

BIOGRAPHICAL SKETCH

NAME: Center, David

eRA COMMONS USER NAME (credential, e.g., agency login): dcenter@bu.edu

POSITION TITLE: Associate Provost, Translational Research; Gordon and Ruth Snider Professor of Pulmonary Medicine; Chief, Allergy, Pulmonary and Critical Care Medicine, Boston University Medical Campus

EDUCATION/TRAINING (*Begin with baccalaureate or other initial professional education, such as nursing, include postdoctoral training and residency training if applicable. Add/delete rows as necessary.*)

INSTITUTION AND LOCATION	DEGREE	Completion Date	FIELD OF STUDY
Boston University School of Medicine, Boston, MA	AB, MD	1972	Medicine
Boston University Medical Center, Boston, MA	Resident	1974	Medicine
Boston University Medical Center, Boston, MA	Fellow	1975	Pulmonary Medicine
Robert B. & Peter B. Brigham Hospital/Harvard Medical School, Boston, MA (K. Frank Austen Lab)	Fellow	1978	Research

A. Personal Statement

David Center has been the PI of NIH funded grants since 1978. He directs the Boston University Clinical and Translational Science Institute since 2008. He is the Chief of Pulmonary, Allergy, Sleep and Critical Care Medicine since 1986. He is the discoverer of Interleukin 16, and has studied its nuclear and extracellular structure and function since 1982. He defined its role as a dual function Alarmin, identifying its immunomodulatory functions on CD4 cells and its structure and function as a nuclear scaffold for transcriptional regulatory elements related to the cell cycle and has obtained over 30 patents for reagents, diagnostics and therapeutics related to IL-16 function in inflammatory diseases. His laboratory described the first non-infectious functions for HIV-1 derived proteins and the mechanism of action of IL-16 repression of HIV-1 replication. He is an experienced mentor for predoctoral students, post-doctoral fellows and junior faculty with successful precepting individuals with F31, F32, K08 and fellow and faculty minority supplements to his R01s and P01s. He has been a DSMB member for national NHLBI sponsored trials and now chairs the NHLBI DLD Pulmonary Pragmatic Trials Cooperative which oversees 4 simultaneous large, multi-site pragmatic trials in COPD and Interstitial Lung Disease. As a result he has extensive experience in complex study design and administration of clinical and translational trials. He is an experienced administrator, laboratory and translational scientist with over 34 years as head of a large multidisciplinary Pulmonary, Allergy, Sleep and Critical Care Section and over 12 years of experience leading a CTSA Hub and interacting with the CTSA Network.

Ongoing projects that I would like to highlight include:

UL54 TR001430

Bair-Merritt; Center Saitz

8/13/08 – 03/31/2025

NIH/NCATS

Boston University Clinical and Translational Science Award (CTSA)

Major Goals: The major goals of this project are to infrastructure for clinical and translational scientists, including pilot awards, junior faculty and trainee positions, assistance in bioinformatics, biostatistics, human subjects research approval, study design and recruitment and engagement of the community in translational research efforts.

B. Positions and Honors**Positions and Employment**

2014 – Professor of Biomedical Engineering, Boston University School of Medicine

2008 – Director, Boston University Clinical and Translational Science Institute, Boston University School of Medicine

2007 – Associate Provost for Translational Research, Boston University School of Medicine

- 2006 – Assistant Provost for Translational Research; Director, Clinical and Translational Science Institute, Boston University School of Medicine
- 2000 – Gordon and Ruth Snider Professor of Pulmonary Medicine, Boston University School of Medicine
- 1989 – Professor of Medicine and Research Professor of Biochemistry, Boston University School of Medicine
- 1987 – Chief, Pulmonary, Allergy and Critical Care Medicine, Boston University School of Medicine
- 1983 – 1989 Associate Professor of Medicine, Boston University School of Medicine
- 1978 – 1983 Chief, Allergy Unit, Boston University School of Medicine
- 1978 – 1983 Assistant Professor of Medicine, Boston University School of Medicine

Other Experience and Professional Memberships

- American Association of Immunologists
- American Thoracic Society
- Elected, American Society for Clinical Investigation
- Elected, Association of American Physicians
- Society for Clinical and Translational Science
- Fellow, American Association for the Advancement of Science
- American Academy of Asthma, Allergy and Immunology

Honors and Notable Leadership Positions

- 2017 – Treasurer, Association for Clinical and Translational Science (ACTS)
- 2017 Breathing for Life Award, American Thoracic Society
- 2015 – Chair, DLD/NHLBI Pulmonary Trials Cooperative
- 2014 – Elected Fellow, American Association for the Advancement of Science
- 2014 – Distinguished Editor, NIH Director's New Innovator Award Review Editorial Board
- 2013 Elected, National Academy of Inventors
- 2013 Edward Livingston Trudeau Medal of the American Thoracic Society
- 2011 Commencement Speaker, Boston University Seven Year Medical Department Graduation
- 2011 Secretary/Treasurer, Society for Clinical and Translational Medicine
- 2009 – 2010 Board Member, Society for Clinical and Translational Science
- 2009 – NIH Director's Pioneer Awards Reviewer
- 2008 – NIH Director's New Innovator Award Reviewer
- 2008 Chadwick Medal of the Massachusetts Thoracic Society
- 2007 – 2013 NHLBI T32 Review Committee
- 2006 – NHLBI LRP Reviewer
- 2003 – 2009 Best of Boston, Pulmonary Physicians
- 2002 Best Physicians in America
- 2002 Distinguished Alumnus Award, Boston University School of Medicine
- 2000 – 2001 Chair, ATS Nomination Committee
- 1999 – 2006 NHLBI Board of Extramural Advisors
- 1999 – 2003 Chair, American Thoracic Society Publications Policy Committee
- 1996 – 2000 Charter Member, NIH Peer Review Oversight Group (PROG)
- 1995 – 1996 NIH DRG Advisory Council
- 1992 – 1995 Chairman, American Lung Association Research Grant Review Committee
- 1992 – 1994 Chairman, NIH Lung Biology Pathology Study Section
- 1989 Co-Chairperson, Fellowship Fellow Review Committee, American Lung Association
- 1987 – 1990 Respiration Study Section, Veteran's Administration Merit Review
- 1987 – 1988 National Secretary/Treasurer, A.F.C.R.
- 1985 – 1990 Career Investigator of the American Lung Association
- 1981 Edward Livingston Trudeau Fellow of the American Thoracic Society

C. Contributions to Science

1. **Chemotactic Factor for Lymphocytes:** These back-to-back papers represent the first description of a naturally derived chemotactic factor for lymphocytes. This represented a complete paradigm shift in the thinking of lymphocyte homing and accumulation, for before these papers lymphocytes were thought not to

respond along concentration gradients as part of accumulation in lymph nodes and tissue sites of inflammation but homed via adhesion molecules. These papers also set the stage for the discovery of the entire family of lymphocyte specific chemotactic cytokines (chemokines). The Lymphocyte Chemoattractant Factor described here was later renamed Interleukin 16 and began a series of over 50 papers describing the functions of IL-16 from which over 30 patents have been issued to BU.

- a) **Center DM**, Cruikshank W. Modulation of lymphocyte migration by human lymphokines. I. Identification and characterization of chemoattractant activity for lymphocytes from mitogen-stimulated mononuclear cells. *J Immunol.* 1982;128(6):2563-8. PMID: 7042840.
- b) Cruikshank W, **Center DM**. Modulation of lymphocyte migration by human lymphokines. II. Purification of a lymphotactic factor (LCF). *J Immunol.* 1982;128(6):2569-74. PMID: 7042841.

2. **Gene Products of HIV-1:** This paradigm shifting work demonstrated for the first time that gene products of HIV-1, known to be present in the circulation of infected individuals, had specific functions independent of viral infection. The study shows that gp120 is responsible for CD4+ T cell homing and activation, making T cells more susceptible to infection. A follow up paper in *Nature* (Mackiewicz C, Levy J, Cruikshank W, Kornfeld H, Center D. Role of IL-16 in HIV Replication. *Nature.* 1996;383:488-9) defined the mechanism of IL-16 repression of HIV1 replication; and a *J. Virology* paper (Green, DS, Center, DM, Cruikshank, WW. HIV-1 gp120 reprogramming of T cell migration provides a mechanism for lymphadenopathy. *J Virol.* 83:5765-5770. PMID: PMC2681967) demonstrated that HIV-1gp120 was capable of heterologous chemotactic factor desensitization and deactivating responses to S1P, therefore explaining the lymphadenopathy in early HIV-1 infection.

- a) Kornfeld H, Cruikshank WW, Pyle SW, Berman JS, **Center DM**. Lymphocyte Activation by HIV-1 Envelope Glycoprotein. *Nature.* 1988;335:445-448. PMID: 2843775.
- b) Mackiewicz CE, Levy JA, Cruikshank WW, Kornfeld H, **Center DM**. Role of IL-16 in HIV Replication. *Nature.* 1996;383:488-9. PMID: 8849720.
- c) Green DS, **Center DM**, Cruikshank WW. Human immunodeficiency virus type 1 gp120 reprogramming of CD4+ T-cell migration provides a mechanism for lymphadenopathy. *J Virol.* 2009;83(11):5765-72. PMID: PMC2681967

3. **Interleukin-16:** Cloning of Interleukin-16 and identification of its function as a CD4 ligand inducing migration and activation of CD4+ cells led to the naming of this cytokine as IL-16 and provided the neutralizing antibodies for all the subsequent studies of function and for the issue of 33 patents related to immune modulation by IL-16 in a variety of diseases, ranging from multiple myeloma to multiple sclerosis and asthma.

- a) Cruikshank WW, **Center DM**, Nisar N, Wu M, Natke B, Theodore AC, Kornfeld H. Molecular and functional analysis of a lymphocyte chemoattractant factor: association of biologic function with CD4 expression. *Proc Natl Acad Sci U S A.* 1994;91(11):5109-13. PMID: PMC43941.
- b) **Center DM**, Kornfeld H, Ryan TC, Cruikshank WW. Interleukin 16: implications for CD4 functions and HIV-1 progression. *Immunol Today.* 2000 Jun;21(6):273-80. PMID: 10825739.
- c) **Center DM**, Kornfeld H, Cruikshank WW. Interleukin 16 and its function as a CD4 ligand. *Immunol Today.* 1996 Oct;17(10):476-81. PMID: 8908813.
- d) **Center DM**, Kornfeld H, Wu MJ, Falvo M, Theodore AC, Bernardo J, Berman JS, Cruikshank WW, Djukanovic R, Teran L, et al. Cytokine binding to CD4+ inflammatory cells: implications for asthma. *Am J Respir Crit Care Med.* 1994 Nov;150(5 Pt 2):S59-62. PMID: 7952594.

4. **IL-16 Selective Chemotactic Activity:** We demonstrated IL-16 selective chemotactic activity for T regulatory cells via CD4 ligands and the induction of the FoxP3+IL-2Rhi phenotype of CD4+ T regulatory cells, thus providing a mechanism for the immunosuppressive effects of IL-16 and a TGFbeta independent pathway for generating T regulatory cells.

- a) McFadden CS, Green D, Yamasaki H, **Center DM**, Cruikshank WW. Preferential migration of T regulatory cells induced by IL-16. *J Immunol.* 2007;179:6439-45. PMID: 17982032.

- b) Lynch EA, Heijens CA, Horst NF, **Center DM**, Cruikshank WW. Cutting edge: IL-16/CD4 preferentially induces Th1 cell migration: requirement of CCR5. *J Immunol.* 2003 Nov 15;171(10):4965-8. PMID: 14607889.
5. **Pro-IL-16:** The first paper listed defined Pro-IL-16 as first (described) PDZ domain containing nuclear scaffold. It is the most highly regulated gene following T cell activation and is the limiting factor in Skp2 transcription, and thus p27 levels permitting T cell exit from G1. It also defines IL-16 as a dual function Alarmin.
- a) Zhang Y, Tuzova M, Xiao Z-XJ, Hanson SK, Xie H, Kornfeld H, Cruikshank WW, **Center DM**. Pro-Interleukin-16 is a scaffold protein which targets Histone Deacetylase 3 to Transcription Factor GABP in the Skp2 Core Promoter. *J Immunol.* 2008;180:402-408. PMID: 18097041.
- b) **Center DM**, Cruikshank WW, Zhang Y. Nuclear pro-IL-16 regulation of T cell proliferation: p27(KIP1)-dependent G0/G1 arrest mediated by inhibition of Skp2 transcription. *J Immunol.* 2004 Feb 1;172(3):1654-60. PMID: 14734747.
- c) Zhang Y, Tuzova M, Xiao ZX, Cruikshank WW, **Center DM**. Pro-IL-16 recruits histone deacetylase 3 to the Skp2 core promoter through interaction with transcription factor GABP. *J Immunol.* 2008 Jan 1;180(1):402-8. PMID: 18097041.

Complete List of Published Work in MyBibliography:

<https://www.ncbi.nlm.nih.gov/myncbi/david.center.2/bibliography/public/>