Commonwealth Health Research Board 2006 Annual Report

"To promote and protect the health of the citizens of the Commonwealth through human health research." Commonwealth Health Research Board Members

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Message From The Chairman

The Commonwealth Health Research Board (CHRB) provides grant funding for creative and innovative research projects that have scientific merit and hold promise for maximizing human health benefits for citizens of the Commonwealth of Virginia. The CHRB supports both new research efforts and the expansion or continuation of existing research efforts.

Since its inception, the CHRB has made 89 grant awards totaling \$5.9 million in grant funding to institutions of higher education and other organizations in Virginia. When the required 33% matching funds are added to the CHRB funded amount, the total project funds amount to \$8.6 million for health research in Virginia.

Grants have been awarded to 16 institutions of higher education and other organizations across the Commonwealth to include: University of Virginia, Eastern Virginia Medical School, Virginia Commonwealth University, Virginia Military Institute, The College of William and Mary, University of Richmond, Longwood University, Norfolk State University, George Mason University, Lynchburg College, James Madison University, Sweet Briar College, Shenandoah University, Virginia Polytechnic Institute and State University, and the Riverside Health System. Grants have been awarded for research on a wide variety of important health conditions including: Alzheimer's disease, diabetes, cancer, asthma, Lyme disease, high blood pressure, renal disease, new chemotherapeutic interventions, and genetic diseases, to mention a few.

The CHRB encourages collaborative research efforts and gives priority to those research efforts where Board support can be leveraged to foster contributions from other entities. CHRB grant recipients, for grant awards made in 1999 through 2002, have leveraged almost \$13 million in additional private and federal grant funds to further their research studies. In addition, numerous publications in peer-reviewed scientific journals and periodicals as well as presentations of the data at regional and national scientific meetings have resulted from CHRB grant funded research projects.

We are proud of the accomplishments of the CHRB and our grant recipients as we work toward a healthier future for all Virginians. As Chairman, I am pleased to present the 2006 CHRB Annual Report.

S. Lawrence Kocot, Chairman Commonwealth Health Research Board July 2006

Commonwealth Health Research Board Evaluation Highlights

Comments from Principal Investigators concerning their research and their success in obtaining additional grant funds from federal or private foundation organizations as a result of initial CHRB grant support.

Margaret Saha, Ph.D.

The College of William and Mary: 2000 CHRB grant recipient for a project entitled, *Imaging the aging brain: In vivo detection of key aging molecules in small animals.*

"The CHRB is an extremely valuable source of funding because it not only allows, but encourages, innovative and risky interdisciplinary projects that could not have received funding elsewhere. This allows investigators to obtain the preliminary data and be successful in future funding endeavors.

Certainly, in part, funding from the CHRB, and the preliminary data obtained from that funding, has allowed me to obtain a grant from the Department of Defense Breast Cancer Research Program (BCRP) of the Office of the Congressionally Directed Medical Research Programs, "In Vivo Molecular Imaging of Mammary Tumorigenesis in Murine Model Systems" (2005-2006), \$107,015."

Image of a live mouse showing expected iodide accumulation in stomach and thyroid and accumulation of signal as a marker of a developing breast tumor (left side of photo).



Introduction



Geoffrey Krystal, M.D., Ph.D. Virginia Commonwealth University: 1999 CHRB grant recipient for a project entitled, Inhibition of P13K as a Novel Therapeutic Strategy for the Treatment of Small cell lung cancer (SCLC).

Preliminary data generated as a result of the CHRB award was used to obtain a Merit Review Award from the Department of Veteran's Affairs Research Service of \$689,800 over the period of 2001-2006. The complete set of data also served as a cornerstone for the renewal of the Merit Review Award that will run from 2006-2010 at total direct cost of \$535,200.

CHRB Introduction

The goals of the Commonwealth Health Research Board (CHRB) include providing grant funding for research to advance the understanding of biological systems, to improve the treatment and control of human disease, and to improve human health services and the delivery of human health care.

The CHRB provides grant funding for research efforts that have the potential of maximizing the health of Virginia's citizens. Research efforts eligible for support include traditional medical and biomedical research related to the causes and cures of human diseases, as well as research related to human health services and the delivery of human health care.

More specifically, in accordance with § 23-279 of the *Code of Virginia*, the Board encourages collaborative research efforts among two or more institutions or organizations, gives priority to those research efforts where Board support can be leveraged to foster contributions from federal agencies or other entities, and supports both new research efforts and the expansion or continuation of existing research efforts.

Background

Legislation in 1997 created the Commonwealth Health Research Fund to provide financial support for research projects that have the potential of maximizing health benefits for the citizens of the Commonwealth. This initiative of the General Assembly and the Governor used the proceeds from the sale of Trigon stock to create the framework and fiscal resources for a research grant program. The funds result from the stock and cash distributed to the Commonwealth of Virginia pursuant to the conversion of Blue Cross and Blue Shield from a mutual insurance company to a stock corporation. Income from the funds is used to make grants. The estimated value of the Fund as of March 31, 2006 was \$29.6 million.

The Commonwealth established the Commonwealth Health Research Board (CHRB) to develop and implement the grant program. The following chart shows the number and amount of grant funds awarded along with the amount of matching funds provided by the grantee institution.

Grant Year	Number of Grant Awards	CHRB Grant Awards	Grantee Matching Funds	Total Project Funds
1999	9	\$ 597,377	\$ 260,151	\$ 857,528
2000	11	\$ 719,442	\$ 429,489	\$ 1,148,931
2001	13	\$ 825,590	\$ 341,680	\$ 1,167,270
2002	12	\$ 718,382	\$ 344,603	\$ 1,062,985
2003	8	\$ 509,806	\$ 199,999	\$ 709,805
2004	14	\$ 887,914	\$ 376,735	\$ 1,264,649
2005	10	\$ 755,436	\$ 305,909	\$ 1,061,345
2006	12	\$ 954,058	\$ 451,983	\$ 1,406,041
Cumulative Total	89	\$ 5,968,005	\$ 2,710,549	\$ 8, <mark>678</mark> ,554



Number of CHRB Grant Awards to Date by Institution or Organization



CHRB Grant Funding to Date by Institution or Organization



Birgit Winther, M.D. University of Virginia: 1999 CHRB grant recipient for a project entitled, Effects of common colds caused by viruses on middle-ear pressure in children.

"I am so grateful for the CHRB award. It had major impact on my research career. In addition, it provided new information on the effect of cold virus on the ears in children. The new information was used as a stepping stone for a NIH award which I have now had for two years."

Dr. Winther received a 5-year grant award from the National Institutes of Health (NIH) in the amount of \$980,000 as a result of the initial CHRB grant award.



Nasal mucus is obtained for laboratory testing of cold viruses by PCR.



Middle ear pressure is obtained with a digital tympanometer.

Grant Awards FY 2006/2007



Michael McVoy, Ph.D.

Virginia Commonwealth University: 2001 CHRB grant recipient for a project entitled, Antiviral mechanisms of herpes virus DNA packaging inhibitors.

"It takes substantial preliminary data and a proven track record in the form of publications for a new investigator to obtain federal research grants. For scientists engaged in research at institutions within the Commonwealth of Virginia, the opportunities to obtain sufficient funds with which to generate this preliminary data and a solid publication record (i.e., > \$50,000) are extremely limited (I know of only two - Jeffress and CHRB). I am therefore very grateful for the CHRB grant that my lab received. It came at a critical time in my career in which I very much needed those funds to move my research to a point where I could be competitive for NIH grants. It also engendered an important and fruitful collaboration with my partner in this grant, Jay Brown at the University of Virginia. I only wish that more funds were available for small grants of this type."

As a result of the CHRB grant award, Dr. McVoy was awarded a grant from the National Institute of Allergy and Infectious Diseases (NIH/NIAID) R21 AIO53768: Analysis of cytomegalovirus DNA cleavage/packaging genes, in the amount of \$347,500 for the period 10-01-03 to 09-30-05. Dr. McVoy also has a grant renewal pending: NIH/NIAID R01 AI46668, Human cytomegalovirus DNA cleavage and packaging in the amount of \$1,489,575 for the period 07-01-06 to 06-30-11.

Commonwealth Health Research Board

Total funding for all grant awards in FY 2006/2007 including second year grant funding for four 2005/2006 grant awards

Submitting Institution/ Organization	Principal Investigator	Grant Title	CHRB Grant Award	Matching Funds	Total Project Funds
Eastern Virginia Medical School	Henri Parson, Ph.D.	Characterization of Diabetic Neuropathy and Genetic Differences in African-Americans	\$96,215	\$52,187	\$148,402
University of Virginia	G. Paul Matherne, M.D.	Evaluation of metabolic remodeling in the prevention of left ventricular hypertrophy with A1 adenosine receptors	\$75,000	\$94,894	\$169,894
Eastern Virginia Medical School	Diane M. Duffy, Ph.D.	Aspirin, Eggs and Pregnancy: A New Method of Birth Control?	\$97,830	\$39,915	\$137,745
University of Richmond	Michael C. Leopold, Ph.D.	Crown Ether Modified Naoparticle Films as Metal Ion Sensing Materials	\$48,000	\$29,595	\$77,595
Virginia Commonwealth University	Shyama Masilamani, Ph.D.	Sex Hormone-Mediated Regulation of Kidney Transporters and Blood Pressure Control	\$98,107	\$32,375	\$130,482
Eastern Virginia Medical School	Michael J. Solhaug, M.D.	Elucidating the Mechanism of IGF- 1's Protective Role on Acute Renal Failure	\$100,000	\$33,000	\$133,000
Virginia Commonwealth University	J. Randy Koch, Ph.D.	Assessing the Utility of Consumer Surveys for Improving the Quality of Behavioral Healthcare Services	\$42,349	\$13,975	\$56,324
Virginia Commonwealth University	Zendra Zehner, Ph.D.	The role of the Oncogene Stat3 and Tumor Suppressor ZBP-89 in Tumor Metastasis	\$100,000	\$33,000	\$133,000
Eastern Virginia Medical School	Yuliya Dobrydneva, Ph.D.	Thrombosis in women undergoing tamoxifen chemoprevention therapy	g \$61,421	\$21,497	\$82,918
University of Virginia	Gary Kupfer, M.D.	Tax 1 induced chemo sensitization of p53 mutant resistant tumors	\$75,000	\$25,000	\$100,000
Eastern Virginia Medical School	Gyorgy Lonart, Ph.D.	The role of SNAP-25 and RIM1a genes in attention-deficit hyperactivity disorder	\$99,869	\$50,349	\$150,218
Norfolk State University	Kara Witzke, Ph.D.	Improving Treatment of Charcot Foot in the Diabetic Patient: Understanding the Etiology of the Disease and its associated Fractures	\$60,267	\$26,196	\$86,453

Grand Total all grant awards for FY 2006/2007



Abstracts for 2006/2007 Grant Awards (July 1, 2006 to June 30, 2007)

#21-06 Henri Parson, Ph.D. Eastern Virginia Medical School

Grant Title: Characterization of Diabetic Neuropathy and Genetic Differences in African-Americans

Brief summary: Diabetes mellitus is the leading cause of non-traumatic lower extremity amputations (LEAs) and ulcerations in the United States. Recent studies paradoxically indicate that while rate of neuropathy in the African-American (AA) population is lower than in the Caucasian population, lower extremity amputations and ulcerations are more prevalent in AA. Our preliminary data indicates that AAs with diabetes may have an altered vascular system that potentiates abnormal blood flow and causes decreased skin nutrition and nerve integrity which may lead to a greater prevalence of ischemia, loss of sensation, foot ulcers and amputations. We suspect that the reasons AA have a higher frequency of amputation, despite lower rates of neuropathy is not only due to differences in access and disparities regarding health care, but also physiologic and possibly genetic differences which may predispose them to severe diabetic neuropathy and its complications. These facts underscore the need to understand the racial differences in the pathophysiology of diabetic neuropathy.

#16-06 Paul Matherne, M.D. University of Virginia

Grant Title: Evaluation of metabolic remodeling in the prevention of left ventricular hypertrophy with A1 adenosine receptors

Brief summary: Myocardial hypertrophy (thickening of the heart) is a serious consequence of congenital heart defects in children (including aortic stenosis), high blood pressure and heart attacks. Hypertrophy of the heart can in turn impair the ability of the heart to function well, causing heart failure. For this reason it is important to develop effective strategies to prevent hypertrophy of the heart. Adenosine is a naturally occurring compound which we think can prevent a heart from becoming enlarged. This study will use state of the art imaging techniques developed at UVA to investigate how adenosine prevents hypertrophy. These findings will support future national grant applications to develop effective treatments to protect the hearts of children born with congenital heart defects, and in adults with cardiovascular disease.

#31-06 Diane Duffy, Ph.D. Eastern Virginia Medical School

Grant Title: Aspirin, Eggs and Pregnancy: A New Method of Birth Control?

Brief summary: Prostaglandins are produced by the ovary around the time of ovulation, and prostaglandin synthesis inhibitors can reduce fertility in women. This project will determine if prostaglandins act directly at the oocyte to stimulate the maturation and fertilization of the oocyte, steps which are essential for successful reproduction. PCR, immunofluorescence, and measurements of intracellular messengers will be utilized to determine if oocytes express functional receptors for prostaglandins. Studies will also test the hypothesis that prostaglandin treatment increases oocyte maturation and fertilization in vitro. If prostaglandins enhance oocyte maturation and fertilization, then prostaglandin synthesis inhibitors such as aspirin and ibuprofen (Advil, Motrin) may prevent these essential steps, resulting in an effective pre-fertilization contraceptive.

John J. Beck, Ph.D.

Sweet Briar College: 2002 CHRB grant recipient for a project entitled, Syntheses and Structure-Activity Relationship Studies of Aromatic Side-Chain (z)-Ligustilide Derivatives: A Natural Product from Ligusticum Species.

"Funds from the CHRB supported research that provided positive results of an initial investigation into a line of antibacterial compounds. These positive results provided evidence for a new line of antibacterial compounds. These new compounds will be the subject of a grant proposal to the NSF."

As a result of the CHRB grant award, the grant was reported as being awarded on an Naional Science Foundation (NSF) MRI grant request for the acquisition of a high-field NMR. The NSF MRI was for \$254,066 and the funding period was for September 2002 to September 2005. Additionally, the results of the project are the basis of a proposal currently under preparation. The proposal will be submitted to the NSF, Organic Chemistry Dynamics (DYN) program for funding consideration.



George Kulik, Ph.D.

University of Virginia: 2001 CHRB grant recipient for a project entitled, *Molecular Targets for Cancer Therapy by Proteomic Analysis of Antiapoptotic Signaling Pathways.*

As a result of the CHRB grant award, Dr. Kulik leveraged grant funds from the Department of Defense Prostate Cancer Research Program Antiapoptotic signaling in prostate cancer cells in the amount of \$330,000 for the period January 1 2002 – December 31, 2005.

Frank Castora, Ph.D.

Eastern Virginia Medical School: 2001 CHRB grant recipient for a project entitled, *Alzheimer Disease linked to a mutation in mitochondrial DNA*.

"The CHRB award can be instrumental in allowing exciting research projects that may lack the preliminary data to warrant national funding to begin to obtain the data necessary for successful application for NIH or similar national funding."

#01-06 Michael Leopold, Ph.D. University of Richmond

Grant Title: Crown Ether Modified Nanoparticle Films as Metal Ion Sensing Materials

Brief summary: Have you improved your mood by eating a banana or watched the television show ER and heard doctors order a "Chem-7" for a patient? These two observations are both concerned with the amount of potassium in your bloodstream. Adequate potassium levels are crucial for many physiological functions, including the ability to concentrate and regulation of the heart. Clinicians must be able to accurately and efficiently determine potassium concentrations. This task is currently performed by extracting blood and sending it to the laboratory for instrumental analysis while the patient and doctor wait. It would be ideal if this measurement could be made bedside, saving crucial time during an emergency. Nanotechnology has illuminated certain materials that may revolutionize the way potassium levels are measured. Networked films of nanoparticles affixed with potassium specific sensors are thought to be a novel sensing material for this application and are the subject of the proposed research.

#32-06 Shyama Masilamani, Ph.D. Virginia Commonwealth University

Grant Title: Sex Hormone-Mediated Regulation of Kidney Transporters and Blood Pressure Control

Brief summary: One out of three Virginians have been told by a health care provider that they have hypertension. The prevalence of hypertension is greater in men than premenopausal women (men=24%, women=13%; 35-44 age group). The long- term goal of this study is to enhance our understanding of the molecular mechanisms that are involved in the sex related difference in the development of hypertension. Our hypothesis is that premenopausal female DOCA-salt hypertensive rats retain less sodium and fluid compared to males due to a lower expression of epithelial sodium channels (ENaC) and aquaporin 2 (AQP2) water channels. Further, this sex difference is mediated by differential regulation of ENaC and AQP2 by gender hormones. This study will use an integrated physiologic approach, utilizing both in vivo and in vitro studies to examine the regulation of ENaC and AQP2 by sex hormones in the DOCA-salt hypertensive rat.

#30-06 Michael Solhaug, M.D. Eastern Virginia Medical School

Grant Title: Elucidating the Mechanism of IGF-1's Protective Role on Acute Renal Failure

Brief summary: The current proposal will investigate whether the protective effect of Insulinlike growth factor (IGF-1) in ischemic acute renal failure (ARF) is mediated through nitric oxide (NO). Ischemic ARF is characterized by decreased filtration capacity of the kidney leading to impaired function. IGF-1 is a growth factor known to regulate filtration capacity through regulation of blood flow. Therefore, vasodilation of microdissected renal vessels in response to IGF-1, mediated through NO, will be investigated. Renal proximal tubular injury is also associated with ARF. Using a proximal tubule cell culture system, the current proposal will also investigate whether IGF-1 directly protects proximal tubular injury or through modulation of NO synthesis. Data from this study would provide the first direct evidence of a mechanism by which IGF-1 can protect against the renal disturbances seen in ARF. This could lead to new therapeutic agents that improve patient outcome or reduce mortality rate in ARF.

#36-06 Randy Koch, Ph.D. Virginia Commonwealth University

Grant Title: Assessing the Utility of consumer Surveys for Improving the Quality of Behavioral Healthcare Services

Brief summary: There has been significant growth in the use of consumer surveys to assess the quality of behavioral healthcare services. A key rationale for using consumer surveys is their presumed value to providers for improving the quality of services they provide. Although there is anecdotal evidence that some providers have successfully used consumer surveys to improve service quality, we have not determined the extent to which this has occurred. Given the widespread use of consumer surveys and their cost, it is critical that this and related questions be addressed. The proposed study will use a combination of focus groups, surveys, and structured interviews with key stakeholders in both private and public sector treatment programs to describe the extent to which consumer surveys are used for QI, the ways in which they are used, to identify barriers to effective use, and to develop strategies to increase their use.

#40-06 Zendra Zehner, Ph.D. Virginia Commonwealth University

Grant Title: The role of the Oncogene Stat3 and Tumor Suppressor ZBP-89 in Tumor Metastasis

Brief summary: There is no doubt that cancer is a major health threat. Statistics show that 1 in 3 Americans will experience some form of cancer in their lifetime. Most, if not all, cancers originate from epithelial cells, which during tumor progression leave the confines of the primary tumor, invade the interstitial tissue, and disseminate to secondary sites. In this process the epithelial cell is modified making it receptive to new directives. It is thought that by understanding the requirements for this transition, new targets will be uncovered, which, if blocked, could prevent tumor progression. We propose one such target is the oncogene Stat3, which we have shown stimulates gene expression by overcoming the tumor suppressor ZBP-89. Ultimately, we propose to restore the repressor activity of the tumor suppressor ZBP-89, in the hopes that this will serve to control cell growth and halt tumor progression.

#30-05 Gyorgy Lonart, Ph.D.

Eastern Virginia Medical School (a two-year grant awarded in FY 2005/2006)

Grant Title: The role of SNAP-25 and RIM1a genes in attention-deficit hyperactivity disorder

Brief summary: Attention-deficit hyperactivity disorder (ADHD) is a heritable, clinically heterogeneous psychiatric syndrome of inattention, impulsivity and variable hyperactivity. While modern research methods have greatly expanded knowledge, its etiology still remains unknown. Thus, there is a need for improved animal models to aid the accuracy of diagnosis and to develop novel therapeutics for this common disorder. We propose to develop a mouse line carrying mutations for Snap25 and RIM1_ genes as a model for ADHD. The coloboma mice that are mutant for Snap25 are spontaneously hyperactive. In contrast, RIM1_ null mutant mice are not spontaneously hyperactive but display excessive arousal in novel environments, a similarity to ADHD. We plan to create double (RIM1_/ Snap25) mutant mice to take advantage of their complementary features for better modeling of ADHD, and we will test whether they also display other fundamental behavioral and neural deficits of ADHD.

Abdelali Haoudi, Ph.D. Eastern Virginia Medical School: 2002 CHRB grant recipient for a project entitled, *Novel Cancer Gene Therapy for Prostate Cancer*.

"Funding from the CHRB was instrumental for establishing the grounds for a novel scientific finding in the area of cancer gene therapy. Further investigations are needed to clearly establish this exciting finding and further clarify its mechanism of action and control therefore opening a new avenue for a potential cancer gene therapy."

As a result of the CHRB award, Dr. Haoudi leveraged additional grant support from the Elsa U. Pardee Foundation for the period of Sept 2004-Sept 2006 for the amount of \$125,000. Dr. Haoudi is in the process of submitting additional grant proposals to the NIH.



Talissa Altes, M.D.

University of Virginia: 2001 CHRB grant recipient for a project entitled, Hyperpolarized Helium-3 Diffusion Weighted MR of the Lung: An New Technology to Assess the Lung Microstructure.

"Thank you very much for your support! It was integral to our getting started in what has turned out to be a very productive and interesting area of research."

As a result of the CHRB award, Dr. Altes was able to leverage additional grant support from the following sources: (1) Assessment of the variability of hyperpolarized helium-3 gas magnetic resonance imaging in patients with chronic obstructive pulmonary disease, GlaxoSmithKline (GSK), 11/1/03-10/31/04, \$363,069; and (2) A New Method to Detect Early Changes of Emphysema in Persons Exposed to Second Hand Cigarette Smoke, Flight Attendant Medical Research Institute (FAMRI), 7/1/04- 6/31/07, \$317,000.

#15-05 Yuliya Dobrydneva, Ph.D. Eastern Virginia Medical School (a two-year grant awarded in FY 2005/2006)

Grant Title: Thrombosis in women undergoing tamoxifen chemoprevention therapy

Brief summary: Tamoxifen is the most widely used drug for treatment and prevention of breast cancer. However, women taking tamoxifen have elevated risk of thrombosis, a severe and often life-threatening adverse effect of this drug. Though it concerns millions of women, the reason for the higher occurrence of thrombosis has not yet been established. We discovered that tamoxifen activates calcium entry into human platelets by novel mechanism, leading to platelet activation and formation of blood clots. We propose to investigate the cellular mechanism of tamoxifen-induced thrombosis. We will identify intracellular signaling pathways that are activated by tamoxifen and investigate if pharmacological agents such as phytoestrogens and antioxidants can counteract tamoxifen-induced platelet activation. We will identify the key structural features of tamoxifen molecule that are responsible for Ca2+ elevation and thrombosis. This project directly addresses a problem in breast cancer chemotherapy and will afford better understanding of novel pharmacological effect of tamoxifen.

#13-05 Gary Kupfer, M.D. University of Virginia (a two-year grant awarded in FY 2005/2006)

Grant Title: Tax 1 induced chemo sensitization of p53 mutant resistant tumors

Brief summary: p53 protein is crucial to maintenance of normal genome structure. Disruption of normal function of p53 is implicated in a majority of all human cancer and is a major cause of resistance to chemotherapy. We have found that Tax1, a protein made by human T cell leukemia virus I, causes increased sensitivity of p53 disrupted cells to DNA damage. The purpose of this grant is to use Tax1 as a tool to chemosensitize resistant, p53 mutated tumors. This approach has the potential to be used as a novel form of cancer therapy, thus impacting the more than 1.2 million Americans and 31,000 Virginia who were diagnosed with cancer in 2004.

#06-05 Kara A. Witzke, Ph.D.

Norfolk State University (a two-year grant awarded in FY 2005/2006)

Grant Title: Improving Treatment of Charcot Foot in the Diabetic Patient: Understanding the Etiology of the Disease and its associated fractures.

Brief summary: Up to 9,000 Virginians with diabetes also suffer from Charcot foot, a condition where the bones of the foot and ankle become weak and dislocate or crush with even minor trauma. This leads to foot deformity that requires surgery or amputation, which adversely affects quality of life. Management of Charcot is difficult because doctors are unsure what causes Charcot. The purpose of this study is to investigate the cellular mechanism responsible for reduced bone strength in the feet in diabetic patients with Charcot disease compared to healthy subjects and diabetic subjects without Charcot and to determine if a simple bone density test can predict those patients at risk for this disease. These results will revolutionize the way doctors treat Charcot, in suggesting new medications that target the underlying mechanism of the disease, and may help to identify patients susceptible to Charcot so that early interventions are possible.





Investment of Funds

The Virginia Retirement System invests the Commonwealth Health Research Fund (CHRF). The estimated value of the CHRF as of March 31, 2006 was \$29.6 million. The Commonwealth Health Research Fund is a very straightforward account that consists of a combination of equity (70%) and bonds (30%). This asset mix results in a relatively low risk profile when viewed over a market cycle. This account has performed very well over time and has benefited from low investment management fees.



Grant funding is calculated by an amount not to exceed six percent of the moving average of the market value of the Fund calculated over the previous five years or since inception, whichever is shorter, on a one-year delayed basis, net of any administrative fee assessed pursuant to subsection E of § 51.1-124.36, may be expended in a calendar year for any purpose permitted by this chapter. The Board shall not be required to expend such amount in a calendar year, and any amount up to such six percent that is not expended in a calendar year may be expended in any other calendar year. As of April 6, 2005, the Board elected to cap available grant funds at 4% per year.

Commonwealth Health Research Board Fiscal Year 2006 budget for the period – January 1, 2006 to December 31, 2006

Grant Budget for Calendar Year 2006							
Calendar Year January 1 - December 31, 2000 January 1 - December 31, 2001 January 1 - December 31, 2002 January 1 - December 31, 2003 January 1 - December 31, 2004	Year 1 Year 2 Year 3 Year 4 Year 5	Market Value @ 12/31/XX \$28,152,982 \$26,150,016 \$22,273,551 \$26,449,255 \$28,010,649					
Avera Funds available for 2006 grants on average market value:	\$26,207,290 <u>4.00%</u> \$1,048,292						
Less Estimated VRS Administra Estimated Mellon Custodial Fee Estimated VRS Management Fee Estimated State Street Management Total Estimated Administrative Expe	\$ 5,100 \$ 4,600 <u>\$ 3,300</u> \$13,000						
Funds available for 2006 grants estimated administrative expe	\$1,035,292						

Methodology:

The valuation date for market values will be 12/31/XX of each year. Each annual calculation will be made based on the previous five calendar years, with a one year delay. Christopher Kramer, M.D. University of Virginia: 2001 CHRB grant recipient for a project entitled, Imaging Inflammation within Atherosclerotic Plaque with magnetic resonance.

"The CHRB grant was instrumental in completing the project, publishing a manuscript, and the data accrued served as preliminary data for an NIH R01 application that was subsequently funded."

As a result of the CHRB grant award, Dr. Kramer was awarded a National Institutes of Health (NIH) National Heart, Lung and Blood Institute (NHLBI) R01 grant, *Comprehensive Magnetic Resonance in Peripheral Arterial Disease* for the period 9/22/03-8/30/08, in the amount of \$3.7 million total costs including \$2.6 million direct costs.



Representative sequential black blood magnetic resonance images (upper left to lower right) obtained with the use of a surface coil from the femoral artery of a subject with mild to moderate peripheral arterial disease with both the luminal and adventitial border clearly delineated. Note the slice to slice variation in plaque morphology. This technique can be used to reduce sample sizes for clinical trials of novel approaches to reducing atherosclerotic plaque burden. 11

Grant Funding



George Kunos, M.D., Ph.D. Virginia Commonwealth University: 1999 CHRB grant recipient for a project entitled, Development of Novel Drugs for the Treatment of High Blood Pressure Disease.

"I am grateful for the CHRB for the support I received. There is considerable foresight on the part of CHRB in supporting research with potential practical implications, such as the development of novel therapeutic agents."

Jennifer Wayne, Ph.D. Virginia Commonwealth University: 2000 CHRB grant recipient for a project entitled, Mechanical function predicted by MRI parameters in cartilage.

"I am truly grateful for the award from the CHRB which allowed me to explore a new avenue of research and establish collaborations with additional colleagues. The CHRB clearly has foresight in advancing science and technology by supporting research within the Commonwealth for the benefit of Virginians and society as a whole."

Dr. Wayne's continuing work was funded by The National Institute of Arthritis and Musculoskeletal and Skin Diseases (NIAMS) of NIH, 9/2002-2005, for \$326,500.

Eligibility for CHRB Grant Funding

The following Virginia-based entities may apply for a grant from the Commonwealth Health Research Board (CHRB):

- State-supported institutions of higher education,
- > Private, not-for-profit institutions of higher education established in Virginia,
- Agencies of the Commonwealth of Virginia, whose mission is to conduct health or health related research, and
- Nonprofit organizations exempt from income taxation under Section 501 c (3) of the Internal Revenue Code and with their principal offices and programs in the Commonwealth of Virginia whose mission is to conduct health or health related research.

CHRB Grant Application Process

- As part of the Commonwealth Health Research Board (CHRB) grants application process, there are three steps that take place in the review process. More details regarding the required information as part of the submission of a concept paper or a full proposal to the CHRB can be found at www.chrb.org under the heading of CHRB Grant Guidelines.
- <u>Step One:</u> Submission of a concept paper. Concept papers are normally due October 1st. Concept papers (excluding the cover page) must be no longer than five typewritten, doublespaced pages. In general, concept papers will provide information on the problem, need or opportunity that the project will address and the anticipated results or impact of the project. Concept papers will also include estimated total project costs and the amount of funding the applicant is seeking from the CHRB as well as a timeframe for conducting the research. Each concept paper undergoes scientific and technical merit review.
- <u>Step Two:</u> Submission of a full proposal. The Board requests those applicants whose concept papers have been judged, in Step one, to have potential for successful research outcomes, to submit full proposals. Only applicants whom the Board has invited to develop a full proposal may submit a full proposal to the Board. Full proposals are normally due February 1st. The full proposal, excluding the cover page, must be no longer than 12 typewritten, double-spaced pages. In general, full proposals will provide similar information as provided in the concept paper except in greater detail. As in Step One, each full proposal receives in-depth review.

<u>Step Three:</u> Presentation to the CHRB. The Board invites finalists from Step two to make a presentation in-person to the Board. Presentations to the Board are normally scheduled for the May meeting. Presentations, including questions and responses, should take no longer than 15 minutes. The presentation should elaborate on the information contained in the concept paper and formal proposal.

CHRB Grant Guidelines

CHRB grant guidelines are updated annually and posted to the CHRB website at www.chrb.org by August 1st of each year. The grant guidelines are designed to help individuals determine if the research project or initiative for which financial support is sought is a good match with the CHRB purposes and criteria. The guidelines also describe the kinds of research projects and activities the CHRB funds - and does not fund, and tells how and when to apply for a grant. The CHRB website also provides a description of past and current CHRB grant awards and grant abstracts.



Maximum CHRB Grant Award

Applicants may request funding to support projects over either a one-year or a two-year period. The maximum amount of a one-year award is \$100,000. The maximum amount for a two-year award is \$200,000. However, no more than \$100,000 will be provided in either the first or second year. The number of one-year and two-year awards that the CHRB anticipates it will make, is dependent upon the amount of funds available and the number of requests received for each category.

Required 33% Matching Funds from Grantee Institution/Organization

The grantee institution must agree to provide 33% in required matching funds for the CHRB grant award. For example, if the grantee institution requests \$100,000 in CHRB grant funds, then the 33% matching funds would amount to \$33,000.

Other Grant Requirements

- The starting date for all projects awarded CHRB grant funds shall be July 1st of the funding year.
- Requests for funding will be reviewed according to criteria that are consistent with the CHRB's purposes and goals.
- Individuals applying for funding may submit no more than one application per funding cycle.
- The CHRB will accept no more than 15 applications from any one organization or institution of higher education per funding cycle.
- Grantees are responsible for meeting federal, state, and local health and safety standards and for establishing and implementing necessary measures to minimize their employees' risk of injury or illness in activities related to CHRB grants. Grantees are further responsible for meeting all applicable federal, state, and local regulations, requirements, and standards related to human subjects and animal welfare.
- Grantees are responsible for obtaining the appropriate approvals by any internal or external institutional review boards when the use of human subjects or vertebrate animals is proposed. Written documentation must be provided that the approvals have been obtained prior to the release of any CHRB grant funding. If these approvals are not in place by July 1 of the funding year, an award will not be made and the applicant will be required to reapply at a subsequent date for CHRB funding.

CHRB Grant Criteria

Requests for funding will be reviewed in terms of the following criteria to evaluate the likelihood that the proposed research will have a substantial impact on the pursuit of the CHRB's purposes and goals:

Significance: Does the research address an important problem? If the aims of the application are achieved, how will scientific or other knowledge be advanced? What will be the effect of this research on the concepts, methods, or practices in this field?

Paul H. Ratz, Ph.D. Eastern Virginia Medical School:

2000 CHRB grant recipient for a project entitled, Regulation of detrusor smooth muscle contraction by CA2+ and CA2- sensitization.

"Funds provided by the Commonwealth Health Research Board of Virginia enabled my laboratory to acquire a substantial amount of high-guality data that was included in an NIH RO1 grant application. Our goal with this research is to provide a cellular mechanistic approach for the design of new therapeutic agents that will reduce the incidence of urinary incontinence, a chronic disorder that is more prevalent than diabetes. Organizations exist to support research on specific lifethreatening disorders such as hypertension, cancer and diabetes, but research on many non-life-threatening disorders is under funded. Support by the CHRB addresses this issue by providing funds of sufficient magnitude and duration for investigators with diverse interest to pursue their medical research problems in a meaningful and significant way."

As a result of work funded by the CHRB, Dr. Ratz was funded for 4 years at \$730,000 by the National Institute of Diabetes and Digestive and Kidney Diseases at the National Institutes of Health.

Grant Funding



Glenda E. Gillaspy, Ph.D. Virginia Polytechnic Institute and State University: 2001 CHRB grant recipient for a project entitled, Isolation of Genes for Transgenic Production of a Diabetes Treatment.

"The CHRB funding I received was critical in starting a new research project in my laboratory in 2002. I really appreciate this opportunity, and hope the CHRB can continue to fund "seed" projects of Virginia scientists. Although we did not accomplish our intended goal of cloning a chiro inositol epimerase gene from plants, we did find a really exciting connection between inositol and Vitamin C. This finding allowed us to pursue studies currently funded by the National Science Foundation. "

Based on preliminary data funded by the CHRB, Dr. Gillaspy was awarded the following grants from the NSF:

National Science Foundation, Sole Principal Investigator, *Inositol Synthesis and Catabolism in Plants*, for the period 9/01/03- 8/31/06 in the amount of \$380,000; and, National Science Foundation, Sole Principal Investigator, *REU: Inositol Synthesis and Catabolism in Plants*, for the period 9/01/03- 8/31/05 in the amount of \$11,250.

- Collaboration: Will the initiative employ useful collaborative arrangements among two or more institutions or organizations either within or outside the Commonwealth of Virginia? (The Board encourages the submission of collaborative applications.) Are there unique Virginia research resources that will be utilized as part of this proposal?
- Leverage: How will funding provided by the CHRB be used to leverage additional contributions from other federal or private organizations? (The CHRB encourages the submission of applications that employ leveraging.)
- Approach: Are the conceptual framework, design, methods and analyses adequately developed, well integrated, and appropriate to the aims of the project?
- Innovation: Does the project employ novel concepts, approaches or methods? Are the aims original and innovative? Does the project challenge existing paradigms or develop new methodologies or technologies?
- Experience and Qualifications of Research Team: Does the principal investigator have the proper training and experience to direct and manage the project? What amount of time will the Principal Investigator contribute to the project? Has the Principal Investigator conducted research related to this project? Through training and experience, is the research team qualified to conduct this research? Is the research team experienced with research evaluation processes?

Conditions for CHRB Grant Acceptance

Conditions for grant acceptance include a grant agreement between the principal investigator and the grantee institution and the CHRB. Each grantee must sign a Grant Agreement with the CHRB that delineates the terms and specific objectives of the project. Each grantee receiving a one-year or two-year CHRB grant award will be required to submit scientific and fiscal reports at specific times. Specific grant reporting dates are specified in the individualized grant agreement. Also provided are general dates for the distribution of CHRB grant funds over the course of the grant project.







Post Award Reporting Requirements

Each grantee and grantee institution must agree to submit any additional requested data and reports on a timely basis, and to participate in other evaluation efforts required by the CHRB. All published journal articles, monographs, or other special reports based on grant-supported projects must carry a standard footnote of acknowledgment as follows: *"This research was supported by grant funding from Virginia's Commonwealth Health Research Board."* Two reprints of any publication resulting from the funded research must be sent to the Board as soon as they are available. For a period up to five years, the recipient organization also agrees to notify the CHRB of any future grant awards that are received as a result of research funded with grant funds from the Commonwealth Health Research Board.

The CHRB submits an annual report to the Governor and the General Assembly on the Board activities to include an executive summary of the grant process. It also provides information on grants funded in prior years and their success in leveraging additional grant funding from federal or private foundation funding sources.



Additional Funds Leveraged based upon Evaluation Responses Received

Cynthia S. Kelly, M.D. Eastern Virginia Medical School: 2000 CHRB grant recipient for a project entitled, EZ Breathers: Partnership for Asthma Awareness and Prevention in Head Start children.

"Funding provided by the Commonwealth Health Research Board not only helped us to improve care for preschool-aged children with asthma in our community but it provided us with the outcome data necessary to successfully compete for national funding from the Robert Wood Johnson Foundation so that we could expand our program to asthmatic children of all ages in Hampton Roads."

As a result of work funded by the CHRB, Dr. Kelly was successful in obtaining one of eight awards for an "Allies Against Asthma" program funded by the Robert Wood Johnson Foundation, in a competition of 250 investigators. The grant is in an amount of \$1,500,000 over four years.

Commonwealth Health Research Board 2006 Annual Report

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