# Author Instructions American Journal Of Medical Genetics

United States. Health Professions Education and Manpower Training Bureau, National Cancer Institute (U.S.). Viral Oncology. Program Analysis and Communication Section

<u>Peer Review in Health Sciences</u> Tom Jefferson, Fiona Godlee, 2003-09-26 This book has established itself as the authoritative text on health sciences peer review. Contributions from the world's leading figures discuss the state of peer review, question its role in the currently changing world of electronic journal publishing, and debate where it should go from here. The second edition has been thoroughly revised and new chapters added on qualitative peer review, training, consumers and innovation.

A Compilation of Journal Instructions to Authors National Cancer Institute (U.S.),1979

Guide for the Care and Use of Laboratory Animals National Research Council, Division on Earth and Life Studies, Institute for Laboratory Animal Research, Committee for the Update of the Guide for the Care and Use of Laboratory Animals, 2011-01-27 A respected resource for decades, the Guide for the Care and Use of Laboratory Animals has been updated by a committee of experts, taking into consideration input from the scientific and laboratory animal communities and the public at large. The Guide incorporates new scientific information on common laboratory animals, including aquatic species, and includes extensive references. It is organized around major components of animal use: Key concepts of animal care and use. The Guide sets the framework for the humane care and use of laboratory animals. Animal care and use program. The Guide discusses the concept of a broad Program of Animal Care and Use, including roles and responsibilities of the Institutional Official, Attending Veterinarian and the Institutional Animal Care and Use Committee. Animal environment, husbandry, and management. A chapter on this topic is now divided into sections on terrestrial and aquatic animals and provides recommendations for housing and environment, husbandry, behavioral and population management, and more. Veterinary care. The Guide discusses veterinary care and the responsibilities of the Attending Veterinarian. It includes recommendations on animal procurement and transportation, preventive medicine (including animal biosecurity), and clinical care and management. The Guide addresses distress and pain recognition and relief, and issues surrounding euthanasia.

Physical plant. The Guide identifies design issues, providing construction guidelines for functional areas; considerations such as drainage, vibration and noise control, and environmental monitoring; and specialized facilities for animal housing and research needs. The Guide for the Care and Use of Laboratory Animals provides a framework for the judgments required in the management of animal facilities. This updated and expanded resource of proven value will be important to scientists and researchers, veterinarians, animal care personnel, facilities managers, institutional administrators, policy makers involved in research issues, and animal welfare advocates.

**Human Genetics and Genomics** Bruce R. Korf, Mira B. Irons, 2012-11-19 This fourth edition of the best-selling textbook, Human Genetics and Genomics, clearly explains the key principles needed by medical and health sciences students, from the basis of molecular genetics, to clinical applications used in the treatment of both rare and common conditions. A newly expanded Part 1, Basic Principles of Human Genetics, focuses on introducing the reader to key concepts such as Mendelian principles, DNA replication and gene expression. Part 2, Genetics and Genomics in Medical Practice, uses case scenarios to help you engage with current genetic practice. Now featuring full-color diagrams, Human Genetics and Genomics has been rigorously updated to reflect today's genetics teaching, and includes updated discussion of genetic risk assessment, "single gene" disorders and therapeutics. Key learning features include: Clinical snapshots to help relate science to practice 'Hot topics' boxes that focus on the latest developments in testing, assessment and treatment 'Ethical issues' boxes to prompt further thought and discussion on the implications of genetic developments 'Sources of information' boxes to assist with the practicalities of clinical research and information provision Self-assessment review questions in each chapter Accompanied by the Wiley E-Text digital edition (included in the price of the book), Human Genetics and Genomics is also fully supported by a suite of online resources at www.korfgenetics.com, including: Factsheets on 100 genetic disorders, ideal for study and exam preparation Interactive Multiple Choice Questions (MCQs) with feedback on all answers Links to online resources for further study Figures from the book available as PowerPoint slides, ideal for teaching purposes The perfect companion to the genetics component of both problem-based learning and integrated medical courses, Human Genetics and Genomics presents the ideal balance between the bio-molecular basis of genetics and clinical cases, and provides an invaluable overview for anyone wishing to engage with this fast-moving discipline.

American Journal of Medical Genetics, 1987

**Psychoeducational Assessment of Students who are Visually Impaired Or Blind** Sharon Bradley-Johnson, Sandra Kanouse Morgan, 2008

**A Compilation of Journal Instructions to Authors** National Cancer Institute (U.S.) Viral Oncology Area. Information Unit, 1971

Author's Handbook of Styles for Life Science Journals Michel Atlas, 1995-11-08 Let the Author's Handbook of Styles

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Suggestions to Medical Authors and A.M.A. Style Book American Medical Association, 1919

The AST Handbook of Transplant Infections Deepali Kumar, Atul Humar, 2011-04-18 Whether you need to manage a post-transplant infection or reduce the possibility of infection, you will find effective guidance in this handbook. The work of the American Society of Transplantation Infectious Diseases Community of Practice, this reference exclusively uses tables and flowcharts to speed up decision making. This distinguished group of investigators and teachers provide point of care information on optimum management of infection in adult and pediatric organ and stem cell transplant patients. The unique tables and flowcharts are devised by the authors, backed up with extensive references, making the book a fully researched yet easy to use guide. The fast growing specialty of transplantation will be well served by this book as increasing numbers of successful procedures mean transplant teams have to be ever more alert to the possibility of and need for action in the event of ensuing infection.

<u>A Compilation of Journal Instructions to Authors</u> United States. Health Professions Education and Manpower Training Bureau, National Cancer Institute (U.S.). Viral Oncology. Program Analysis and Communication Section, 1971

Handbook of Physical Measurements Karen W. Gripp, Anne M. Slavotinek, Judith G. Hall, Judith E. Allanson, 2013-09 This handbook provides a practical collection of reference data on a variety of physical measurements for use in the evaluation of children and adults with dysmorphic features and/or structural anomalies. Each chapter concentrates on a specific body area.

Neurocutaneous Disorders E. Steve Roach, Van S. Miller, 2004-01-08 'The reader is most definitely in for a treat provides

the essential clinical and genetic data which points the way to the future. From the foreword by Roger N. RosenbergThe neurocutaneous disorders comprise a group of neurological disorders featuring skin lesions and often eye lesions, central and peripheral nervous system tumors, brain malformations, mental retardation, and psychiatric syndromes or seizures. This book provides an authoritative, illustrated review of the recognition, investigation, treatment and genetics of these disorders. It will be essential reading for neurologists as well as dermatologists, geneticists and pediatricians.

**Developmental Neuropsychiatry** James C. Harris, 1998 James Harris's two volume work on developmental neuropsychiatry sets the agenda for this emerging clinical specialty. Written by an individual with the developmental expertise of a pediatrician, the behavioral sophistication of an adult and child psychiatrist, and a deep appreciation of neuroscience, these two books offer an integrated yet comprehensive approach to developmental neuropsychiatry. Grounded in neuroscience but enriched by clinical realities, Volume II provides a comprehensive review of the developmental neuropsychiatric disorders. Throughout the text current DSM-IV diagnostic criteria are provided. Part I outlines the diagnostic process and the genetic history, provides details on the conduct of neuropsychological testing, and offers a detailed review of brain imaging techniques, moving from CT and MRI scanning to the most recent developments in functional MRI and PET scanning. Part II discusses mental retardation, cerebral palsy, the learning disorders, the pervasive developmental disorders, and traumatic brain injury. Part III describes behavioral phenotypes in cytogenetic and other genetic disorders, genetic metabolic disorders, and disorders that result from gestational substance abuse. Part IV is devoted to developmental psychopathology and includes Attention Deficit/Hyperactivity disorder, schizophrenia, Tourette's disorder, sleep disorders, and the syndromes of aggression and self-injury primarily occurring in mentally retarded persons. Part V covers treatment and includes detailed descriptions of psychotherapy, behavior therapy, pharmacological interventions, genetic counseling, and gene therapy. Finally, Part VI deals with legal and ethical issues as they pertain to developmentally disabled persons.

The Carriers Anne Skomorowsky,2022-05-03 A tiny mutation on the X chromosome can shape a family's history. Passed down from a "carrier" parent to a child, fragile X syndrome is the most common inherited cause of intellectual disability and autism. Beyond that—and a rarity among genetic disorders—some fragile X carriers not only transmit the mutation but also experience related conditions themselves. In such cases, carriers can have tremors, infertility, and psychiatric disorders that complicate raising children with fragile X syndrome—and all too often, they suffer in silence. The Carriers investigates this common but still little-known genetic condition and its life-altering consequences. Anne Skomorowsky reveals how this disorder afflicts families across generations, telling the stories of the mothers and grandparents of fragile X patients and considering how genes interact with family dynamics. She interweaves the personal narratives and family histories of the people affected by fragile X disorders with clear and accessible explanations of the science behind them. Skomorowsky

unpacks the latest research on the fragile X mutation and explores the history of its discovery. She highlights the roles of women as carriers, caregivers, and researchers who have made astonishing scientific breakthroughs over the last three decades. The Carriers is an essential book for fragile X families, including those just learning they are carriers, and for all readers interested in the complexities of heredity, the ethical dilemmas of genetic medicine, and the relationship between genes and personality.

<u>Renal Vascular Disease</u> Andrew C. Novick, J. Scoble, G. Hamilton, 1996 This is a reference text covering all aspects of renal disease, including: pathology, clinical features, imaging, hypertension, atherosclerotic disease, medical and surgical treatment.

The Obstetric Hematology Manual Sue Pavord, Beverley Hunt, 2018-02-08 Understand the rapidly growing complexities of obstetric hematology and high-risk pregnancy management, with experts in the field. Now in its second edition, this comprehensive and essential guide focuses on providing the best support for patients and clinical staff, to prevent serious complications in pregnancy and the post-partum period for both mother and baby. Wide-ranging and detailed, the guide offers discussions on basic principles of best care, through to tackling lesser-known hematological conditions, such as cytopenias and hemoglobinopathies. Updated with color illustrations, cutting-edge research, accurate blood film reproductions, and practical case studies, the revised edition places invaluable advice into everyday context. This unique resource is essential reading for trainees and practitioners in obstetrics, anesthesia, and hematology, as well as midwives, nurses, and laboratory staff. Clarifying difficult procedures for disease prevention, the guide ensures safety when the stakes are high. Reflecting current evidence-based guidelines, the updated volume is key to improving pregnancy outcomes worldwide.

Life Histories of Genetic Disease Andrew J. Hogan, 2016-10-30 A richly detailed history that "uncovers the challenges and limitations of our increasing reliance on genetic data in medical decision making" (Shobita Parthasarathy, author of Building Genetic Medicine). Medical geneticists began mapping the chromosomal infrastructure piece by piece in the 1970s by focusing on what was known about individual genetic disorders. Five decades later, their infrastructure had become an edifice for prevention, allowing expectant parents to test prenatally for hundreds of disease-specific mutations using powerful genetic testing platforms. In this book, Andrew J. Hogan explores how various diseases were "made genetic" after 1960, with the long-term aim of treating and curing them using gene therapy. In the process, he explains, these disorders were located in the human genome and became targets for prenatal prevention, while the ongoing promise of gene therapy remained on the distant horizon. In narrating the history of research that contributed to diagnostic genetic medicine, Hogan describes the expanding scope of prenatal diagnosis and prevention. He draws on case studies of Prader-Willi, fragile X, DiGeorge, and velo-cardio-facial syndromes to illustrate that almost all testing in medical genetics is inseparable from the larger—and

increasingly "big data"-oriented—aims of biomedical research. Hogan also reveals how contemporary genetic testing infrastructure reflects an intense collaboration among cytogeneticists, molecular biologists, and doctors specializing in human malformation. Hogan critiques the modern ideology of genetic prevention, which suggests all pregnancies are at risk for genetic disease and should be subject to extensive genomic screening. He examines the dilemmas and ethics of the use of prenatal diagnostic information in an era when medical geneticists and biotechnology companies offer whole genome prenatal screening—essentially searching for any disease-causing mutation. Hogan's analysis is animated by ongoing scientific and scholarly debates about the extent to which the preventive focus in contemporary medical genetics resembles the aims of earlier eugenicists. Written for historians, sociologists, and anthropologists of science and medicine, as well as bioethics scholars, physicians, geneticists, and families affected by genetic conditions, Life Histories of Genetic Disease is a profound exploration of the scientific culture surrounding malformation and mutation.

<u>Psychiatric Mental Health Nursing</u> Patricia G. O'Brien, Winifred Z. Kennedy, Karen A. Ballard, 2012-02-15 A comprehensive, easy-to-read introductory text for nursing students. The book is organized into three sections: Introduction to Psychiatric-Mental Health Nursing, Mental Health Disorders, and Nursing Management of Special Populations. This unique text is the most comprehensive psychiatric mental health resource available.

Issues in Genetic Medicine: 2011 Edition ,2012-01-09 Issues in Genetic Medicine / 2011 Edition is a ScholarlyEditions™ eBook that delivers timely, authoritative, and comprehensive information about Genetic Medicine. The editors have built Issues in Genetic Medicine: 2011 Edition on the vast information databases of ScholarlyNews.™ You can expect the information about Genetic Medicine in this eBook to be deeper than what you can access anywhere else, as well as consistently reliable, authoritative, informed, and relevant. The content of Issues in Genetic Medicine: 2011 Edition has been produced by the world's leading scientists, engineers, analysts, research institutions, and companies. All of the content is from peer-reviewed sources, and all of it is written, assembled, and edited by the editors at ScholarlyEditions™ and available exclusively from us. You now have a source you can cite with authority, confidence, and credibility. More information is available at http://www.ScholarlyEditions.com/.

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