

CURRICULUM VITAE

Name: Richard E. Gregg, M.D.

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Address: 7 Linden Lane
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Education:

Feb. 1970 - B.S. Biochemistry, Iowa State University

Aug. 1971 - M.S. Biochemistry, Iowa State University

June 1976 - M.D. Stanford University

Board Certification:

Internal Medicine, 1979

Endocrinology and Metabolism, 1981

Brief Chronology of Employment:

- 1976 - 1977 Intern, Internal Medicine, Strong Memorial Hospital, Rochester, NY
- 1977 - 1978 Resident, Internal Medicine, Strong Memorial Hospital, Rochester, NY
- 1978 - 1982 Clinical Associate, Molecular Disease Branch, NHLBI, NIH, Bethesda, MD
- 1982 - 1984 Medical Staff Fellow, Molecular Disease Branch, NHLBI, NIH, Bethesda, MD
- 1984 - 1988 Senior Investigator, Molecular Disease Branch, NHLBI, NIH, Bethesda, MD
- 1988 - 1996 Executive Director, Department of Metabolic Diseases, Bristol-Myers Squibb Pharmaceutical Research Institute, Princeton, NJ
- 1996-1998 Vice President, Metabolic Diseases, Drug Discovery Bristol-Myers Squibb Pharmaceutical Research Institute, Princeton, NJ
- 1999-2001 Vice President, Metabolic and Cardiovascular Drug Discovery Bristol-Myers Squibb Pharmaceutical Research Institute, Princeton, NJ
- 2001-2007 Vice President, Clinical Discovery Bristol-Myers Squibb Pharmaceutical Research Institute, Princeton, NJ

Military Service:

Commissioned Corps, United States Public Health Service, 1978 – 1988

Societies:

American Association for the Advancement of Science
American College of Physicians
American Heart Association, Council on Atherosclerosis
American Diabetes Association

Honors:

Undergraduate

Phi Kappa Phi (Scientific Honorary Society), Iowa State University, 1970
Award for the Most Outstanding Student in One's Major Field (Biochemistry),
Iowa State University, 1970
Graduated with Honors and Distinction, Iowa State University, 1970

Medical School

It was the policy of Stanford University School of Medicine during the time of my enrollment not to give scholastic honors or awards.

Western Student Medical Research Forum Award for Meritorious Research,
1975 and 1976

Professional

Fellow of the Council on Atherosclerosis, American Heart Association, 1982
Fellow of the American College of Physicians, 1985
Pfizer Traveling Lectureship, Montreal Clinical Research Institute, 1985
Visiting Professor, Shandong Medical College, Jinan, China, 1986

Member of Board of Directors

American Federation for Aging Research (1999-2005)
Robert Wood Johnson University Hospital at Hamilton (2001-present)

Research Experience:

- 1969 - 1971 Mechanism of protein biosynthesis in eukaryotic organisms
- 1974 - 1976 Metabolism of triglycerides in experimentally induced acute and chronic uremia
- 1978 - 1988 Metabolism of lipoproteins and apolipoproteins in human subjects
- 1988 - 2001 Led Drug Discovery activities for BMS in Cardiovascular and Metabolic Diseases; regulation of lipoprotein metabolism; regulation of carbohydrate and fatty acid metabolism; regulation of weight; aging and chronic degenerative diseases; atherosclerosis and vessel wall biology; thrombosis. Twenty-five compounds nominated for clinical development; seven compounds still in clinical development.
- 2001-2007 Led all Exploratory Development and Clinical Pharmacology activities for BMS; Phase I&II, mechanism of action, and proof of concept studies; Clinical Pharmacology studies in all phases; development and use of novel biomarkers; pharmacogenetic and genomic studies; developed modeling and simulation capabilities.

BMS Special Activities:

- 1995 - 1998 Led the Exploratory Development working groups for squalene synthase inhibitors and for MTP inhibitors
- 1996 Drug Discovery Productivity Initiative
Member of Task Force
Responsible for writing Drug Discovery Handbook
Change leader; responsible for planning and rolling out Productivity Initiative to all of Drug Discovery
- 1997 Selected to participate in the inaugural CEO's Forum (two day meeting with CEO for high potential mid-level executives)
- 1998 Led the team that developed the vision for Applied Genomics and initiated the efforts for BMS in genetics/genomics/biotechnology
- 2001 Core member of Drug Discovery Redesign Task Force
Member of Clinical Development Redesign Task Force
Developed vision, organizational structure, and processes for Clinical Discovery
Led the team to develop structure and processes for Exploratory Development Teams (strategic planning and development coordination for Exploratory Development compounds)
Responsible for selection and assimilation of members of DuPont Pharma's Drug Metabolism and Clinical Pharmacology Departments into BMS Clinical Discovery
- 2002 Led the team to develop the structure and processes for Exploratory Clinical Research Teams (exploratory development clinical operations)
Led the team to coordinate closure of the Wilmington Drug Discovery site and assimilate employees into other BMS sites and projects
Led the Drug Discovery Strategic Portfolio and Staffing team (set strategies and processes for BMS Drug Discovery)
- 2002 - 2007 Co-leader of Exploratory Development Operating Committee

Member Brand Development Operating Committee (governance body for Full Development)

Member Institute Executive Committee

Member Medical Review Committee (most senior medical review group for BMS)

Responsible for approving First in Human studies

BIBLIOGRAPHY

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3. Gregg, R.E., Diamond, A., and Reaven, G.M.: Effect of chronic uremia and glucocorticoid therapy on triglyceride kinetics in rat. Metabolism 26:875-882, 1977.
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5. Ghiselli, G., Schaefer, E.J., Zech, L.A., Gregg, R.E., Brewer, H.B. Jr.: Increased prevalence of apolipoprotein E₄ in type V hyperlipoproteinemia. J. Clin. Invest. 70:474-477, 1982.
6. Ghiselli, G., Gregg, R.E., Zech, L.A., Schaefer, E.J., Brewer, H.B., Jr.: Phenotype study of apolipoprotein E isoforms in hyperlipoproteinemia patients. The Lancet ii:405-407, 1982.
7. Schaefer, E.J., Zech, L.A., Gregg, R.E., Brewer, H.B., Jr.: Metabolism of high density lipoproteins. In: Proceedings of the USA-USSR First Lipoprotein Symposium. U.S. Department of Health and Human Services, NIH Publication No. 83-1966, Washington, D.C., U.S. Government Printing Office, pp. 105-122, 1982.
8. Goldstein, D.S., Dionne, R., Sweet, J., Gracely, R., Brewer, H.B., Jr., Gregg, R.E., and Keiser, H.R.: Circulatory, plasma catecholamine, cortisol, lipid and psychological responses to a real-life stress (Third molar extractions): Effects of diazepam sedation and of inclusions of epinephrine with the local anesthetic. Psychosomatic Medicine 44:259-272, 1982.
9. Brown, R.E., Gregg, R.E., and Hood, J.C.: Droperidol treatment of streptozocin-induced nausea and vomiting. Drug Intell. and Clin. Pharm. 16:775-776, 1982.
10. Cogan, D.G., Chu, F.C., Barringer, J. and Gregg, R.E.: Maculo Halo Syndrome. Tr. Am. Ophth. Soc. 80:184-192, 1983.
11. Gregg, R.E., Zech, L.A., and Brewer, H.B., Jr.: Apolipoprotein E alleles in severe hypertriglyceridemia. The Lancet 1:353, 1983.
12. Gregg, R.E.: Molecular and metabolic defects. In: Type III hyperlipoproteinemia: Diagnosis, molecular defects, pathology, and treatment. Brewer, H.B., Jr. Moderator

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