



Linking Research and Researchers



Boston University Chobanian & Avedisian School of Medicine

Boston University Research
Interdisciplinary Biomedical Research Office

IBRO <https://rb.gy/ybd8r>



www.bumc.bu.edu/evanscenteribr





MISSION STATEMENT

Affinity Research Collaboratives (ARCs) consist of faculty and trainees from different disciplines across campuses, and are organized around foci of common research interests. The extraordinary strength in biomedical and physical sciences at Boston University, and the support and development of the ARCs create opportunities for new interdisciplinary approaches to both research and training in biomedical research. Basic science discovery promoted by ARCs are also available to the Clinical and Translational Science Institute and to other centers for collaborative translational applications.

FOUNDED and SUPPORTED BY

The Evans Center for Interdisciplinary Biomedical Research (**Dr. Katya Ravid, Founding Director**), open to faculty across BU campuses, is supported by the Department of Medicine (past recent chair Dr. David L. Coleman; current chair, Dr. Anthony N. Hollenberg), BU Clinical & Translational Science Institute, BU vice chair for research office and by the Evans Medical Foundation (2009-present). The Boston University Interdisciplinary Biomedical Research Office (BU IBRO) (2015-present) within the office of BU Associate Provost for Research provides additional collaborations and support.

ARC PATHWAYS TO IMPACT (with achievement of ≥ 3 deemed as success)

1. **Publish Science** to build a scientific knowledge base to make an impact in research
 2. **Obtain grants** to advance scientific knowledge to make an impact in research
 3. Engage with corporations to implement **Sponsored Research Agreements** to make an impact in research
 4. Engage with corporations to get **licensing agreements** to make an impact in research
 5. Engage with corporations and investors to launch **start-up companies** to make an impact in research
 6. **Present** science at meetings to catalyze scientific knowledge and collaborators to make an impact in research
 7. **Use Science dissemination strategies** to fast-track timely uptake of research by investigators, policy groups, news media and the public to increase visibility and to make an impact in research
 8. **Engage in advocacy and community-engagement** to translate research into guidelines, practices & policy to make an impact in research
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The Evans Center for Interdisciplinary Biomedical Research (ECIBR)
and BU Interdisciplinary Biomedical Research Office (BU IBRO)
Founding and current Director: Katya Ravid, Barbara E. Corkey Professor of Medicine
Professor of Biochemistry, Biology, Health Sciences; Fulbright Research Scholar

BY THE NUMBERS

ARC Outcomes and Types of Metrics for Monitoring and Evaluating Success (2009-2023). IBRO, established in 2015, serves to promote team science university-wide).

1. 24 multidisciplinary ARCs have received funding to investigate novel research topics since 2009, with about 309 core faculty participants, and **261 predocs** & **198 postdocs**. Minority researchers have similar percent representation as in the faculty body.
2. 4 multidisciplinary **ARCs have engaged with 4 hubs in CTSA network**; an ARC-driven Thrombotic Microangiopathy regional Consortium; Adoption of the ARC Model by 2 hubs in the CTSA network.
3. **1,183 collaborative publications** catalyzed by ARCs, to date; 75% of co- authors had never published together prior to the ARC.
4. **428 external grants** awarded to ARC affiliated projects out of 762 applied for to date (totaling **\$299,107,317**), with at least two ARC faculty in the co-PI or co-I roles.
5. ARCs catalyzed **investigators from 14 schools, 38 departments & industry** to form multidisciplinary research projects.
6. Average of 12 seminars, 3 workshops, a retreat, and a symposium per each ARC per year, attracted a range of 25 – 60 investigators & trainees per event. *Exception 2023: Precision Medicine in Alzheimer Disease ARC International Symposium with 400+ attendees.
7. Dr. Ravid co-developed Master of Science Programs in Nanomedicine and in Biomedical Research Technologies, currently instructed by some ARC members.
8. 8 ARCs developed during 2015-2023, including a new initiative focused on Tuberculosis and one on Musculoskeletal Health; Multidisciplinary Program to Identify Predictors of Efficacy and Resistance to Cancer Checkpoint inhibition; COVID-19; Tobacco Sciences; Thrombosis and Hemostasis; Fibrosis: Connecting Tissues and Investigators; Mobile and Electronic Health; Precision Medicine for Alzheimer Disease
9. 4 publications about the influence of the ARC Model and best practices shared with the CTSA network.
10. We rapidly respond to newly emerging biomedical research and technology challenges, as exemplified by a new ARC focused on Biomedical Innovation Technologies (**BIT-ARC**), now in its inaugural year (developed in 2023; see below).

NEW INITIATIVES!!

- page 29: Biomedical Innovation & Technology ARC
- page 30: Department of Medicine Researchers Forum

KEYS FOR SUCCESS: THE FIVE Cs (K. Ravid)

- 1 **Capability**
Creative research ideas
- 2 **Cooperation**
Willingness to learn
- 3 **Communication**
Effective scientific exchanges
- 4 **Coaching**
Generous mentoring/insights by the Center's Director and ARC leadership
- 5 **Conditions**
Supportive resources and academic culture

PUBLICATIONS

"Thrombotic Microangiopathy: A Multidisciplinary Team Approach." Gordon CE, Chitalia VC, Sloan JM, Salant DJ, Coleman DL, Quillen K, Ravid K, Francis JM. Am J Kidney Dis. 2017 Nov;70(5):715-721.

View at <http://bit.ly/2k9qLG8>

"Catalyzing Interdisciplinary Research and Training: Initial Outcomes and Evolution of the Affinity Research Collaboratives Model." Ravid K, Seta F, Center D, Waters G, Coleman D. Acad Med. 2017 Oct;92(10):1399-1405.

View at <http://bit.ly/2kdXH0g>

"Promoting interdisciplinary research in departments of medicine: results from two models at Boston University School of Medicine." Coleman DL, Spira A, Ravid K. Trans Am Clin Climatol Assoc. 2013;124:275-82.

View at <http://bit.ly/2kdYgXW>

"Building interdisciplinary biomedical research using novel collaboratives." Ravid K, Faux R, Corkey B, Coleman D. Acad Med. 2013 Feb; 88(2):179-84.

View at <http://bit.ly/2AKOJhU>

Musculoskeletal Health (MHet)

Directors: Drs. Paola Divieti Pajevic, David Felson, and LaDora Thompson

(ARC, 2023-present)



SYNOPSIS

(This ARC includes 20 faculty from various disciplines)

The mission of the Musculoskeletal Health (MHet) Affinity Research Center (ARC) is to bring together investigators across specialties and campuses (Medical and Charles River Campuses) to conduct multidisciplinary studies related to the musculoskeletal system.

The MHet ARC will comprise investigators from Boston University Chobanian & Avedisian School of Medicine, Boston University Goldman School of Dental Medicine, Boston Medical Center, Boston University Sargent College, and Boston University College of Engineering.

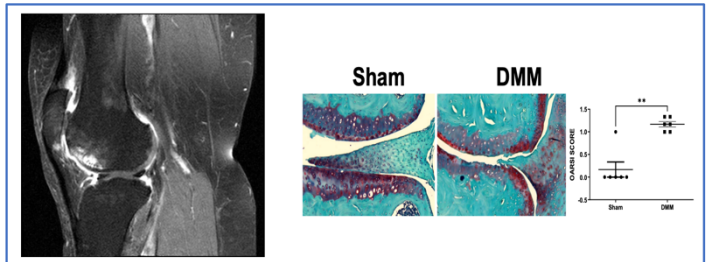


Image by Divieti Pajevic, Felson, Thompson

In addition to specific research projects, the following aims are proposed:

- 1) Create a university-wide community of investigators interested in researching musculoskeletal conditions;
- 2) Generate pilot data using animal models and organ systems to support NIH grant proposals;
- 3) Train the next generation of researchers (pre- and post-doctoral fellows) and;
- 4) Organize monthly meetings, seminars, and an annual mini symposium with invited speakers.

HIGHLIGHTS

Core members are Drs. Louis Gerstenfeld and Beth Bragdon (project 1), Drs. Brianne Connizzo and Katherine Zhang (project 2), Beth Bragdon (project 3) and Drs. Louis Gerstenfeld and Paul Tornetta (Aim2). The MHet will comprise numerous trainees and more than 20 faculty members committed to and interested in musculoskeletal research.

EXAMPLE OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Perazza LR, Gower AC, Brown-Borg HM, Pajevic PD, Thompson LV. Protectin DX as a therapeutic strategy against frailty in mice. *Geroscience*. 2023 Apr 14. doi: 10.1007/s11357-023-00789-3. Epub ahead of print. PMID: 37059838.

WHERE ARE THEY NOW?

ARC launched in 2023

Tuberculosis Interdisciplinary Approach to Research Alliance (TIARA)

Directors: Drs. Karen Jacobson, Charles Horsburgh, and Laura White

(ARC, 2023-present)



SYNOPSIS

(This ARC includes 25 faculty from various disciplines)

Tuberculosis (TB) remains the leading infectious disease killer globally, with an estimated 1.5 million deaths and 10 million new infections annually. In 2020, case notifications plummeted because of the COVID-19 pandemic related disruptions of services, with subsequently the first rise in TB mortality in more than a decade. We are falling gravely short of the WHO's End TB Strategy goal of reducing TB incidence by 80%, TB deaths by 90% and eliminating catastrophic costs for TB-affected households by 2030. It is now clear that



progress towards TB elimination will require focused attention on addressing comorbidities, improving diagnostics and biomarkers of treatment response, and awareness of post-TB sequelae, from understanding biology through potential intervention impacts. The **Boston University Tuberculosis Interdisciplinary Approach to Research Alliance (TIARA) Affinity Research Collaborative (ARC)** has as its mission *the identification and promotion of a cohort of diverse TB clinical and translational research and researchers at Boston University who will accelerate ending the TB epidemic through innovative, interdisciplinary, and creative approaches*. The ARC builds on existing TB core faculty with strong research portfolios, but as of recently with minimal coordinated efforts to advance as consortium interdisciplinary approaches to this research topic.

HIGHLIGHTS

This ARC consists of researchers from Boston Medical Center, Boston University Chobanian & Avedisian School of Medicine, Boston University Sargent College, Boston University National Emerging Infectious Diseases Laboratories, and Boston University School of Public Health. Devoted to the pursuit of translational research to reduce the burden of TB worldwide, there is special focus on key populations including persons who use alcohol and/or illicit drugs, persons with malnutrition and diabetes, children/adolescents, and persons with drug resistant TB; *all reflective of cases in the BMC patient populations*.

EXAMPLE OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Myers B, Carney T, Rooney J, Malatesta S, Ragan EJ, White LF, Natcheva H, Bouton TC, Weber SE, Farhat M, McIlhleron H, Theron D, Parry CDH, Horsburgh CR, Warren RM, Jacobson KR. Smoked drug use in patients with TB is associated with higher bacterial burden. *Int J Tuberc Lung Dis* 2023;27:444-50.

WHERE ARE THEY NOW?

ARC launched in 2023

A Multi-Disciplinary Program to Identify Predictors of Efficacy and Resistance to Cancer Checkpoint Inhibition (PIPER-C)

Directors: Drs. Matthew Kulke, Evan Johnson, David Sherr, and Gerald Denis

(ARC, 2021-present)



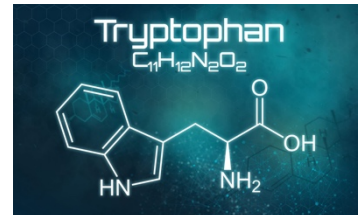
SYNOPSIS

(This ARC includes 26 faculty from various disciplines)

Immune checkpoint inhibitors (ICIs) target inhibitory receptors on T cells and antigen-presenting cells and have transformed cancer treatment. Because the efficacy of immune checkpoint inhibitors is dependent on T cell function, factors that affect the host immune environment can have a significant impact on treatment efficacy. Early studies have shown that levels of immuno-suppressive metabolites, the host microbiome, and tumor-derived exosomes may play a role in mediating the efficacy of checkpoint inhibitors. All these factors can be influenced, either directly or indirectly, by socioeconomic status. These associations, however, are essentially unstudied: only 4% of enrollees in clinical trials of immune checkpoint inhibitors have been from under-represented minority populations. To address this knowledge gap, this broad-based study will investigate associations between circulating metabolites, AhR activity, the microbiome, and tumor-derived exosomes on treatment efficacy in patients at Boston Medical Center receiving treatment with immune checkpoint inhibitors. Our studies will utilize an expanding cancer biobank within the BU-BMC Cancer Center and will bring clinical investigators together with laboratory-based investigators in a broad range of fields.

HIGHLIGHTS

The findings from our study should shed light not only on how specific biomarkers are associated with immune status and treatment outcomes, but also on how these biomarkers may differ in diverse patient populations. As such, our study has broad implications for understanding cancer biology and optimizing cancer treatment.



EXAMPLE OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Based on work done on ARC, we received funding for two new investigator-initiated studies on breast and prostate cancers in collaboration with Bristol Myer Squibb.

- Multidimensional Markers of Response and Resistance in advanced breast cancer: CDK4/6 Inhibitors (HR+/HER2-)
- Multidimensional Markers of Response and Resistance to Therapies in Metastatic Castration Resistant Prostate Cancer

Amr Radwan, Chinmay Jani, ..., Matthew Kulke, Umit Tapan. The association between the development of immune-related adverse events and survival outcomes in a racially diverse patient population. *The Oncologist*. June 2023, under review

WHERE ARE THEY NOW?

ARC Program with newly developed platforms for clinical trials

Respiratory Viruses: A Focus on COVID-19

Directors: Drs. Markus Bosmann and Mohsan Saeed

(ARC, 2020-2023; ARC Program, 2023-present)



SYNOPSIS

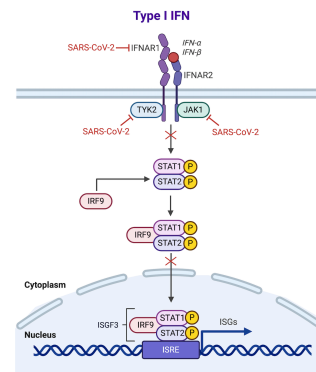
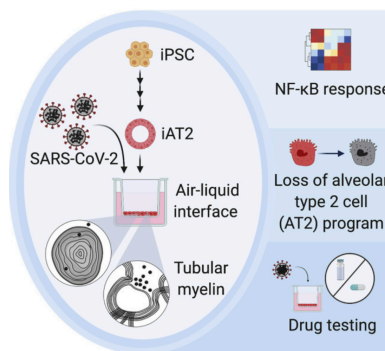
(This ARC includes 46 faculty from various disciplines)

COVID-19 emerged as a new human disease in November 2019 and it swept through the whole world at a lightning speed, claiming over a million human lives and bringing the global economy to a standstill. The purpose of this ARC was to assemble a team of scientists with expertise in various disciplines of biomedical sciences with the goals to (1) generate tools for the investigation of COVID-19, (2) delineate the molecular mechanisms that underlie the COVID-19 pathophysiology, and (3) to develop therapeutic options for the treatment of COVID-19.

HIGHLIGHTS

A collaborative work by several members of this ARC led to the development of a stem cell-derived model of SARS-CoV-2 infection to interrogate the virus-host interface in a physiologically relevant setting. Using this infection model that comprises the lung type II pneumocytes, the team dissected the mechanisms by which SARS-CoV-2 impairs the normal lung functions.

Also, we established a panel of highly infectable human cell lines representing various SARS-CoV-2 target organs, such as lungs, liver, intestine, brain, heart, and kidneys. We used this panel of cell lines to uncover the molecular details of how SARS-CoV-2 blocks antiviral immune signaling and creates a favorable environment for its replication.



TWO EXAMPLES OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Huang J, Hume AJ, ..., Kotton DN. SARS-CoV-2 infection of pluripotent stem cell-derived human lung alveolar type 2 cells elicits a rapid epithelial-intrinsic inflammatory response. *Cell Stem Cell*. 2020 Sep 18.

Chen DY, Chin CV, ..., Saeed M. Spike and nsp6 are key determinants of SARS-CoV-2 Omicron BA.1 attenuation. *Nature*. 2023 Mar;615(7950):143-150. doi: 10.1038/s41586-023-05697-2.

WHERE ARE THEY NOW?

ARC Program focused on researching emerging pathogens

Connecting Tissues and Investigators: Fibrosis ARC

Directors: Drs. Xaralabos Varelas, Maria Trojanowska, and Irving Bigio

(ARC, 2017-2020; ARC Program 2020-Present)



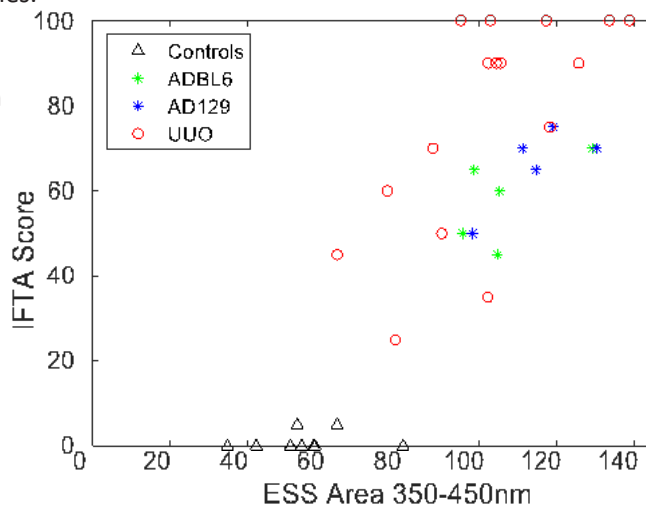
SYNOPSIS

(This ARC includes 26 faculty from various disciplines)

Aberrant fibrogenesis or fibrosis is a common response to chronic tissue injury of almost every organ, contributing to about 45% of all deaths in the U.S. Fibrosis at the phenotypic level shows remarkable similarities across different organ systems, but it is not understood whether cell populations or molecular mechanisms that contribute to the excessive deposition of extracellular matrix are conserved across organs. The central hypothesis is that there are both shared and tissue-specific factors in organ fibrosis that can be utilized to develop improved diagnostics and therapies.

HIGHLIGHTS

- Completed single cell RNA sequencing of fibrosis in a variety of organs, including skin, lungs, adipose tissue, oral cavity, and kidneys.
- Developed new portable spatial-frequency domain imaging to measure collagen content in the skin of scleroderma patients.
- Optimized elastic-scattering spectroscopy to quantify interstitial fibrosis/tubular atrophy in kidney
- Identified candidate metabolites to differentiate between diseased and healthy lungs in preclinical model of pulmonary hypertension.



TWO EXAMPLES OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Mostafa E. Belghasem, Ousama A'amar, Daniel Roth, Joshua Walker, Nkiruka Arinze, Sean M. Richards, Jean M. Francis, David J. Salant, Vipul C. Chitalia, Irving J. Bigio. Towards minimally-invasive, quantitative assessment of chronic kidney disease using optical spectroscopy. *Sci Rep.* 2019; 9: 7168.

Pilvar A, Mehendale AM, Karrobi K, El-Adili F, Bujor A, Roblyer D. Spatial frequency domain imaging for the assessment of scleroderma skin involvement. *Biomed Opt Express.* 2023 May 25;14(6):2955-2968. doi: 10.1364/BOE.489609. PMID: 37342706; PMCID: PMC10278615.

WHERE ARE THEY NOW?

ARC Program in collaboration with BU Center for Multiscale & Translational Mechanobiology

Tobacco Regulatory Sciences

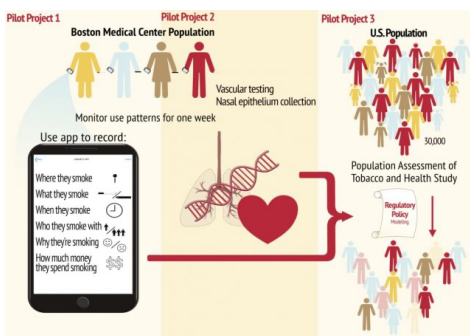
Directors: Drs. Jessica Fetterman, Naomi Hamburg, Andrew Stokes, and Stine Grodal
(ARC, 2017-2020; ARC Program 2020-2022)



SYNOPSIS

(This ARC includes 32 faculty from various disciplines)

Tobacco Regulatory Science (**TRS**) ARC brings together a multidisciplinary investigative team (Pulmonary/Cardiovascular Medicine; Computational Biology; Global Health; Community Health Sciences;



Health Policy & Health Services; Epidemiology; Health Law, Policy, and Management; Biomedical Engineering; Strategy and Innovation; Public Relations, Communication, Computer Sciences) with the mission of understanding complex tobacco use patterns and health impacts in vulnerable populations across the life course.

The safety of e-cigarette use has been called into question with the emergence of EVALI and younger individuals who vape appear to be at greater risk for more severe symptoms and hospitalization with COVID-19 for reasons that are not yet understood. E-cigarettes have been on the market for nearly 10

years but have evolved rapidly, outpacing scientific inquiry into the health effects.

The majority of studies to date evaluating the cardiopulmonary effects of e-cigarettes have focused on e-cigarette products (first- and second- generation devices) no longer in use.

HIGHLIGHTS

This collaboration developed an innovative computer vision approach to detect health warning labels in influencer promotions of cigar products on social media. The presence of a health warning on an Instagram cigar post was associated with lower social media engagement (fewer likes and comments). This study provides support for the implementation of health warning requirements on social media tobacco promotions.

TWO EXAMPLES OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Fetterman JL†, Keith RJ, ..., Hamburg NM. Alterations in vascular function associated with the use of combustible and electronic cigarettes. *JAHA*. 2020. 9(9):e014570.

Wu J, Origgi JM, ..., Fetterman JL. Compliance With the US Food and Drug Administration's Guidelines for Health Warning Labels and Engagement in Little Cigar and Cigarillo Content: Computer Vision Analysis of Instagram Posts. *JMIR Infodemiology*. 2023 Mar 14;3:e41969. doi: 10.2196/41969. PMID: 37113379; PMCID: PMC10132024.

WHERE ARE THEY NOW?

Morphed into an American Heart Association-funded Strategically Focused Research Network

Precision Medicine for Alzheimer Disease & Related Disorders

Directors: Drs. Rhoda Au, Alice Cronin-Golomb, and Lindsay Farrer

(ARC, 2017-2019; ARC Program 2020-Present)



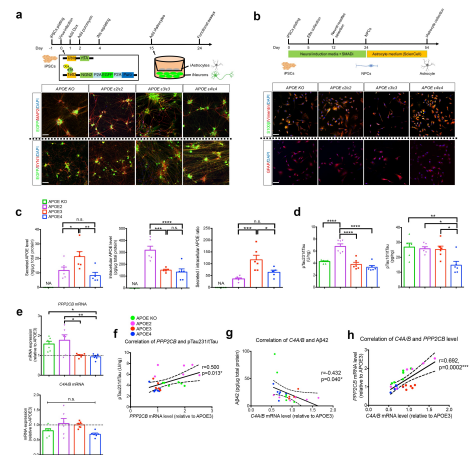
SYNOPSIS

(This ARC includes 33 faculty from various disciplines)

The primary aims of this ARC are to identify subtypes of AD within the Framingham Heart Study (FHS) dataset, validate these subtypes using other available data from the national AD Centers database and other public databases, investigate the biological underpinnings of these subtypes, and identify new therapeutic targets specific for these subtypes. This project introduces a very novel approach for extracting key information from very large and complex datasets that will define subtypes and risk profiles for AD.

HIGHLIGHTS

The mechanism(s) underlying the protective effect of the *APOE* $\epsilon 2$ allele against Alzheimer disease (AD) is poorly understood. To evaluate the contribution of other genetic factors to the protective effect of $\epsilon 2$, we conducted a genome-wide association study for AD among $\epsilon 2$ carriers and applied a systems biology approach involving generating and analyzing various 'omic data and performing immunohistochemistry experiments in brain tissue from 761 AD cases and controls, as well as validation experiments in *APOE* allele-specific iPS cells. Collectively, our findings demonstrated for the first time a molecular link between a tau phosphatase and the classical complement pathway, especially C4, and AD-related tau pathology.



TWO EXAMPLES OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Qiu S, Miller MI,..., Kolachalama VB. Multimodal deep learning for Alzheimer's disease dementia assessment. Nat Commun. 2022 Jun 20;13(1):3404. doi: 10.1038/s41467-022-31037-5. PMID: 35725739; PMCID: PMC9209452.

Zhang X, Tong T,..., Au R, Farrer LA. Midlife measures of high-density lipoprotein and fasting glucose are associated with late-onset Alzheimer disease. Alzheimer Dement. 2023; 19:181-193. PMID: 35319157

WHERE ARE THEY NOW?

ARC Program with NIH-funded U19 center of excellence

Systems Biology Approaches to Microbiome Research

Directors: Drs. Daniel Segrè and W. Evan Johnson

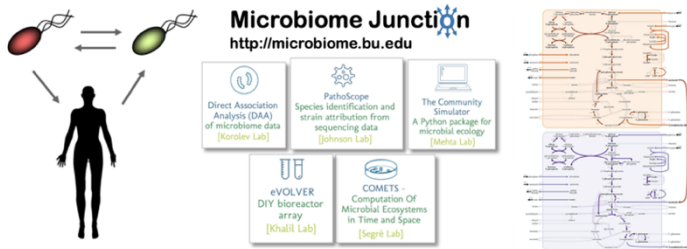
(ARC, 2017-2019; ARC Program 2019; Formalized BU-supported Microbiome Program, 2019)



SYNOPSIS

(This ARC includes 47 faculty from various disciplines)

Microbial communities play a crucial role in the health of plants, animals, and humans, and of marine and terrestrial ecosystems. Understanding these communities can have great impact in many areas, including agriculture and food production, climate change, immune system function and infectious disease. The goal of our ARC is to develop a new, multi-level mechanistic understanding of how microbe-microbe, microbe-environment, and microbe-host interactions determine microbial community dynamics, diversity and stability, and use this knowledge to understand how to control and engineer microbial communities for defined purposes.



HIGHLIGHTS

We have made significant progress towards the construction of the Microbiome Junction, a hub of computational tools to help conduct quantitative microbiome research. This serves as the point of convergence

of multiple data types generated by different investigators, helping them interpret the data in the form of microbial and host interaction networks. We are actively applying these methods and tools in multiple contexts, including cancer (lung, pancreatic, immunotherapies, etc) and infectious diseases (HIV, TB, etc).

TWO EXAMPLES OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Faits T, Walker ME, ... , Matthan NR. Exploring changes in the human gut microbiota and microbial-derived metabolites in response to diets enriched in simple, refined, or unrefined carbohydrate-containing foods: a post hoc analysis of a randomized clinical trial. *Am J Clin Nutr.* 2020 Sep 16.

Yue Zhao, Anthony Federico, Tyler Faits, Solaiappan Manimaran, Daniel Segrè, Stefano Monti and W. Evan Johnson: animalcules: interactive microbiome analytics and visualization in R, *Microbiome* (2021) 9, 76.

WHERE ARE THEY NOW?

BU-supported Program on System Biology Approaches to Microbiome Research: Boston University Microbiome Initiative
[<https://www.microbu.org/>]

Mobile and Electronic (ME) Health

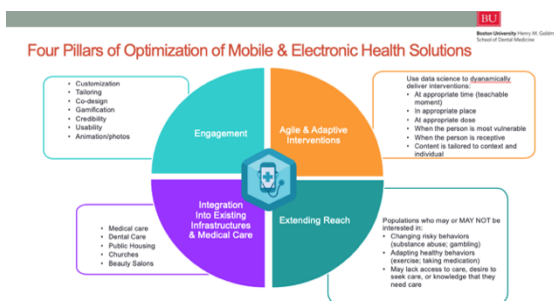
Director: Dr. Belinda Borrelli & Co-Directors: Drs. Joanne Murabito and Daniel Fulford
(with Dr. Tibor Palfai, thru 2022 and Dr. Lisa Quintiliani, thru 2023)
(ARC, 2015-2019; ARC Program, 2020-Present)



SYNOPSIS

(This ARC includes 175 faculty from various disciplines)

The mission the ME-ARC is to conduct research and training on mobile health with an emphasis on underserved populations and transdisciplinary research. Since, 2017, the ME-ARC has held monthly seminars, annual symposiums, reduced digital silos at BU, and provided mentorship to post-docs, K awardees, and consultation to BU researchers. The ME-ARC has received external funding across a wide variety of areas (e.g., smoking, diet, physical activity, obesity, alcohol, oral health) and digital platforms (Virtual Reality, mobile apps, web, social media, text messaging, connected devices) that have been deployed in real-world clinical and public health settings. The ME-ARC has built strong collaborations with other BU groups (Tobacco-ARC and Framingham Heart Study) resulting in additional publications and external grant award funding. In addition to the leadership team above, the ME-ARC has a steering committee, external advisory board, trainees, and >176 BU members.



HIGHLIGHTS

Dr. Borrelli received a \$4.2M five-year award from the NIH that resulted from one of the ME-ARC pilots, entitled, “Delivery of a Smoking Cessation Induction Intervention Via Virtual Reality Headset During a Dental Cleaning.” She also received a \$1.8M award from the American Heart Association to study cessation of adolescent vaping using Virtual Reality and texting. This project is part of a multi-project center grant led by Naomi Hamburg of the Tobacco-ARC. In addition, Dr. Lisa Quintiliani was awarded an NIH R01 to develop and test a weight management intervention for public housing residents that uses text messaging and connected devices. ME-ARC members are Co-Is on all the above projects.

TWO EXAMPLES OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Quintiliani, LM, Foster, M. Oshry, LJ. Preferences of mHealth app features for weight management among breast cancer survivors from underserved populations. *Psycho-Oncology*. 2019. 28(10):2101-2104.

Shankar, D., Borrelli, B., Cobb, V., Quintiliani, L.M., Palfai, T., Weinstein, Z., Bulekova, K., & Kathuria, H. (2022). Text-messaging to promote smoking cessation among individuals with opioid use disorder: quantitative and qualitative evaluation. *BMC Public Health*, 22(1), 1-12. PMID: 35387648

WHERE ARE THEY NOW?

CTSI and university Program focused developing digital solution to health-inequities.

Thrombosis and Hemostasis in Health and Disease

Directors: Drs. Vipul Chitalia, Katya Ravid, and Jean Francis

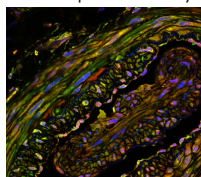
(ARC, 2015-2019; ARC Program, 2020-present)



SYNOPSIS

(This ARC includes 34 faculty from various disciplines)

The Thrombosis and Hemostasis in Health and Disease ARC had evaluated several facets of thrombosis spanning from novel mediators to the generation of the animal model up to validating the hypotheses in human cohorts. This ARC has investigated thrombosis using emerging techniques such as machine-learning techniques in the context of organ pathologies (e.g., renal failure), infectious disease (e.g., Shigella-Toxin mediated) and Thrombotic Microangiopathy Collaborative (TMA), a highly complex disease mediated primarily by the aberrant hyperactivation of complement system. The ARC established the first Boston University TMA initiative with BU CTSI.



Vasa vasorum: tissue factor (red) and its cognate ubiquitin ligase, STUB1 (green); On the Cover of *Science Translational Medicine*

HIGHLIGHTS

This ARC has successfully brought together basic and translational research through a cross-disciplinary approach to gain deeper understanding of thrombosis pathogenesis and to develop novel personalized predictive tools and therapeutic targets. As reported earlier, ARC members have initiated and developed the first Thrombotic Microangiopathy (TMA) program in BUSM. An additional unique, innovative contribution of the TtoH ARC has been a focus on *organ pathology-induced changes in thrombosis and vascular hemostasis*, which in the past year has been expanded to initiate studies related to mechanisms of cancer-induced thrombosis. Thrombotic mechanisms related to chronic kidney disease and bone marrow myelofibrosis have been probed, both used as model systems to examine *this newly studied paradigm*, resulting in notable publications and new grant support. Our collaborative pursuits led to identifying certain human plasma metabolites retained in chronic kidney disease, such as Kynurenine, as thrombogenic. Further, the secreted enzyme in myelofibrosis, lysyl oxidase, was found to enhance platelet activation and thrombosis in vivo in mice, and more recently new collaborative proteomic approaches led to identifying in bone marrow malignant cells modified integrins known to participate in promoting thrombosis.

TWO EXAMPLES OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Matsuura S, Thompson CR, Belghasem ME, Bekendam RH, Piasecki A, Leiva O, Ray A, Italiano J, Yang M, Merrill-Skoloff G, Chitalia VC, Flaumenhaft R, Ravid K, Platelet Dysfunction and Thrombosis in JAK2V617F-Mutated Primary Myelofibrotic Mice. *Arterioscler Thromb Vasc Biol.* 2020 Oct;40(10):e262-e272.

Ravid JD, Leiva O, Chitalia VC. Janus Kinase Signaling Pathway and Its Role in COVID-19 Inflammatory, Vascular, and Thrombotic Manifestations, *Cells*, 2022 Jan 17;11(2):306.PMID: 35053424, PMCID: PMC8773838

WHERE ARE THEY

Morphed into an American Heart Association-funded Strategically Focused Research Network (Cardio-Oncology)

Protein Trafficking and Neurodegenerative Disease

Director: Dr. Lindsay Farrer

(ARC, 2009-2012; ARC Program 2013-2021)

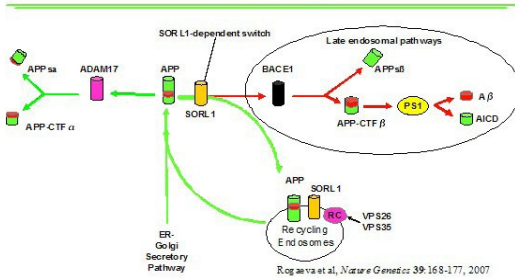


SYNOPSIS

(This ARC includes 20 faculty from various disciplines)

Recycling of the amyloid precursor protein (APP) from the cell surface via the endocytic pathways plays a key role in the generation of amyloid β -peptide ($A\beta$), the accumulation of which is thought central to the pathogenesis of Alzheimer disease (AD).

SORL1 is a sorting receptor for APP



This ARC explores the role of vesicular sorting proteins and other genes involved in protein trafficking in the etiology and pathophysiology of AD and other neurodegenerative disorders. The power of this ARC lies in its diverse, interdisciplinary expertise, and ability to validate any finding using independent approaches of genetic epidemiology, cell biology, model systems and pathology.

HIGHLIGHTS

Notably, cognitive decline has been observed disproportionately in persons affected with MS and to a lesser extent with celiac disease, suggesting that these three diseases share a common immune or inflammatory pathway. Future studies of the protective mechanism of these *HLA* alleles may provide insight into the neurobiology of cognitive decline and dementia and novel therapeutic approaches for these disorders.

TWO EXAMPLES OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Li M, Reisman J, Morris-Eppolito B, Qian SX, Kazis LE, Wolozin B, Goldstein LE, Xia W. Beneficial association of angiotensin-converting enzyme inhibitors and statins on the occurrence of possible Alzheimer's disease after traumatic brain injury. *Alzheimers Res Ther.* 2020; 12(1):33.

Panitch R, Hu J, Xia W, Bennett DA, Stein TD, Farrer LA, Jun GR. Blood and brain transcriptome analysis reveals APOE genotype-mediated and immune-related pathways involved in Alzheimer disease. *Alzheimers Res Ther.* 2022 Feb 9;14(1):30. doi: 10.1186/s13195-022-00975-z. PMID: 35139885; PMCID: PMC8830081.

WHERE ARE THEY NOW?

ARC Program with platforms for researching genetic biomarkers

Mitochondrial Dynamics in Health and Disease (mtARC)

Directors: Drs. Orian Shirihai and Andrea Havasi

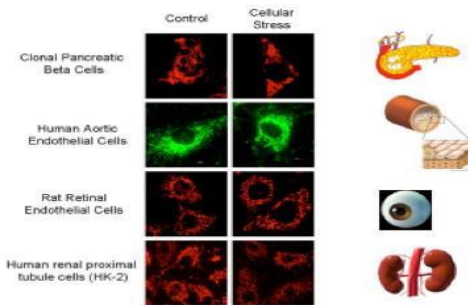
(ARC, 2009-2012; ARC Program, 2013-2016; The ARC gave rise to a Bioenergetics Core)



SYNOPSIS

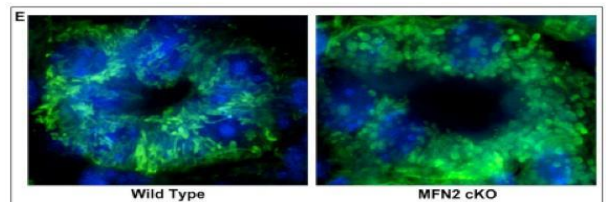
(This ARC included 28 faculty from various disciplines)

The mtARC focused on the role of mitochondria in physiology and pathophysiology. Diverse membership encompassed 12 sections/departments with representation from both the Medical and Charles River Campuses. In addition, membership included labs from other universities and from industry. This ARC also provided training and tools for investigating mitochondrial bioenergetics, dynamics and reactive oxygen species (ROS) in various systems.



Data from the labs of Drs Shirihai, Vita, Roy & Borkan

Mitochondrial morphology in Mfn2 conditional KO mice



HIGHLIGHTS

The mitochondrial ARC collaboratively discovered a role for mitofusin 2 (Mfn2) and mitochondrial dynamics in neurodegeneration and metabolic diseases in different organ systems and was shown to have an important role in organ survival after ischemia. This ARC gave rise to an institutional Research Core focused on Bioenergetics.

EXAMPLE OF A COLLABORATIVE DISCOVERY AMONG SEVERAL

Liesa M, Luptak I, Qin F, Hyde BB, Sahin E, Siwik DA, Zhu Z, Pimentel DR, Xu XJ, Ruderman NB, Huffman KD, Doctrow SR, Richey L, Colucci WS, Shirihai OS. Mitochondrial transporter ATP binding cassette mitochondrial erythroid is a novel gene required for cardiac recovery after ischemia/reperfusion. *Circulation*. 2011 Aug16;124(7):806-13.doi: 10.1161/ CIRCULATIONAHA.110.003418. Epub 2011 Jul 25.

WHERE ARE THEY NOW?

Leading faculty recruited to chair a department at UCLA, where ARC principles were implemented

BU iPS Cell Bank

Directors: Drs. Darrell Kotton, Gustavo Mostoslavsky, and George Murphy

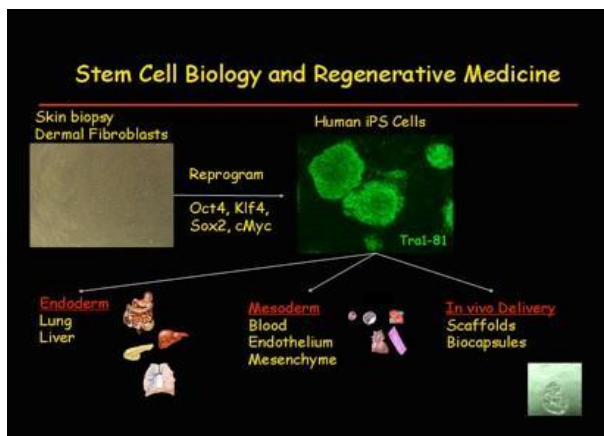
(ARC, 2009-2012; ARC Program, 2013-2014; CReM (2014-Present))



SYNOPSIS

(This ARC included 15 faculty from various disciplines)

The discovery that mature somatic cells can be reprogrammed into induced pluripotent stem (iPS) cells provides unprecedented opportunities to better understand and treat a variety of human diseases.



ARC members developed in the past a novel reagent, the lentiviral 'stem cell cassette' vector (STEMCCA) that allows the most efficient generation of iPS cells.

This ARC utilized this vector to develop an extensive bank of human iPS cells from individuals with diseases commonly found in the Boston Medical Center patient population: Sickle Cell Anemia, Amyloidosis, Emphysema, Inflammatory Bowel Disease, Scleroderma, and Diabetes.

HIGHLIGHTS

The iPS Cell Bank, Regenerative Medicine ARC initiative, among the first ARCs to be awarded at Boston University, facilitated the launch of an Open-Source stem cell bank that has grown over the years into one of the most widely shared NIH-sponsored stem cell repositories in the US. Today, this repository continues to serve as the heart of a new ARC-facilitated Center for Regenerative Medicine and has become the largest bank of disease-specific induced pluripotent stem (iPS) cells for studying sickle cell disease, amyloidosis, and genetic lung diseases including alpha-1 antitrypsin deficiency, cystic fibrosis, and a wide variety of genetic neonatal lung disorders.

EXAMPLE OF A COLLABORATIVE DISCOVERY AMONG SEVERAL

Somers A1, Jean JC, ..., Kotton DN. Generation of transgene-free lung disease-specific human induced pluripotent stem cells using a single excisable lentiviral stem cell cassette. *Stem Cells*. 2010 Oct. 28 (10):1728-40.

WHERE ARE THEY NOW?

Launched a BU center for Regenerative Medicine (CReM)

Sex Differences in Adipose Tissue Biology and Related Metabolic Disease

Directors: Drs. Susan K. Fried and Paul Pilch

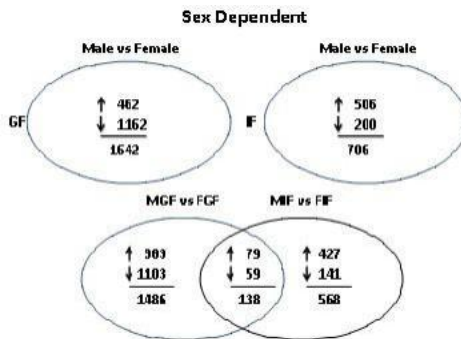
(ARC, 2009-2012; ARC Program, 2013-2015; continued BNORC collaboration)



SYNOPSIS

(This ARC included 21 faculty from various disciplines)

This ARC focused on the role of adipose tissue biology in the metabolic complications of obesity in men and women. Members have complementary expertise in biochemistry, cell biology, immunology and translational research in obesity, diabetes, and cardiovascular disease. This ARC had a close relationship to an existing NIH-funded center, the Boston Nutrition and Obesity Research Center, and has initiated novel collaborations resulting in interdisciplinary grant proposals, including 'brite' adipocytes, adiporedoxin, a novel adipocyte dysfunction in insulin resistance. The ARC also engaged in fruitful ARC-ARC collaborations with four other ARCs: Mitochondria, Arterial Stiffness, Cancer and Inflammation, and Metabolic Disease and Insulin Resistance: Studies in Patients Undergoing Bariatric Surgery.



HIGHLIGHTS

The ARC team found that glucocorticoids (GC) have profound effects on adipose tissue, adipogenesis and adipose tissue metabolic and endocrine function. With chronic excess GC, produced systemically or through local adipose tissue conversion, fat accumulates in central adipose depots and contributes to metabolic derangements.

EXAMPLE OF A COLLABORATIVE DISCOVERY AMONG SEVERAL

Jedrychowski MP, Liu L, Laflamme CJ, Karastergiou K, Meshulam T, Ding SY, Wu Y, Lee MJ, Gygi SP, Fried SK, Pilch PF. Adiporedoxin, an upstream regulator of ER oxidative folding and protein secretion in adipocytes. *Mol Metab*. 2015 Sep 18;4(11):758-70.

WHERE ARE THEY NOW?

Leading faculty recruited to build program at Columbia University, where collaborative ARC principles are implemented

Cardiovascular Consequences of Metabolic Disease

Director: Dr. Kenneth Walsh

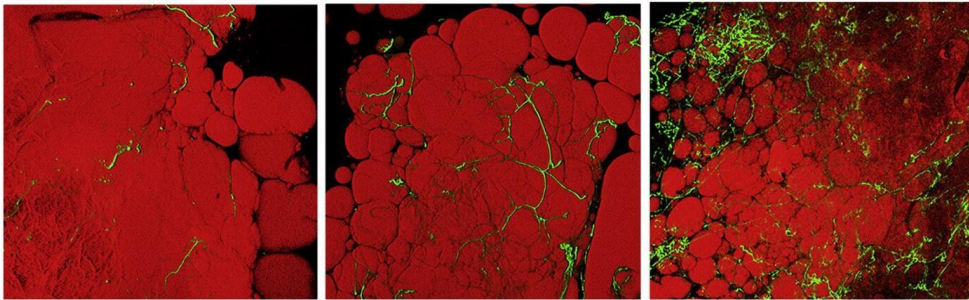
(ARC, 2009-2010; Successful completion with achievement of an NHLBI-funded PPG)



SYNOPSIS

(This ARC included 14 faculty from various disciplines)

Obesity promotes a chronic inflammatory state, which contributes to the development of insulin resistance and cardiovascular disease. This ARC intended to promote translational research by bringing basic and clinical research laboratories together to study the cardiovascular consequences of obesity and diabetes. Activities of basic science labs and clinical research labs were coordinated to assess the interrelationship between metabolic dysfunction and vascular disease.



Angiogenesis in adipose tissue (Walsh and Aprahamian et al.)

HIGHLIGHTS

The ARC identified and studied the role of angiogenesis in adipose tissue biology, with focus on fat cell expansion.



EXAMPLE OF A COLLABORATIVE DISCOVERY AMONG SEVERAL

Bretón-Romero R1, Feng B1, ..., Hamburg NM2. Endothelial Dysfunction in Human Diabetes Is Mediated by Wnt5a-JNK Signaling. *Arterioscler Thromb Vasc Biol.* 2016 Mar;36(3):561-9.

WHERE ARE THEY NOW?

Launched an NIH-funded Program Project before the ARC leader was recruited to lead a section at the University of North Carolina at Chapel Hill

Biomarkers of Disease

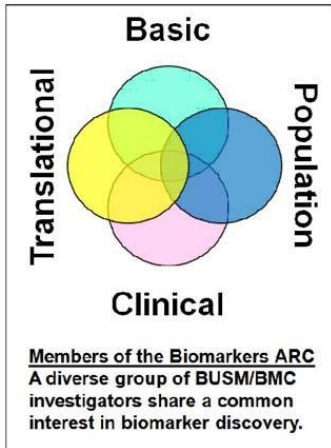
Directors: Drs. Mark McComb, Richard A. Cohen, and Catherine Costello

(ARC, 2009-2012; ARC Directors re-directed efforts into a newly funded NIH grant to support a National Proteomic Center at BU)



SYNOPSIS

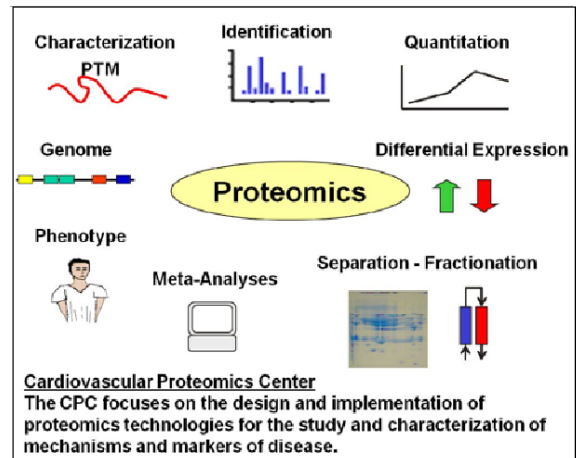
(This ARC included 17 faculty from various disciplines)



Projects designed by this ARC were to build on existing expertise and to take advantage of preliminary research studies performed at BUMC in transgenic mice and animal models of metabolic disease. A discovery-based proteomics approach was designed to identify candidate biomarkers related to metabolic disease. The studied hypothesis was that metabolic changes in detected by changes in plasma protein abundances that betray diseased tissues may be detected by changes in plasma protein abundances that betray leakage of tissue-specific proteins, and bypost-translational modifications (PTMs) that reflect abnormal tissue metabolism.

HIGHLIGHTS

This ARC team discovered changes in metabolic function that may be maladaptive, leading to a situation in which substrate supplied to the heart for energy generation is adequate in amount but cannot be fully utilized. They provided a model to offer specific molecular insights regarding the cellular and physiological mechanisms that lead to metabolic heart disease.



TWO EXAMPLES OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Papers that received support from the COVID ARC

Amraei, R., Yin, W., ... Costello, C. E.; Rahimi, N. CD209L/L-SIGN and CD209/DC-SIGN Act as Receptors for SARS-CoV-2. *ACS Cent. Sci.*, 2021, 7, 1156-1165 (Epub 2021 Jun 30). [cited approx. 300 times]

Amraei, R., Xia, C., ..., Costello, C. E.; Rahimi, N. Extracellular Vimentin is an Attachment Factor that Facilitates SARS-CoV-2 Entry into Human Endothelial Cells. *Proc. Natl Acad. Sci. USA*, 2022, 119(6):e2113874119. doi: 10.1073/pnas.2113874119. (Epub 2022 Jan 25). [cited more than 50 times]

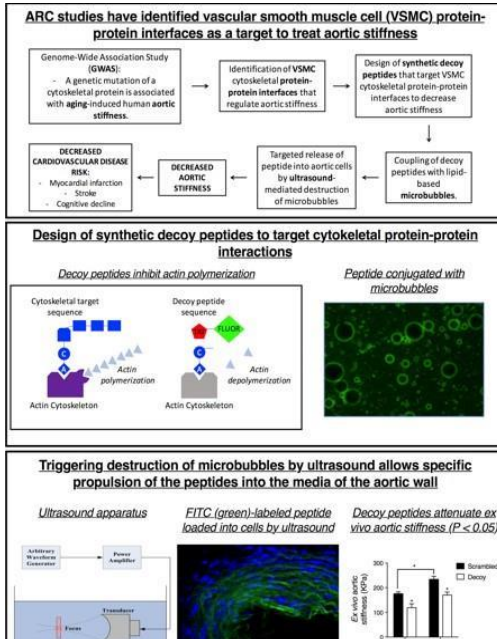
WHERE ARE THEY NOW?

Launched an NIH-funded Program Project and later fused with BUSM Proteomic Center

Molecular, Biomechanical and Genetic Mechanisms of Arterial Stiffness

Directors: Drs. Richard Cohen, Kathleen Morgan, and Francesca Seta

(ARC, 2010-2013; ARC Program, 2014-2018; Arterial Function Core)



SYNOPSIS

(This ARC included 16 faculty from various disciplines)

Arterial stiffness, a vascular condition characterized by loss of elastic compliance of large arteries, is an independent predictor of, and probable cause of, subsequent adverse cardiovascular events. Targeting arterial stiffness could represent a novel approach to decrease the risk of developing cardiovascular diseases, which remain the major cause of mortality and morbidity in US. The Arterial Stiffness ARC was conceived as an interdisciplinary collaborative group of basic scientists, epidemiologists, and bioengineers, from BUSM/BMC, The Framingham Heart Study and CRC, to tackle the complex question of what causes arterial stiffness with the common goal of identifying therapeutic targets.

HIGHLIGHTS

This ARC team-initiated efforts to identify therapeutic target molecules to prevent or reverse aortic stiffness, discovered that activation of the lysine deacetylase sirtuin-1 has potent anti-inflammatory and antioxidant effects on the vascular wall decreasing arterial stiffness in a model of diet-induced obesity, and successfully applied GWAS data to identify candidate molecules to develop cell permeant, microbubble targeted decoy peptides that are effective in decreasing aortic stiffness in an aged rodent model.

EXAMPLE OF A COLLABORATIVE DISCOVERY AMONG SEVERAL

Nicholson CJ, Singh K, ..., Morgan KG. Reversal of Aging-Induced Increases in Aortic Stiffness by Targeting Cytoskeletal Protein-Protein Interfaces. J Am Heart Assoc. 2018 Jul 18;7(15):e008926. doi: 10.1161/JAHA.118.008926. PMID: 30021807; PMCID: PMC6201469.

WHERE ARE THEY NOW?

Gave rise to a research core focused on Arterial Stiffness

Calcium Homeostasis in Health and Disease

Directors: Drs. Victoria Bolotina and Michael Kirber

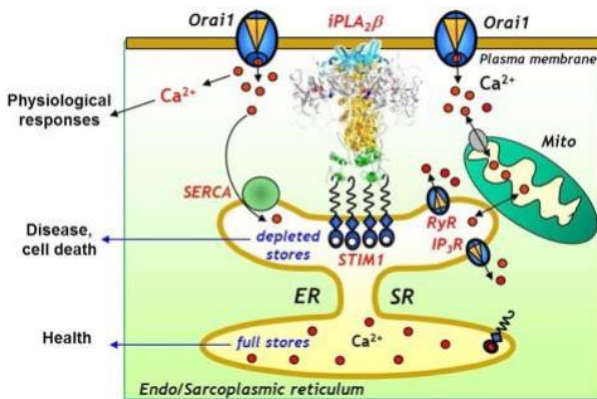
(ARC, 2010-2013; ARC Program, 2014-2015)



SYNOPSIS

(This ARC included 26 faculty from various disciplines)

Molecular determinants of ER / SR Ca^{2+} homeostasis



Comprised of experts in complementary fields from 18 laboratories in 15 Sections and Departments from Boston University Medical and Charles River campuses, this ARC looked at the mechanisms of cellular function from many different perspectives, which enabled the group to identify common molecular determinants and mechanisms of Ca^{2+} signaling in diverse cell types, and allowed their translation to human disease to address the mechanisms of impairment in Ca^{2+} homeostasis in cardiovascular, neurological, pancreatic, and other systems.

HIGHLIGHTS

Calcium ARC supported a ground-breaking discovery of a novel Ca^{2+} signaling mechanism of Parkinson's disease and has provided school-wide expertise in the study of calcium homeostasis.

EXAMPLE OF A COLLABORATIVE DISCOVERY AMONG SEVERAL

Zhou Q, Yen A, Rymarczyk G, Asai H, Trengrove C, Aziz N, Kirber MT, Mostoslavsky G, Ikezu T, Wolozin B, Bolotina VM. Impairment of PARK14-dependent Ca^{2+} signaling is a novel determinant of Parkinson's disease. Nat Commun. 2016 Jan 12; 7:10332.

WHERE ARE THEY NOW?

Leading faculty developed a Biotech Start up focused on Calcium Signaling in Parkinson Disease

Obesity, Inflammation and Cancer

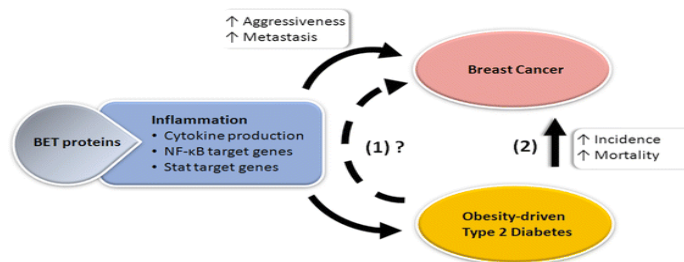
Directors: Drs. Gerald V. Denis and Barbara Nikolajczyk
(ARC, 2010-2013)



SYNOPSIS

(This ARC included 9 faculty from various disciplines)

Population studies identify cohorts of high body mass index (BMI) subjects with unexpectedly reduced risk for breast and colon cancer, and normal BMI subjects with unexpectedly elevated risk for breast cancer, provoking hard thinking about cellular and molecular mechanisms that most strongly couple obesity to cancer occurrence or progression. Emerging work suggests that abnormal metabolism and its associated chronic inflammation make the difference.



Nichols et al, *Cell Molecular Life Science* **74**(2): 231-233, 2017

HIGHLIGHTS

Discoveries made by this ARC support a new hypothesis: metabolic disease in obesity promotes breast cancer incidence and metastasis.



EXAMPLE OF A COLLABORATIVE DISCOVERY AMONG SEVERAL

Nicholas DA1,2, Andrieu G1, Strissel KJ1, Nikolajczyk BS2, Denis GV3,4. BET bromodomain proteins and epigenetic regulation of inflammation: implications for type 2 diabetes and breast cancer. *Cell Mol Life Sci.* 2017 Jan;74(2):231-243.

WHERE ARE THEY NOW?

ARC Projects and members joined BUSM Cancer Center

Nanotheranostics

Directors: Drs. Victoria Herrera, Mark Grinstaff, Joyce Wong, and Karl Karlson

(ARC, 2012-2015; ARC Program, 2015-2017 giving rise to Master of Science and research programs in Nanomedicine)

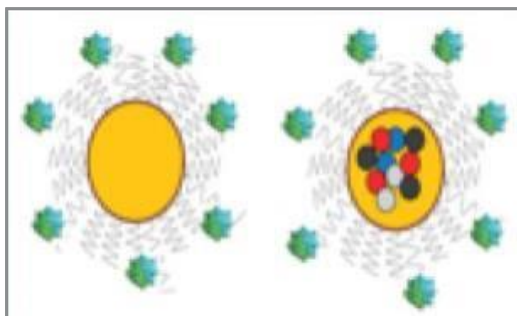


SYNOPSIS

(This ARC included 26 faculty from various disciplines)

Nanotheranostics is defined as the integrated combination of target-specific diagnostics and delivery of therapeutics based on nanotechnology platforms. The ARC-program has provided the opportunity to utilize prototype nanotechnology platforms and proof of concept imaging formats.

Operationalization of the experiments to test the diagnostics, therapeutic and intraoperative survival visualization and resection required the inter-disciplinary expertise and capabilities of several departments: Chemistry, Biomedical Engineering, Molecular Medicine, Pathology, and Surgery.



HIGHLIGHTS

The ARC discovered and proved that low pH-responsive expansile nanoparticles (eNPs) home to and selectively localize intracellularly in pancreatic cancer cells in xenograft model of pancreatic peritoneal carcinomatosis, while sparing peri-tumoral normal cells – endothelium, mesothelium, adipocytes, pancreatic tissue, and all normal intra-abdominal organs including the liver and spleen where all other nanoparticles typically end-up by default.

EXAMPLE OF COLLABORATIVE DISCOVERY AMONG SEVERAL

Whitaker, R.D., Decano, J.L., Gormley, C., Beigie, C.A., Meisel, C., Tan, G.A., Moran, A.M., Giordano, N.J., Park, Y., Huang, P., Andersson, S., Gantz, D., Grant, A.K., Ruiz-Opazo, N., Herrera, V.L.M., Wong, J.Y. (2022). Janus USPIO modular platform (JUMP) for theranostic ultrasound-mediated targeted intratumoral microvascular imaging and DNA/miRNA delivery. *Theranostics*, 12(18), 7646-7667. <https://doi.org/10.7150/thno.78454>.

WHERE ARE THEY NOW?

A formalized new BU Program in Nanomedicine, hosting a MS of Sciences program on the topic

Metabolic Diseases and Insulin Resistance: Studies in Patients Undergoing Bariatric Surgery

Directors: Drs. Neil Ruderman, Konstantin Kandror, and Caroline Apovian

(ARC, 2012-2014; 2015-on, members joined other metabolic-related ARCs)



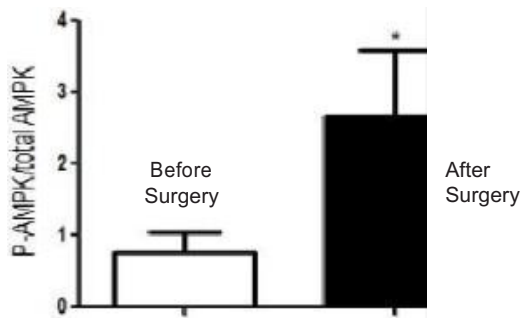
SYNOPSIS

(This ARC included 4 faculty from various disciplines)

Morbidly obese individuals are predisposed to a wide range of diseases including type 2 diabetes, atherosclerotic cardiovascular disease (ASCVD), fatty liver disease, and certain cancers, all of which can be improved or prevented by bariatric surgery. This ARC team along with a few other groups, have observed that approximately 25% of morbidly obese individuals are insulin sensitive (IS) and the remainder are insulin resistant (IR). Intriguingly, compared to the IR group, the IS patients are less likely to develop ASCVD, and presumably other obesity- associated comorbidities.

HIGHLIGHTS

The ARC found that the activity of fuel sensing enzyme AMP-activated protein kinase (P-AMPK) in adipose tissue is markedly diminished (70%) in the insulin resistant patients at the time of surgery. Likewise, 3-months post-operatively decreased AMPK activity was eliminated.



Comparison of AMPK phosphorylation in the subcutaneous fat of 11 matched pre- and post-bariatric surgery patients(*p<0.05).

EXAMPLE OF A COLLABORATIVE DISCOVERY AMONG SEVERAL

Xu XJ, Apovian C, Hess D, Carmine B, Saha A, Ruderman N. Improved Insulin Sensitivity 3 Months After RYGB Surgery Is Associated With Increased Subcutaneous Adipose Tissue AMPK Activity and Decreased Oxidative Stress. Diabetes. 2015 Sep;64(9):3155-9.

WHERE ARE THEY NOW?

Lead faculty retired; members joined other ARCs

Computational Genomics Models of Environmental Chemical Carcinogenicity

Directors: Drs. Stefano Monti and David Sherr

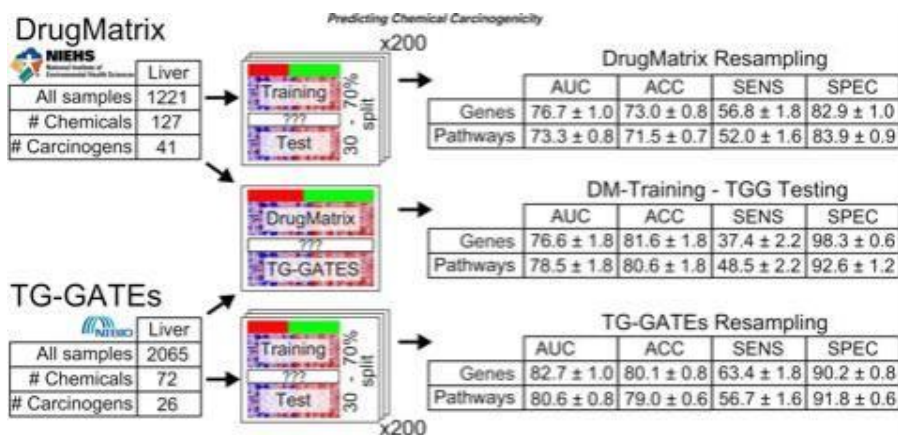
(ARC, 2013-2016; ongoing discussions related to a new Environmental Toxicants Program)



SYNOPSIS

(This ARC included 12 faculty from various disciplines)

This project was centered around the over-arching goal of developing genomic models of carcinogenicity for cancer prevention and tailored treatment, the goal being to develop accurate and cost-effective methods for the identification of threats to our health from exposure to chemical and environmental carcinogens. An essential component of this ARC is the research into development and application of novel computational approaches to the analysis and integration of multi- dimensional, multi-omics data.



HIGHLIGHTS

The ARC team confirmed the hypothesis that the genomic profile of human cells generated in response to short term exposure with environmental chemicals in vitro can predict, with up to 83% accuracy, a long-term biologic consequence, in vivo cancer development. This discovery paves the way for predicting which of the 85,000 chemicals in consumer and industrial use are human carcinogens. To date, only ~2.5% of those chemicals have been tested for carcinogenicity.

EXAMPLE OF A COLLABORATIVE DISCOVERY AMONG SEVERAL

Wang Z, Snyder M, Kenison JE, Yang K, Lara B, Lydell E, Bennani K, Novikov O, Federico A, Monti S, Sherr DH. How the AHR Became Important in Cancer: The Role of Chronically Active AHR in Cancer Aggression. Int J Mol Sci. 2020 Dec 31;22(1):387. doi: 10.3390/ijms22010387. PMID: 33396563; PMCID: PMC7795223.

WHERE ARE THEY NOW?

Faculty and related research contributed to founding a new Computational Biomedicine Section

Etiology & Pathogenesis of Oral Cancer

Directors: Drs. Maria Kukuruzinska, Maria Trojanowska and Avrum Spira

(ARC, 2014-2017; ARC Program, 2017-2019)

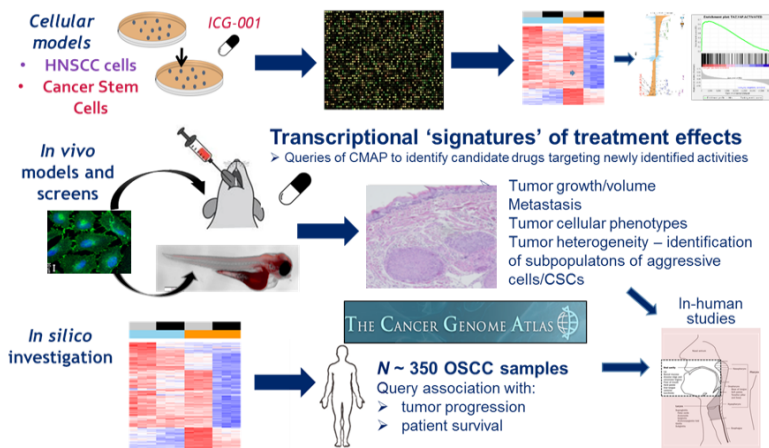


SYNOPSIS

(This ARC included 14 faculty from various disciplines)

Studies conducted by the Etiology & Pathogenesis of Oral Cancer (EPOC) ARC have been CON ducted by an array of faculty from different schools, aiming at developing new pathways and understanding of this pathology.

Identifying New Druggable Targets and Treatments for Head and Neck Cancer



HIGHLIGHTS

EPOC activities have led to significant new findings related to: 1) the mechanisms of oral squamous cell carcinoma (OSCC) development and progression with a focus on the early pathways involving the N-glycosylation/ β -catenin/TAZ-YAP signaling axis, as well as the role of aryl hydrocarbon receptor in tumor initiation in collaboration with the oral microbiome; 2) the remodeling and activation of OSCC tumor stroma and its role in cancer progression; and 3) personalized early detection and treatment of oral cancer, with emphasis on early detection and amelioration of radiation treatment-induced fibrosis.

EXAMPLE OF A COLLABORATIVE DISCOVERY AMONG SEVERAL

Reed, E.R., Jankowski, S.A., Spinella, A.J., Noonan, V., Nomoto, K., Matsui, J., Varelas, X.*, Kukuruzinska, M.A.*, and Monti, S.* (2023). Multi-resolution scRNAseq Subtyping Reveals that β -catenin/CBP Activation of mTORC1 Signaling Promotes Partial EMT Cellular States in Head and Neck Cancer. Translational Research *Corresponding authors. June 21, 2023. Epub ahead of print.

WHERE ARE THEY NOW?

Formalized Cross-disciplinary Program focused on Head and Neck Cancer

PneumoniARC

Director: Dr. Joseph 'Jay' Mizgerd

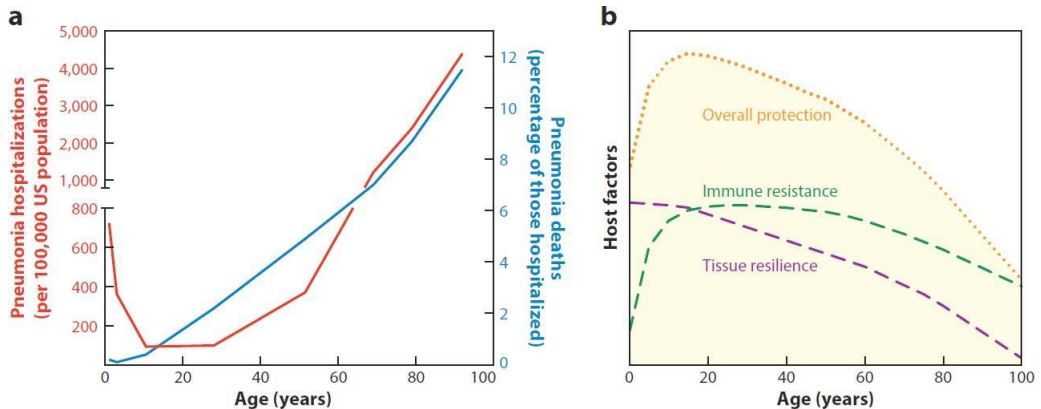
(ARC, 2015-2017)



SYNOPSIS

(This ARC included 9 faculty from various disciplines)

A multidisciplinary team was gathered, integrating critical distinct perspectives into this complex disease process, including the basic biology disciplines of Lung Biology, Immunology, and Microbiology, combined with clinical research realms from those caring for children (Pediatrics) or older adults (Pulmonary & Critical Care Medicine), to form the PneumoniARC. In order to attack the problem of high-susceptibility to pneumonia, this ARC is coordinating the knowledge bases, research tools, and investigational activities of 9 principal investigators from 4 academic units across 6 different buildings, all of whom have studies that are complementary with others in the group and relevant to pneumonia.



From Annual Rev Physiol, 2015

HIGHLIGHTS

The ARC identified the B cell repertoire as a window into the nature and impact of the lung “Virome.”

EXAMPLE OF A COLLABORATIVE DISCOVERY AMONG SEVERAL

Barker KA, Etesami NS, ..., Mizgerd JP. Lung-resident memory B cells protect against bacterial pneumonia. J Clin Invest. 2021 Jun 1;131(11):e141810. doi: 10.1172/JCI141810.PMID: 34060477

WHERE ARE THEY NOW?

ARC members contributed to rapid development of a timely (2020) ARC on *Respiratory Viruses: A focus on COVID-19*

Biomedical Innovation Technologies ARC (BIT-ARC)

Initiated, 2023



SYNOPSIS

Biomedical Innovation Technologies ARC (BIT-ARC)

Initiated by the Evans Center for Interdisciplinary Biomedical Research Center, the BIT-ARC is supported by the Department of Medicine (DoM), Clinical & Translational Science Institute (CTSI), Interdisciplinary Biomedical Research Office (IBRO) and the Office of Technology Development (OTD).

The ARC is designed to include at least 3 investigators representing different disciplines. Physical, chemical and data science investigators/engineers and biomedical researchers identify a technology to collaboratively develop to address a biomedical and technology need. Technologies include the development and/or application of new devices, data and analysis infrastructure, materials or innovative chemical or biological therapeutics, that address unmet needs for the diagnosis, treatment, or rehabilitation of patients, especially those served by Boston University and Boston Medical Center faculty and staff. Boston University guidance towards licensing and commercialization is made available to successful projects. The goal is for such projects to be developed by the pharmaceutical, materials or device development industries under license to Boston University or Boston Medical Center.

HIGHLIGHTS

In its inaugural year, 7 dynamic projects submitted proposals for consideration. Those not selected are on a trajectory to develop new patents and technologies. The inaugural awardee was announced as denoted in the next page.

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Additional information is found at
Evans Center for Interdisciplinary Biomedical Research Center web page: bumc.bu.edu/evanscenteribr/

Biomedical Innovation Technologies ARC (BIT-ARC)

Developing an artificial intelligence digital pathology diagnostic tool for proteinuric kidney disease

Directors: Drs. Weining Lu, Chao Zhang, Joel Henderson, William Tomlinson, and Vijaya Kolachalama

(ARC, 2023-present)



SYNOPSIS

(This ARC includes 10 faculty from various disciplines)

Newly funded in September 2023, this BIT-ARC project plans to develop an artificial intelligence digital biopsy glomerular ultrastructure diagnostic prototype and product for proteinuric kidney disease in drug development and clinical diagnosis. Proteinuric kidney disease (e.g., podocytopathy) is a group of chronic kidney diseases (CKD) with significant proteinuria/albuminuria (i.e., excess serum albumin in the urine). CKD affects 13% of the population (~37 million in the US and over 850 million people worldwide) and costs the US at least \$50 billion annually. Many CKD patients with proteinuria progress to kidney failure quickly and need dialysis or kidney transplantation to survive. Proteinuria/albuminuria is an early biomarker, risk factor, and surrogate outcome of CKD progression. Proteinuria/albuminuria is often caused by podocyte injury and loss in the kidney glomeruli. There are no kidney podocyte-specific anti-proteinuria therapies to halt CKD progression to kidney failure in patients with severe proteinuria and podocyte loss, which poses significant unmet medical needs worldwide.

We propose to develop an artificial intelligence (AI)/machine learning/deep learning digital biopsy prototype/product, which can measure and calculate kidney glomerular ultrastructure automatically. This AI digital kidney biopsy tool will facilitate novel drug development for proteinuric kidney disease by improving new drug pharmacodynamic and efficacy studies in clinical trials and pre-clinical animal models. This new AI digital biopsy tool may also revolutionize clinical diagnosis in pathology for kidney and other organ diseases to improve patient outcomes by enabling better treatment decisions.

Supported by the Evans Center for Interdisciplinary Biomedical Research (via the Department of Medicine) and BU Office of Technology Development (via the office of BU Associate Provost for Research)

Department of Medicine Researchers Forum

Co-Directors: Drs. Karen Jacobson, George Murphy, and Francesca Seta

(Initiated Winter, 2022)



SYNOPSIS

This Forum, initiated by the Evans Center for Interdisciplinary Biomedical Research, includes faculty from various disciplines within the Department of Medicine.

Department of Medicine Researchers Forum

GLOBAL MISSION STATEMENT

Create a dynamic space for research facilitation, knowledge exchange, wall-breaking, and to connect with and build our community. *A forum to create change.*

Priorities of this group are to expand researchers' voice and build community, not complaints, through collaborations. Participants are encouraged to promote catalyzing new ideas and fostering interactions among and between centers, groups, and innovative programs to enhance community engagement for translational research and improved infrastructure toward overall support such as technology and biosafety improvements.

Pre-ARC

Bridging Basic and Clinical Science to Reduce the Harms and Consequences of Persistent or Risky Substance Use

Directors: Drs. Kathleen Kantak, Alexander Walley, and Venetia Zachariou



SYNOPSIS

(This ARC included 9 faculty from various disciplines)

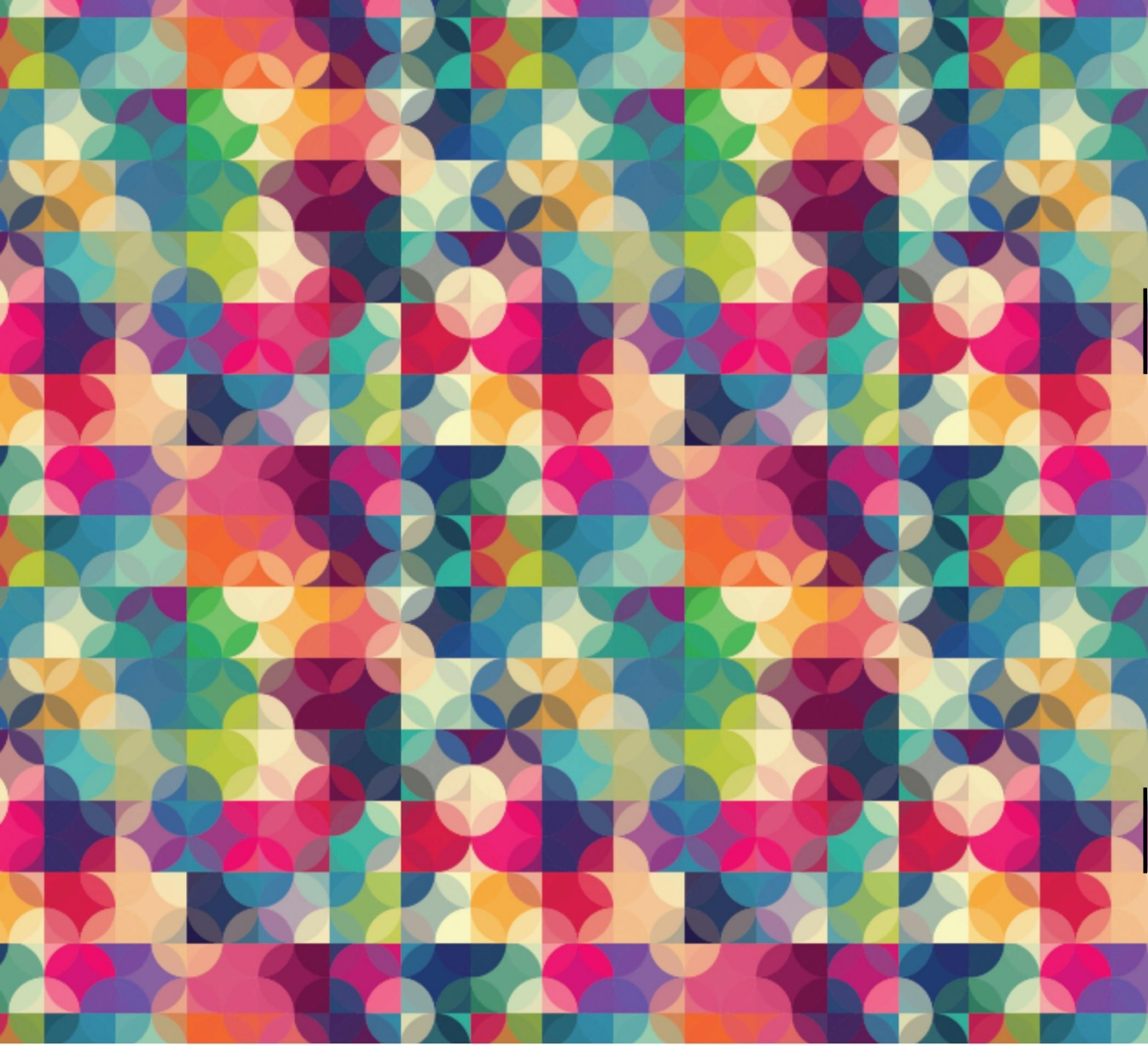
The formation of this pre-ARC stems from the opportunity to foster collaboration among the continuum of nationally recognized clinical and basic science researchers housed in various schools and departments across Boston University campuses, but with not enough collaboration between these often-times separate worlds.

This Pre-ARC intends to explore ideas for building an ARC geared towards understanding the persistent or risky use of addictive substances (i.e., alcohol or other licit and illicit substances) despite negative consequences to the individual, including impairments in cognitive and emotional health, long-term medical complications, and overdose death. Through the discovery of genetic, neurobiological, and psychosocial underpinnings of persistent or risky substance use, the hope is to discover new pathways for prevention and harm reduction.

Currently, Pre-ARC leadership is recruiting additional members and forming faculty teams interested in utilizing resources from the Brain Tissue Repository, the Animal Science Center Breeding Programs, and existing clinical human cohorts, data, and sample repositories. Long-term objectives for an ARC include fostering back-translational collaborations among clinical and preclinical substance use researchers across campuses and to advance new collaborations with, for example, researchers from Computer and Data Sciences and the Center for Systems Neuroscience to broaden the approaches used to mitigate the individual and societal burdens of persistent and risky substance use.

Additional information is found at

Evans Center for Interdisciplinary Biomedical Research Center web page: bumc.bu.edu/evanscenteribr/



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Interdisciplinary Biomedical Research Office

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