

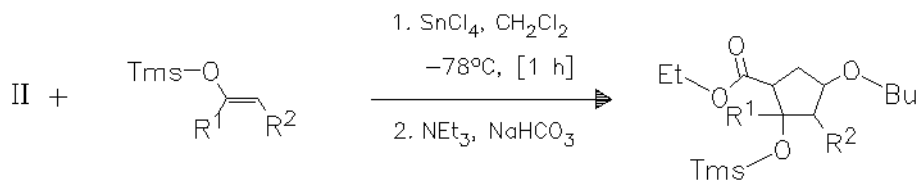
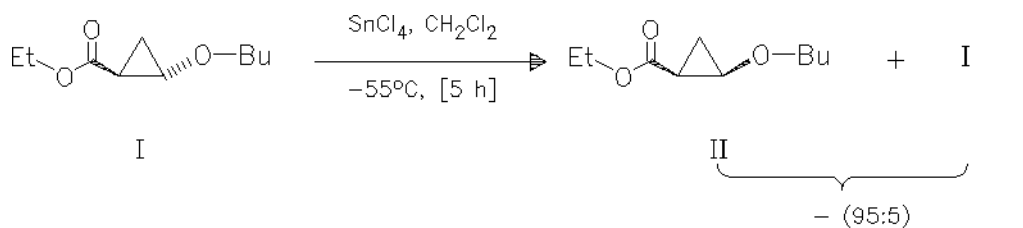
cycloaddition reactions

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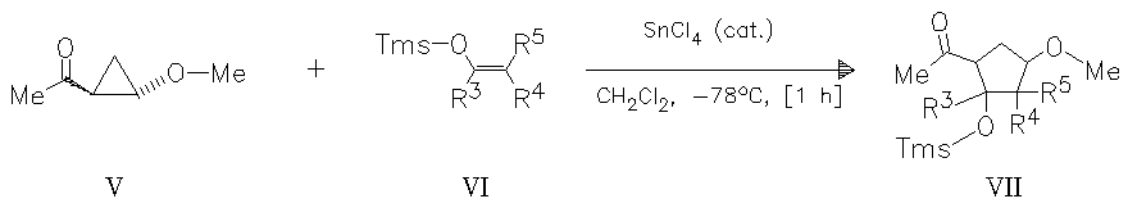
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**(3 + 2)Cycloaddition of 2-Alkoxypropyl Carbonyl Compounds with Enol Silyl Ethers for Functionalized Cyclopentane Formation.**

— Epimerization of the trans-ethyl 2-butoxycyclopropanecarboxylate (I) to give the cis-isomer (II) is achieved using tin(IV) chloride in methylene chloride. The cyclopentane derivatives (IV) are generated as a mixture of diastereomers by cycloaddition reaction of (II) with the enol silyl ethers (III) in the presence of tin(IV) chloride. Only catalytic amounts of SnCl<sub>4</sub> are required to induce the cycloaddition reaction of 2-methoxycyclopropyl methyl ketone (V) with the enol silyl ethers (VI), forming the cyclopentane derivatives (VII). A mechanism is discussed. — (KOMATSU, M.; SUEHIRO, I.; HORIGUCHI, Y.; KUWAJIMA, I.; Synlett (1991) 11, 771-773; Dep. Chem., Tokyo Inst. Technol., Meguro, Tokyo 152, Japan; EN)



III	IV
a R <sup>1</sup> : -Ph; R <sup>2</sup> : -H	66%
b R <sup>1</sup> -R <sup>2</sup> : -(CH <sub>2</sub> ) <sub>3</sub> -	67%
c R <sup>1</sup> -R <sup>2</sup> : -(CH <sub>2</sub> ) <sub>4</sub> -	70%
d R <sup>1</sup> -R <sup>2</sup> : -(CH <sub>2</sub> ) <sub>5</sub> -	69%



a R <sup>3</sup> : -CH=CH <sub>2</sub> ; R <sup>4</sup> , R <sup>5</sup> : -H	34%
b R <sup>3</sup> : -Bu; R <sup>4</sup> , R <sup>5</sup> : -H	60%
c R <sup>3</sup> : -Ph; R <sup>4</sup> , R <sup>5</sup> : -H	78%
d R <sup>3</sup> -R <sup>4</sup> : -(CH <sub>2</sub> ) <sub>4</sub> -; R <sup>5</sup> : -H	74%
e R <sup>3</sup> : -Et; R <sup>4</sup> : -H; R <sup>5</sup> : -Me	41%