In considering the book as a guide to laboratory practice, the unwillingness of the author to editorialize might hamper a practitioner of gas chromatography in selecting equipment or techniques. For example, Dr. Littlewood clearly describes eight ionization detectors, but prefaces the discussion with only a mild statement that the argon and flame ionization detectors are receiving much more attention than the others. In the United States, at least, the flame detector essentially monopolizes the field today, electron capture detectors are of increasing interest, and argon detectors are rapidly disappearing from use. Such information might be of more value to the reader looking to the text for guidance in practice. An analogous situation exists in the chapter on catharometry in which it implied that ion gages are equivalent to catharometers for low sensitivity work, are superior for high sensitivity work, and are gradually replacing them. Again, in the United States, at least 80% of all work is still performed on catharometer-equipped chromatographs, and their lower cost and operating convenience will probably perpetuate this situation.

Some minor errors were noted in the text. For example, in the coating of capillary columns (p. 208), methyl chloride is not a practical choice as a solvent; methylene chloride was probably used. Further, the oven evaporation system described as one technique is never used in practice. The author also states technique is never used in practice. The author also states rather dogmatically (p. 209) that capillary columns coated with polar stationary phases are far less efficient than those coated with non-polar phases. While early results tended to show this distinction, the difference has been shown to be a result of failure to produce equivalently thin coating films with polar liquids. Where the coating procedure is modified to yield suitably thin coatings, polar columns of very high efficiency have been pro-

In summary, Dr. Littlewood's book is highly recommended for clarity and completeness and should serve as an excellent general text in gas chromatography.

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nermodynamische Elektrochemie. Von Dr.-Ing. Erich Lange, ordentlicher Professor für Physikalische Chemie, Thermodynamische Elektrochemie. Vorstand des Instituts für Physikalische Chemie der Universität Erlangen-Nürnberg, in Gemeinschaft mit Dr. Phil Nat. HERMANN GÖHR, Assistent am Institute für Physikalische Chemie der Universität Erlangen-Nürnberg. Hüthig Verlag G.m.b.H., Postfach 727, Heidelberg, Germany. 1962. 429 pp. 16.5 × 23.5 cm. Price, DM. 39.

The outstanding characteristic of this book is that it presents electrochemistry in a series of brief chapters arranged in order of increasing complexity with respect to the thermodynamic phases under consideration. Beginning with a single phase in the absence of current, it proceeds to discuss a single phase with current passing (conductance, transference, diffusion), transport processes in a non-isothermal single phase, a simple electrode at equilibrium, the Volta potential of a two-phase electrochemical system, the electrochemical double layer, phase transformations with flow of current at simple electrodes, Peltier heat at simple electrodes, some electrochemical three-phase systems at equilibrium, the ideal galvanic cell at equilibrium, the real galvanic cell, the galvanic thermocell, a general thermodynamic-kinetic view of polarized electrodes a general thermodynamic-kinetic view of polarized electrodes, a general thermodynamic-kinetic view of polarized galavanic cells, the twofold electrode, the surface-layer equilibrium electrode, the surface-layer mixed electrode, the corrosion surface-layer electrode, the double surfacelayer electrode and abrasion effects at electrodes.

This approach will undoubtedly appeal more to the electrochemical pragmatist than to the thermodynamic purist. pealing as the idea is to proceed gradually from simplicity to complexity, an inevitable result is that single ion activities are introduced before mean activities of strong electrolytes (p. 66).

An exceptionally clear distinction is made between Volta and

galvani potentials, and the treatment in general is orderly, sys-

tematic and logical. The notation unfortunately becomes cumbersome, and the reader finds himself making frequent reference to the five-page list of symbols at the front of the book. Even simple equations become loaded down, as for example the equation for the oxygen electrode reaction in acid solution (p. 70)

$$_{1}\Theta + _{11}H^{+} + _{1}/_{4} _{111}O_{2} \rightarrow _{1}/_{2} _{11}H_{2}O$$

where the subscripts I, II and III refer to the metal, solution and

gas phases, respectively

A more serious criticism though, is that the kinetics which is allowed to permeate a "thermodynamic" book is at best a treatment of the steady state, and the variables that affect the steady state are not adequately handled nor is the rate of attainment of steady state considered. Thus the Nernst diffusion layer concept (p. 196) is used to describe limiting currents, but the inadequacy of this treatment is not pointed out, nor are transient currents upon sudden polarization considered. It is true that an adequate discussion of transient and steady state processes would have lengthened the book considerably, but an incomplete and out-of-date treatment can hardly be defended in the light of the emphasis on transient phenomena in modern electrochemical research.

On p. 198, in considering the effect of supporting electrolyte on limiting currents, the charge-types but not the individual ion

mobilities are taken into account.

The book is relatively free from typographical errors, and the sality of paper, printing and binding are adequate. The Gerquality of paper, printing and binding are adequate. man is easily understood even by a reader whose native language is English. Nevertheless, it is difficult to find sufficient advantage in this treatment over material available in English to justify purchase by the average American electrochemist.

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Magnetische Kernresonanz und Chemische Struktur-Fortschritte der Physikalischen Chemie, Band 7. Low, Max-Planck-Institut für physikalische Chemie, Göttingen. Dr. Dietrich Steinkopff Verlag, Darmstadt, Germany. 1962. xii + 179 pp. 15.5 × 23 cm. Price, DM. 36. 1962.  $xii + 179 pp. 15.5 \times 23 cm.$ 

The author states in the preface that this book is intended as an introduction to nuclear magnetic resonance (n.m.r.) for chemists and students of chemistry, and should enable the reader to evaluate possible applications of n.m.r. in his own research. statement characterizes the scope of the book well. It is not a reference book, and neither does it present any original approaches in theoretical development or in chemical applications. It does provide a clear exposition of those aspects of n.m.r. which are of interest to the chemist, at a level which presupposes only average knowledge of mathematics or physics.

The book is divided into two about equal parts. The first, "physical principles," contains an introduction to relaxation and line width theory, a discussion of the Bloch equations and their solution, the principles of high resolution spectrum interpretation in terms of chemical shift and spin-spin coupling constants, and a short section on experimental methods. In the second part, "chemical applications," the use of n.m.r. in structural analysis is illustrated by a number of examples, both from organic and inorganic chemistry. The effect of hydrogen bonding and complexing on the n.m.r. spectrum is discussed, and the use of n.m.r. in fast chemical kinetics is treated in some detail. A short section on n.m.r. in solids follows.

The book is well written, and very few errors in printing or in fact have been noted by this reviewer (the statement on page 19 that for liquids  $T_1 \approx T_2$  is one of the few). As a short introduction to a large and rapidly growing field, it should prove very useful.

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