Cycloaddition Reactions of 1,1-Diphenylethylene Triplets

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Summary Irradiation of 1,1-diphenylethylene in the presence of certain olefins leads to 2 + 2 cycloadducts, sometimes accompanied by products of hydrogen abstraction, by a process apparently involving diphenylethylene excited triplets.

Rosenberg and Serve,1 have shown that the excited triplet state of 1,1-diphenylethylene has hydrogen-abstracting capability similar to that of the lowest triplet of benzophenone.2 We now report results on the cycloaddition of excited 1,1-diphenylethylene (1) to simple olefins, a process analogous to oxetan formation from benzophenone lowest triplets.3

Irradiation (0.02 mol, Corex filter, 450 W mediumpressure mercury arc) of (1) in the presence of a 30-50-fold excess of isobutene for 80 h resulted in the isolation, after distillation and preparative g.c., of 1,1-dimethyl-2,2diphenylcyclobutane (2)† in 63% yield, based on unrecovered starting material [33% conversion of (1)]. Compound (2) was identified by its n.m.r. $[\tau 3.0 (10H, narrow m)]$ 7.30 (t, J 7.4, 2H), 8.23 (t, J 7.4, 2H), and 9.00 (s. 6H)] and mass spectral $[m/e \ 236 \ (rel. intensity \ 29), \ 208 \ (15), \ 193 \ (11),$ 180 (100), 165 (53), and 115 (30)] data.

Irradiation of (1) in cyclopentene (32 h) gave 56% of a mixture of (3), 3-(1,1-diphenylethyl)cyclopentene, [n.m.r. τ 3.04 (m, 10H), 4.3—4.8 (2H, br m), 7.8—8.7 (4H, m) 8.48 (3H, s); m/e 248 (20) 180 (100), 171 (73)] and (4), 7,7diphenylbicyclo[3,2,0]heptane,† [\tau 2.9 (10H, m) and 7.8-8.5 (10H, m); m/e 248] in a 3:1 ratio, separable by g.c. on SE-30.

From irradiation through Corex for 32 h of a mixture of

† All compounds so designated give acceptable analytical data.

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³ D. R. Arnold, Adv. Photochem., 1969, 6, 301.

*Compare data for 3,3-diphenylcyclobutene: M. Pomerantz, J. Amer. Chem. Soc., 1966, 88, 5349.

H. M. Rosenberg, P. Servé, and R. Rondeau, Canad. J. Chem., 1969, 47, 4295.

(1) and 1,2-dichloroethylene, followed by treatment of the reaction mixture with zinc dust, there was obtained, in addition to much recovered (1), small amounts ($\leq 10\%$) of 3,3-diphenylcyclobutene identified by its spectral properties.4 Other olefins tried as substrates (2-methylbut-2-ene

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and tetramethylethylene) only gave hydrogen abstraction products analogous to (3). Irradiation of (1) in isobutene in the presence of sufficient xanthone to absorb >99% of the incident light resulted in formation of (2), at a slower rate. The addition of (1) (0.2m) to isobutene was completely quenched by piperylene (0.4M). The additions must therefore proceed via an excited triplet. The similarities in chemical behaviour of triplet (1) and benzophenone thus include the 2 + 2 cycloaddition to olefins, as well as hydrogen abstraction1,5

Added in proof. Addition of triplet (1) to furan gives 55% of a 1:1 adduct, m.p. 83-84°.

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