AN APPLICATION OF THE ANGULAR HYDROXYLATION USING BENZENE-SELENINIC ANHYDRIDE TO THE SYNTHESIS OF 108-HYDROXYFURANO-EREMOPHILANES AND RELATED COMPOUNDS

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The total syntheses of eremophilane type sesquiterpenes having furan ring, furanceremophilanes, have been achieved by the Diels-Alder reaction of <u>1</u> and <u>2</u>. In the case of the syntheses of highly oxygenated furanceremophilanes the hydroxylation of the angular (C-10) position of furanceremophilanes with benzene-seleninic anhydride [(PhSeO)₂O; BSA] was a critical step.

The Diels-Alder adduct (<u>3</u>) was converted to <u>4</u> in three steps [a) $AcOH-H_2O$; b) isomerization of the C-4 methyl group to β orientation with p-TsOH; c) ketalization]. The angular hydroxylation of <u>4</u> with BSA gave <u>5a</u> and <u>5b</u> in excellent yield. The synthesis of decompositin (<u>6</u>) and other $\Delta^{1,10}$ -furanceremophilanes were achieved from <u>5b</u>, stereoselectively. The furanceremophilanes having 10 β hydroxyl group (<u>8</u> and <u>9</u>) were synthesized from 10 β -hydroxy compound (<u>5a</u>) in good yield via epoxide (<u>7</u>) as a key intermediate. Synthesis of other highly oxygenated furanceremophilanes will be discussed.

