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Announcements¹

EMPLOYMENT AND FELLOWSHIP OPPORTUNITIES

Genetic Epidemiologist/Director. — Strang Cancer Prevention Center seeks genetic epidemiologist/director for a large, high-risk breast-cancer registry with over 15,000 families. Strang is well-funded and affiliated with Cornell Medical College. The position would carry an appointment with that institution. Candidate must have an M.D. or Ph.D., experience, and ability to work closely with clinicians and basic scientists. Excellent stipend, all benefits. The position will go to the candidate with the strongest research agenda. Write to Dr. D. G. Miller, Strang Cancer Prevention Center, 428 East 72d Street, New York, NY 10021.

Eye Research Institute of Canada. — The Eye Research Institute of Canada is searching for a scientist interested in the molecular biology of vision. A concern for the development of technology for molecular diagnosis of human disease would be considered an asset. The appointee will establish an independent research program and will be eligible for appointment in the graduate faculty at the University of Toronto. Collaboration with other scientists studying molecular biology of eye disease is anticipated. The institute provides

1. This service is offered free of charge to members of The American Society of Human Genetics. To place an announcement, write to The American Journal of Human Genetics, Department of Pediatrics, U-587, University of California, San Francisco, CA 94143-0748. Submission must be made at least *four months* before month of issue in which publication is requested and must be typed double-spaced on a separate sheet of paper. Maximum length is *100 words*, not including address for correspondence.

salary support, excellent research facilities, and start-up funds. Basic qualifications will include Ph.D., M.D., or D.V.M. Interested candidates should send, to the address below, a letter outlining their current and future research interests, a curriculum vitae, and the names of three individuals who can provide a reference. The deadline for receipt of application is 1 November 1992. Send application to Dr. B. L. Gallie, Eye Research Institute of Canada, 399 Bathurst Street, Toronto, Ontario M2T 2S8 Canada.

Molecular biologist. — The Cytogenetics Laboratory, part of the Medical Genetics Birth Defects Center at Cedars-Sinai Medical Center in Los Angeles (a UCLA-affiliated teaching hospital), has a position at the post-doctoral level for an accomplished molecular biologist interested in training in clinical cytogenetics. The position will involve initiating molecular studies of clinical cases, as well as participation in ongoing projects involving pediatric leukemia and solid tumors, while learning cytogenetic techniques as part of an ABMG-accredited training program. For more information, please contact Rhona Schreck, Ph.D., Director, Cytogenetics Laboratory, Cedars-Sinai Medical Center, Steven Spielberg Building 3, 8700 Beverly Boulevard, Los Angeles, CA 90048; phone (310) 855-6451; FAX (310) 967-0112.

Search Reopened for Assistant Director of Cytogenetics. — The Medical Genetics Birth Defects Center at Cedars-Sinai Medical Center in Los Angeles is cur-

rently recruiting an assistant director for the cytogenetics laboratory. The laboratory, under the direction of Rhona Schreck, Ph.D., provides prenatal diagnosis, assessment of congenital abnormalities and reproductive failure, and evaluation of malignancies. As the laboratory is processing over 2,000 patient samples per year, we are recruiting a board-certified (or eligible) cytogeneticist to serve as assistant director and to share the clinical service with the director. The candidate should meet the requirements of, and will be eligible for, a faculty appointment at UCLA. Independent research in any area of cytogenetics is strongly encouraged. If interested, please contact Rhona Schreck, Ph.D., Director, Cytogenetics Laboratory, Division of Medical Genetics, Cedars-Sinai Medical Center, Steven Spielberg Building 3, 8700 Beverly Boulevard, Los Angeles, CA 90048; phone (310) 855-6451; FAX (310) 967-0112.

Postdoctoral Fellowships.— Available in the general areas of physical mapping of human chromosome 21, research into the etiology of Down syndrome, and research into the etiology of human neurodegenerative diseases such as Alzheimer disease and ALS. Methods include molecular biology, somatic cell genetics, and transgenic animal studies. Send curriculum vitae, statement of research interests, and names of three references to Dr. David Patterson, President, Eleanor Roosevelt Institute, 1899 Gaylord Street, Denver, CO 80206. Equal Opportunity/Affirmative Action Employer; women and minorities are encouraged to apply.

BOARD REVIEW COURSE

Board Review Course.— A 2½-day course in medical genetics, designed as a review for the 1993 American Board of Medical Genetics examination, will be sponsored by Baylor College of Medicine and held in Houston on May 14–16, 1993. The course will follow the format used in 1990 and will include local and guest faculty, a graded pretest, and workshops on sample questions. For information, write to Genetics Board Review Course, Office of Continuing Education, Room S104, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030-3498; phone (713) 798-6020; FAX (713) 798-6600.

GRANTS AVAILABLE

Grants for Travel and Research at the Rockefeller Archive Center.— The Rockefeller Archive Center, a division of The Rockefeller University, invites applications for its program of grants for travel and research at the Rockefeller Archive Center for 1993. The competitive program makes grants of up to \$1,500 to researchers in any discipline, usually graduate students or postdoctoral scholars, who are engaged in research that requires use of the collections at the center, which include the records of the Rockefeller family, the Rockefeller Foundation, The Rockefeller University, and other philanthropic organizations and associated individuals. The deadline for applications is December 31, 1992; grant recipients will be announced in March. Inquiries about the program and requests for applications should be addressed to Darwin H. Stapleton, Director, Rockefeller Archive Center, 15 Dayton Avenue, North Tarrytown, NY 10591-1598.

Information for Contributors

The American Journal of Human Genetics is a record of research and review relating to heredity in humans; to the applications of genetic principles in medicine, psychology, anthropology, and social services; and to areas of molecular and cell biology relevant to human genetics. It appears 12 times a year. The *Journal* is owned and controlled by The American Society of Human Genetics and is edited, in conjunction with the publisher, by a staff appointed by the board of directors of the society.

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It is assumed that authors whose research is published in the *Journal* will make cell lines, antibodies, cloned DNA sequences, and similar materials available to other investigators so as to permit the reported research to be reproduced.

The Human Genetics Education section of the *Journal* serves as an international medium for the exchange of ideas in human genetics education. The *Journal* welcomes the submission of manuscripts concerned with any aspect of education in human genetics at the undergraduate and postgraduate levels, and for service providers of genetic health care. These include articles on research in human genetics education, innovations for its improvement, development and review of educational resources, conceptual frameworks for topic and course development, educationally useful historical perspectives, letters to the editor, and activities of The American Society of Human Genetics relative to education. All manuscripts and correspondence regarding the Human Genetics Education section should be addressed to the Editor, Human Genetics Education Section, Kenneth L. Garver, MD., Ph.D., Department of Medical Genetics, Western Pennsylvania Hospital, 4800 Friendship Avenue, Pittsburgh, PA 15224.

Manuscript Submission

Original articles, review articles of approximately 3,000 words or less, and minireviews of 1,500 words or less on timely subjects concerning all aspects of human genetics and related areas will be considered for publication. In addition, brief commentaries on previously published material, editorials, and book reviews will also be included in the *Journal*. We have adopted a policy of not publishing descriptions of single mutations unless the mutation is of highly unusual significance.

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All manuscripts must be written in English, and all manuscript components (including references, tables, and figure legends) typed double spaced, preferably on heavyweight bond. The page margins should be 1½ inches on top, bottom, and left- and right-hand sides to allow for corrections and manuscript editor's and typesetter's instructions and queries. Word-processing output is acceptable as long as it is letter quality. Handwritten items (e.g., Greek letters) should be identified in the margin. Each manuscript component should begin on a separate page, but pages should be numbered consecutively, beginning with the title page and continuing through the summary (abstract), text, references, tables, and figure legends.

Title page. — The title page should contain a concise title, the names and current affiliations of all authors, and the name and complete mailing address of the contact author. A running head of no more than 40 characters (including spaces) should be identified on the title page.

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Text. — The text should be as concise as possible and include the following sections: Introduction, Material and Methods, Results, Discussion, and Acknowledgments. Long articles, case studies, and editorials may require subheadings within some sections or different formats to clarify their content.

The Introduction should summarize the rationale for the study without reviewing the subject extensively. Clearly describe (using past tense) your selection of observational or experimental subjects and identify the methods, apparatus, and procedures in detail. Provide references for established methods, with brief descriptions for published but not well-known methods; describe new or modified methods, giving reasons for their use and evaluating their limitations.

Present your results in logical sequence in the text, tables, and illustrations. Do not repeat in the text all the data in the tables and illustrations. Detailed statistical analyses may be presented in the form of an appendix.

Terminology and abbreviations. — Authors should make an effort to present articles in a precise, comprehensible manner. In general, all material should conform to the style of the *CBE Style Manual: A Guide for Authors, Editors, and Publishers in the Biological Sciences* (5th ed., Council of Biology Editors, Inc., Bethesda, MD).

Nomenclature, abbreviations, and symbols. — Authors should use genetic notation and symbols for human genes which are approved by Human Gene Mapping (HGM). See Shows et al.'s "Guidelines for Human Gene Nomenclature" (Cytogenet. Cell Genet. 46:11–28, 1987), McAlpine et al.'s "The 1988 Catalog of Mapped Genes and Report of the Nomenclature Committee" (Cytogenet. Cell Genet. 49:4–38, 1988), and Kidd et al.'s "Report of the Committee on Human Gene Mapping by Recombinant DNA Techniques" (Cytogenet. Cell Genet. 49:132–218, 1988). Symbols for human genes not included in the above may be obtained from P. J. McAlpine (cochair, HGM Nomenclature Committee), Department of Human Genetics, University of Manitoba, T250-770 Bannatyne Avenue, Winnipeg, Manitoba R3E 0W3, Canada; phone (204)788-6393; FAX (204)786-8712; E-mail GENMAP@UOFMCC.

Mathematical equations must be carefully typed with spacing in final format. Measurements of length, height, weight, and volume should appear in metric units or other internationally accepted units.

Information for Contributors

Authors should use the following standard abbreviations (for others, generally follow the *CBE Style Manual*):

AMP = adenosine monophosphate	NAD (NADP) = nicotinamide adenine dinucleotide phosphate
ATP = adenosine triphosphate	PBS = phosphate-buffered saline
BSA = bovine serum albumin	PAGE = polyacrylamide-gel electrophoresis
BrdUrd = 5-bromodeoxyuridine	PCR = polymerase chain reaction
df = degree(s) of freedom	PIC = polymorphic information content
DEAE = diethylaminoethyl	RFLP = restriction-fragment-length polymorphism
DTT = dithiothreitol	rRNA = ribosomal RNA
DZ = dizygotic	SSC = saline sodium citrate
EDTA = ethylenediaminetetraacetate	SDS = sodium dodecyl sulfate
FCS = fetal calf serum	SD = standard deviation
FSH = follicle-stimulating hormone	SEM = standard error of the mean
mRNA = messenger RNA	TSH = thyroid-stimulating hormone
mtDNA = mitochondrial DNA	tRNA = transfer RNA
MZ = monozygotic	UV = ultraviolet
NAD (NADH) = nicotinamide adenine dinucleotide	VNTR = variable number of tandem repeats

Acknowledgments.—Acknowledge those who have made substantive contributions to the study, as well as the institution(s) or grant sponsor(s) that supported the research. Authors are advised to obtain permission from those acknowledged by name, since the readers may infer endorsement of the data and conclusion. Grants and/or institutions supporting research presented in the article should be mentioned in this section.

References.—Reference citations in the text should be in parentheses and include author name(s) and year of publication. Text citations of two or more works at the same time should be given in chronological order. When citing a paper written by three or more authors, write the name of the first author plus “et al.” (however, the names of the first *seven* authors—plus, if there are more than seven, “et al”—must be given in the Reference section). The References section appears after the acknowledgments and should be arranged alphabetically by author name(s) and then chronologically. To use “personal communication” as a reference citation, you must include written permission from the scientist(s) named.

Following are examples of the proper reference style for various sources:

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Meeting Paper:	Novacek MJ, Wyss AR (1985) Morphology, molecules, and eutherian phylogeny: the search for congruence. Paper presented at the Fourth International Theriological Congress, Edmonton, Alberta, August 13–20
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Tables. — Type each table double-spaced on separate sheets of paper. Exceptionally large tables and pedigree diagrams should be constructed in sections whenever possible to avoid the need for folded inserts. Number tables consecutively and give each a brief title. Give each column a short heading. Footnotes should be used to explain all nonstandard abbreviations that are used in each table and should be denoted by letter. Further guidelines for table format may be found in the *CBE Style Manual* and *The Chicago Manual of Style*.

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