Synthesis of Montroumarin

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Z. Naturforsch. **58c**, 691–696 (2003); received November 12, 2002/April 14, 2003 A simple stereoselective synthesis of montroumarin [(3S)-6,8-dihydroxy-3-phenyl-3,4-dihydroisocoumarin] isolated from *Montrouziera sphaeroidea* has been achieved. Condensation

of benzoyl chloride with 3,5-dimethoxyhomophthalic acid afforded 6,8-dimethoxy-3-phenylisocoumarin (3) which on sequential saponification and esterification yielded the keto ester 5. Enantioselective reduction of the latter with baker's yeast directly furnished the (3S)-6,8-

dimethoxy-3-phenyl-3,4-dihydroisocoumarin (6) in good enantioselectivity which on demethviation provided montroumarin. All of the synthesized compounds were examined in vitro for antifungal activity. Key words: Montroumarin, Dihydroisocoumarin, Antifungal