatives bearing electron withdrawing groups

Richard D. Chambers,* Mandy Parsons, Graham Sandford,* Emmanuelle Thomas, Jelena Trmcic and John S. Moilliet

The effect of each functionality upon the conversion of unactivated carbon-hydrogen bonds to carbon-flourine bonds at sites within an alkyl chain is established.

Highly enantioselective fluorination reactions of β-ketoesters and β-ketophosphonates catalyzed by pp 7168–7179 chiral palladium complexes

Yoshitaka Hamashima, Toshiaki Suzuki, Hisashi Takano, Yuta Shimura, Yasunori Tsuchiya, Ken-ichi Moriya, Tomomi Goto and Mikiko Sodeoka*



Structural and stereochemical aspects of the enantioselective halogenation of 1,3-dicarbonyl compounds catalyzed by Ti(TADDOLato) complexes

Mauro Perseghini, Massimo Massaccesi, Yanyun Liu and Antonio Togni*



Selenium-catalyzed oxidative halogenation Shelli R. Mellegaard-Waetzig, Chao Wang and Jon A. Tunge*



pp 7162–7167

pp 7191-7198

Radical trifluoromethylation of ketone Li enolates

Yoshimitsu Itoh and Koichi Mikami*



Highly basic Li enolates are shown to be applicable to radical trifluoromethylation. The reaction is extremely fast and the minimum reaction time is ~ 1 s.

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COVER

The cover graphic depicts examples of selenium-promoted halogenations. © 2006 T. Lectka. Published by Elsevier Ltd.



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pp 7199-7203