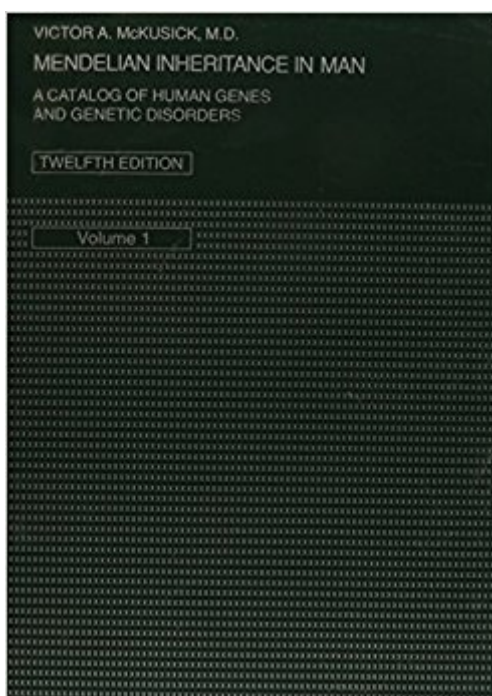


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Mendelian Inheritance In Man: A Catalog Of Human Genes And Genetic Disorders



Synopsis

The twelfth edition of this classic reference work includes:

- More than 2,000 new entries
- A total of more than 9,000 entries
- New features and enhancement of the familiar old features
- Mapping information on more than 4,000 genes of known function
- Information on specific point mutations responsible for more than 700 genetic disorders or neoplasms

Mendelian Inheritance in Man (MIM) is a genetic knowledgebase that serves clinical medicine and biomedical research, including the Human Genome Project. It aims to be comprehensive (not only complete, but also collated, integrated, and interpreted), authoritative (not only accurate but also sound in its interpretations and judgements), and timely (not only up-to-date but also historically dimensioned). From a review of the eleventh edition, *Reproductive Toxicology*: "Even the convenience of computer-based forms of MIM cannot eliminate the need for MIM in book form. The preface provides a wonderful synopsis of human genetics. The information contained in this text serves as a concise review for those with a genetics background." From a review of the tenth edition, *New England Journal of Medicine*: "[Victor McKusick] has been for all these years the shepherd of the development of the field [of clinical genetics]. Perhaps his most important pragmatic achievement has been the 10 editions of *Mendelian Inheritance in Man*, which rapidly became and has remained the principal source of information on inherited diseases for all clinical geneticists." In addition to the erudite entries in the books, the references given with each description represent a magnificent bibliography of clinical genetics. With McKusick's leadership and continued interest in gene mapping, the book also represents an important compendium of the location of genes on specific chromosomes. "The book is a magnificent security blanket for the clinical geneticist and should be in the libraries not only of these specialists, but also of all others who see patients with diseases that have genetic components."

Book Information

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Victor A. McKusick, M.D., is University Professor of Medical Genetics at the Johns Hopkins University School of Medicine. He was formerly William Osler Professor and Chairman of the Department of Medicine at that institution and Physician-in-Chief of the Johns Hopkins Hospital. He is co-founder and co-director of the Short Course in Medical and Experimental Mammalian Genetics, held annually in Bar Harbor, Maine, since 1960.

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