

ORGANIC CHEMISTRY.

1. INTRODUCTION.

THE following report on organic chemistry covers work published during the year except for the theoretical topics which are better reviewed at longer intervals than one year, so that the work discussed can be considered critically in relation to contemporary views. The section on theoretical organic chemistry, therefore, deals with the more salient features of the chemistry of free radicals in solution, since that rapidly advancing field has not been reviewed since 1948. The mechanism of the ozonisation of aromatic compounds, which was last reported in 1947, now merits special attention in view of the wider application of recent theories to other contemporary studies of the reactivity of polynuclear aromatic systems.

Perhaps the most novel discoveries of the year are the recognition of iron dicyclopentadienyl as a new type of symmetrical structure showing typical aromatic behaviour, and the use of clathrate compounds for optical resolutions. In synthetic organic chemistry the year has been notable for a stereospecific total synthesis of cortisone, a synthesis of flavin-adenine-dinucleotide, a synthesis of morphine, and a synthesis of all-*trans*-methylbixin; each of these considerable achievements exemplifies in some way or other the high degree of specificity of modern techniques in organic chemistry, to which has been added a welcome catalytic hydrogenation method for reducing triple bonds to double bonds. The structure of lanosterol has now been established and the stereochemistry of the β -amyrin and lupeol groups of triterpenes has been elucidated, while appreciation of the rich variety of types of substance found to occur naturally is increased by the isolation of an antibiotic incorporating every known type of carbon-carbon unsaturation.

The increasing use of enzymic methods for the study of organic chemical problems is seen in recent work on nucleotides and macromolecules, while expanding facilities for infra-red spectroscopy are reflected in the increasing use being made of this tool in structural problems.

J. W.
W. A. W.

2. THEORETICAL ORGANIC CHEMISTRY.

Bromonium Cations.—Though iodonium salts have long been known, stable bromine and chlorine analogues (I) and (II) have been prepared by



R. B. Sandin and A. S. Hay¹ only in the year under review. The corresponding iodonium salts were made by L. Mascarelli in 1907.² Previously,

¹ *J. Amer. Chem. Soc.*, 1952, **74**, 274.

² *Atti R. Accad. Lincei*, 1907, **16**, II, 562; 1908, **17**, II, 580; 1912, **21**, II, 617; *Gazzetta*, 1908, **38**, 624.

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Christian G. Meyer



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