

Condensed Conference Schedule

Thursday, May 16

11:00 am-6:00 pm

Registration

1:50 pm

Opening Remarks

2:00-4:00 pm

Platform Session 1: Environmental Session

5:00-6:00 pm

Keynote Talk- Chuck Murry

6:00-7:30 pm

Reception

7:30-10:00 pm

Poster Preview and Social

Friday May 17

7:30-9:00 am

Continental Breakfast

8:00-10:15 am

Poster Session 1: (180 posters up for entire meeting; Odd

Numbered Posters Present)

8:30-9:30 am

Moderated Poster Discussion 1

9:00-10:00 am

Moderated Poster Discussion 2

10:20-12:00 pm

Platform Session 2 :CHD Models

12:00-1:30 pm

Box Lunch; Business Meeting

1:30-3:30 pm

Platform Session 3: Genomics and Regulation

3:30-3:50 pm

Coffee Break

3:50-5:50 pm

Platform Session 4: Coronaries, Epicardium, Conduction System

Free evening for dinner and discussions

8:30-10:00 pm

Posters and Social

Saturday May 18

7:30-9:00 am

Continental breakfast

8:00-10:15 am

Poster Session 2 (Even Numbered Posters Present)

8:30-9:30 am

Moderated Poster Discussion 3

9:00-10:00 am

Moderated Poster Discussion 4

10:20-12:00 pm

Platform Session 5: Valve Development

12:00-1:30 pm

Box lunches, NIH Presentations

1:30-3:10 pm

Platform Session 6: Heart Formation and Progenitors

3:10-3:30 pm

Coffee Break

3:30-5:10 pm

Platform Session 7: Cardiac Growth and Regeneration

6:00 pm - ?

Banquet with awards and music

Detailed Conference Schedule

Thursday, May 16, 2013

11:00am - 6:00pm

Registration - Foyer of the Sonoran Ballroom

1:50pm

Opening Remarks, Sonoran Ballroom

Thursday Afte	ernoon Platform Session 1: Environmental Session - Sonoran Ballroom
Session Chai	Todd Camenisch and Ching-Pin Chang
2:00-2:20pm	S1.1. Ambient air pollution and traffic exposures and congenital heart defects in the San Joaquin Valley of California Amy Padulai, Ira B Tager2, Suzan L Carmichaeli, SK Hammond2, FW Lurmanni, Gary M. Shawi and Edward J. Lammer3i Department of Pediatrics, Stanford University, Palo Alto, CA, 2University of California, Berkeley, CA, 3Children's Hospital Oakland Research Institute, Oakland, CA
2:20-2:40	S1.2. Environmental risk factors for congenital heart defects, results from the national birth defects prevention study Jennita Reefhuis, PhD, for the National Birth Defects Prevention Study, Birth Defects Branch, Division of Birth Defects and Developmental Disabilities, Centers for Disease Control and Prevention, Atlanta, GA
2:40-3:00	S1.3. Nicotine Inhibits endocardial cushion development through Nfatc and Vegf Yiqin Xiongi, Jin Yangi, Calvin T. Hangi, Richard M. Cheni, Ching-Pin Changi Department of Medicine, Cardiovascular Medicine, Stanford University School of Medicine, Stanford, CA
3:00-3:20	S1.4. Mouse model of hypoplastic left heart syndrome shows multigenic etiology and defects in energy Metabolism Xiaoqin Liui, Shazina Saeedi, Brian Gibsi, You Lii, Richard Francisi, George Gabrieli, Linda Leatherburyz, Kimimasa Tobitai, Dennis Kostkai, Cecilia.W. Loi iDept of Developmental Biology, University of Pittsburgh School of Medicine, Pittsburgh, PA, 2Children's National Medical Center, Washington, DC
3:20-3:40	S1.5. The epicardium as a sensor of AHR-mediated heart malformation Peter Hofsteeni, Jess Plavickii, Richard E. Petersoni, and Warren Heidemani Division of Pharmaceutical Sciences, School of Pharmacy University of Wisconsin, Madison, WI
3:40-4:00	S1.6. Depletion of Retinoic Acid Receptors Initiates a Novel Positive Feedback Mechanism that Promotes Teratogenic Increases in Retinoic Acid Enrico D'Aniello and Joshua S. Waxmanı*

Department of Molecular Cardiovascular Biology and Developmental Biology Divisions, Cincinnati Children's Hospital Medical Center, Cincinnati, OH

Thursday Evening Keynote Address and Reception

5:00 - 6:00pm Keynote Address: Charles Murry, PhD, University of Washington Cardiac Regeneration: Lessons from Development

6:00 - 7:30pm Reception, Sonoran Ballroom Rooftop

7:30-10:00pm Poster Sessions & Social, Santa Catalina Ballroom

Friday, May 17, 2013

7:30-9:00am

Continental Breakfast, Sonoran Ballroom Rooftop

8:00-10:15 am

Poster Session 1: Odd Numbered Posters Present

8:30 - 9:30 am

Moderated Poster Discussion 1

Moderator- Heather Bartlett, MD, University of Iowa

M1.1. The cell cycle kinase Nek2 modulates ciliogenesis and resorption in the development of left right asymmetry

Basudha Bas, Samuel Endicott, Shiaulou Yua, Mustafa Khokha, and Martina Brueckner

Yale University School of Medicine

M1.2. Identification of a role for phosphofructokinase-1 in heterotaxy

Jason Cowan, Ashley Cast, Stephanie M. Ware Cincinnati Children's Hospital Medical Center

M1.3. Forward genetic screen with mouse ENU mutagenesis reveals a central role for the cilia and left-right patterning in congenital heart disease.

You Li, Nikolas Klena, Richard Francis, Xiaoqin Liu, Kristi Lemke, George Gabriel, Bishwanath Chatterjee, Rama Damerla, Yu Chen, Shane Anderton, Caroline Lawhead, Laura Reinholdt, Kimimasa Tobita, Linda Leatherbury, Gregory Pazour, Cecilia W. Lo

Department of Developmental Biology, University of Pittsburgh School of Medicine, Pittsburgh, PA, 2Jackson Laboratory, Bar Harbor, ME, 3Children's

National Medical Center, Washington, D.C., 4Program in Molecular Medicine, University of Massachusetts Medical School, Worcester, MA

M1.4. Genetic variant located in an intron of SCN10A functionally affects an SCN5A enhancer

Malou van den Boogaard¹, David E Arnolds^{2,3,*} Scott Smemo⁴, Ozanna Burnicka-Turek⁴, Petra Klous⁵, Harmen J G van de Werken⁶, Wouter de Laat⁵, Marcelo A Nobrega⁴, Ivan P Moskowitz^{2,3,#}, Phil Barnett¹, and Vincent M Christoffels¹, #

1Department of Anatomy, Embryology and Physiology, Heart Failure Research Center, Academic Medical Center, University of Amsterdam, The Netherlands; 2,3,4Departments of Pediatrics2, Pathology3 and Human Genetics4, University of Chicago, Illinois; 5Hubrecht Institute, KNAW and University Medical Center Utrecht, The Netherlands; 6Department of Cell Biology, Erasmus MC Rotterdam, The Netherlands *,# equal contribution

M1.5. Common genetic variant for conduction system function at SCN10A

modulates SCN5A enhancer activity.

Ozanna Burnicka-Turek*1.2, Malou van den Boogaard,4, Scott Smemo*3, David E. Arnolds*1.2 Harmen J.G. van de Werken5, Petra Klous, Wouter de Laaf, Vincent M. Christoffels#4, Marcelo A. Nobrega#3 Phil Barnett#4, and Ivan P. Moskowitz#1,2

Departments of Pediatrics1, Pathology2, and Human Genetics3, The University of Chicago, Chicago, IL, USA; Department of Anatomy, Embryology and Physiology₄ Academic Medical Center, University of Amsterdam, Amsterdam, The Netherlands; Institute-KNAW & University Medical Center Utrechts, Utrecht, The Netherlands. * authors contributed equally to this work; # authors jointly directed this work

M1.6. A strict lineage boundary between the first and second heart fields is defined by the contribution of the Tbx5 lineage.

Joshua D. Wythe, W. Patrick Devine, Kazuko Koshiba-Takeuchi, Kyonori Togi, Benoit G. Bruneau.

Gladstone Institute of Cardiovascular Disease, University of California, San Francisco, San Francisco, CA, USA.

9:00-10:00 am

Moderated Poster Discussion 2

Moderator- Sylvia Evans. PHD, University of California, San Diego

M2.1. Regional expression of Cyp26 enzymes in the anterior lateral plate mesoderm is required to balance cardiac and vascular lineages

> Ariel Rydeen^{1,2,3} and Joshua Waxman^{2,3} Cincinnati Children's Hospital

M2.2. IGF2 signaling regulates cardiomyocyte proliferation

Hua Shen', Susana Cavallero', Peng Li', Kristine Estrada', Miguel Constancia2, and Henry M. Sucov1 University of Southern California

M2.3. Arterial and venous progenitors of the major axial vessels originate at distinct locations

Vikram Kohli¹, Jennifer A. Schumacher¹, Sharina Palencia Desai^{1,2}, Kira Rehn¹ and Saulius Sumanas1'

Cincinnati Children's Hospital

M2.4. The role of hippo and Pitx2 signaling in cardiac regeneration

Ge Tao1, Todd Heallen1, Yuka Morikawa2, John Leach1, Min Zhang1,5, Ashley Benham², James T. Willerson², Robert J. Schwartz^{2,6}, Randy L. Johnson^{3,4} and James F. Martin^{1,2,3,5}

Baylor College of Medicine

M2.5. Fzd2 mediated non-canonical Wnt signaling in cardiac progenitor development

Ethan David Cohen University of Rochester

M2.6 YAP1 promotes adult cardiomyocyte proliferation and improves outcome after myocardial infarction

Zhiqiang Lin^{1*}, Alexander von Gise^{1*}, Pingzhu Zhou¹, Qing Ma¹, Jianming Jiang², Allan Yau¹, Jessica Buck¹, Jinghai Chen¹, Da-zhi Wang¹ and William Pu^{1#}, Children's Hospital Boston

Friday Morning

Platform Session 2: CHD Models, Sonoran Ballroom
Session Chairs

Michiko Watanabe and Kristy Red-Horse

10:20-10:40am S2.1. Mouse and human CRKL is dosage sensitive for heart morphogenesis

SE Racedoi, J Chungi, DM McDonald-McGinni, A Baileyi, N Meyeri, E Goldmuntzi, E Zackaii, BS Emanueli, B Zhoui, B Funkei and BE Morrowi
Department of Genetics, Albert Einstein College of Medicine, Yeshiva University,
Bronx, NY, 2Children's Hospital of Philadelphia, University of Pennsylvania,
Philadelphia, PA, 3Laboratory for Molecular Medicine, Partners HealthCare Center for Personalized Genetic Medicine, Cambridge, MA

10:40-11:00 S2.2. A Dominant Negative JUN mutant is associated with human congenital heart disease and alters a physical and functional interaction with TBX1

Hua Pani, Tao Zhangi, Julie De Mesmaekerz, Anuja Nevez, Matthew A. Bensonz, Brande C. Latneyz, Petra Wernerz, Elizabeth Goldmuntzz, Shoumo Bhattacharyaz, and Jason Z. Stolleri

¹Division of Neonatology, Department of Pediatrics, Children's Hospital of Philadelphia, Philadelphia, PA, ²Department of Cardiovascular Medicine, The Wellcome Trust Centre for Human Genetics, University of Oxford, Roosevelt Drive, Oxford, UK, ³Department of Pharmacology and the Institute for Translational Medicine and Therapeutics, Perelman School of Medicine at the University of Pennsylvania, Philadelphia, PA, ⁴Division of Cardiology, Department of Pediatrics, Children's Hospital of Philadelphia, PA

11:00-11:20 S2.3. Mechanisms of Gene Haploinsufficiency: Chromatin factors and phenotypic rescue in a mouse model of DiGeorge syndrome

F. Gabriella Fulcolii, Monica Franzesez, Claudia Angeliniz, Antonio Baldinii.3

1) Institute of Genetics and Biophysics, CNR; 2) Istituto Applicazioni del Calcolo, CNR; 3).University Federico II, Napoli, Italy

11:20-11:40 S2.4. Composition and function of the TBX5 cardiac transcriptional complex

Lauren Waldroni, 2, Kerry Dorri, 2, Ileana M. Cristea 4, Ivan Moskowitz 5 and Frank L. Conloni, 2,3

- (1) McAllister Heart Institute, School of Medicine, UNC-Chapel Hill, Chapel Hill, NC,
- (2) Department of Genetics, School of Medicine, UNC-Chapel Hill, Chapel Hill, NC
- (3) Department of Biology, UNC-Chapel Hill, Chapel Hill, NC, (4) Department of Molecular Biology, Princeton University, NJ, (5) Departments of Pediatrics and Pathology, The University of Chicago, Chicago, IL

11:40-12:00 S2.5. Foxf genes integrate Tbx5 and Hedgehog pathways in second heart field cardiac progenitors for atrial septation

Andrew Hoffmanni, Holly Yangi, Ozanna Burnicka-Tureki, Josh Bosmani, Yufang Zhang², StevenVokes³, Andrew McMahon⁴, Vladimir Kalinichenko², Ivan Moskowitzi iDepartments of Pediatrics and Pathology, The University of Chicago, Chicago, IL, 2Cincinnati Children's Hospital Medical Center, Cincinnati, OH ₃Section of Molecular Cell & Developmental Biology, Institute for Cellular and Molecular Biology, Austin, TX, ₄Department of Stem Cell and Regenerative Biology, Department of Molecular and Cellular Biology and Harvard Stem Cell Institute, Harvard University, Cambridge, Massachusetts

12:00 - 1:30

Box Lunch; Weinstein Conference Business Meeting

Friday Afternoon

Platform Session 3: Genomics & Regulation, Sonoran Ballroom
Session Chairs

Tom Doetschman and Kryn Stankunas

1:30-1:50pm S3.1. Whole-exome Sequencing Identifies Novel Genes for Autosomal Recessive Heterotaxy

Muhammad Tariq₁, Ashley E. Cast₁ and Stephanie M. Ware_{1, 2} Divisions of ₁Molecular Cardiovascular Biology, ₂Human Genetics, Cincinnati Children's Hospital Medical Center, Cincinnati, OH

1:50-2:10 S3.2. Altered enhancer usage between fetal and adult cardiac transcriptional programs identified by genome-wide GATA4 ChIP-seq

Aibin Hei, Fei Gui, Qing Mai, Len A. Pennacchio3, William T. Pui.2.

Department of Cardiology, Boston Children's Hospital and Harvard Medical School, Boston, MA, 2Harvard Stem Cell Institute, Harvard University, Cambridge, MA, 3Genomics Division, MS 84-171, Lawrence Berkeley National Laboratory, Berkeley, CA

2:10-2:30 S3.3. microRNA-1 is required for normal cardiac sarcomere formation and function

Amy Heidersbach, Karen Carver-Moore, Yu Huang, Kathryn Ivey, and Deepak Srivastava Gladstone Institute of Cardiovascular Disease, University of California, San Francisco CA

2:30-2:50 S3.4. Direct differentiation from pluripotent stem cells tocardiac lineages by defined factors

Yuika Morita1.2, Kazuko Koshiba-Takeuchi2 and Jun K. Takeuchi1.2.3

1:Graduate School of Biological Sciences, the University of Tokyo, 2:Division of Cardiovascular Regeneration, IMCB, the University of Tokyo, 3:JST PRESTO, Understanding life by iPS cells technology

2:50-3:10 S3.5. Heart field origin of great vessel precursors relies on nkx2.5-mediated vasculogenesis

Noëlle Paffett-Lugassy1.2.3, Reena Singh4, Burcu Guner-Ataman1.2, Richard P. Harvey4, C. Geoffrey Burns1.2.3 * & Caroline E. Burns1.2.3 *

1Cardiovascular Research Center, Massachusetts General Hospital, Charlestown, MA, 2Harvard Darlinghurst, New South Wales, Australia

3:10-3:30 S3.6. ETS factors regulate the Vegf-dependent, arterial-specific expression of DII4.

Joshua D. Wythe1,2.3, Lan T. H. Dang4,5,6, W. Patrick Devine1,3.7, Emilie Boudreau4,5,6, Stanley T. Artap8, William Schachterle3, Didier Y.R. Stainier3,9, Peter Oettgen8, Brian L. Black3, Benoit G. Bruneau1,2.3, Jason E. Fish4,5,6

1. Gladstone Institute of Cardiovascular Disease, San Francisco, CA, 2. Department of Pediatrics, University of California, San Francisco CA, 3. Cardiovascular Research Institute, University of California, San Francisco, CA, 4. Toronto General Research Institute, University Health Network, Toronto, Canada, 5. Department of Laboratory Medicine and Pathobiology, University of Toronto, Toronto, Canada, 6. Heart and Stroke Richard Lewar Centre of Excellence in Cardiovascular Research, Toronto, Canada, 7. Department of Pathology, University of California, San Francisco, CA, 8. Divisions of Cardiology and Molecular and Vascular Medicine, Center for Vascular Biology Research, Beth Israel Deaconess Medical Center, Harvard Medical School, Boston, MA. 9. Department of Developmental Genetics, Max Planck Institute for Heart and Lung Research, BadNauheim, Germany

3:30 - 3:50pm coffee break

Friday Afternoon Platform Session 4: Coronaries, Epicardium, Conduction System Sonoran Ballroom Session Chairs Stacey Rentschler and Chris Brown

3:50-4:10pm S4.1. Epicardial CXCL12 and Coronary Vessel Development Susana Cavallero, Hua Shen, Henry M. Sucov, Broad Center for Regenerative Medicine and Stem Cell Research, Keck School of Medicine, University of Southern California, Los Angeles, CA

4:10-4:30 S4.2. Efficient Differentiation of Human Pluripotent Cells into Wt1 Epicardium through an IsI1+ Primitive Mesoderm Intermediate

Miranda Hayworthi,2, David Reynoldsi,2, Laura Menendezi,2, Tatiana Yatskievych3,4, Parker Antin3,4 and Stephen Daltoni,2

1Paul D. Coverdell Center for Biomedical and Health Sciences and 2Department of Biochemistry and Molecular Biology, Franklin College of Arts and Sciences, University of Georgia, Athens, GA, 3Department of Cellular and Molecular Medicine and 4Department of Molecular and Cellular Biology, College of Medicine, The University of Arizona, Tucson AZ

4:30-4:50 S4.3. Coronary artery stems are patterned by aorta specific myocardial development

Heidi Chena, Aruna Poduria, Brian Raftreya, Andrew S. McKaya, Kirk Willmarthb, Jareda Churko, Joseph Wuc, and, Kristy Red-Horsea,d
Department of Biological Sciences, bDepartment of Developmental Biology, aDepartment of Medicine, Division of Cardiology, Stanford University, Stanford CA

4:50-5:10 S4.4. Locus-confined combinatorial regulatory elements control Tbx3 expression in the cardiac conduction system

Henk van Weerdi*, Ileana Badii*, Malou van den Boogaardi, Corrie de Gier-de Vriesi, Harmen J van de Werkenz, Wouter de Laatz, Phil Barnetti, Vincent M

Christoffels:*equal contribution
Dept. of Anatomy, Embryology & Physiology, Academic Medical Center, Amsterdam,
The Netherlands, 2Hubrecht Institute-KNAW and University Medical Center Utrecht,

Utrecht, The Netherlands

5:10-5:30 S4.5. A Pitx2-microRNA pathway that delimits sinoatrial node development and inhibits predisposition to atrial fibrillation

Jun Wangi, Yan Baii.4, Na Lii, Stephanie B. Greenei, Min Zhangi.4, Ye Taoi, Xander H.T. Wehrens 1.5, and James F. Martin#,1,2,3,4

Department of Molecular Physiology and Biophysics, Baylor College of Medicine, Houston, Texas, 2 Texas Heart Institute, Houston, Texas, 3 Program in Developmental Biology, Baylor College of Medicine, Houston, Texas, 4 Institute of Biosciences and Technology, Texas A&M Health Science Center, Houston, Texas, 5 Department of Medicine, Division of Cardiology, Baylor College of Medicine, Houston, Texas

5:30-5:50 S4.6. Engineered Epicardium Activates Cardiac Regeneration

Vahid Serpooshani, Ke Weiz, Mingming Zhaoi, Scott A. Metzleri, Parisha B. Shahi, Paul J. Kimi, Yuka Matsuurai, Wenqing Caiz, Alex Savtchenkoz, Andrew Wangi, Marta Diez-Cuñadoz, Wenhong Zhuz, Morteza Mahmoudis, Manish J. Buttei, Phillip C. Yangi, Daniel Bernsteini, Mark Mercolaz, and Pilar Ruiz-Lozanoi

1 Stanford University School of Medicine, Stanford, CA, 2 Sanford-Burnham Medical

Research Institute, La Jolla, CA, 3 University of Illinois at Urbana-Champaign, Urbana, IL, 4 Department of Bioengineering, University of California, San Diego, La Jolla, CA

Free Evening for Dinner and Discussions

8:30 - 10:00pm Posters and Social, Santa Catalina Ballroom

Saturday, May 18, 2013

7:30-9:00 am	Continental Breakfast, Sonoran Ballroom Rooftop
8:00-10:15 am	Poster Session 2 Even Numbered Posters Present
8:30 - 9:30 am	Moderated Poster Discussion 3 Moderator- Christopher Wendler, PhD., University of Florida

M3.1. TGFβ and Hedgehog signaling in flow-induced EndoMT;Role of endothelial primary cilia.

Beerend P Hierck¹, Gonzalo Sánchez-Duffhues³, Marie-José Goumans³, Peter ten Dijke³, Robert E Poelmann¹, Anastasia D Egorova^{1,2} Leiden University Medical Center

M3.2. Endocardial Brg1 loss-of-function studies distinguishbetween origins of semilunar valve defects and mechanisms of disease progression

Brynn N. Simek¹, Maithri Sarangam¹, Vidusha Devasthali¹, Ching-Pin Chang², Bin Zhou³, Kryn Stankunas_University of Oregon

M3.3. TGFβ2 non-canonical effectors, Src and p38 MAPK, can mediate epicardial EMT.

Patrick Allison¹, Derrick Broka¹, Todd Camenisch^{1,2,3,4} University of Arizona

M3.4. A novel role for Class II Small Leucine Rich Proteoglycans (SLRP) in maturation of the Cardiovascular ECM During Valvulogenesis

Loren E. Dupuis¹, Matthew G. Berger¹, Vennece Fowlkes¹, Shukti Chakravarti², Marian F. Young³, and Christine B. Kern¹ Medical University of South Carolina

M3.5. YAP1 is required for endothelial-mesenchymal transition of the atrioventricular cushion

Hui Zhang¹, Alexander von Gise^{2,6}, Tianyuan Hu¹, Qiaozhen Liu¹, Xueying Tian¹, Xiuzhen Huang¹, Liang He¹, Bin Zhao³, Chen-Leng Cai⁴, Fernando D. Camargo⁵, William Pu², Bin Zhou¹
Chinese Academy of Sciences, Shanghai

M3.6. Bmp signaling and epicardial contribution to the atrioventricular function

Marie M. Lockhart¹, Aimee L. Phelps¹, Maurice J. van den Hoff², John B. Burch³, Andy Wessels¹
Medical University of South Carolina

9:00-10:00 Moderated Poster Discussion 4

Moderator- Tony Firulli, PhD, Indiana University

M4.1. Role of PINK1 in mitochondrial homeostasis and cardiac aging

Sreehari Kalvakuri, Takeshi Akasaka and Rolf Bodmer Sanford-Burnham Medical Research Institute

M4.2. A Bmp-responsive *Hand1 cis*-regulatory element drives gene expression in the developing outflow tract

Joshua W. Vincentz and Anthony B. Firulli Indiana University

M4.3. Venous endothelium-derived endothelin directs sympathetic innervation of the heart

Eleana Manousiouthakis, Monica Mendez, and Takako Makita UCLA Keck School of Medicine

M4.4. Cardiac laterality, migration and morphogenesis depend upon small heat shock proteins

Mark Springel¹, Jonathan Wosen¹, Yongchang Ji³, Jonah Zuflacht¹, Paloma Marin¹, Jamie Lahvic¹, Lara Hutson¹, Jeff Amack³ and Martha Marvin²

¹Department of Biology and ²Neuroscience Program, Williamstown, MA ³ Department of Cell and Developmental Biology, State University of New York Upstate Medical University, Syracuse, NY

M4.5 Interrogating translating RNAs and cell-type specific gene expression using Cre- activated translating ribosome affinity purification

Pingzhu Zhou¹, Yijing Zhang¹, Qing Ma¹, Fei Gu¹, Daniel S. Day^{2,3}, Aibin He¹, Bin Zhou⁴, Jing Li⁵, Sean Stevens¹, Daniel Romo⁵ and William T. Pu^{1,6}
Boston Children's Hospital

M4.6 Conserved requirements for Hsp23/Hspb1 in cardiac tolerance to ischemia/reperfusion

Sarah Piloto and Rolf Bodmer Sanford-Burnham Medical Research Institute

Saturday Morning Platform Session 5: Valve Development, Sonoran Ballroom

Session Chairs Mohamad Azhar and Andrea Ladd

10:20-10:40 S5.1. Genetic and Cellular Determinants of Mitral Valve Prolapse in Humans

Kimberly Sauls1, Annemarieke de Vlaming1, Katherine Williams1, Roger Markwald1, Ken Irvine2, David Peal3, David Milan3, Ronen Durst4, Robert Levine5, Susan Slaugenhaupt6, Chip Norris1

1- Department of Regenerative Medicine and Cell Biology, College of Medicine, Medical University of South Carolina, Charleston, SC,2- Department of Molecular Biology and Biochemistry, Waksman Institute & Howard Hughes Medical Institute, Rutgers University, Piscataway, NJ, 3- Cardiovascular Research Center, Cardiology Division, Boston, Massachusetts General Hospital, Boston, Massachusetts, , 4- Department of Cardiology, Hadassah Hospital, Israel, 5- Department of Cardiology, Massachusetts General Hospital, Boston, Massachusetts, 6- Center for Human Genetic Research, Massachusetts General Hospital/Harvard Medical School, Boston, Massachusetts

10:40-11:00 S5.2. Endocardial Loss of Hand2 Function Disrupts Notch Signaling Causing Hypotrabeculation and Tricuspid Atresia

Nathan J. VanDuseni, Joshua W. Vincentzi, Beth A. Firullii, Joaquim Grego Bessas, Hanying Cheni, Wenjun Zhangi, Bin Zhouz, José Luis De La Pompa4, Weinian Shoui, and Anthony B. Firullii

Riley Heart Research Center Department of Pediatrics, Wells Center for Pediatric Research, Indiana University, Indianapolis, IN, 2Department of Genetics, Albert Einstein College of Medicine, New York, NY, 3Department of Developmental Biology, Memorial Sloan Kettering Cancer Center, New York, NY, 4Cardiovascular

Developmental Biology Program, Cardiovascular Development and Repair Department, Centro Nacional de Investigaciones Cardiovasculares (CNIC), Madrid, Spain

11:00-11:20 S5.3. MEF2C regulates expression of components of the Nodal signaling pathway in the outflow tract and is required for proper alignment of the aorta and pulmonary artery

Ralston M. Barnesi, Ian S. Harrisi, Eric J. Jaehnigi, William Schachterlei, David J. McCulleyi, Anabel Rojasi, Linh H. Vongz, John J. Schwartzz, and Brian L. Blacki Cardiovascular Research Institute, University of California, San Francisco, San Francisco, CA, 2Albany Medical College, Albany, NY USA

11:20-11:40 S5.4. The bHLH transcription factor Twist1a functions to limit cardiomyocyte production in zebrafish

Kristina M. Garske, Yocheved L. Schindler, and Deborah Yelon Division of Biological Sciences, University of California, San Diego, La Jolla, CA

11:40-12:00 S5.5. Olfactomedin-1 activity identifies a cell invasion checkpoint during epithelial-mesenchymal transition in the embryonic heart

Alejandro Lencinas 1.2, Danny C. Chhuni, Kelvin P. Dani, 3, Kristen D. Rossi, Elizabeth A. Hooveri, Parker B. Antini and Raymond B. Runyani. Departments of Cellular and Molecular Medicinei, Pharmacology and Toxicology and Program in Physiological Sciences. The University of Arizona, Tucson Arizona

12:00 - 1:30 pm Box lunch with NIH Presentation by Charlene Schramm (NHLBI)

Sonoran Ballroom

Saturday Afternoon Platform Session 6: Heart Formation and Progenitors

Sonoran Ballroom

1:30-1:50pm S6.1. Numb family proteins are essential for cardiac morphogenesis and progenitor differentiation

Hua Guoi; Chen Zhaoi; Thomas Myinti; William Pittmani; Le Yangi; Sheng Weiz; Duan Maz; John Schwarzi; Weimin Zhongi; Robert J. Schwartzi; Harold Singeri; Michelle Tallquists; and Mingfu Wui.

Cardiovascular Science Center, Albany Medical College, Albany N, 2 Key Laboratory of Molecular Medicine, Fudan University, Shanghai, China; 3Molecular, Cellular and Developmental Biology, Yale University, New Haven, CT; 4Biology and Biochemistry, University of Houston, Houston, Texas; 5JABSOM, Center for Cardiovascular Research, University of Hawaii, Honolulu, HI

1:50-2:10 S6.2. Notch Activation by Delta4 and Jagged1 Ligands in the Heart: A Story through Time and Space

Gaetano D'Amatoi, Gonzalo del Montez, Stanislao Travisanoi, Guillermo Luxáni and José Luis de la Pompai

Department of Cardiovascular Developmental Biology, Centro Nacional de Investigaciones Cardiovasculares (CNIC), Madrid SPAIN, 2Department of Developmental and Stem Cell Biology, The Victor Chang Cardiac Research Institute, Darlinghurst NSW

2:10-2:30 S6.3. Gaint11, identified in congenital heart disease, is a novel glycosylator of Notch and specifies cilia type in left-right patterning

Marko T. Boskovskii*, Shiaulou Yuani*, Nis Borbye Pedersenz, Christoffer Knak Gothz Svetlana Makovai, Henrik Clausenz, Martina Brueckneri, Mustafa K. Khokhai Program in Vertebrate Biology, Department of Pediatrics and Genetics, Yale University, School of Medicine, 333 Cedar Street, New Haven CT, 3Copenhagen Center for Glycomics, Department of Cellular and Molecular Medicine, Faculty of Health Sciences, University of Copenhagen, Copenhagen, Denmark

2:30-2:50 S6.4. Unilateral Dampening of Bmp Activity by Nodal Generates Cardiac Left-Right Asymmetry

Justus Veerkampi, Franziska Rudolphi, Zoltan Cseresnyesi.2, Florian Prilleri, Cécile Otteni, Marc Renzi and Salim Abdelilah-Seyfriedi
Cardiovascular and Metabolic Disease Program, Max Delbrück Center for Molecular Medicine, Berlin, Germany, 2German Rheumatism Research Center (DRFZ), a Leipniz Institute, Berlin, Germany

2:50-3:10 S6.5. hERG/seizure K+-channel mutants exhibit Wnt/Pygopusassociated cardiac remodeling in *Drosophila*

Karen Ocorri, Alex Zambonz, Yoav Nudelli, Min Tang3, Xiushan Wu3, Rolf Bodmeri Development and Aging Program, Sanford-Burnham Medical Research Institute, 2Dept. of Pharmacology, University of California San Diego, La Jolla, CA, 3The Center for Heart Development, College of Life Science, Hunan Normal University, Changsha, P.R.China.

3:10-3:30 Coffee Break

Saturday Afternoon Platform Session 7: Cardiac Growth & Regeneration, Sonoran Ballroom

Session Chairs Caroline Burns and Chuck Murry

3:30-3:50pm S7.1. miR-17-92 cluster is required for and sufficient to induce cardiomyocyte proliferation in postnatal and adult hearts

Jinghai Cheni, Zhan-Peng Huangi, Hee Young Seoki, Jian Dingi, Masaharu Kataokai, Zheng Zhangi, Xiaoyun Hui, Gang Wangi, Zhiqiang Lini, Si Wangi, Willam T. Pui,2, Ronglih Liao2,3 and Da-Zhi Wangi,2* Department of Cardiology, Boston Children's

Hospital, Harvard Medical School, Boston, 2Harvard Stem Cell Institute, Harvard University Cambridge, 3Brigham and Women's Hospital, Boston

3:50-4:10 S7.2. Notch Signaling Regulates Cardiomyocyte Proliferation during Zebrafish Heart Regeneration

Long Zhaoi, Burcu Guner-Atamani, Kazu Kikuchiz, Kenneth D. Possz, Caroline E. Burns*1,3,4, & C. Geoffrey Burns*1,3,4

1 Cardiovascular Research Center, Massachusetts General Hospital, Charlestown, MA

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- 4:10-4:30 **S7.3. Identification of chemical modifiers of heart regeneration**Wen-Yee Choi, Jennifer Holdway, and Kenneth Poss Department of Cell Biology,
 Howard Hughes Medical Institute, Duke University Medical Center, Durham, NC
- 4:30-4:50 S7.4 The Mesp1 Lineage of Cardiac Progenitor Cells Contribute to Neovascularization and Functional Improvement in Post-MI Mouse Hearts

Yu Liui, Li Cheni, Andrea Diaz Diazz, Xueping Xu3, Austin Cooney3, Bradley McConnell2, and Robert Schwartz1, 4

¹Department of Biology and Biochemistry, ²Department of Pharmacological and Pharmaceutical Sciences, University of Houston, Houston TX, ³Department of Molecular and Cellular Biology, Baylor College of Medicine, Houston TX, 4Stem Cell Engineering, Texas Heart Institute, Houston TX

4:40-5:10 S7.5 Wnt6 and sfrp1 regulate heart development and cardiomyocyte differentiation via a negative-feedback regulatory loop

Natalie Gibb and Stefan Hoppler
Aberdeen Developmental Biology Group and Cardiovascular Research Programme,
Institute of Medical Sciences, Foresterhill Health Campus, University of Aberdeen,
Scotland, UK

Banquet w/ Awards and Music by Code Orange

6:00 -