Metodi Di Separazione Nella Chimica Inorganica. MICHAEL LEDERER, Editor. Consiglio Nazionale Delle Ricerche, Rome, Italy. Vol. 1. 1963. 260 pp. 18 × 25.5 cm. Price, \$12.00.

This book, the first of two volumes, is a collection of some of the lectures given in Rome at a postgraduate summer school course in chemistry. The course was held in September, 1962 and was sponsored by the National Research Council of Italy and the F. Giordani Foundation.

The book contains ten chapters written in four different languages. Four chapters are in Italian, three in English, two in French, and one in German. The different languages cause a minimum of difficulty because each chapter is well written and very well illustrated. The most important contribution of this volume is that it focuses attention on the application of modern separation methods to inorganic systems. The theoretical basis, the useful applications, and the limitations of each technique are presented in a readable fashion by experts in the field. A total of 450 references are given which can be most helpful to one who wants more detailed information on a particular topic.

The topics, the authors, the language, and the number of pages are listed here and a brief comment is made to indicate the content of each chapter. "Liquid–Liquid Extraction" by H. Irving and R. J. P. Williams is in Italian, 50 pages. This is an excellent review of the theory and technique of separations by lquid–liquid extraction methods. The chapter is largely a translation of a chapter written by the same authors in "Treatise on Analytical Chemistry," Vol. I, Interscience, New York, N. Y., 1961. "Liquid Ionic Exchangers" by E. Cerrai is in Italian, 26 pages. This describes the separations of metal ions on inert supports impregnated with anionic and cationic liquid exchangers. "Adsorption Chromatography in Inorganic Chemistry" by L. Sacconi is in Italian, 20 pages. Most of the discussion deals with the properties and applications of alumina. "The Adsorption of Inorganic Substances on Paper" by M. Lederer, is in English, 8 pages. Examples are given of specific systems that show changes in  $R_t$  values with changes in solvent and added HCl or LiCl. "Chromatographic Separations on Paper Impregnated with Synthetic Inorganic Exchangers" by G. Alberti is in Italian, 12 pages. Examples are given of separations on paper impregnated with zirconium phosphate and with ammonium phosphomolybdate. "Gas-Liquid Chromatography" by A. T. James is in English, 2 pages. This is an abstract and has no references. "High-Voltage Electrophoresis in Inorganic Chemistry" by D. Grass is in English, 10 pages. The apparatus and technique are described in some detail. "Chromatography and Complex Chemistry" by E. Blasius is in German, 58 pages. This is a very complete treatment providing also details of the author's research on the halogeno and mixed halogeno complexes of the platinum metals. "Chromatography of the Condensed Phosphates and Other Polyanions' by J. P. Ebel is in French, 50 pages. Only a very short survey, 2 pages, is given of the other iso- and hetero-polyanions. "Radiochromotography in the Gas Phase" by J. P. Adloff is in F

The printing in the book, on glossy paper, is excellent. Its illustrations are good and it has relatively few typographical errors. This is a valuable reference book on inorganic separations, which with the next volume promised should provide a fairly

complete treatise on the subject.

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Physical Adsorption of Gases. By D. M. Young, Research Department, Dow Chemical of Canada, Limited, and A. D. Crowell, Professor of Physics, University of Vermont. Butterworth, Inc., 7235 Wisconsin Avenue, Washington 14, D. C. 1962. 426 pp. 14 × 22 cm. Price, \$13.00.

In 1945, "The Adsorption of Gases and Vapors," by Brunauer, was published. His book, which was Volume I of a projected set, was devoted to physical adsorption. The book under review is an overdue replacement for Brunauer's valuable work. Many of the questions left unanswered in the earlier work have now been solved, and the new book appears to give a definitive treatment of the subject.

The authors have found a reasonable balance between an exhaustive literature survey and a readable treatment that can be used profitably without having to look up the original papers. They appear not only to have read, but understood the large number of papers that they have chosen to digest in whole or part. The book is rich in useful figures and tables.

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On the whole, they have chosen a logical scheme of presentation. A chapter on intermolecular forces, with specific emphasis on forces at a surface, is adequate, although necessarily fragmentary. A proper consideration of the fundamentals of this subject would involve considerably more molecular quantum

mechanics than is appropriate in a book of this type, and would take the reader far afield from the specific topic of surfaces. The next chapter presents a consistent account of the thermodynamics of adsorption. The vexed subject of the relationship of pure thermodynamic results to particular molecular models is now at least well-aired and the authors treat this problem thoroughly.

In the next two chapters the authors take up specific adsorption isotherms. A random sampling shows that the treatment of the Hüttig isotherm is correct, although somewhat less severe than the reviewer would have liked. On the other hand, the treatment of the Volmer isotherm, although not incorrect, fails to expose the exact nature of the approximation that makes it inapplicable at high coverages. The authors imply that a comparison between it and the Langmuir equation is valid near  $\theta=1$ , which is certainly wrong.

It is perhaps wise to have a separate chapter on surface area estimation, although the subject is inextricably mixed up with isotherm equations or thermodynamics. This arrangement is probably justified, however, because of the large number of readers who will be concerned almost solely with this chapter. A chapter follows on the measurement of isotherms.

Although there is a chapter on the properties of the adsorbed layers, these and related topics are relatively slighted in the book. Noticeably absent are the sorts of things that physicists and what may be called vacuum engineers study in the range of very low pressures and temperatures. Perhaps this omission arises from the arbitrary line that is drawn between physical and chemical adsorption. Certainly, Becker's study of the intermediate state that precedes the chemisorption of gases on tungsten would be relevant to the material in this book.

In sum, the book is very good, and should certainly become the standard work in this area for years to come.

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G. D. HALSEY, JR.

Mass Spectral Correlations. By Fred W. McLafferty. Eastern Research Laboratory, Dow Chemical Company, Framingham, Mass. American Chemical Society, Applied Publications, 1155 Sixteenth Street, N.W., Washington 6, D. C. 1963. 117 pp. 15.5 × 23.5 cm. Price, \$4.75.

This book consists of a tabulation (85 pages, preceded by a 13-page introduction) listing the most intense peaks in the approximately 4000 mass spectra (a collection larger than any other one publicly available at the present time) existing in the files of the Dow Chemical Company and of those spectra that have appeared in the literature prior to 1960. The three or five most intense peaks are listed up to mass 200, while peaks down to the tenth most abundant are listed up to mass 400.

The table lists first the mass of the ion, the empirical formula assigned to it, the structural characteristics of the compounds in whose spectrum it appeared, and finally the number of spectra (in that particular collection) in which it was found to be the most intense peak, or the second or the third one, etc. From these tabulations one can conclude in which type of compound a peak at a certain mass would be a very significant one.

The table thus aids in the identification of important peaks in a mass spectrum. For example, if one encounters an intense peak at m/e 74, one finds that such a fragment is easily formed from methyl esters,  $\alpha$ -methyl carboxylic acids, or amino alcohols; that m/e 31 is most probably due to primary alcohols, ethers, or fluorocarbons; that m/e 95 and 105 are indicative of furoates and benzoates, respectively; and that m/e 149 is formed from phthalates, to mention only a few. Obviously, many different but related compounds give rise to a peak at a certain mass, and their higher (or lower) homologs will lead to peaks 14 m $\mu$ higher (or lower). This is indicated by "etc.," suggesting to the user to search also up and down the homologous series to find other possibilities.

Because of the origin of the table, one is, of course, not guaranteed to find for each peak in question the correct answer, as this depends on the occurrence of this particular compound type in the Dow collection. In such a case it is important that the user realizes this fact and starts thinking of other possibilities not listed. Sometimes the suggestions in the book may lead him at least on the right track. For example, at mass 89 there is listed the group  $C_3H_3CH_2$ , but not  $C_2H_6SCH(CH_3)$ . This should not imply that the latter group does not give rise to an intense peak at mass 89 (it is, in fact, the most intense peak of ethyl (2-butyl) sulfide) but only that this particular type of compounds is not represented in the Dow collection and thus also not mentioned in the tables (even not indicated by ''etc.'').

The book will be most useful to those who are using mass spectrometry only occasionally and are therefore not very familiar with the interpretation of mass spectra. The more experienced mass spectrometrist may sometimes get a clue if faced with a spectrum of a compound type he has not previously encountered