[Heterocycles, 38, 1751-1756 (1994)]

[Lab. of Pharm. Synthetic Chemistry]

## Diastereoselective Addition of Allyltriphenylstannane to 3-Sulfinylfurfural Mediated by Titanium (IV) Tetrachloride and Tin (IV) Tetrachloride.

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The addition of allyltriphenylstannane to 3-sulfinylfurfural in the presence of titanium(IV) tetrachloride proceeded with high diastereoselectivity to give the furyl alcohol, whereas the similar treatment with tin(IV) tetrachloride afforded the other diastereoisomeric alcohol, exclusively.

[J. Chem. Soc., Perkin Trans. 1, 1994, 15-24]

[Lab. of Pharm. Synthetic Chemistry]

Enantioselective Synthesis of (+)-Indolizine, (+)-Laburnine and (+)-Elaeokanines A and C Using the Diels-Alder Reaction of  $\alpha$ -(2-exo-Hydroxy-10-bornylsulfinyl) maleimides.

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The Diels-Alder adduct derived from N-butynylmaleimide and cyclopentadiene was transformed into the tetracyclic lactams and via a common precursor. The lactams were converted into (+)-indolizine and (+)-laburnine, respectively, via retro-Diels-Alder reaction. Similar methodology was successfully applied to the synthesis of (+)-elaeokanine A and (+)-elaeokanine C.

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[Lab. of Pharm. Synthetic Chemistry]

Diels-Alder Reactions of Optically Active  $\alpha$ -(2-exo-Hydroxy-10-bornylsulfinyl)-maleimides and its Application to Optically Active 5-Functionalised Pyrrolidines via Retro-Diels-Alder Reaction.

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Optically pure sulfinylmaleimides were synthesized. The Diels-Alder reactions of the sulfoxides with various dienes showed high diastereoselectivity. Regioselective reduction of the adducts followed by desulfinylation afforded the  $\gamma$ -hydroxy lactams. N-Acyliminium additions using these comounds proceeded diastereoselectively to give  $\gamma$ -alkyl lactams by virtue of its conformationally rigid, bicyclo [2.2.1]- and 7-oxabicyclo [2.2.1] heptene moiety.