RING-TRANSFORMATION REACTIONS OF AZAHETEROMODIFIED ADAMANTANES AND RELATED DERIVATIVES

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We have been exploring new synthetic routes to heteromodified polycyclic skeletons as one of novel new-typed heterocycles. We have now developed the following routes to 2-azaadamantyl system, 4-azahomoadamantano-imidazoles and -isoquinolinones.

(i) Oxaziranes 1 were prepared by m-CPBA oxidation of 4-azahomoadamant-4-enes 2.1 Compound 1 in cyclohexane was irradiated with a low pressure Hg lamp through a quartz filter under argon. The initial product in each cases was isolated by chromatography to afford N-acy1-2-azaadamantane 3 (44-67%). This ring-expansion and ring-contraction sequence of the adamantane skeleton thus provides a novel and convenient route to 2-azaadamantyl system. Thermal reaction of 1 with DMAD and methyl propiolate will be discussed also.

(ii) Dihydropyrazines 5 prepared from 4,5-dione 4 rearranged to homoadamantanoimidazoles 6 in high yields on photolysis (a high pressure Hg lamp, pyrex filter).

(iii) The enamide derivatives 7 were readily obtainable from 2b by acylation in high yields. Irradiation of 7 in ether through a quartz filter with a low pressure Hg lamp afforded photocyclization products 8 in 66-83% yields after chromatography. Dehydrogenation of 8 with DDQ gave the corresponding 4-azahomoadamantano [4,5-b]1(2H) isoquinolinones 9 in good yields.

References

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