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

μ-Dichlorotetraethylenedirhodium 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 coördinated ethylene of I is displaced by a number of ligands. For example, when suspensions in methanol were treated with cycloöctadiene, ethylene was displaced within 2 min., and the yellow μ-dichlorobis-(cycloöctadiene)-dirhodium(I) described by Chatt[6] 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 than the ethylene complex in solution or when heated.

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

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.
$Ba_3Fe_2ReO_9$	5.81	14.10
$Ba_3Cr_2ReO_9$	5.70	13.8
Ba_2MnOsO_6	5.82	14.2
Ba_2FeOsO_6	5.76	14.1

DEPARTMENT OF CHEMISTRY ROLAND WARD
UNIVERSITY OF CONNECTICUT
STORRS, CONNECTICUT

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-

ter on gas analysis logically includes the handling of organic gases and vapors. American readers may be interested to know that the author describes the use of as much, or more, apparatus of American, as of British, manufacture.

A feature which should merit the commendation of most readers is the inclusion of realistic chapters on the theory and practice of such important separation techniques as ion exchange, chromatography, and solvent extraction. An excellent short chapter on inorganic microgravimetric and microtitrimetric analysis is frosting on the cake. As with the book of Charlot and Bezier, a *very* brief chapter on errors in quantitative analysis "exists"; in a minimal sense it is acceptable, but unexceptional. In a work of this magnitude, the inclusion of an extensive and well organized table of contents, appendix, and index are worthwhile features.

Objection may be raised to the author's conservation of space by omitting specific literature references throughout the text, though selected bibliographies follow each chapter. A more serious objection, related to use as a textbook, concerns the absence of any numerical problems. But some instructors prefer to dispense their own problem material, and the textual exposition of theory often does include typical calculations.

A good treatment of procedures for the analysis of complex materials (ferrous and non-ferrous alloys, limestone, silicate mineral, etc.) is included. The procedure given for the analysis of a silicate mineral provides one example of the struggle between modernism and classicism. Classicists may resent the abbreviated description given the old approach to this analysis and the greater attention given a streamlined substitute. Neither the NaOH fusion nor the Berzelius (HF + H_2SO_4) decompostion specified have quite the universality of applicability of the corresponding Na_2CO_3 fusion or J. L. Smith method. But the use of a Berzelius dissolution with the EDTA determinations of Ca and Mg and flame photometry for Na and K must certainly conform with more typical modern practice.

This reviewer is conservative enough, however, to join the classicists' club in bemoaning the use of a silica assay in a silicate mineral *via* a low temperature drying form of quinoline silicomolybdate. But the majority of Dr. Vogel's innovations are not this drastic.

Anyone concerned with quantitative inorganic analyses should be interested in this book. Those concerned with the teaching of this subject could seriously consider this text as a worthy guide for two semesters of work. Dr. Vogel is to be congratulated on his uniquely "modern" approach *with* due regard to the basic and the "classic."

DEPARTMENT OF CHEMISTRY
UNIVERSITY OF MICHIGAN
ANN ARBOR, MICHIGAN

CHARLES L. RULFS

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