

Emory+Children's Pediatric Research Center

Update June 2014

Research Resources

Research Resources:

The resources to the right are available to all investigators affiliated with Children's Healthcare of Atlanta (CHOA), including medical staff, Emory Department of Pediatrics (DOP) faculty and staff, and those outside of the DOP and CHOA who are members of our research centers. We encourage involvement of all those interested in research throughout our system, and provide this as a guide to resources along with our research website www.pedsresearch.org. Our goals are to build infrastructure and programs that serve a broad community of scientists and clinicians engaged in pediatric research, and provide training in grant writing and grant opportunities that enhance our extramural funding for all child health investigators affiliated with Children's Healthcare of Atlanta. For suggestions and comments on any of the initiatives and resources, please contact Paul Spearman, MD (paul.spearman@emory.edu).

Grant and Manuscript Support

➤ **Stacy Heilman, PhD**
Grants Advocate (404-727-4819,
stacy.heilman@emory.edu)

- Assistance with finding grant opportunities and connecting to collaborators
- Core laboratory assistance, supervision

Grants & Manuscript Editing

- Prioritized for extramural funding opportunities, program projects
- Experienced at program project management, grant and scientific paper editing
- Request form on pedsresearch.org; send to Stacy Heilman.

Biostatistics Core

➤ Traci Leong, PhD
 ➤ Courtney McCracken, PhD
 ➤ Scott Gillespie, MS

Procedure: Request form located at <http://www.pedsresearch.org/core/detail/biostats>

Priorities: analysis for grant applications and publications

Clinical studies/ coordinators

➤ **Kris Rogers, RN, CRA**
Director, Clinical Research:
 (404-785-1215,
Kristine.rogers@choa.org)

➤ **Manager, Egleston campus:**
Allison Wellons (404-785-6459,
Allison.wellons@choa.org)

➤ **Manager, Hughes Spalding/Scottish Rite campuses:** **Beena Desai**
 (404-785-2269,
beena.desai@choa.org)

➤ **Nurse Manager, Pediatric Research Unit (Egleston):**
Stephanie Meisner, RN
Stephanie.Meisner@choa.org
 (404-785-0400-main number)

➤ **Pediatric Research Unit (Egleston):**
Services: The Research Department manages clinical coordinators and research nurses centrally, and provides training in research procedures and compliance. As needs grow or new grants are obtained, new personnel are hired who report to Kris Rogers and to the natural supervisor (grant PI, service line chief, division director).

Common Equipment/ Specimen Processing Core

2nd floor ECC 260 lab:
Technical Director:
 ➤ **Yelena Blinder**
ybesnov@emory.edu

Equipment: Biosafety cabinet, incubators, clinical centrifuge, real-time PCR machine, standard PCR machine, multilabel plate reader, gel documentation system on order

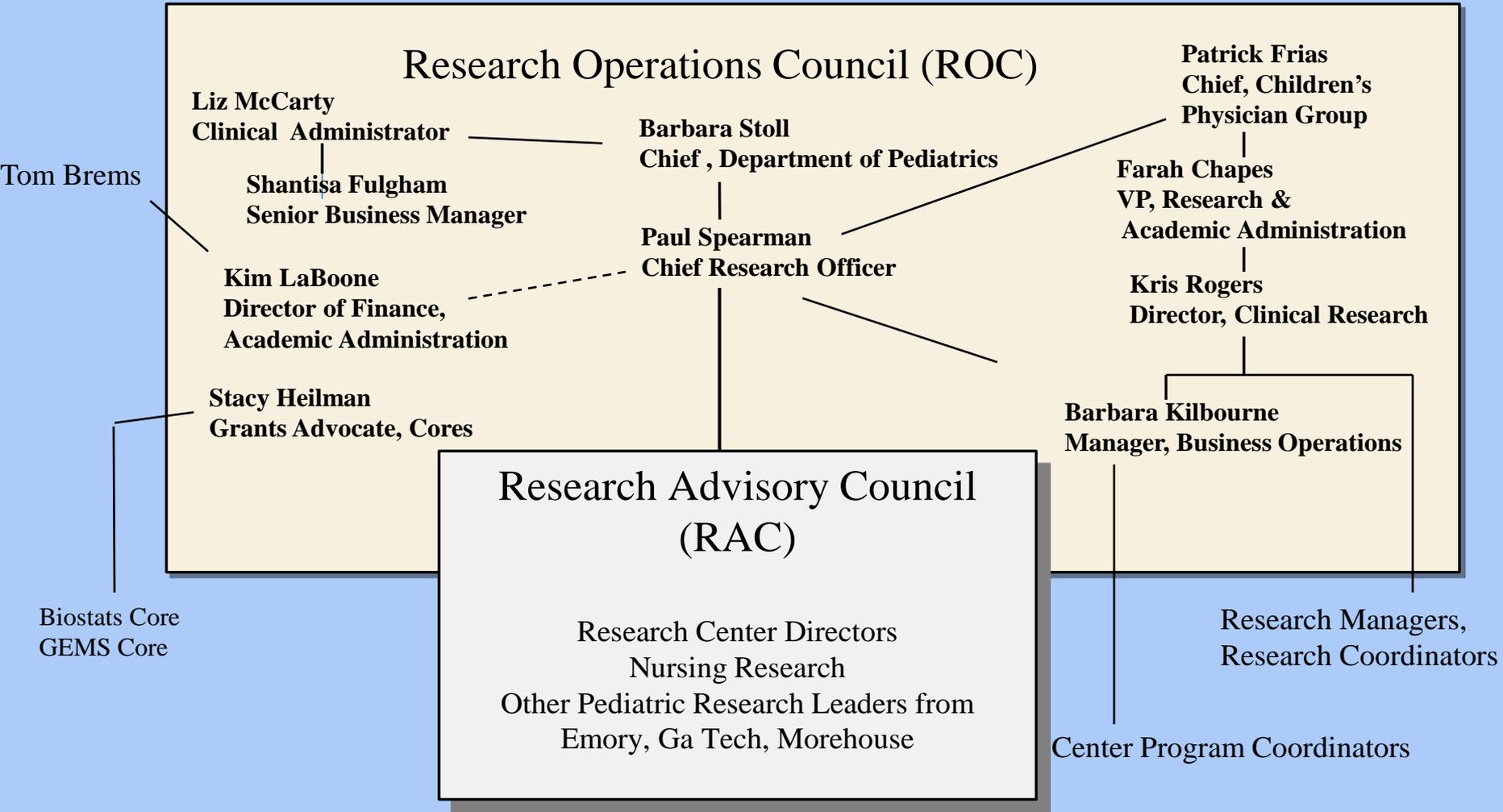
Services: this core provides common equipment for investigator's use, including access to benchtop space and hood space, centrifuges for clinical specimen processing

Laboratory Specimen Processing: Egleston

Manager: Diana Worthington-White (404-785-1721
diana.worthington-white@choa.org)

- Clinical trials specimen processing, shipping, limited storage
- ACTSI processing lab
- Laboratory inventory management system (LIMS) available

Research Leadership:



Emory+Children's Pediatric Research Centers*

Neurosciences

Pediatric
Nanomedicine

Cardiovascular
Biology

Marcus
Autism
Center

Pediatric
Innovation

Drug
Discovery

Immunology
and Vaccines

Clinical/
Translational
Research
Center

Clinical
Outcomes
Research and
Public Health

Aflac Cancer
Center

Center for CF
and Airways
Disease
Research

Transplantation
& Immune-
mediated
Disorders

*For more information, please see
center WebPages

Emory+Children's Pediatric Research Center Contacts

Center Directors:

Aflac Cancer and Blood Disorders Center

Center Director: Bill Woods, MD

william.woods@choa.org

Program Coordinator: Linda Campbell

linda.campbell@emory.edu

Center for Cardiovascular Biology

Center Director: Mike Davis, PhD

michael.davis@bme.gatech.edu

Program Coordinator: Kristen Herzegh, BA,

MPH kcoshau@emory.edu

Children's Center for Clinical and Translational Research

Center Director: Cynthia Wetmore, MD, PHD

Program Coordinator: Kristen Herzegh, BA,

MPH kcoshau@emory.edu

Center for Cystic Fibrosis & Airways Disease Research

Center Director: Nael McCarty, PhD

namccar@emory.edu

Program Coordinator: Jennifer Kenny

(acting)

Center for Drug Discovery

Center Director: Baek Kim, PhD

Baek.kim@emory.edu

Program Coordinator: Kristen Herzegh, BA,

MPH kcoshau@emory.edu

Center for Immunology and Vaccines

Center Director: Paul Spearman, MD

paul.spearman@emory.edu

Program Coordinator: Kristen Herzegh, BA,

MPH kcoshau@emory.edu

Center for Neurosciences Research

Center Director: Ton deGrauw, MD, PhD

ton.degrauw@choa.org

Program Coordinator: Jennifer Kenny

jkenny@emory.edu

Center for Pediatric Innovation

Center Directors: Bob Guldberg, PhD and Kevin Maher, MD

robert.guldberg@me.gatech.edu and

maherk@kidsheart.com

Program Coordinator: Hazel Stevens

hazel.stevens@me.gatech.edu

Center for Pediatric Nanomedicine

Center Director: Gang Bao, PhD

gang.bao@bme.gatech.edu

Senior Manager: Amy Tang

amy.tang@bme.gatech.edu

Program Coordinator: Erin Kirshtein

Erin.kirshtein@bme.gatech.edu

Center for Transplantation & Immune-mediated Disorders

Center Director: Subra Kugathasan, MD

skugath@emory.edu

Program Coordinator: Jennifer Kenny

jkenny@emory.edu

Clinical Outcomes

Research and Public Health

Center Director: Paul Spearman, MD (Acting)

paul.spearman@emory.edu

Program Coordinator: TBN

Marcus Autism Center

Center Director: Ami Klin, PhD

Director of Research: Warren Jones, PhD

ami.klin@emory.edu or

ami.klin@choa.org and

warren.r.jones@choa.org

Program Coordinator: TBN

Research Center Administration:

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Medicine, CEO, The Emory Children's Center
Executive Director, The Pediatric Center of Georgia

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Children's Healthcare of Atlanta

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Children's Healthcare of Atlanta

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Manager, Business Operations
Research Strategy Leadership, Children's Healthcare of
Atlanta

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Research-sponsored events/meetings:

(This is an overview, for specific dates/events, go to: <http://www.pedsresearch.org/calendar>)

MONDAYS	TUESDAYS	WEDNESDAYS	THURSDAYS	FRIDAYS	VARIOUS DAYS
<p>Research Operations Council (ROC) meetings: occurs weekly at Egleston, 1st Floor Admin Boardroom. Designed for central team to discuss detailed operations and issues.</p>		<p>Research Brainstorming Sessions: Help as needed to allow development and exploration of special research topics. For suggested topic nominations, contact (Stacy.heilman@emory.edu)</p>		<p>PerCS: 10 AM coffee social every 1st and 3rd Friday, usually held 3rd floor break area, E-CC</p>	<p>Research Advisory Council (RAC) meetings: twice monthly; restricted to RAC membership, contact Paul Spearman for inquiries or suggestions paul.spearman@emory.edu</p>
<p>K club: Monthly discussions/lectures for K award training, other grants training/education. Typically 2nd Monday, September to May, Contact Stacy Heilman (Stacy.heilman@emory.edu) for more information. <i>Sponsored by Departments of Pediatrics and Medicine and ACTSI.</i></p>		<p>Research Grand Rounds: 3rd Wednesday of month, Egleston, 7:30 AM</p>		<p>Research Seminars: Fridays (Egleston Classrooms). Contact Barbara Kilbourne for suggestions or needs (barbara.kilbourne@choa.org)</p>	<p>Invited speakers through seminar series sponsored by centers; contact Center Directors or Barbara Kilbourne at barbara.kilbourne@choa.org if interested in upcoming events. Center Directors are listed on pedsresearch.org website.</p>

Specialized Research Equipment/Service Cores:

CORE	SCIENTIFIC DIRECTOR	TECHNICAL DIRECTOR/CONTACT	EQUIPMENT	LOCATION	SERVICES
<u>Animal Physiology Core</u>	Mary Wagner, PhD mary.wagner@emory.edu 404-727-1336	Rong Jiang, MD rjiang2@emory.edu	Small animal surgical equipment	Emory-Children's Center, 3 rd Floor Lab	This core assists with and provides the surgical expertise and equipment for small animal survival surgery, including IACUC protocol assistance. Currently, the core offers pulmonary banding, aortic banding, coronary ligation and intramyocardial injections for mice, rats and rabbits and is available for development of other surgical procedures.
<u>Biomarkers Core</u>	Lou Ann Brown, PhD lou.ann.brown@emory.edu 404-727-5739	Janine Ward janine.ward@emory.edu	Agilent gas chromatography/mass spectrometer and Waters high performance HPLC with fluorescence detector	Emory-Children's Center, 3 rd Floor Lab	This core analyzes markers of oxidative stress and markers of alcohol exposure. Speak to Scientific Director about other chromatography/mass spec assays available.
<u>Cardiovascular Imaging Research Core (CIRC)</u>	Ritu Sachdeva, MD sachdevar@kidshcart.com 404-785-CIRC	Cynthia Mott, MPH, CCRC, PMP Cynthia.mott@choa.org	-Echocardiograms - Flow Doppler -3-D Imaging -Upright Bicycle -VO2 Analysis -Electrocardiogram -Cardiac MRI	Outpatient Cardiac Services, 2 nd Floor, Tower 1	This core provides non-invasive cardiac support for investigators involved in clinical research involving infants, children and adolescents. The CIRC has dedicated space, equipment and staff to provide you with quality cardiovascular imaging data that is collected in a meticulous, systematic, detail-orientated manner. Because of our unique set-up, we are able to utilize state-of-the-art imaging modalities not typically seen in the clinical setting.

Specialized Research Equipment/Service Cores (continued)

CORE	SCIENTIFIC DIRECTOR	TECHNICAL DIRECTOR/CONTACT	EQUIPMENT	LOCATION	SERVICES
Flow Cytometry/Cell Sorting	David Archer darcher@emory.edu	Aaron Rae aaron.j.rae@emory.edu	FACSCanto, LSRII, FACSAria, AutoMACS	Health Sciences Research Building, E-362	This core offers access to several state of the art analytical flow cytometers as well as high-speed cell sorting. We also offer training as well as expert help to enable our users to improve the quality and scope of their research.
Immunology Core	Larry Anderson larry.anderson@emory.edu 404-712-6604	Sujin Lee, PhD sujin.lee@emory.edu	Specimen processing (hood, centrifuges, Coulter counter), Zeiss ELISPOT reader, ELISAs, assay design for intracellular cytokine staining (ICS), luminex 200 assays for protein quantitation, real-time PCR	Emory-Children's Center, Room 510	This core provides equipment and technical expertise for the performance of immunologic assays and diagnostic assays for infectious pathogens. Our mission is to enhance the ability of investigators at Children's and affiliated institutions to perform research in the areas of immunology, vaccine testing, and infectious diseases.
Radiology Core	Radiologists at Children's are board certified with additional training in pediatric imaging and are available for consultation upon request. This operation also includes physicists with imaging expertise and other staff experts.	Melinda Dobbs, RN, BSN, CCRC melinda.dobbs@choa.org	<ul style="list-style-type: none"> • Access to clinical CT (4), PET (1), Bone Densitometry (2), Fluoroscopy (8), Nuclear Medicine (4), Ultrasound (9) and X-ray. • Access to 6 clinical MRI scanners including a 1.0T intraoperative, 1.5T and 3T systems. • Access to 2 fMRI systems. • Sedation Services • Access to radiology investigators specializing in radiology, neuroradiology and interventional radiology. • Access to MRI physicists (3). • Access to research professionals including administrators and research coordinators. • Administrative services including scheduling, archival of images 		The is an interdisciplinary research core that recognizes the importance of medical imaging in the diagnosis and treatment of diseases in children and young adults. PIRC provides investigators with modern imaging technology and collaborating imaging researchers to achieve research goals. Our team consults with investigators to enhance their research through access to state-of-the-art technology and enables the conduct of standard imaging associated with large clinical trials. Services include MRI, CT, PET, Bone Densitometry, Fluoroscopy, Nuclear Medicine, Ultrasound and X-ray.

CORE in Development	EQUIPMENT/LOCATION	DESCRIPTION
Specimen Repository (which will enhance the Specimen Processing Core)	LIMS, freezers (-80, LN2) Sync with freezer space in new building; temporary space until then being identified	The specimen repository will offer organized storage of blood and body fluids and nucleic acids. Tissue repository services are under further discussion. Specimen processing can be coordinated to link with the specimen repository. Bar-coded standard vial storage and a dedicated LIMS will offer automated tracking and organized retrieval of specimens.

Partnership Core

CORE	SCIENTIFIC DIRECTORS	EQUIPMENT	LOCATION	SERVICES
Integrated Cell Imaging Core	Adam Marcus, PhD Director, ICI aimarcu@emory.edu Alexa Mattheyses, PhD Associate Director, ICI mattheyses@emory.edu Neil Anthony, PhD neil.anthony@emory.edu 404-969-CORE	The rates for the microscopes included in this effort can be found at: http://ici.emory.edu/document/ICI%20Pediatrics%20Rates.pdf . Pediatric researchers will benefit from a 40% subsidy when using any of the ICI equipment and technologies. ICI also provides expert consultation, training, and assistance on all technologies. More information on the microscopes and services available, locations, and how to become a user is available at ici.emory.edu	A partnership facilitated by the Emory School of Medicine and includes the Emory+Children's Pediatric Research Center Cellular Imaging Core along with other cellular imaging sites on campus including Winship Cancer Institute, Emory NINDS Neuroscience Core Facilities (ENNCF), and the Department of Physiology	This core provides training and access to advanced cellular imaging systems, including confocal and TIRF microscopy. For more information: http://www.pedsresearch.org/cores/detail/cell-imaging

Funding Opportunities:

Funding Opportunity	Funding Limit	Funding Term	Deadline	Eligibility	Post Award Expectations	Additional Information
Friends	\$25,000	12-18 months	3rd Friday in Sept	<ol style="list-style-type: none"> Children's professional staff who do not also have a compensated faculty appointment Must be for clinical or outcomes research taking place in Children's facilities 	<ol style="list-style-type: none"> Must provide annual and final reports. Must be willing to present findings to Friends groups, Children's leadership, etc. 	Fund does not provide for investigator salary support
EECRSeed: Engaging Emory & Children's Researchers Seed Grant Program	\$50,000	12 months	3rd Friday in Sept	<ol style="list-style-type: none"> Regular faculty in clinical departments at Emory. Applicants outside of Dept. of Peds must have clinical privileges at Children's. Must not have an active R01 or P01. Must provide agency and proposed date they will submit for extramural funding. Priority given to faculty with New Investigator status. 	Must submit a grant to an extramural agency.	<p>\$25,000 of total award may be directed to investigator salary.</p> <p>This seed grant is sponsored by Children's Healthcare of Atlanta and Emory University</p>
Research Center Pilot Grants (including Emory & GA Tech based centers)	\$50,000 (some GA Tech are \$60K)	12 months	Usually mid-winter; Emory-based are due roughly every other year and GA Tech-based offered every year	<ol style="list-style-type: none"> Must include a member of the center and/or member of Children's medical staff GA Tech-based centers (CPN, CPI and IPaT) must also include member of GA Tech faculty 	<ol style="list-style-type: none"> Must provide annual report specifying related publications, grant applications submitted and extramural funding received. Must apply for extramural funding within one year of project conclusion date. 	https://pediatricconnect.gtri.gatech.edu/grants

Funding Opportunities (continued):

Funding Opportunity	Funding Limit	Funding Term	Deadline	Eligibility	Post Award Expectations	Additional Information
Dudley Moore Nursing and Allied Health Research Fund	\$15,000	6-18 months	Usually 1st Friday in May and Oct	<ol style="list-style-type: none"> 1. All Children's nursing and allied health staff who provide services at one of Children's locations are eligible. 2. Excludes those with regular faculty appointments or who are employed by Emory 3. Projects must have an impact on enhanced patient care, priority is given to projects that will provide evidence to change practice. 	Must be willing to present findings by request.	Fund restricted by donor to support nursing and allied health research at Children's
Quick Wins	varies	12-24 months	ongoing	<ol style="list-style-type: none"> 1. Project proposals must be submitted by teams comprised of individuals from each organization, Children's and Georgia Tech. 2. The proposals must address a project that provides an answer to an unmet business or clinical need as identified by a clinician, technologist, or Children's leader. 	The project must be capable of delivering a workable solution (at minimum a validated "prototype") into the hands of a clinician or team within 18 months from the receipt of funds and project start.	https://pediatriconnect.gtri.gatech.edu/grants

Additional Resources/Updates:

Research listserv:

Contact barbara.kilbourne@choa.org to be added to this listserv used to disseminate all pediatric research related announcements including seminars, funding opportunities, such as BiRD (Bringing in the Research Dollars), and the Weekly PREP (Pediatric Research Events and Programs)

Website:

www.pedsresearch.org

This is the central resource for research seminar info, contacts, cores, calendars, forms

Health Sciences Research Building:

1760 Haygood Road

Atlanta, GA 30322

190,000 ft²; 115,000 for pediatric research

Dry and wet lab research

For floor plans go to: http://pedsresearch.org/_files/HSRB_FloorPlans.pdf

Go to: <http://www.pedsresearch.org/about-us> for more info

Research Recruitment Update*:

NAME	PHOTO	CENTER	TITLE	START DATE	RECRUITED FROM	RESEARCH INTERESTS
Changwon Park, PhD		Center for Cardiovascular Biology		April 2014	Department of Pharmacology, College of Medicine, University of Illinois Chicago, IL	FLK1 (VEGFR2), a receptor tyrosine kinase, plays a critical role for blood and vessel development. Fate mapping studies have demonstrated that FLK1+ mesoderm contributes to the development of the cardiovascular system consisting of hematopoietic, endothelial, cardiac muscle and smooth muscle cells. FLK1 continues to play a critical role in (pathological) angiogenesis in the adult. Therefore, understanding molecular mechanisms that regulate Flk1 expression is essential for delineating the pathways involved in blood and vessel differentiation during embryogenesis as well as postnatal angiogenesis. We have demonstrated that Bone Morphogenetic Protein (BMP) 4 is a major factor to generate FLK1 expressing mesoderm which can subsequently differentiate into endothelial and hematopoietic cells. Furthermore, we reported that ER71, a novel member of the ETS transcription factor family, is the direct upstream regulator of FLK1 expression and that ER71 is indispensable for vessel and blood development in mouse embryogenesis. Extending from our previous findings, we are currently studying the role of ER71 for the establishment of the cardiovascular system and for pathological angiogenesis. Outcome from the proposed studies will provide a new and detailed insight on the role of ER71 in vascular development and pathological angiogenesis, which can provide a new research venue for the development of specific targets for the cardiovascular diseases. In addition, we are investigating mechanisms which can induce direct reprogramming of somatic cells to functional endothelial cells.
Cynthia Wetmore, MD, PhD		Center for Clinical & Translational Research (CTTR)	Director	April 2014	St. Jude's Research Hospital	Basic science: Developmental neurobiology, genetic control of normal and neoplastic proliferation in the nervous system, neural stem cells, gene expression in the nervous system, repair of DNA damage in the nervous system. Clinical science: Developmental therapeutics for pediatric oncology, neuro-oncology; design and conduct of Phase I/II clinical studies; translation of basic science discoveries to improving clinical care of patients.
Dmitry M. Shayakhmetov, Ph.D.		Center for Transplantation and Immune-Mediated Disorders (CTID)	Professor, Division of Rheumatology, Department of Pediatrics	April 2014	Department of Medicine, Division of Medical Genetics, University of Washington, Seattle	<ul style="list-style-type: none"> • Molecular mechanisms of a novel type of pro-inflammatory necrotic cell death in vivo. • Identification of molecular sensors triggering transcriptional and functional activation of macrophages in vivo. • Defining the role of pro-inflammatory types of cell death in the disruption of tissue homeostasis and triggering the systemic inflammatory host response • Modification of adenovirus interaction with circulating antibodies for cancer therapy.

*Recruits for the past year

Research Update June 2014

Research Recruitment Update*:

NAME	PHOTO	CENTER	TITLE	START DATE	RECRUITED FROM	RESEARCH INTERESTS
Chris Gunter, PhD		Marcus Autism Center (MAC)	Associate Director for Research	February 2014	Nature—Senior Editor University of Alabama in Birmingham—Adjunct Professor ASHG—Chair, Communications Committee	Spokesperson for science.
Paul A. Dawson, PhD		Center for Transplantation and Immune-Mediated Disorders (CTID)	Professor	February 2014	Department of Internal Medicine, Section on Gastroenterology, Wake Forest School of Medicine, Medical Center Boulevard	BILE ACIDS, CHOLESTEROL METABOLISM, MOLECULAR CLONING, GENE EXPRESSION AND REGULATION, MOLECULAR GENETICS Molecular Genetics of Ileal Bile Acid Transporter. My lab identified and cloned the human ileal bile acid transporter cDNA and gene. These probes are being used to identify dysfunctional mutations in patients with bile acid malabsorption. Various classes of dysfunctional mutations in the ileal bile acid transporter gene have been identified. In addition to null mutations (i.e., splicing defects), we have also identified missense mutations that interfere with bile acid transporter processing and mechanism of action. The Class 2 mutations cause misfolding and ER retention of the transporter. More interesting are the Class 3 and 4 mutations that block bile acid transport at the substrate binding and solute translocation steps. The actions of these mutations are being studied to gain insight into the molecular mechanism of sodium-coupled solute transport. The association of these mutations with other gastrointestinal and lipid metabolism disorders including gallstone disease, irritable bowel syndrome, hypocholesterolemia, and hypertriglyceridemia is currently being investigated.
Cheng-Kui Qu, MD, PhD		Aflac Cancer and Blood Disorders Center (Aflac)	Associate Professor	January 2014	Case Comprehensive Cancer Center Case Western Reserve University	His specific interests are in myeloid malignancies, with an emphasis on PTPN11/SHP-2 and cell signaling mechanisms that control hematopoietic stem cell function. Also focusing on the role of protein phosphatases in normal hematopoietic cell development and in leukemogenesis. Works closely with Kevin Bunting and Himalee Sabnis.

*Recruits for the past year

Research Recruitment Update*:

NAME	PHOTO	CENTER	TITLE	START DATE	RECRUITED FROM	RESEARCH INTERESTS
Elizabeth "Beth" Stenger, MD		Aflac Cancer and Blood Disorders Center (Aflac)	Assistant Professor	August 2013	Children's Hospital of Pittsburgh, University of Pittsburgh	Enhanced IL-12 Production by mTOR-inhibited DC and Protection from GVHD
Brandon Aylward, PhD		Children's Center for Neurosciences/ Children's Center for Cardiovascular Biology (CCNR and CCB)	Assistant Professor	July 2013	Cincinnati Children's Hospital Medical Center	He received his doctoral degree in clinical child psychology with a minor in quantitative psychology from the University of Kansas and completed his predoctoral residency program at Cincinnati Children's. His research interests encompass a broad range of health-related issues for children and adolescents within the context of pediatric psychology. To this end, his work has focused on three main areas: (1) predictors and correlates of children's psychosocial, developmental and physical functioning in various chronic illness populations; (2) trends and correlates of adherence and self-management behaviors; and 3) use of advanced statistical methodology and innovative technology to examine predictors and outcomes for chronic health issues.

*Recruits for the past year