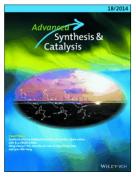
Cover Pictures

Synthesis of (E)-5-Aryl(halo)methylenebicyclo[2.2.2]oct-2-enes and -[2.2.1]hept-2-

Ming-Chang P. Yeh, Hsin-Hui Lin, Siou-Fu Kuo, Pei-Jie Chen and Jyun-Wei Hong Article first published online: 10 NOV 2014 | DOI: 10.1002/adsc.201401045



The back cover picture, provided by Ming-Chang P. Yeh and co-authors, illustrates an efficient synthesis of (E)-5-halomethylenebicyclo[2.2.2]oct-2-enes and -[2.2.1]hept-2-enes from the simple and inexpensive FeX₃ (X=Cl or Br) and C-4 propargyl-tethered cyclic 2-enols under air at room temperature in minutes. The fluoro-analogs are available from the reaction of TBS-protected cyclic 2,6-enynols with BF₃·OEt₂ under the similar reaction conditions. Details of this work can be found in the communication on pages xxxx–yyyy (M. C. P. Yeh, H. H. Lin, S. F. Kuo, P. J. Chen, J. W. Hong, *Adv. Synth. Catal.* **2014**, *356*, xxxx–xxxx; DOI:

10.1002/adsc.201400671).

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