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# GENETIC EPIDEMIOLOGY OF CORONARY HEART DISEASE PAST, PRESENT, AND FUTURE

EDITORS: D.C. Rao
Robert C. Elston
Lewis H. Kuller
Manning Feinleib
Christine Carter
Richard Havlik

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## GENETIC EPIDEMIOLOGY OF CORONARY HEART DISEASE

## Past, Present, and Future

Proceedings of a Workshop held in St. Louis, Missouri August 10–12, 1983

#### **Editors**

#### D.C. Rao

Washington University School of Medicine St. Louis, Missouri

#### Robert C. Elston

Louisiana State University Medical Center New Orleans, Louisiana

#### Lewis H. Kuller

University of Pittsburgh Pittsburgh, Pennsylvania

#### Manning Feinleib

National Center of Health Statistics Hyattsville, Maryland

#### **Christine Carter**

National Heart, Lung and Blood Institute National Institutes of Health Bethesda, Maryland

#### Richard Havlik

National Heart, Lung and Blood Institute National Institutes of Health Bethesda, Maryland

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### **Contributors and Participants**

Ronald T. Acton, Departments of Microbiology, Medicine, and Epidemiology, University of Alabama School of Medicine, Birmingham, AL 35294 [479]

John Albers, University of Washington, Mail Stop ZA-36, Seattle, WA 98104

S. Anderson, Boston University Medical Center, Boston, MA 02118 [407]

K. Owen Ash, Department of Pathology, University of Utah, Salt Lake City, UT 84132 [419]

Bruce O. Barger, Departments of Microbiology, Medicine, Epidemiology, University of Alabama School of Medicine, Birmingham, AL 35294 [479]

**Terri H. Beaty,** Department of Epidemiology, John Hopkins University School of Hygiene and Public Health, Baltimore, MD 21205 [501]

Kåre Berg, Institute of Medical Genetics, University of Oslo, Oslo 3, Norway [85,549]

D. Timothy Bishop, Department of Human Genetics, University of Utah, Salt Lake City, UT 84132 [271]

Michael Boehnke, Department of Biostatistics, University of Michigan, Ann Arbor, MI 48109 [173] **George E. Bonney,** Department of Biometry, Louisiana State University Medical Center, New Orleans, LA 70112 [349]

Ingrid Borecki, Division of Biostatistics, Washington University School of Medicine, St. Louis, MO 63110

R.H. Bradford, Oklahoma Medical Research Foundation, Oklahoma City, OK 73104

John D. Brunzell, Department of Medicine, Division of Metabolism, University of Washington, Seattle, WA 98195 [403]

Clareann H. Bunker, Department of Epidemiology, University of Pittsburgh, Pittsburgh, PA 15213

Pamela J. Byard, Department of Anthropology, Case Western Reserve University, Cleveland, OH 44106

Chris Cannings, Department of Probability and Statistics, University of Sheffield, Sheffield S3 7RH, England [271]

Rita M. Cantor, Division of Medical Genetics, Harbor-UCLA Medical Center, Torrance, CA 90509

Christine Carter, Epidemiology Branch, Clinical and Genetic Epidemiology, National Heart, Lung and Blood Institute, National Institutes of Health, Bethesda, MD 20205 [79]

The number in brackets is the opening page number of the contributor's article.

- **Gregory Cary**, Department of Psychiatry, The Jewish Hospital of St. Louis, St. Louis, MO 63108
- W. Castelli, Framingham Heart Study, National Heart, Lung and Blood Institute, Framingham, MA 01701 [407]
- Ranajit Chakraborty, Center for Demographic and Population Genetics, University of Texas Health Science Center at Houston, Houston, TX 77030 [155,213,324]
- **Aravinda Chakravarti**, Department of Biostatistics, University of Pittsburgh, Pittsburgh, PA 15261 [293]
- Joseph Christian, Department of Medical Genetics, Indiana University, Indianapolis, IN 46223
- C.S. Chung, Department of Public Health Sciences, University of Hawaii, Honolulu, HI 96822 [135]
- Susanna F.M. Clarkson, Division of Biostatistics, Washington University School of Medicine, St. Louis, MO 63110
- C. Robert Cloniger, Department of Psychiatry, The Jewish Hospital of St. Louis, St. Louis, MO 63108
- Bernice H. Cohen, Department of Epidemiology, Johns Hopkins University School of Hygiene and Public Health, Baltimore, MD 21205 [501]
- Robert B. Copeland, Georgia Heart Clinic, LaGrange, GA 30240 [479]
- Nancy J. Cox, Department of Psychiatry, The Jewish Hospital of St. Louis, St. Louis, MO 63108
- Gary Cutter, Department of Epidemiology, University of Alabama School of Medicine, Birmingham, AL 35294 [479]
- Mary M. Dadone, Department of Internal Medicine, Cardiology Division, University of Utah, Salt Lake City, UT 84132 [419]

- Thomas F. Deuel, Department of Internal Medicine, The Jewish Hospital of St. Louis, St. Louis, MO 63108
- **Eric J. Devor,** Department of Psychiatry, The Jewish Hospital of St. Louis. St. Louis, MO 63108
- Elaine D. Eaker, Epidemiology and Biometry Program, National Heart, Lung and Blood Institute, National Institutes of Health, Bethesda, MD 20205 [481]
- Robert C. Elston, Department of Biometry, Louisiana State University Medical Center, New Orleans, LA 70112 [349,539]
- Richard R. Fabsitz, Epidemiology Branch, National Heart, Lung and Blood Institute, National Institutes of Health, Bethesda, MD 20205
- Manning Feinleib, National Center for Health Statistics, Hyattsville, MD 20782 [1]
- **B. Finizola**, Boston University Medical Center, Boston, MA 02118 [407]
- Wilford Forbush, Health and Human Services, Washington, DC 20201
- Charles J. Glueck, General Clinical Research Center, University of Cincinnati, Cincinnati, OH 45267 [149,193,525]
- Rodney C.P. Go, Department of Epidemiology, University of Alabama School of Medicine, Birmingham, AL 35294 [479]
- **Lynn R. Goldin**, Biological Psychiatry Branch, National Institute of Mental Health, Bethesda, MD 20205
- Irving I. Gottesman, Department of Psychiatry, Washington University School of Medicine, St. Louis, MO 63110
- Samuel B. Guze, Vice Chancellor for Medical Affairs, Washington University School of Medicine, St. Louis, MO 63110

Craig L. Hanis, Center for Demographic and Population Genetics, University of Texas Health Science Center at Houston, Houston, TX 77030 [213]

Emily L. Harris, Department of Epidemiology, University of Pittsburgh, Pittsburgh, PA 15261 [499]

Sandra J. Hasstedt, Department of Human Genetics, University of Utah, Salt Lake City, UT 84132 [271]

Richard Havlik, Epidemiology Branch, Clinical and General Epidemiology, National Heart, Lung and Blood Institute, National Institutes of Health, Bethesda, MD 20205

Hans Ole Hein, University Hospital, DK-2200 Copenhagen N, Denmark

Ivar Heuch, Department of Biostatistics, University of North Carolina, Chapel Hill, NC 27514

Millicent W. Higgins, Departments of Epidemiology and Internal Medicine, School of Public Health, University of Michigan, Ann Arbor, MI 48109 [43,51]

Paul N. Hopkins, Department of Internal Medicine, Cardiology Division, University of Utah, Salt Lake City, UT 84132 [419]

Steven C. Hunt, Department of Internal Medicine, Cardiology Division, University of Utah, Salt Lake City, UT 84132 [419]

Lynn B. Jorde, Department of Pediatrics, University of Utah, Salt Lake City, UT 84132 [419]

Candace M. Kammerer, Genetics Department, Southwest Foundation for Research and Education, San Antonio, TX 78284 [297]

Samuel Karlin, Department of Mathematics, Stanford University, Stanford, CA 94305 [317]

Kathe Kelly, Lipid Research Clinics Program, National Heart, Lung and Blood Institute, National Institutes of Health, Bethesda, MD 20205 [149] Muin J. Khoury, Department of Epidemiology, Johns Hopkins University School of Hygiene and Public Health, Baltimore, MD 21205 [501]

Philip Khoury, General Clinical Research Center, University of Cincinnati, Cincinnati, OH 45267 [149]

H. Kuida, University of Utah, Salt Lake City, UT 84132 [419]

**Lewis H. Kuller**, Department of Epidemiology, University of Pittsburgh, PA 15261 [533]

Jean-Marc Lalouel, Department of Biological Anthropology, University of Paris, Paris 75005, France [217,267,549]

Kenneth Lange, Department of Biomathematics, University of California, Los Angeles, CA 90024 [173]

Peter M. Laskarzewski, General Clinical Research Center, University of Cincinnati, Cincinnati, OH 45267 [149]

**G.M. Lathrop**, Department of Biological Anthropology, University of Paris, Paris 75005, France [267]

Namsoo Lee, Lipid Research Center, Washington University School of Medicine, St. Louis, MO 63110

**D. Levy**, Boston University Medical Center, Boston, MA 02118 [407]

C.C. Li, Graduate School of Public Health, University of Pittsburgh, Pittsburgh, PA 15261 [159]

Ira M. Longini, Department of Biostatistics, University of Michigan, Ann Arbor, MI 48109 [43]

Jean W. MacCluer, Genetics Department, Southwest Foundation for Research and Education, San Antonio, TX 78284 [297]

Martha M. McCrate, Division of Biostatistics, Washington University School of Medicine, St. Louis, MO 63110

- Matthew K. McGue, Division of Biostatistics, Washington University School of Medicine, St. Louis, MO 63110 [23]
- Lynne McMahan, Department of Pathology, Louisiana State University, New Orleans, LA 70112
- F. John Meany, Department of Medical Genetics, Indiana University, Indianapolis, IN 46223
- Margot J. Mellies, General Clinical Research Center, University of Cincinnati, Cincinnati, OH 45267 [149]
- Beverly G. Mellon, Department of Human Genetics, Medical College of Virginia/Virginia Commonwealth University, Richmond, VA 23298
- Harold A. Menkes, Department of Environmental Health Sciences, Johns Hopkins University School of Hygiene and Public Health, Baltimore, MD 21205 [501]
- J. Philip Miller, Division of Biostatistics, Washington University School of Medicine, St. Louis, MO 63110
- Judy Z. Miller, Hypertension Research Center, Indiana University, Indianapolis, IN 46233 [499]
- Patricia P. Moll, Department of Human Genetics, University of Michigan, Ann Arbor, MI 48109 [37]
- John A. Morrison, General Clinical Research Center, University of Cincinnati, Cincinnati, OH 45267 [149]
- Newton E. Morton, Population Genetics Laboratory, University of Hawaii, Honolulu, HI 96822 [117,245,270]
- **Arno G. Motulsky**, Departments of Medicine and Genetics, University of Washington, Seattle, WA 98195 [403,541]
- R. Myers, Boston University Medical Center, Boston, MA 02118 [407]

- Kadambari K. Namboodiri, Department of Biostatistics, University of North Carolina, Chapel Hill, NC 27514 [65,155]
- Walter E. Nance, Department of Human Genetics, Medical College of Virginia/Virginia Commonwealth University, Richmond, VA 23298 [325]
- Carol A. Newill, Department of Epidemiology, Johns Hopkins University School of Hygiene and Public Health, Baltimore, MD 21205 [501]
- Thomas W. Nielsen, Division of Heart and Vascular Diseases, Cardiac Functions Branch, National Institutes of Health. Bethesda, MD 20205
- James J. Nora, Departments of Genetics, Preventive Medicine, and Pediatrics, University of Colorado School of Medicine and Rose Medical Center, Denver, CO 80262 [47]
- Gary Olson, Division of Cardiovascular Disease, Dalton Cardiology, Dalton, GA 30720 [479]
- **Dennis H. O'Rourke,** Department of Anthropology, University of Utah, Salt Lake City, UT 84112
- S. Padgett, Epidemiology Branch, National Heart, Lung and Blood Institute, National Institutes of Health, Bethesda, MD 20205 [407]
- William A. Peck, Department of Internal Medicine, The Jewish Hospital of St. Louis, St. Louis, MO 63108
- H. Mitchell Perry, Jr., Hypertension Division, Washington University School of Medicine, St. Louis, MO 63110
- J. Plehn, Boston University Medical Center, Boston, MA 02118 [407]
- Michael A. Province, Division of Biostatistics, Washington University School of Medicine, St. Louis, MO 63110 [193]

- Leslie J. Raffel, Division of Medical Genetics, Departments of Medicine and Pediatrics, Harbor-UCLA Medical Center, Torrance, CA 90509 [445]
- D.C. Rao, Division of Biostatistics, Washington University School of Medicine, St. Louis, MO 63110 [149,193]
- **Theodore Reich**, Department of Psychiatry, The Jewish Hospital of St. Louis, St. Louis, MO 63108
- George G. Rhoads, Epidemiology Branch, National Institute of Child Health and Human Development, National Institutes of Health, Bethesda, MD 20205 [11]
- John P. Rice, Department of Psychiatry, The Jewish Hospital of St. Louis, St. Louis, MO 63108
- Basil Rifkind, National Heart, Lung and Blood Institute, National Institutes of Health, Bethesda, MD 20205
- **David L. Rimoin**, Division of Medical Genetics, Departments of Medicine and Pediatrics, Harbor-UCLA Medical Center, Torrance, CA 90509 [445]
- Lee N. Robins, Department of Psychiatry, Washington University School of Medicine, St. Louis, MO 63110
- Jeff M. Roseman, Department of Epidemiology, University of Alabama School of Medicine, Birmingham, AL 35294 [479]
- **Peter A. Rosenbaum**, Departments of Biometry and Medicine, Louisiana State University Medical Center, New Orleans, LA 70112
- Jerome I. Rotter, Division of Medical Genetics, Departments of Medicine and Pediatrics, Harbor-UCLA Medical Center, Torrance, CA 90509 [445]
- Daniel D. Savage, Framingham Heart Study, National Heart, Lung and Blood Institute, Framingham, MA 01701 [407]

- Kenneth Schechtman, Division of Biostatistics, Washington University School of Medicine, St. Louis, MO 63110
- Gustav Schonfeld, Departments of Preventive Medicine and Medicine, Washington University School of Medicine, St. Louis, MO 63110 [375]
- Sue Seyeanhan, Department of Human Biology, John Curtin School of Medical Research, Canberra 2601, Australia
- Edwin K. Silverman, Washington University School of Medicine, St. Louis. MO 63110
- Mark H. Skolnick, Department of Human Genetics, University of Utah, Salt Lake City, UT 84132 [271]
- Jean B. Smith, Department of Pathology, University of Utah, Salt Lake City, UT 84132 [419]
- **George Sopko**, Department of Internal Medicine, St. Louis University School of Medicine, St. Louis, MO 63103
- Edward L. Spitznagel, Department of Mathematics and Division of Biostatistics, Washington University, St. Louis, MO 63130
- Michael P. Stern, Department of Medicine, University of Texas Health Science Center at San Antonio, San Antonio, TX 78284 [93]
- **Brian K. Suarez**, Departments of Psychiatry and Genetics, Washington University School of Medicine, St. Louis, MO 63110 [105]
- **John R. Taylor,** Department of Psychiatry, Washington University School of Medicine, St. Louis, MO 63110
- Constance M. Vadheim, Division of Medical Genetics, Departments of Medicine and Pediatrics, Harbor-UCLA Medical Center, Torrance, CA 90509 [445]

#### xvi / Contributors and Participants

Paul Van Eerdewegh, Department of Psychiatry, The Jewish Hospital of St. Louis, St. Louis, MO 63108

Diane K. Wagener, Western Psychiatric Institute and Clinic, University of Pittsburgh, Pittsburgh, PA 15261 [443]

Carolyn Walden, University of Washington, Mail Stop Gl-22, Seattle, WA 98195

Robert B. Wallace, Department of Preventive Medicine and Environmental Health, University of Iowa, Iowa City, IA 52242 [125] Reimut Wette, Division of Biostatistics, Washington University School of Medicine, St. Louis, MO 63110 [193]

Paul T. Williams, Department of Mathematics, Stanford University, Stanford, CA 94305 [317]

Roger R. Williams, Department of Internal Medicine, Cardiology Division, University of Utah, Salt Lake City, UT 84132 [89,419]

Wick R. Williams, Department of Genetics, Division of Human Genetics, M.D. Anderson Hospital and Tumor Institute, Houston, TX 77030 [121]













Participants at the Workshop













Participants at the Workshop

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Participants at the Workshop



#### **Preface**

Considerable attention has been paid to the role of familial factors in the etiology of coronary heart disease (CHD) in man. Several conferences have been held over the past decade that specifically highlighted research priorities and characterized the relevant phenotypes that are known or suspected precursors of a variety of CHD endpoints. The major aim of two such conferences, "Task Force on Genetic Factors in Atherosclerotic Disease" in 1974 and "Genetic Analysis of Common Diseases: Application to Predictive Factors in Coronary Disease" in 1978, was to summarize the state of the art findings on the clinical, genetic, and epidemiological aspects of CHD. This format effectively disseminated a large amount of information to a wide audience. However, disciplinary isolation continues to be a limitation despite the emergence of collaborative efforts. So long as geneticists, epidemiologists, and clinicians continue with their parallel approaches, none is likely to resolve the etiological basis of complex human diseases such as CHD. A sound strategy should forge an interdisciplinary approach. We have deliberately designed this workshop for this purpose, by inviting experts in various disciplines to examine and constructively criticize the scientific value of each others' studies.

These proceedings are the result of an attempt to maximize interdisciplinary communication among geneticists, epidemiologists, and clinicians. On the first day of the conference four major epidemiological studies, chosen as examples of representative, prospective cohort studies containing data on a variety of CHD risk factors, were discussed. Each was reviewed by two experts who had not participated in the study, first by an epidemiologist and then by a geneticist. A formal discussion of these reviews was then conducted by one of the major contributors to the study, which generated additional discussion of similar studies. This format was designed to identify the extent to which earlier contributions have been received by the scientific community and to recognize the potential opportunities that still exist. Only by recognizing the limitations and strengths of past studies can we hope to design future investigations more efficiently. We believe this format succeeded in meeting these objectives, was successful in initiating discussions, and set the stage for a continuous and effective exchange of ideas throughout the conference period and beyond. Using these four studies as case models, the two subsequent days were dedicated to a review of current methodological approaches

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and clarification of intervening phenotypes and endpoint definitions. In addition, and perhaps most importantly, specific genetic hypotheses and optimal statistical procedures were discussed that might best serve as guidelines for future research. Some attention was paid to the general utility of these recommendations for a wide spectrum of common diseases in man including CHD, cancer, behavioral phenotypes, and a variety of lung and blood disorders.

It is hoped that this volume represents another significant, albeit small, step toward the ultimate goal of understanding the etiological basis of coronary heart disease. While we were unable to be comprehensive, we believe we have encapsulated important features of past epidemiological studies, covered the most recent clinical findings, and indicated some optimal approaches to examining the role of genetic epidemiology in the manifestation of CHD. The more successful we become at communicating progressive ideas, revealing methodological or conceptual frustrations, and clarifying joint hypotheses and analytical strategies, the closer we will be to understanding, managing, and eventually controlling coronary heart disease. This workshop will have served its major purpose if it succeeds in convincing clinicians, epidemiologists, and geneticists that real progress on the genetic epidemiology of CHD can only be made through meaningful interactions among these disciplines.

The Editors

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