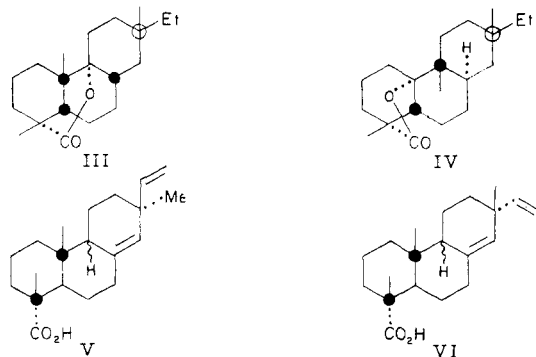


13-substituent in an axial configuration. Thus pimaric acid is V and isopimaric acid VI.<sup>8</sup>



The thus-derived C-13 stereochemistry of the pimaric acids implies that the biosynthesis of ring C takes place in a non-specific manner. Hence, the stereochemistry of tetracyclic diterpenoids, phyllocladene, the aconite alkaloids, etc., cannot be predicted with safety on biogenetic grounds.<sup>9</sup>

(8) The present chemical data corroborate the structural conclusions from surface tension measurements of the pimaric acids [H. H. Brunn, *Acta Acad. Aboensis, Math. et Phys.*, **19**, (3), 7 (1954)]. The authors are most grateful to Mr. L. J. Gough for first drawing their attention to this work.

(9) The authors express their gratitude to Drs. Lawrence and Levan-Thoi and Professor Jeger for gifts of the resin acids and to the Institute for Atomic Research, Ames, Iowa, for the use of a Baird infrared spectrophotometer.

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#### ACETYLENIC $\pi$ -COMPLEXES OF CHROMIUM IN ORGANIC SYNTHESIS<sup>1</sup>

Sir:

The concept of the  $\pi$ -complexed intermediate originating from the interaction of sufficiently activated  $\pi$ -electron systems, exemplified previously by aromatic Grignard compounds,<sup>2</sup> with chromium has led to the examination of other  $\pi$ -electron systems and of the prospects of isolating intermediates and final products arising from their reaction with the transition metals. Evidence of the ability of chromium to complex acetylenes and of the fusion of these complexes to aromatic structures has been obtained.<sup>3</sup> The description by Sternberg, Markby and Wender<sup>4</sup> of the use of iron pentacarbonyl in preparing duroquinone constitutes another example of this phenomenon. We now wish to report a specific one-step synthesis of hexamethylbenzene and 1,2,3,4-tetramethylnaphthalene from 2-butyne via a  $\pi$ -complex synthesis on chromium as a prototype of a general synthesis of aromatic molecules from acetylenes.

Triphenylchromium tri-tetrahydrofuranate<sup>1</sup> undergoes heterogeneous reaction smoothly with 2-butyne at 20° within a period of several minutes.

(1) Paper V of " $\pi$ -Complexes of the Transition Metals"; Paper IV, W. Herwig and H. H. Zeiss, *THIS JOURNAL*, **79**, 6561 (1957).

(2) H. H. Zeiss and W. Herwig, *THIS JOURNAL*, **78**, 5959 (1956); *Ann.*, **606**, 209 (1957).

(3) H. H. Zeiss, "Handbook, XVth International Congress of Pure and Applied Chemistry," July, 1957, Paris, p. 134.

(4) H. W. Sternberg, R. Markby and I. Wender, *THIS JOURNAL*, **80**, 1009 (1958).

Hydrolysis and ether extraction of the mixture are employed to obtain a mixed crystalline product which may be separated by fractional crystallization of the picrates. Under heterogeneous conditions about equal amounts of hexamethylbenzene, whose identity was confirmed by mixed m.p. and ultraviolet-infrared spectral comparisons with authentic substance, and 1,2,3,4-tetramethylnaphthalene, m.p. 107–108.5°, picrate m.p. 183.5–184.5° [*Anal.* Calcd.: C, 91.25; H, 8.75; and N, 58.11; H, 4.63; N, 10.17, respectively. Found: 91.45; H, 8.59; and C, 58.13; H, 4.34; N, 10.55], hitherto available by a five-step synthesis from prehnitene,<sup>5</sup> are formed.

The condensation of 2-butyne on chromium may also be carried out homogeneously in tetrahydrofuran solution and in this case product control has been found to be possible. For example, if the temperature of the solution is maintained at 0° for 3 days or at room temperature for 3 hours, the sole aromatic product is the naphthalene derivative; and if the solution is maintained at room temperature for 24 hours or heated to reflux for 2 hours, hexamethylbenzene is produced admixed with the naphthalene. However, the formation of the tetramethylnaphthalene in this reaction may be suppressed entirely by the use of triethylchromium in tetrahydrofuran solution with 2-butyne, and in this case the only aromatic product is hexamethylbenzene.

The product, hexamethylbenzene, arising from the reaction between triphenyl- or triethylchromium and 2-butyne clearly requires the intervention of a  $\pi$ -complexed intermediate such as  $R_3Cr(CH_3C\equiv CCH_3)_3$ , this being formed by replacement of the coordinating tetrahydrofuran molecules in  $R_3Cr(THF)_3$  by 2-butyne, since this acetylene is unaffected either by chromic trichloride (or its tetrahydrofuranate) or by phenylmagnesium bromide. The participation of the phenyl groups of triphenylchromium in this condensation reaction leading to the naphthalene by interaction of phenyl with two molecules of 2-butyne in the  $\pi$ -complex and consequent *ortho* ring closure opens the way for the synthesis of various condensed aromatic ring systems by an appropriate choice of triarylchromium and acetylene. These reactions will be reported later in detail.

(5) M. C. Kloetzel, R. P. Dayton and H. L. Herzog, *ibid.*, **72**, 273 (1950). Ultraviolet and infrared spectral comparisons confirmed the identity of the two samples.

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#### PHOTOCHEMICAL REACTIONS OF KETONES IN SOLUTION

Sir:

The mechanism of photolysis of ketones has been a subject of current interest,<sup>1-6</sup> and we wish to

(1) W. Davis, Jr., and W. A. Noyes, Jr., *THIS JOURNAL*, **69**, 2153 (1947).

(2) A. J. C. Nicholson, *Trans. Faraday Soc.*, **50**, 1067 (1954).

(3) T. W. Martin and J. N. Pitts, *THIS JOURNAL*, **77**, 5465 (1955).

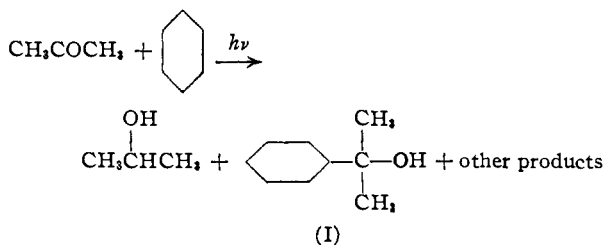
(4) P. P. Manning, *ibid.*, **79**, 5151 (1957).

(5) J. R. McNesby and A. S. Gordon, *ibid.*, **80**, 261 (1958).

(6) P. Ausloos, *Can. J. Chem.*, **36**, 400 (1958).

report our results on the light induced reactions of simple aliphatic ketones in saturated hydrocarbon solvents.<sup>7</sup> Ketones having no  $\gamma$ -hydrogen were found to give the corresponding alcohol and the addition product with the saturated hydrocarbon. Ketones with  $\gamma$ -hydrogens were found to give acetone, an olefin and a cyclobutanol derivative. These reactions demonstrate new types of photochemical reactions markedly different from those previously observed in gas phase. The formation of cyclobutanols indicates a simple, readily accessible synthetic route to a group of compounds otherwise difficult to prepare.

When a mixture of acetone and cyclohexane (1:8) was irradiated in a quartz immersion irradiator,<sup>8,9,10</sup> a small amount of methane and carbon monoxide was obtained. The products isolated by fractional distillation and gas chromatography were 2-propanol (53%), pinacol (15%), acetylacetone (8%), and cyclohexyldimethylcarbinol (I, 12%).<sup>11,12</sup> The recovered cyclohexane showed slight unsaturation. Cyclohexylcyclohexene and hydrocarbon polymers also were obtained.



Under similar condition, irradiation of 2-pentanone (IIa) in cyclohexane gave acetone, ethylene and 1-methylcyclobutanol (IIIa, 12%); phenylurethane of IIIa, m.p. 141° (Found: C, 70.14; H, 7.58; N, 6.84), identical with that of an authen-

(7) For a review on photochemical reactions, see C. R. Masson, V. Boekelheide and W. A. Noyes, Jr., "Technique of Organic Chemistry," Vol. II, Interscience Publishers, New York, N. Y., 1956, p. 257.

(8) M. S. Kharasch and H. N. Friedlander, *J. Org. Chem.*, **14**, 245 (1949).

(9) See ref. 7, p. 274.

(10) All reactions were carried out at 10–15° and the percentage conversions of the ketones ranged from 25 to 30%.

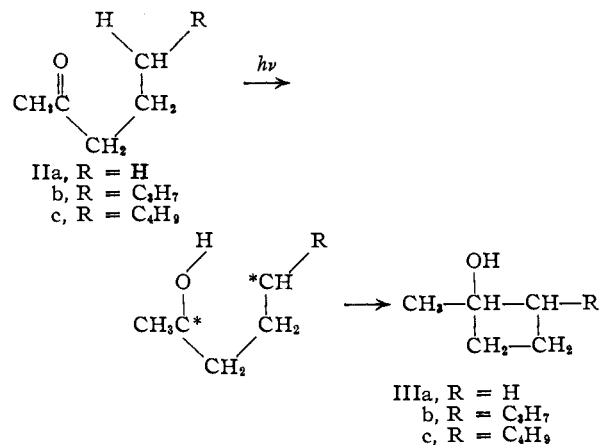
(11) P. Sabatier and A. Mailhe, *Ann. chim.*, [8] **10**, 539 (1907).

(12) E. J. Bowen and A. T. Horton, *J. Chem. Soc.*, 1685 (1936).

tic sample.<sup>13</sup> Compound (IIIa) and the authentic sample also exhibit identical infrared spectra and retention time in gas chromatography.

Irradiation of 2-octanone (IIb) in isoöctane gave, in addition to equimolar quantities of acetone and 1-pentene (67%), a saturated cyclic tertiary alcohol (IIIb, 17%) isomeric with 2-octanone, b.p. 67–68° (9 mm.),  $n_D^{20}$  1.4420 (Found: C, 75.00; H, 12.90); phenylurethane, m.p. 78° (Found: C, 72.75; H, 8.55; N, 5.72). The n.m.r. spectrum of IIIb indicates that it is 1-methyl-2-propylcyclobutanol.<sup>14</sup> Gas chromatography and infrared spectral analysis showed that the 1-pentene obtained was not contaminated by other isomers. Similarly, irradiation of 2-nonanone (IIc) in isoöctane yielded acetone (60%), 1-hexene (60%) and 1-methyl-2-butylcyclobutanol (IIIc, 10%), b.p. 49–50° (0.8 mm.),  $n_D^{20}$  1.4441 (Found: C, 76.21; H, 13.05); phenylurethane, m.p. 91° (Found: C, 73.42; H, 9.06; N, 5.73).

An intramolecular mechanism is formulated for the formation of cyclobutanols.



(13) Prepared from the reaction of cyclobutanone and methylmagnesium bromide.

(14) The authors wish to thank Dr. G. V. D. Tiers of Minnesota Mining and Manufacturing Company for the interpretation of the n.m.r. spectra.

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## BOOK REVIEWS

**Experimental Crystal Physics.** By W. A. WOOSTER, Department of Mineralogy and Petrology, Cambridge. Oxford University Press, 114 Fifth Avenue, New York 11, N. Y. 1957. viii + 115 pp. 14.5 × 22 cm. Price, \$2.90.

This book is based on a course in experimental crystal physics given at Cambridge. The topics covered are the optical, magnetic, thermal, plastic, piezoelectric, pyroelectric and elastic properties of crystals. The experiments described are "only those . . . which can be carried out in about two hours." For this reason the experiments are of necessity either qualitative or semiquantitative and deal only with the less complicated aspects of crystal physics. This is as it should be in a beginning course, although the

reviewer feels that a few four or six hour experiments should be (and probably are) thrown in among those described in the text. The course is excellent and should be emulated by all schools of crystallographers, crystal chemists and crystal physicists.

In reading the book, a certain amount of imagination is required in the reconstruction of the course. A piece of apparatus is described in terms of schematic diagrams which would be invaluable to the student faced with it in the laboratory but which would be of little help to one trying to build it. Circuit diagrams are also somewhat vague, giving resistances, capacitances and impedances, but often failing to mention voltages and occasionally giving nothing more than  $V_1$  or  $V_2$  to tell one what tubes should be included in a circuit.