

Physicists will find this volume useful in making range-energy measurements, evaluating particle momentum by multiple scattering techniques, and making charge identification by grain and delta-ray counting. The theory and systematics of alpha decay processes are considered in great detail, together with quantitative theory on the formation of alpha stars in emulsion. The evaluation of neutron energy spectra is particularly thorough. About one-third of this scholarly work is devoted to applications of the nuclear emulsions in high-energy physics, both with collimated beams of particles from accelerators and in the study of the cosmic radiation. By comparison, coverage of other types of applications, such as that in the fields of geology and histautoradiography, while adequate in view of general principles developed in earlier sections, may appear scanty, occupying only 13 pages of fine-printed text.

In an age where individual scientists take pride in being expert on some small facet of nature, it is exhilarating to find a comprehensive work covering a broad vista of science skillfully integrated by a single mind. Demers is to be congratulated for his arduous efforts in providing a greatly needed, authoritative work on the use of photographic emulsions in nuclear physics.

HERMAN YAGODA  
*Ionospheric Physics Laboratory,  
Geophysics Research Directorate,  
Air Force Cambridge Research Center*

#### Effect of Radiation on Human Heredity.

Report of a study group convened by the World Health Organization, together with papers presented by various members of the group. World Health Organization, Geneva, 1957 (order from Columbia University Press, New York). 168 pp. Illus. \$4.

The purpose of the study group whose report is presented in this volume was, according to the preface, twofold: (i) to obtain the opinions of authorities on genetics from countries other than Great Britain and the United States, whose national committees on radiation hazards reported in 1956; and (ii) to probe certain untouched aspects of the problem of genetic hazards, especially "the lines of research which should be followed, in the light of our present knowledge, to increase our understanding of the genetic effects of ionizing radiations on man." At the meeting of the first International Congress of Human Genetics in Copenhagen during August 1956, a group of 20 highly expert investigators in various aspects of human genetics was assembled. The report actually comprises only ten pages of the volume, and while it

will undoubtedly stimulate some research to bring about the solution of critical problems and to supply key information, and while it may serve to increase financial support of research in human and other branches of genetics, this section is hardly the most valuable or interesting part of the book. It is hard not to mimic those generals who are forever fighting the last war.

The volume contains a collection of a dozen papers, however, every one of which is worth reading and rereading. The happy juxtaposition of the first two papers (by H. J. Muller and T. C. Carter, respectively) spotlights one of the most controversial matters in genetics today—one that is related to the whole problem of estimating genetic damage and hazard. This is the problem of the frequency of deleterious mutant genes that are always, or almost always, harmful no matter what the conditions, in comparison with the frequency of mutant genes that are seriously detrimental only when present in a person in double dose (homozygous) and which, when present in single dose (heterozygous), may, at least under certain conditions, be selectively advantageous. If the first situation predominates, then most mutant genes are held in the population in simple equilibrium between input (mutation) and outflow (elimination through failure to be passed on because of death or infertility). But if the second situation obtains, the interplay of forces is far more complex and consequently less predictable. Muller adopts the former view; Carter, and after him Wallace, the latter.

R. M. Sievert, of Sweden, gives a masterly summary of known human exposures to ionizing radiation. One may note the fact that he had no premonition of the recently discovered zone of extremely heavy radiation at high altitudes, for the curves drawn in 1956 all flatten out at about 50,000 feet. The Swedish data in general are in good agreement with the conclusions of the British and American committees. J. Lejeune discusses the kinds of data needed and the practicability of detecting induced mutations in the offspring of radiated parents. There is wide misapprehension on this subject. Lejeune is properly cautious. Court Brown describes methods being developed for more accurately determining the genetically all-important gonad dose delivered by various types of exposure.

Measurement of the spontaneous mutation rate in man by direct and indirect methods, the effects of induced mutations, the differential sensitivity of human loci to radiation, and the load of abnormal genes per person in the population are topics discussed by L. S. Penrose. There follows the first report of the radiation readings in inhabited areas with

particularly high background radiation, such as the region of monazite sands in Travancore. Gopal-Ayengar reports readings that are high, but hardly as high as rumor had previously made them. He estimates a total gamma dose of about 10 to 30 roentgens over the reproductive span of 30 years in that locality. The world will be understandably interested in the prompt investigation of the populations living on such soils. Is their genetic burden detectably higher than that of similar populations not so exposed to radiation?

Stevenson and Neel probe the difficulties and possibilities of error in estimating spontaneous mutation rates in human populations. Freire-Maia considers the effect of inbreeding in bringing more mutant recessive genes to the surface and exposing them more rapidly to selection. The frequency of genetic defect *manifested* in a population is *not* a simple function of the frequency of the genetically defective genes. Howard Newcombe, of Canada, completes the series of papers by describing the sort of vast genetic health registration of the population that is really needed if many of these problems are to be solved.

BENTLEY GLASS  
*Department of Biology,  
Johns Hopkins University*

**Electronic Instrumentation for the Behavioral Sciences.** Clinton C. Brown and Rayford T. Saucer. Thomas, Springfield, Ill., 1958. xiv + 160 pp. Illus. \$5.50.

In 160 pages this book provides "a simplified presentation of basic electronic theory required for instrumentation problems." The volume is specifically oriented toward fields of experimental psychiatry, psychophysiology, and physiology where electronic instrumentation is required for stimulation or measurement.

The first chapter offers a brief, lucid discussion of the physical theory underlying electronic phenomena, followed by an elementary review of methods of electrical measurement. Tube types and basic circuitry are next considered. This section is followed by chapters on power supplies, amplifiers, oscillators, timing devices, and switching circuits. Valuable suggestions are made regarding input and output transducers, including devices for tracing and displaying physiological changes. There is commendable emphasis on various methods for the protection of human subjects of experimentation. A chapter is devoted to test instruments, with suggestions regarding kits available for their economical construction. Recommendations are made regarding specific instruments that are