The Reaction of 1,2,4-Triazoline-3,5-diones with Mono-olefins

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ONE of the most reactive dienophiles is 4-phenyl-1,2,4-triazoline-3,5-dione (I); solutions of this red dione are decolorized instantaneously at -78° by cyclopentadiene. 1,2° We now report that 1,2,4-triazoline-3,5-diones are also extremely reactive toward mono-olefins having allylic hydrogens. A methylene chloride solution (0.01m) in (II)3 and 1.0m in cyclohexene is decolorized at 25° within 6 min. after mixing. Ethyl azodicarboxylate also reacts with olefins having allylic hydrogens4 but far more slowly; a comparable solution of this ester in an evacuated sealed tube is incompletely reacted even after 3 weeks at 50°. Thus dione (II) is at least thirty thousand times more reactive toward cyclohexene than is ethyl azodicarboxylate.

The olefin-dione reaction results in the quantitative formation of a 1:1 adduct of the additive-substitution type (III-VI) first noted by Diels and

Alder^{5,6} and more recently by Thaler and Franzus.⁷ The structures of the adducts arising from the aliphatic olefins, (III) and (IV), indicate that a

shift in the position of the double bond has occurred; however, one cannot conclude from the structures of the alicyclic olefin addition products, (V) and (VI), whether or not a migration of the double bond has occurred.

The Table indicates the structures of the adducts arising from the various mono-olefins. All adducts have the correct elemental composition and have i.r. and n.m.r. spectra consistent with the suggested structures.

TABLE

Products from the reaction of 4-methyl-1,2,4-triazoline-3,5-dione with mono-olefins

Mono-olefin	Product	M.p. (°c)
2,3-Dimethylbut-2-ene	(III)	95.0 - 96.0
cis-But-2-ene	(IV)	$72 \cdot 0 - 73 \cdot 0$
trans-But-2-ene	(IV)	$71 \cdot 0 - 72 \cdot 0$
Cyclohexene	(\mathbf{V})	149.0-149.6
Cyclopentene	(VI)	118.5 - 119.5

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