

expect that  $(C_2H_4)_4Rh_2Cl_2$  will be found to be structurally similar to  $(CO)_4Rh_2Cl_2$ .

$\mu$ -Dichlorotetraethylenedirrhodium is only sparingly soluble in such solvents as ethanol, acetone, dioxane, chloroform, or methylene chloride and generally cannot be recovered pure from these solvents. The solid is moderately stable. Samples occasionally darken after storage for several days at room temperature but are unaltered for months at 5°. Upon slow heating in an evacuated tube there is little change below 100°, but at 115° the surface of the crystals becomes black, and further heating causes rapid destruction without evidence of melting.

The coordinated ethylene of I is displaced by a number of ligands. For example, when suspensions in methanol were treated with cyclooctadiene, ethylene was displaced within 2 min., and the yellow  $\mu$ -dichlorobis-(cyclooctadiene)-dirrhodium(I) described by Chatt<sup>6</sup> precipitated. A similar displacement of ethylene occurred when I was treated with triphenylphosphine, hydrogen cyanide, acrylonitrile, or pyridine.

A propylene analog of I has been prepared from propylene and hydrated rhodium(III) chloride. It is more soluble and dissociates more readily

(6) J. Chatt and L. M. Venanzi, *J. Chem. Soc.*, 4735 (1957).

than the ethylene complex in solution or when heated.

CONTRIBUTION No. 782 RICHARD CRAMER  
CENTRAL RESEARCH DEPARTMENT  
EXPERIMENTAL STATION  
E. I. DU PONT DE NEMOURS AND COMPANY  
WILMINGTON, DELAWARE

RECEIVED JUNE 15, 1962

### A Correction for the Lattice Constants of Mixed Metal Oxides of Rhenium and Osmium with the Hexagonal Barium Titanate Structure

Sir:

The correct lattice constants for the following compounds erroneously reported in *Inorganic Chemistry*, 1, 245 (1962) are

	a, Å.	c, Å.
Ba <sub>3</sub> Fe <sub>2</sub> ReO <sub>9</sub>	5.81	14.10
Ba <sub>3</sub> Cr <sub>2</sub> ReO <sub>9</sub>	5.70	13.8
Ba <sub>2</sub> MnOsO <sub>6</sub>	5.82	14.2
Ba <sub>2</sub> FeOsO <sub>6</sub>	5.76	14.1

DEPARTMENT OF CHEMISTRY  
UNIVERSITY OF CONNECTICUT  
STORRS, CONNECTICUT

ROLAND WARD

RECEIVED JUNE 21, 1962

## Book Review

**A Text-Book of Quantitative Inorganic Analysis Including Elementary Instrumental Analysis.** Third Enlarged Edition. By ARTHUR I. VOGEL, Woolwich Polytechnic, London. John Wiley and Sons, Inc., 440 Park Avenue South, New York 16, N. Y. (and Longmans, Green and Co., Ltd.), 1961. xxx + 1216 pp. 16 × 23 cm. Price, \$12.00.

Here, indeed, is a book! The comprehensive versatility of this author, whose texts on practical organic chemistry and qualitative analysis are equally well known, is admirably demonstrated in this expanded version of his work on quantitative inorganic analysis. One cannot resist the temptation to compare this work with another having similar aims and scope (though, only 700 pp.), namely, Charlot and Bezier's "Quantitative Inorganic Analysis." It is the latter work which tends to suffer by the comparison. The less critical choice of some procedures and their abbreviated descriptions, and the superficiality with which the theory and practice of some techniques are outlined, comprise possible objections to any such ambitious a coverage in a one-volume treatise. Though using 1246 pages in the process, Dr. Vogel has, by and large, succeeded in avoiding both criticisms.

Represented as being a student text suitable for his

entire training in this area and as being of value to practicing analytical chemists, this book is fairly described as being *all of that*. The theory and practice of classical gravimetric and titrimetric methods are adequately treated. Advantageous and proven newer reagents and techniques applicable to "wet chemical" procedures are described for *use*, and not merely added as a weak appendage indiscriminately enumerating *all* that is new. A similar comment applies with respect to the inclusion of good methods for such elements as molybdenum, zirconium, uranium, etc. An excellent chapter on complexometric titrations is largely devoted to the applications of ethylenediaminetetraacetic acid (EDTA).

The practical and theoretical coverage of "instrumental" measuring techniques is very good and embraces most of the more familiar and consequential electrical and optical themes. Certain specialized techniques (such as the use of X-rays, Raman spectroscopy, radiochemical methods, nuclear magnetic resonance spectroscopy, etc.) are, understandably, not treated. Details on separations *via* distillation and gas (adsorption) chromatography are not given, on the grounds that these are of more general value in organic work. Gas-evolution techniques are concisely treated, however, and a detailed chap-