

Synthesis of 1,2,5,6,10a,10b-Hexahydroindeno[7,1,2-*hij*]quinolizine-3,10-dione: a New Heterocyclic Ring System

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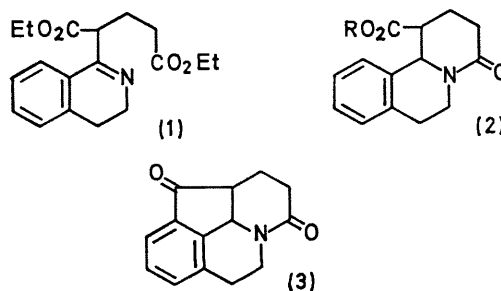
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Summary Diethyl 1-(3,4-dihydroisoquinol-1-yl)glutarate is cyclised reductively to give ethyl 4-oxo-1,3,4,6,7,11b-hexahydro-2*H*-benz[*a*]quinolizine-1-carboxylate: this ester and the corresponding acid are further cyclised by polyphosphoric acid forming 1,2,5,6,10a,10b-hexahydroindeno[7,1,2-*hij*]quinolizine-3,10-dione.

1,2,5,6,10a,10b-hexahydroindeno[7,1,2-*hij*]quinolizine-3,10-dione (3), m.p. 188—190° (ethanol) in 70% yield. The structures of this compound and of the acid (2; R = H) and the ester (2; R = Et) were confirmed by chemical tests and by i.r. and n.m.r.

peri-CYCLISATIONS of derivatives of 1-carboxyalkyltetrahydroisoquinoline into an aromatic ring activated by methoxy-groups are known.¹ We report a comparable *peri*-cyclisation of a benzoquinolizine derivative into an unactivated nucleus and leading to a previously unreported indenoquinolizine ring system.

Diethyl 1-(3,4-dihydroisoquinol-1-yl)glutarate (1) was prepared by a modification of the method of Agbalyan *et al.*² in which dimethylformamide was used as solvent. Hydrogenation at 60° and normal pressure using 5% palladium-charcoal gave ethyl 4-oxo-1,3,4,6,7,11b-hexahydro-2*H*-benz[*a*]quinolizine-1-carboxylate (2; R = Et)† m.p. 98—100° (ether-40/60 petroleum). The ester was easily hydrolysed to the corresponding acid (2; R=H), m.p. 229—230° (water). When heated with polyphosphoric acid at 170° for 10—15 min. both the ester and acid gave



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† Satisfactory elemental analyses have been obtained for all new compounds.

¹ G. C. Morrison and J. Shavel, *J. Org. Chem.*, 1964, **29**, 2486; K. Bernauer, *Experientia*, 1964, **20**, 380; R. M. Carlson and R. K. Hill, *J. Org. Chem.*, 1966, **31**, 2385; K. Bernauer, *Helv. Chim. Acta*, 1968, **51**, 1119.

² S. G. Agbalyan, L. A. Nersesyan, and Zh. A. Khanamiryan, *Armenian. khim. Zhur.*, 1967, **20**, 45.