

## Amidoalkylation of Olefins with 3-Benzyl-5-methoxyhydantoin

By D. BEN-ISHAI\* and G. BEN-ET

(Department of Chemistry, Technion—Israel Institute of Technology, Haifa, Israel)

**Summary** 3-Benzyl-5-methoxyhydantoin alkylates olefins, in the presence of an acid catalyst, to give 5-alkylhydantoin with an unsaturated side chain.

in the presence of boron trifluoride-ether complex as catalyst, isobutene reacted with 5-butoxyhydantoin to give a 44% yield of a product with a C<sub>8</sub> side-chain (IV, m.p.

THE reactions of 5-alkoxyhydantoin with aromatic compounds<sup>1</sup> and conjugated dienes<sup>2</sup> has now been extended to mono-olefins. It was found that 3-benzyl-5-methoxyhydantoin alkylates various olefins in boiling benzene and in the presence of naphthalene-2-sulphonic acid, to give one of the isomeric products (I), (II), or (III), or a mixture of products (see Table):

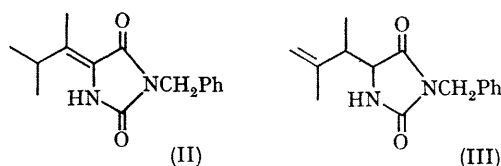
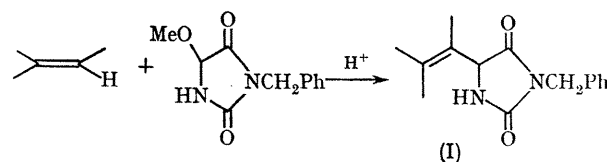
TABLE

| Olefin                   | Product     | M.p.            | Yield (%)          |
|--------------------------|-------------|-----------------|--------------------|
| 1,1-Diphenylethylene ..  | (I)         | 175°            | 78                 |
| 1,1-Diphenylpropylene .. | (I)         | 190°            | 67                 |
| α-Methylstyrene ..       | (I)         | 172°            | 36                 |
| β-Methylstyrene ..       | (I) + (II)  | oil             | 46 <sup>a</sup>    |
| Styrene ..               | (II)        | 174°            | 36                 |
| 2-Phenylbut-2-ene ..     | (III)       | 137°            | 68                 |
| 2-Methylbut-2-ene ..     | (I) + (III) | 73 <sup>a</sup> | 73 <sup>a</sup>    |
| Tetramethylethylene ..   | (III)       | 132°            | 75                 |
| Isobutene ..             | (I) + (III) | oil             | 33 <sup>a, b</sup> |
| Indene ..                | (I)         | 167°            | 35                 |

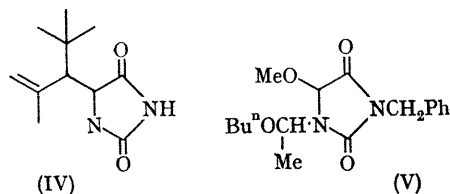
<sup>a</sup> The two isomers were eluted together from an alumina column.

<sup>b</sup> The major product was (I), according to the n.m.r. spectrum of the mixture.

The structure of the products obtained depends on the nature of the olefins used. Tetraphenyl- and triphenylethylene as well as 2-methyl-1,1-diphenylpropene failed to react with the methoxyhydantoin under the conditions used. Isobutene reacted with 3-benzyl-5-methoxyhydantoin in benzene solution, in the presence of the acid catalyst, at 80° (sealed tube) to give a mixture of products (see Table). In methylene chloride, at room temperature and



177°). Butyl vinyl ether and dihydropyran on the other hand, alkylated the hydantoin on the nitrogen to give oily products (e.g. V).



The assignment of structures to the substituted hydantoin are based on their i.r., n.m.r., and mass spectra (molecular peaks).

(Received, September 22nd, 1969; Com. 1423.)

<sup>1</sup> G. Ben-Et and D. Ben-Ishai, *Chem. Comm.*, 1969, 376.

<sup>2</sup> E. Goldstein and D. Ben-Ishai, *Tetrahedron Letters*, 1969, 2631.