



International Conference on Organic Synthesis

*July 1-4, 2012
Tallinn, Estonia*

*Balticum
Organicum
Syntheticum*

**Program
and Abstracts**



**OXY-GROUP DIRECTED DIASTEREOSELECTIVITY IN THE CYCLIZATION OF
ALLYLIC BIS-TRICHLOROACETIMIDATES FOR THE SYNTHESIS OF
C-QUATERNARY VINYLGLYCINOLS**

Klimovica, K.; Rasina, D.; Jirgensons, A.

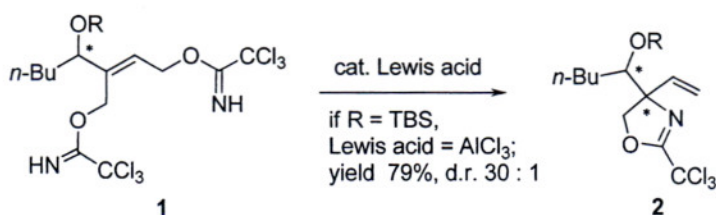
Aizkraukles 21, Riga, Latvia

Latvian Institute of Organic Synthesis

Latvia

dacerasina@osi.lv

C-Quaternary vinylglycinols are structural elements of a number of pharmaceutically relevant compounds and natural products. Previously, we have reported the synthesis of 4-vinyl-2-trichloromethyloxazolines as precursors of C-quaternary vinylglycinols by Lewis acid catalyzed cyclization of allylic bis-trichloroacetimidates.¹ Here we present the investigation of diastereoselectivity for the cyclization of chiral bis-imidates **1** bearing oxy group.



Synthetic routes toward substrates **1** having different *O*-substituents were developed and the stereoselectivity of the reaction was explored with a range of Lewis acid catalysts. The best diastereoselectivity and yield for the formation of oxazoline **2** was achieved with *O*-TBS protected substrate **1** and AlCl₃ as a catalyst.

1. Klimovica, K.; Grigorjeva, L.; Maleckis, A.; Popelis, J.; Jirgensons, A. *Synlett* **2011**, 19, 2849 – 2851.