THOMAS L. PETTY ASPEN LUNG CONFERENCE

66th Annual Meeting "Pulmonary Fibrosis – Focusing on the Future" June 4-7, 2024

Monday, June 3, 2024 -- Evening

5:00-7:00 PM Evening Registration

Gant Conference Center

Tuesday, June 4, 2024 – Morning

8:00-8:20 AM Welcome/Introduction

Elizabeth Redente, Ph.D., Chair David Schwartz, M.D., Co-Chair

8:20-8:30 AM The Thomas L. Petty Aspen Lung Conference: A Historical Perspective

Dennis E. Doherty, M.D., FCCP Professor of Medicine/University of Kentucky Secretary/Treasurer, National Lung Health Education Program

Theme 1/Session 1: Common Thread of ILA Relationship to Established Lung Fibrosis Moderators--

8:30-9:05 AM

THOMAS L. PETTY LECTURE "WHAT CAN WE LEARN FROM ILAS AND DISEASE DIAGNOSIS AND PROGRESSION?"

Fernando J. Martinez, M.D., M.S. Chief, Division of Pulmonary and Critical Care Medicine **Bruce Webster Professor of Medicine** Joan and Sanford I. Weill Department of Medicine Weill Cornell Medicine in New York New York, New York

9:05-9:30 AM **Discussion**

9:30-9:45 AM

GENOME-WIDE ASSOCIATION STUDY OF IDIOPATHIC PULMONARY FIBROSIS AMONG ASIAN ANCESTIRES. Anna L. Peljto^{1*}, Deepa Puthenvedu¹, Haruhiko Furusawa², Jonathan Cardwell¹, Masaki Hirose³, Yoshikazu Inoue³, Dong Soon Kim⁴, Yasunari Miyazaki², Ken Ohta⁵, Shin Ohta⁶, Tsukasa Okamoto², Jong Sun Park⁷, Moo Suk Park⁸, Jin Woo Song⁴, Ivana V. Yang¹, Tasha E. Fingerlin⁹, David A. Schwartz¹, ¹University of Colorado Anschutz Medical Campus, Aurora, CO; ²Tokyo Medical and Dental University, Tokyo, Japan; ³NHO Kinki Chuo Chest Medical Center, Osaka, Japan; ⁴Asan Medical Center, University of Ulsan, Seoul, Republic of Korea; ⁵National Hospital Organization Tokyo National Hospital, Tokyo, Japan; ⁶Showa University, Tokyo, Japan; ⁷Seoul National University College of Medicine, Seoul National University Bundang Hospital, Seongnam, Republic of Korea; ⁸Severance Hospital, Yonsei University College of Medicine, Seoul, Republic of Korea; ⁹National Jewish Health, Denver, CO.

9:45-9:10:00 AM CELL SPECIFIC MOLECULAR PROFILING OF SCLERODERMA ASSOCIATED INTERSTITIAL LUNG DISEASE SUBTYPES. Monica Yang^{1*}, Fred Deiter², Emily Flynn¹, Jessica Neely³, Seoyeon Lee², John Greenland², Marina Sirota⁴, Paul Wolters², ¹Division of Rheumatology, Department of Medicine, University of California, San Francisco; ²Division of Pulmonary, Critical Care, Allergy and Sleep Medicine, Department of Medicine, University of California San Francisco; ³Division of Pediatric Rheumatology, Department of Pediatrics, University of California, San Francisco; ⁴Bakar Computational Health Sciences Institute, University of California, San Francisco.

10:00-10:30 AMCoffee Break MEET THE PROFESSOR SESSION (by Registration table) (Refreshments for conference participants only)

Tuesday, June 4, 2024 -- Morning

Theme 2: Exploring the Dynamic Cellular and Structural Biology of Lung Fibrosis

Session 2: Biological Interface and Interaction in Interstitial Lung Disease

Moderators--

10:30-11:05 AM

PARKER B. FRANCIS LECTURESHIP
"DEVELOPMENT OF LUNG FIBROSIS:
HOW CELLS SIGNAL, CHANGE AND EMERGE"
Harold A. Chapman, M.D.
Professor of Medicine
Department of Pulmonary and Critical Care Medicine
University of California San Francisco
San Francisco, California

11:05-11:30 AM Discussion

11:30-11:45 AM ELUCIDATING THE FUNCTIONAL ROLE OF FIBROBLAST PROLIFERATION IN LUNG FIBROSIS VIA MURINE MODELS AND PRECISION-CUT HUMAN LUNG SLICES. Christopher Molina*, Dean Sheppard, Department of Pulmonary and Critical Care Medicine, University of California San Francisco, CA.

11:45-12:00 Noon ROLE OF GATA6 IN ALVEOLAR FIBROBLAST FUNCTION. J. Green, M.G. Ushakumary, C. Na, Anna-Karina Perl*, Division of Pulmonary. Biology, Cincinnati Children's Hospital; Department of Pediatrics, Univ. of Cincinnati College of Medicine, Cincinnati, OH.

12:00-1:30 PMLunch (lunch not provided by conference)

Tuesday, June 4, 2024 -- Afternoon

<u>Session 3: Biological Interface and Interaction in Interstitial Lung Disease – Re-Building the Matrix</u> Moderators--

1:30-2:05 PM STATE OF THE ART

Kristi S. Anseth, Ph.D.

University of Colorado at Boulder, Boulder, Colorado

"The Development of Biomaterials to Serve as Synthetic Extracellular Matrix (ECM) to Rebuild the Lung"

2:05-2:30 PM Discussion

2:30-2:45 PM

TRPV4 IS A KEY MECHANOSENSOR IN MACROPHAGES THAT DRIVES MYOFIBROBLAST DIFFERENTIATION THROUGH THE SECRETION OF ACTIVE TGF-β. Rachel G. Scheraga^{1,2}*, L.M. Grove², S. Abraham², B.D. Southern^{1,2}, A. Reinhardt², E.M. Orsini¹, M.A. Olman^{1,2}, ¹Departments of Pulmonary and Critical Care and ²Inflammation and Immunity, Cleveland Clinic, Cleveland, OH.

2:45-3:00 PM PIEZO2 IS AN IMPORTANT MECHANO-RECEPTOR IN PULMONARY FIBROSIS.

Margaret A.T. Freeberg*, S.V. Camus, T.H. Thatcher, P.J. Sime Department of Internal Medicine, Virginia Commonwealth University, Richmond, VA.

3:00-3:30 PMBreak (Refreshments for conference participants only)

<u>Session 4: Biological Interface and Interaction in Interstitial Lung Disease – Re-Building the Endothelium</u> Moderators--

3:30-4:05 PM STATE OF THE ART

Tatiana V. Kalin, M.D., Ph.D.

Phoenix Children's Center for Cancer and Blood Disorders University of Arizona College of Medicine – Phoenix, Arizona

"Regeneration of the Vascular Niche During Fibrosis Repair"

4:05-4:30 PM Discussion

4:30-4:45 PM

THE ROLE OF ENDOTHELIAL MECHANOTRANSDUCTION IN PULMONARY FIBROSIS. Patricia Brazee*, Shatruhan Rajput, Ilyaas Sugal, Trong Nguyen, Katharine Ference, Rachel Knipe, Division of Pulmonary and Critical Care Medicine, Center for Immunology and Inflammatory Diseases, Massachusetts General Hospital, Boston, MA.

4:45-5:00 PM

VASCULAR BED TRANSCRIPTIONAL ACTIVATION CHARACTERIZES RAPID PROGRESSORS IN IDIOPATHIC PULMONARY FIBROSIS. Sari Ezgi¹*, N.S. Sharma².³, K. Patel⁴, S. Shankar⁴, M.G. Gastanadui⁵, D. Moncada Giraldo⁶, Y. Soto-Vazquez⁵, D. Stacks¹, L. Hecker¹, K. Dsouza⁴, M. Banday², E. OʻNeill⁴, P. Benson¹, G. Payne⁵,
ß, A. Gaggar⁵,
ß, C. Margaroli¹, ¹Department of Pathology, University of Alabama at Birmingham, Birmingham, Alabama; ²Department of Medicine, Brigham and Women's Hospital, Boston, Massachusetts; ³West Roxbury VA Medical Center, Boston, Massachusetts; ⁴Department of Medicine, University of South Florida, Tampa, Florida; ⁵Department of Medicine, University of Alabama at Birmingham, Birmingham, Alabama; ⁶Department of Pediatrics, Emory University, Atlanta, Georgia; ⁶Birmingham VA Medical Center, Birmingham, Alabama.

5:00-7:00 PM POSTER VIEWING (Refreshments for conference participants only)

Wednesday, June 5, 2024 -- Morning

<u>Session 5: Biological Interface and Interaction in Interstitial Lung Disease – Re-Building Stem Cell Niches</u> <u>Moderators--</u>

8:00-8:35 AM

ROGER S. MITCHELL LECTURE "REGENERATION OF THE STEM CELL NICHE TO DRIVE REPAIR"

Xin Sun, Ph.D.
Professor of Pediatrics
Department of Cell and Development Biology
School of Biological Sciences
University of California San Diego
San Diego, California

8:35-9:00 AM Discussion

- 9:00-9:15 AM TYPE 2 INNATE IMMUNITY PROMOTES THE DEVELOPMENT OF PULMONARY FIBROSIS IN HERMANSKY-PUDLAK SYNDROME. Parand Sorkhdini¹, Kiran Klubock-Shukla¹, Dongqin Yang¹, Alina Xiaoyu Yang¹, Carmelissa Norbrun¹, Wendy J. Introne², Bernadette R. Gochuico², Yang Zhou¹*, Department of Molecular Microbiology and Immunology, Brown University, Providence, R.I. ²Medical Genetics Branch, National Human Genome Research Institute, Bethesda, Maryland.
- 9:15-9:30 AM MOVEMENT OF EPITHELIAL CELLS IS ASSOCIATED WITH THE EXTENT OF LUNG FIBROSIS IN IDIOPATHIC PULMONARY FIBROSIS (IPF) Andrey Krivoy^{1*}, Evgenia Dobrinskikh¹, Seyedeh Zahra Fotook Kiaei¹, Ian Stancil^{1,2}, Janna Brancato¹, Ivana V. Yang¹, David A. Schwartz¹, ¹Division of Pulmonary Sciences and Critical Care Medicine; University of Colorado, Denver, CO; ²Division of Pulmonary and Critical Care Medicine, Stanford University, Palo Alto, CA.

9:30-10:00 AMCoffee Break MEET THE PROFESSOR SESSION (by Registration table) (Refreshments for conference participants only)

Wednesday, June 5, 2024 -- Morning

<u>Theme 3: Diagnostic Challenges and Novel Approaches – The Road to Understanding Interstitial Lung Abnormalities, UIPs and Disease Progression</u>

Session 6: AI Integration to Enhance Early Detection Using Integrated Omics Moderators--

10:00-10:35 AM STATE OF THE ART

Casey S. Greene, Ph.D.

University of Colorado School of Medicine, Aurora, Colorado "The Integration of Omics Data to Model and Understand the Biology of ILDs"

10:35-11:00 AM Discussion

11:00-11:15 AM SKIN TRANSCRIPTOMICS ARE ASSOCIATED WITH LUNG FUNCTION IMPAIRMENT AND DISTINCT CELL TYPE ENRICHMENT IN SYSTEMIC SCLEROSIS-RELATED LUNG DISEASE. Jana Zielonka*, Ningshan Li, Zuoheng Wang, Xiting Yan, Jose L. Gomez, Yale University School of Medicine, Division of Pulmonary, Critical Care & Sleep Medicine, New Haven, CT.

11:15-11:30 AM LRP1 DEFICIENCY CHANGES LIPID METABOLISM AND AGGRAVATES LUNG FIBROSIS. F. Schramm¹, L. Wujak², S. Hadzic³, K. Rubio⁴, T.O. Eichmann⁵, W. Sattler⁵, S. Günther⁶, J. Wilhelm^{1,7}, I. Alexopoulos^{1,7}, B. Kojonazarov³, G. Barreto⁴, Malgorzata Wygrecka^{1,7*}, ¹Center for Infections and Genomics of the Lung, German Center for Lung Research, Justus Liebig University, Giessen, Germany; ²MedComms Warsaw, Warsaw, Poland; ³Excellence-Cluster Cardio-Pulmonary Institute, Justus Liebig University, Giessen, Germany; ⁴Universite de Lorraine, Nancy, France. ⁵Medizinische Universität Graz, Graz, Austria; ⁶German Centre for Cardiovascular Research, Bad Nauheim, Germany; ⁷Institute for Lung Health, Justus Liebig University, Giessen, Germany.

12:00-3:00 PM Picnic – T Lazy 7 - The Ranch (for conference participants and their family)

Thursday, June 6, 2024 -- Morning

<u>Session 7: Common Threads to Usual Interstitial Pneumonitis: Tying it All Together</u> <u>Moderators--</u>

8:00-8:35 AM

MARVIN I. SCHWARZ LECTURE
"ROLE OF TRANSITIONAL EPITHELIAL CELLS
IN LUNG REGENERATION AND FIBROSIS"

Rachel L. Zemans, M.D.
Henry Sewall Research Professor of
Pulmonary and Critical Care Medicine
Professor of Internal Medicine and
Cellular & Molecular Biology
University of Michigan
Ann Arbor, Michigan

8:35-9:00 AM Discussion

9:00-9:15 AM

HOW SHOULD PATIENTS WITH PROGRESSIVE PULMONARY FIBROSIS BE IDENTIFIED? CONSENSUS FINDINGS FROM A MODIFIED DELPHI STUDY. Athol U Wells^{1,2*} Simon LF Walsh,² Ayodeji Adegunsoye,³ Vincent Cottin,⁴ Sonye Danoff,⁵ Anand Devaraj,^{1,2} Kevin R Flaherty,⁶ Peter M George,^{1,2} Martin Kolb,⁷ Yasuhiro Kondoh,⁸ Andrew G Nicholson,¹ Sara Tomassetti,⁹ Elizabeth R Volkmann,¹⁰ Kevin K Brown¹¹, ¹Royal Brompton and Harefield Hospitals, London, UK; ²National Heart and Lung Institute, Imperial College London, London, UK; ³University of Chicago, Chicago; ⁴National Reference Center for Rare Pulmonary Diseases, Louis Pradel Hospital, Claude Bernard University Lyon 1, Lyon, France; ⁵Johns Hopkins Medicine, Baltimore; ⁶University of Michigan, Ann Arbor; ⁷McMaster University and St. Joseph's Healthcare, Hamilton, Canada; ⁸Tosei General Hospital, Aichi, Japan; ⁹Florence University, Florence, Italy; ¹⁰University of California, David Geffen School of Medicine, Los Angeles; ¹¹National Jewish Health, Denver.

9:15-9:30 AM

LUNG MICROENVIRONMENT LABEL-FREE *QUANTITATIVE* **PROTEOMICS** IDENTIFIES DISTINCT ENDOPHENOTYPES IN IDIOPATHIC PULMONARY FIBROSIS. L. T. Ngo¹, M. Rekowski², D. Koestler³, I. Azeem¹, A. Harrison¹, M.K. Demoruelle⁴, J. Boomer¹, B.R. England⁵, P. Wolters⁶, P. Molyneux⁷, M. Castro¹, J.S. Lee⁸, J.J. Solomon⁹, K. Koronuma¹⁰, M.P. Washburn², Scott M. Matson^{1*}, ¹Division of Pulmonary, Critical Care and Sleep Medicine, University of Kansas School of Medicine; ²Department of Cancer Biology, KUMC; ³Department of Biostatistics KUMC; ⁴Division of Rheumatology, University of Colorado; ⁵Division of Rheumatology & Immunology, University of Nebraska Medical Center; ⁶Division of Pulmonary and Critical Care Medicine, UCSF; ⁷National Heart and Lung Institute, Imperial College London; ⁸Division of Pulmonary Sciences and Critical Care Medicine, University of Colorado: 9National Jewish Health Hospital: 10Department of Respiratory, Sapporo Medical University School of Medicine, Sapporo, Japan.

9:30-10:00 AMCoffee Break MEET THE PROFESSOR SESSION (by Registration table) (Refreshments for conference participants only)

Thursday, June 6, 2024 -- Morning

<u>Session 8: Non-Invasive Molecular Imaging and Quantifying Lung Fibrosis</u> <u>Moderators--</u>

10:00-10:35 AM STATE OF THE ART

Sydney B. Montesi, M.D.

Massachusetts General Research Institute, Harvard Medical School, Boston, Massachusetts "Harnessing the Power of Early Detection Through Molecular Imaging by PET/CT and PET/MRI"

10:35-11:00 AM Discussion

11:00-11:15 AM

QUANTIFYING AND EVALUATING TGFβ1 NON-INVASIVELY USING THE NOVEL ⁸⁹Zr DFO-TGFβRII PET IMAGING. Yujun Zhang^{1,2*}, Jessy Deshane³, Tejaswini Kulkarni³, Benjamin Larimer^{1,4}, ¹Graduate Biomedical Sciences, The University of Alabama at Birmingham, AL; ²Department of Radiology, The University of Alabama at Birmingham, AL; ³Division of Pulmonary, Allergy and Critical Care Medicine, The University of Alabama at Birmingham, AL; ⁴O'Neal Cancer Center, AL.

11:15-11:30 AM

DATA-DRIVEN TEXTURAL ANALYSIS (DTA) FIBROSIS SCORES FROM BASELINE TO 1-YEAR HRCT PREDICT SUBSEQUENT DISEASE PROGRESSION. Matthew Koslow^{1*}, J.J. Swigris¹, D. Baraghoshi³, M. Strand³, J. Solomon¹, E. Fernandez Perez¹, Z.X. Yunt¹, R. Keith¹, M.P. Mohning¹, T.J. Huie¹, K.K. Brown¹, D.A. Lynch², S.M. Humphries², Center for Interstitial Lung Disease, Division of Pulmonary, Critical Care and Sleep Medicine; National Jewish Health, Denver, CO, Department of Radiology, National Jewish Health, Denver, CO, Denver, CO, Division of Biostatistics and Bioinformatics, National Jewish Health, Denver CO.

11:30-1:30 PMLunch (lunch not provided by conference)

Thursday, June 6, 2024 -- Afternoon

Theme 4: Transformational Interventions for Lung Fibrosis
Session 9: Regenerative Capacity and Scar Removal in the Lungs
Moderators-

1:30-2:05 PM STATE OF THE ART

Kamran Atabai, M.D.

University of California San Francisco, San Francisco, California

"Age-Dependent Regulation of Cell-Mediated Collagen Turnover"

2:30-2:45 PM

DEXTROMETHORPHAN INHIBITS SECRETION OF PRO-FIBROTIC CARGOES ELICITING AN ANTI-FIBROTIC RESPONSE IN IN-VIVO, EX-VIVO AND IN-VITRO MODELS OF PULMONARY FIBROSIS. Muzamil M Khan^{1,2*}, Joanna Zukowska¹, Juan Jung¹, George Galea¹, Nadine Tuechler¹, Aliaksandr Halavatyi¹, Beate Neumann¹, Thomas Muley², Hauke Winter², Julia Duerr³, Marcus A Mall³, Ernesto de la Cueva¹, Mikhail Savitski¹, Rainer Pepperkok^{1,2}, ¹Cell Biology and Biophysics Unit, European Molecular Biology Laboratory, Heidelberg, Germany; ²Translational Lung Research Center Heidelberg, German Center for Lung Research (DZL), Heidelberg, Germany; ³Department of Pediatric Respiratory Medicine, Immunology and Critical Care Medicine, Charité-Universitätsmedizin Berlin, Germany.

2:45-3:00 PM

COLLAGEN PROLYL-3-HYDROXYLASE 1 IS INCREASED IN IDIOPATHIC PULMONARY FIBROSIS AND CONTROLS COLLAGEN QUALITY AND COMPOSITION. Claudia A. Staab-Weijnitz^{1*}, Juliane Merl-Pham², Vivek Sarohi³, Rachel Z. Blumhagen⁴, Leonhard Binzenhöfer¹, Karolina Pijadina¹, Marleen Stremlau¹, Elisabeth Hennen¹, Natalia Cabeza-Boeddinghaus¹, Ceylan Onursal¹, Jürgen Behr⁵, Anne Hilgendorff¹, Hans Peter Bächinger⁶, David A. Schwartz⁷, Ivana Yang⁷, Naftali Kaminski⁶, Stefanie M. Hauck², Oliver Eickelberg¹, Roberto Vanacore⁶, Trayambak Basak³, ¹Comprehensive Pneumology Center, Institute of Lung Health and Immunity, Helmholtz Munich, Germany; ²School of Biosciences and Bioengineering (SBB), Indian Institute of Technology (IIT), Mandi, Mandi, India; ⁴Center for Genes, Environment and Health, National Jewish Health, Denver, CO; ⁵University Hospital of the LMU, Munich, Germany; ⁶Department of Biochemistry and Molecular Biology, Oregon Health & Science University, Portland, OR; ¬Department of Medicine, University of Colorado Anschutz Medical Campus, Aurora, CO; ⁶Pulmonary, Critical Care and Sleep Medicine, Yale School of Medicine, New Haven, CT; ⁰Division of Nephrology and Hypertension, Department of Medicine, Vanderbilt University Medical Center, Nashville, TN.

3:00-3:30 PMBreak (Refreshments for conference participants only)

Thursday, June 6, 2024 -- Afternoon

<u>Session 10: Regenerative Capacity and Scar Removal in the Lung</u> <u>Moderators--</u>

3:30-4:05 PM

THOMAS A. NEFF LECTURE

"CELLULAR SENSING AND REMODELING OF THE LUNG MATRIX ENVIRONMENT"

Daniel J. Tschumperlin, Ph.D.
Professor and Vice Chair
Department of Physiology and Biomedical Engineering
Mayo Clinic
Rochester, Minnesota

4:05-4:30 PM Discussion

4:30-4:45 PM MECHANOSENSORY PRIMARY CILIA REWIRE LUNG FIBROBLAST METABOLISM TO

PROMOTE FIBROSIS. G.S. Ozcebe#¹, C.S. Trempus#¹, B. Papas¹, J.L. Li¹, M. Prates Mori¹, R. Snyder¹, R. Vancini¹, J. Watts¹, J. Santos¹, C.M. Hogaboam², D.A. Schwartz³, A. Mora⁴, M. Rojas⁴, Stavros Garantziotis^{1*}, ¹National Institute of Environmental Health Sciences; ²Cedars Sinai Medical Center; ³University of Colorado; ⁴Ohio State University. #equal contribution as first

authors

4:45-5:00 PM RECOGNITION OF CELL DEATH METABOLICALLY PRIMES LUNG FIBROBLASTS FOR

COLLAGEN SYNTHESIS AND PROLIFERATION. Hope Chatwin, Shannon McManus, Elizabeth Redente, Alexandra McCubbrey*, Department of Medicine, National Jewish Health,

Denver CO.

5:00-7:00 PM POSTER VIEWING (Refreshments for conference participants only)

Friday, June 7, 2024 -- Morning

<u>Session 11: Pharmacologic Targets Lead to Cures</u> <u>Moderators--</u>

8:00-8:35 AM

REUBEN M. CHERNIACK LECTURE
"LESSONS FROM CF: GENETIC TARGETING FOR
SUCCESSFUL PHARMACOLOGIC INTERVENTION
AND PARALLELS TO ILD DRUG DEVELOPMENT"

Paul A. Negulescu, Ph.D. Senior Vice President, Research Vertex Pharmaceuticals, Incorporated San Diego, California

8:35-9:00 AM Discussion

9:00-9:15 AM

ELEVATED SINE OCULIS HOMEOBOX HOMOLOG 1 (SIX1) IN FIBROTIC LUNG DISEASES, A POTENTIAL TARGET FOR GENE SILENCING? Cory Wilson¹, Sarah Shin², Scott Collum², Nancy Wareing³, Howard J Huang⁴, Bindu Akkanti⁵; Bela Patel⁵, **Harry Karmouty-Quintana**^{2,5*}, Department of Internal Medicine, University of Iowa; Iowa City, Iowa; Department of Biochemistry and Molecular Biology, UTHealth, Houston, TX; Department of Internal Medicine, Emory University, Atlanta, GA; Department of Pulmonary Critical Care, Houston Methodist, Houston TX; Divisions of Critical Care, Pulmonary and Sleep Medicine, Department of Internal Medicine, UTHealth, Houston, TX.

9:15-9:30 AM

SYNERGISTIC EFFECTS OF NINTEDANIB AND ABT-199 ON FIBROBLAST APOPTOSIS. **Joseph Cooley**^{1*}, J. Wilson², N. Javkhlan², D. A. Schwartz³, D. W. Riches², E. F. Redente², ¹Pulmonary and Critical Care, National Jewish Health, Denver, CO; ²Pediatrics, National Jewish Health, Denver, CO, United States; ³Dept of Med, Univ of Colorado, Aurora, CO.

9:30-10:00 AMCoffee Break (Refreshments for conference participants only)

Friday, June 7, 2027 – Morning

<u>Session 12: Cures Tissue Repair Through Rebuilding the Lung</u> Moderators:

10:00-10:35 AM

GILES F. FILLEY LECTURE
"THE FUTURE OF LUNG REGENERATION
AND TRANSPLANTATION"

Thomas Petersen, M.D., Ph.D.
Vice President, Regenerative Medicine
United Therapeutics
Durham, North Carolina

10:35-11:00 AM Discussion

11:00-11:15 AM

MUC5B SUPPRESSES ALVEOLAR REGENERATION. Yan Hu^{1*}, Naoko Liu¹, James Needell¹, Melanie Königshoff², Christopher Evans¹, ¹Division of Pulmonary Sciences and Critical Care Medicine, University of Colorado School of Medicine, Aurora, CO, ²Division of Pulmonary, Allergy, and Critical Care Medicine; Department of Medicine, University of Pittsburgh, Pittsburgh, PA.

11:15-11:30 AM

LUNG CELL TRANSPLANTATION FOR PULMONARY FIBROSIS. Irit Milman Krentsis^{1*}, Yangxi Zheng¹, Christa Blagdon¹, Sarah Y. Shin³, Sandeep K. Yadav¹, Esther Bachar Lustig¹, Chava Rosen^{1,2}, Eli Shezen¹, Einav Shoshan¹, Burton F. Dickey⁴, Harry Karmouty-Quintana³, Yair Reisner¹, ¹Department of Stem Cell Transplantation and Cell Therapy, MD Anderson Cancer Center, Houston, TX; ²Department of Neonatology, Children's Hospital, Sheba Medical Center, Tal Hashomer, Israel; ³Department of Internal Medicine, The University of Texas Health Science Center, Houston, TX; ⁴Department of Pulmonary Medicine, MD Anderson Cancer Center, Houston, TX.

11:30-12:30 PM

CONFERENCE SUMMARY

Patricia J. Sime, M.D.
Chair, Department of Internal Medicine
William Branch Porter Professor of Medicine
Division of Pulmonary Diseases and Critical Care Medicine
Virginia Commonwealth University
Richmond, Virginia

12:30-1:00 PM Discussion and Adjourn

POSTER VIEWING

Tuesday, June 4, 2024 5:00-7:00 PM

POSTERS

ALDEHYDE DEHYDROGENASE 2 MITIGATES LUNG FIBROSIS AFTER INJURY. Yael Aschner^{1*}, Elisabeth Murphy², Kelly A Correll², Keriann Beke², Paul R Reynolds², Gregory P Downey^{1,2}, Dept of Medicine, University of Colorado Anschutz Medical Campus, Aurora, CO, Dept of Academic Affairs, National Jewish Health, Denver, CO.

DEVELOPMENT OF NOVEL TECHNOLOGY FOR THE VISUALIZATION AND QUANTITATION OF MICROVASCULAR PROGENITOR DRIVEN ADAPTIVE ANGIOGENESIS DURING CHRONIC LUNG DISEASE. Hannah Thorndyke¹*, Evan Lundberg¹, Maggie Dawson¹, Edwin Ortiz Gaxon², Emma Mason¹, Eszter Vladar³, David Coronado Escobar⁴, and Susan Majka¹,³,¹Department of Medicine, Division of Pulmonary & Critical Care Medicine, National Jewish Health, Denver CO; ²Cell Biology, Stem Cells & Development Graduate Program, University of Colorado Anschutz Medical Campus; ³Department of Medicine, Division of Pulmonary & Critical Care Medicine, University of Colorado Anschutz Medical Campus; ⁴Onimagin Technologies SCA, Cordoba, Spain.

PGK1 MEDIATES DEFICIENT PYRUVATE UTILIZATION AND BIOENERGETIC COMPROMISE IN LUNG FIBROBLASTS IN AGE-RELATED LUNG FIBROSIS. Pilar Londono, Emily Turner, Gavriel Roda, Christopher M. Evans and Sunad Rangarajan*, Division of Pulmonary Sciences and Critical Care, University of Colorado Anschutz Medical Campus, Aurora, CO.

LONG NON-CODING RNA IN IPF: REGULATORY PLAYERS IN LUNG FIBROSIS. Aileen C. Button^{1*}, I. V. Yang¹, D. A. Schwartz¹; ¹Department of Medicine, University of Colorado, Aurora, CO.

TARGETING A NOVEL TRPV4-PI3Kγ INTERACTION BLOCKS FIBROGENIC MYOFIBROBLAST DIFFERENTIATION. Lisa M. Grove¹, M.L. Mohan², K.D. Singh², S. Abraham¹, A. Reinhardt¹, H. Mao¹, R.G. Scheraga^{1,3}, B.D. Southern^{1,3}, S.V. Naga Prasad², S.S. Karnik², and **Mitch A. Olman^{1,3*}**, ¹Departments of Inflammation and Immunity and ²Cardiovascular and Metabolic Sciences of Lerner Research Institute and ³Respiratory Institute, Cleveland Clinic, Cleveland, Ohio.

CELL-CELL AND CELL-MATRIX INTERACTIONS IN BIOENGINEERED 3D MODELS OF LUNG FIBROSIS. Rachel Blomberg^{1*}, Mikala C. Mueller¹, David W.H. Riches^{2,3,4,5}, and Chelsea M. Magin^{1,3,6}, ¹Department of Bioengineering, University of Colorado Denver | Anschutz, ²Program in Cell Biology, Department of Pediatrics, National Jewish Health, ³Division of Pulmonary Sciences and Critical Care Medicine, Department of Medicine, University of Colorado Anschutz, ⁴Department of Research, Veterans Affairs Eastern Colorado Health Care System, ⁵Department of Immunology and Microbiology, University of Colorado Anschutz, ⁶Department of Pediatrics, University of Colorado Anschutz, Aurora, CO.

ENGINEERING TUNABLE STIFFNESS HYDROGELS TO MODEL FIBROTIC ALVEOLAR TRANSITIONAL CELLS AND STUDY HUMAN PULMONARY FIBROSIS. Alicia E. Tanneberger^{1*}, Rachel Blomberg¹, Amy L. Ryan², Chelsea M. Magin^{1,3,4}. ¹Department of Bioengineering, University of Colorado Denver Anschutz; ²Department of Anatomy and Cell Biology, Carver College of Medicine, University of Iowa, Iowa City, IA; ³Department of Pediatrics, University of Colorado, Anschutz Medical Campus, Aurora, CO; ⁴Division of Pulmonary Sciences & Critical Care Medicine, Department of Medicine, University of Colorado, Anschutz Medical Campus, Aurora, CO.

POSTERS – Tuesday, June 4, 2024 – continued

REPETATIVE O₃ EXPOSURE IN HYPERGLYCEMIC INSULIN RESISTANT MICE EXACERBATES LUNG INJURY AND FIBROSIS. **Robert M. Tighe^{1*}**, A.V. Vose¹, J.G. Wagner² and J.R. Harkema², ¹Duke University, Durham, NC ²Michigan State University, East Lansing, MI.

DEFINING THE EFFECT OF PDGFRa LUNG FIBROBLASTS ON ALVEOLAR EPITHELIAL CELLS USING TRANSCRIPTOMINC PROFILING. Carol S. Trempus^{1*}, Brian N. Papas², Erica Scappini³, Charles J. Tucker³, Deloris Sutton⁴, and Stavros Garantziotis¹, ¹Immunity, Inflammation, and Disease Laboratory, NIEHS, Research Triangle Park, NC; ²Biostatistics & Computational Biology Branch, NIEHS, Research Triangle Park, NC; ³Signal Transduction Branch, NIEHS, Research Triangle Park, NC; ⁴Comparative & Molecular Pathogenesis Branch, NIEHS, Research Triangle Park, NC.

SOX9 UPREGULATION IN THE PATHOGENESIS OF SEVERE FIBROTIC LUNG DISEASE. **Priyanka** Singh^{1*}, P. R. Gajjala¹, H. H. Ediga¹, V. Sontake², C. P. Vemulapalli¹, P.K. Patel¹, H. Miyazaki², D. Popov², S. Alisher¹, S. K. Huang³, M. S. Walters⁴, and S. K. Madala¹, ¹Division of Pulmonary, Critical Care and Sleep Medicine, University of Cincinnati, Cincinnati, OH; ²Gordian Biotechnology, South San Francisco; ³Division of Pulmonary and Critical Care Medicine, University of Michigan Medical School, Ann Arbor; ⁴Department of Medicine, Section of Pulmonary, Critical Care and Sleep Medicine, University of Oklahoma Health Sciences Center, Oklahoma.

CELL COMPETITION DRIVES BRONCHIOLIZATION AND PULMONARY FIBROSIS. Rachel Warren¹, Kylie Klinkhammer¹, Handeng Lyu¹, Changfu Yao², Barry Stripp² and **Stijn P. De Langhe^{1*}**, ¹Department of Medicine, Division of Pulmonary and Critical Medicine, Mayo Clinic, Rochester, MN; ²Women's Guild Lung Institute, Department of Medicine, Cedars-Sinai Medical Center, Los Angeles, CA.

GENETIC LOCUS ASSOCIATED WITH BLEOMYCIN INDUCED LUNG INJURY IN MICE. **Yingping Wang^{1*}**, Corinne Hennessy¹, Kristina Hatakka¹, Stephen Humphries², Evgenia Dobrinskikh², David Clouthier^{3†}, Ivana V. Yang^{1†}, David A. Schwartz^{1†}, ¹ Department of Medicine, University of Colorado School of Medicine, Aurora, Colorado; ² Department of Radiology, National Jewish Health, Denver, Colorado; ³ Department of Craniofacial Biology, University of Colorado School of Medicine, Aurora, Colorado. [†]Authors contributed equally.

'SILICOSARCOIDOSIS': IMPORTANCE OF SILICA IN GRANULOMATOUS LUNG FIBROSIS. Jeremy T. Hua^{1*}, Carlyne D. Cool², Einat Fireman Klein⁴, Lukas J. Lee⁵, Lauren M. Zell-Baran¹, Robert A. Cohen⁶, Richard Kraus¹, Charles Van Hook³, Cecile S. Rose¹, ¹Division of Environmental and Occupational Health Sciences, National Jewish Health, CO; ²Department of Pathology and ³Department of Medicine, University of Colorado, CO; ⁴Pulmonary Division, Lady Davis Carmel Medical Center, Faculty of Medicine Technion Institute of Technology, Haifa, Israel, ⁵Tao-Yuan General Hospital, Taiwan; ⁶Environmental and Occupational Health Sciences, School of Public Health, University of Illinois Chicago, IL.

ROLE OF αTAT1 IN THE MECHANOBIOLOGY OF LUNG FIBROSIS. **Ingo Ganzleben**^{1,2,3}*, Alyce Segal^{1,3}, Benjamin D. Medoff^{1,2,3}, ¹Division of Pulmonary and Critical Care Medicine – Massachusetts General Hospital; ²Center for Immunology and Inflammatory Diseases – Massachusetts General Hospital; ³Harvard Medical School, Boston, Massachusetts.

APOPTOTIC RESISTENCE IN COLLAGENIA1-EXPRESSING FIBROBLASTS DRIVES SILICA-INDUCED PULMONARY FIBROSIS. **Daniel G. Foster***, N. Javkhlan, J. Wilson, B. L. Edelman, D. W. H. Riches, E. F. Redente; Pediatrics, National Jewish Health, Denver, CO.

POSTERS – Tuesday, June 4, 2024 – continued

ENDOTHELIAL SIPRI SUPPORTS ALVEOLAR EPITHELIAL REPAIR. **Patricia L Brazee**^{1*}, K.G. Ference ¹, A. Pickering², T.G. Kooistra¹, T. Hla³, B.D. Medoff¹, R.S. Knipe¹, ¹Massachusetts General Hospital, Division of Pulmonary and Critical Care Medicine, Boston, MA; ²Harvard Medical School, Department of Biomedical Informatics; ³Boston Childrens Hospital, Department of Surgery, Boston, MA.

SPATIAL AND TRANSCRIPTIONAL FEATURES IN IDIOPATHIC PULMONARY FIBROSIS (IPF) ASSOCIATED WITH THE MUC5B PROMOTER VARIANT. Rachel Z. Blumhagen^{1*}, Jonathan S. Kurche^{2,3}, Carlyne D. Cool^{4,5}, David Heinz⁵, David A. Schwartz^{2,**}, Ivana V. Yang^{6**}, ¹Center for Genes, Environment and Health, National Jewish Health, ²Rocky Mountain Regional Veteran's Administration Medical Center, ³Division of Pulmonary Sciences and Critical Care Medicine, University of Colorado Anschutz Medical Campus, ⁴Department of Pathology, University of Colorado Anschutz Medical Campus, ⁵Pathology Laboratory, National Jewish Health, ⁶Department of Biomedical Informatics, University of Colorado Anschutz Medical Campus **authors contributed equally

REDOX HETEROGENEITY IN FIBROBLASTS DURING FIBROSIS PROGRESSION. Patrick A. Link^{1*}, Jeffrey A. Meridew¹, Nunzia Caporarello², Ashley Y. Gao¹, Victor Peters³, Mauricio Rojas³, Daniel J. Tschumperlin¹, ¹Department of Physiology and Biomedical Engineering, Mayo Clinic, Rochester, MN; ²Department of Medicine, Loyola University, Chicago, IL; ³Department or Internal Medicine, Ohio State University, Columbus, OH.

SIGLECF KNOCKOUT MICE ARE PROTECTED FROM DEVELOPMENT OF LUNG FIBROSIS. Marika Orlov^{1*}, Naoko Liu¹, Kenny Ngo¹, James Needell¹, Fan Jia², Brian Vestal², Rachel Blumhagen², Jazalle McClendon³, William Janssen³, and Christopher Evans¹, ¹Division of Pulmonary Science and Critical Care Medicine, University of Colorado, Aurora, CO, ²Center for Genes, Environment and Health, National Jewish Health, Denver, CO ³Department of Medicine, National Jewish Health, Denver, CO.

UPREGULATION OF MARCKS ACTIVITY IN MACROPHAGE REPROGRAMMING AND ITS POTENTIAL AS A THERAPEUTIC TARGET IN PULMONARY FIBROSIS. Ching-Hsien Chen*, So-Yi Chang, Wen-Hsien Chang, Angela Linderholm, David C. Yang, Ssu-Wei Hsu, Reen Wu, Division of Pulmonary, Critical Care, and Sleep Medicine, Department of Internal Medicine, University of California Davis, Davis, CA.

POSTER VIEWING

Thursday, June 6, 2024 5:00-7:00 PM

POSTERS

BOOKMARKING DYSPNEA IN FIBROTIC INTERSTITIAL LUNG DISEASE. **Jeff Swigris***, Kerri Aronson, Michelle Kam, National Jewish Health, Center for Interstitial Lung Disease, Denver, CO.

WNT SIGNALING IN PULMONARY MICROVASCULAR PROGENITOR CELLS (MVPC) DRIVES ADAPTIVE ANGIOGENESIS DURING FIBROSIS AND REGULATES REPAIR OF THE ALVEOLAR-CAPILLARY UNIT. **Emma C Mason**^{1*}, Benjamin R. Schneider¹, Maggie Dawson¹, Evan Lundberg¹, David Cingel¹, Peter Kim¹, Kevin Kim¹, Edwin Ortiz Gaxon¹, Hannah Thorndyke¹, Elizabeth Redente², Patrick Geraghty³, M. Mark Taketo⁴ and Susan M Majka^{1, 5}, ¹ Department of Medicine, National Jewish Health, Denver, CO; ²Division of Cell Biology and Department of Pediatrics, National Jewish Health, Denver, CO; ³ State University of New York, Downstate Health Science University, Brooklyn, New York; ⁴ Kyoto University, Sakyo, Kyoto, Japan; ⁵ Gates Center for Regenerative Medicine & Stem Cell Biology, University of Colorado, Aurora, CO.

PATIENT'S VIEWS ON THE ASSESSMENT OF INTERSITIAL LUNG DISEASE-RELATED DYSPNEA. **Joseph B. Pryor**^{1*}, Dolly Kervitsky², Jeffrey J. Swigris³, ¹Division of Pulmonary and Critical Care, University of Colorado, Denver, CO; ²PFWarriors, ³Center for Interstitial Lung Disease, National Jewish health, Denver, CO.

INTERSTITIAL LUNG DISEASE EDUCATION IN PULMONARY FELLOWSHIP: SUPPORTING PATIENTS THROUGH THEIR ILLNESS JOURNEY. Samantha King^{1*}; Anna Neumeier^{1,2}; Bridget Graney^{1,2}; Tristan Huie^{1,3}, ¹Department of Medicine, University of Colorado School of Medicine, Aurora, CO; ²Department of Medicine, Denver Health Medical Center, Denver, CO; ³Department of Medicine, National Jewish Health, Denver, CO.

CIRCULATING BIOMARKERS IN CHILDREN AND ADOLESCENTS WITH FIBROSING INTERSTITIAL LUNG DISEASE (ILD). Robin Deterding^{1*}, Kevin K. Brown², Steven Cunningham³, Emily M. DeBoer¹, Matthias Griese⁴, Nicolaus Schwerk⁵, Lisa R. Young⁶, Carina Ittrich⁷, Thomas Schlange⁷, Martina Gahlemann⁸, David Warburton⁹, ¹Section of Pediatric Pulmonary and Sleep Medicine, Department of Pediatrics, University of Colorado Denver, Denver, CO and The Children's Hospital Colorado, Aurora, CO; ²Department of Medicine, National Jewish Health, Denver, CO; ³Centre for Inflammation Research, University of Edinburgh, Edinburgh, United Kingdom; ⁴Hauner Children's Hospital Ludwig Maximilians University, German Center for Lung Research (DZL), Munich Germany; ⁵Clinic for Pediatric Pulmonology, Allergology and Neonatology, Hannover Medical School, Hannover, Germany; ⁶Division of Pulmonary and Sleep Medicine, The Children's Hospital of Philadelphia, Philadelphia, PA; ⁷Boehringer Ingelheim Pharma GmbH & Co. KG, Biberach an der Riss, Germany; ⁸Boehringer Ingelheim (Schweiz) GmbH, Basel Switzerland; ⁹Children's Hospital Los Angeles, Los Angeles, CA and Keck School of Medicine, University of Southern California, Los Angeles, CA.

MUC5B PROMOTER VARIANT IS ASSOCIATED WITH A UNIQUE PROTEIN SIGNATURE IN EARLY LUNG FIBROSIS. Jeremy A. Herrera^{1*}, Mark Maslanka², Rachel Blumhagen¹, Janna Brancato¹, Jonathan Huber¹, Carlyne Cool¹, Kirk C. Hansen², Ivana V. Yang¹, David A. Schwartz¹, ¹Division of Pulmonary Sciences and Critical Care Medicine; ²Department of Biochemistry and Molecular Genetics; University of Colorado Denver.

POSTERS – Thursday, June 6, 2024 – continued

WILMS TUMOR 1 IMPAIRS FIBROBLAST CLEARANCE IN SEVERE FIBROTIC LUNG DISEASE. Harshavardhana H. Ediga^{1*}, C. P. Vemulapalli¹, V. Sontake³, H. Miyazaki³, D Popov³, P.K. Patel¹, S. Paranthaman¹, S. K. Huang² and S. K. Madala¹, ¹Division of Pulmonary, Critical Care and Sleep Medicine, Department of Internal Medicine, University of Cincinnati, Cincinnati, OH; ²Division of Pulmonary and Critical Care Medicine, University of Michigan Medical School, Ann Arbor; ³Gordian Biotechnology, South San Francisco, CA.

NON-TYPICAL LUNG FIBROSIS: IMAGING PREDICTS SURVIVAL. **David A Lynch***, Stephen M Humphries, Department of Radiology, National Jewish Health, Denver, CO.

SPUTUM LEVELS OF EXTRACELLULAR DNA ARE INCREASED IN RHEUMATOID ARTHRITIS-ASSOCIATED INTERSTITIAL LUNG DISEASE. M. Kristen Demoruelle^{1*}, Melissa Griffith¹, Timothy M. Wilson², Marie L. Feser¹, Kevin D. Deane¹, Stephen Humphries³, Joshua J. Solomon⁴, ¹University of Colorado Denver, Division of Rheumatology, Aurora, CO; ²Thomas Jefferson University, Division of Rheumatology, Philadelphia, PA; ³National Jewish Health, Department of Radiology, Denver, CO; ⁴National Jewish Health, Division of Pulmonary, Critical Care and Sleep Medicine, Denver, CO.

PCSK6 AND RESPIRATORY-RELATED OUTCOMES IN PATIENTS WITH PULMONARY FIBROSIS EN-ROLLED IN THE PULMONARY FIBROSIS FOUNDATION PATIENT REGISTRY. **Kristin N. Berger^{1*}**, Will Whalen¹, Will Simmons¹, John S. Kim², Imre Noth², Justin M. Oldham³, Anna J. Podolanczuk¹, ¹Department of Medicine at ¹Weill Cornell Medicine, New York, NY; ²University of Virginia, Charlottesville; ³University of Michigan, Ann Arbor, MI.

CT FINDINGS ASSOCIATED WITH MYOSITIS RELATED INTERSITIAL DISEASE. Joseph B Pryor^{1*}, Joshua Solomon², Jeffrey Swigris², Rebecca Keith², Tami Bang³, Andrea Fuentealba³, David A Lynch³, Liudmila Kastsianok⁴, Zulma Yunt², ¹Division of Pulmonary and Critical Care, University of Colorado, Denver, CO; ²Center for Interstitial Lung Disease, National Jewish Health, Denver, CO; ³Department of Radiology, National Jewish Health, Denver, CO; ⁴Division of Rheumatology, National Jewish Health, Denver, CO.

EFFICACY OF ENSIFENTRINE, A DUAL PDE3/PDE4 INHIBITOR, IN THE RAT MODEL OF BLEOMYCIN INDUCED PULMONARY FIBROSIS. **Margot MacDonald-Berko***, Verona Pharma plc, NC; Joanne Kilgour, Regulatory Science Associates, Inverkip, UK; Tara Rheault, Verona Pharma plc, NC. Lab work was completed at Labcorp Early Development Laboratories Ltd, UK.

RECRUITED MACROPHAGES PRODUCE COAGULATION FACTOR XIII-A WHICH ENHANCES COLLAGEN DEPOSITION AFTER BLEOMYCIN. **Peter Moore***, Raleigh Garner, Emily King, Shannon Hott, Katrina Kopf, William Janssen, Alexandra McCubbrey. Departments of Medicine, University of Colorado Anschutz, Aurora, CO and National Jewish Health, Denver, CO.

UNIQUE PHOSPHOPROTEINS TRIGGER SENESCENCE IN RADIATION-INDUCED PULMONARY FIBRO-SIS (RIPF). Sadiya Bi Shaikh^{1*}, Eric Hernady², Brian Marples² and Irfan Rahman¹, ¹Department of Environmental Medicine, University of Rochester Medical Center, Rochester, NY; ²Department of Radiation Oncology, University of Rochester Medical Center, Rochester, NY.

LUNG TRANSPLANT OUTCOMES FOR PATIENTS WITH CONNECTIVE TISSUE DISEASE-RELATED IN-TERSTITIAL LUNG DISEASE. **Sarah L. Khan^{1*}**, Samuel J. Minkove², Kevin J. Psoter³, Sonye K. Danoff¹, Pali D. Shah¹, ¹Division of Pulmonary and Critical Care Medicine, Johns Hopkins University School of Medicine, Baltimore, MD; ²Pulmonary Care and Sleep Medicine, St. Joseph's Medical Center, Towson, MD; ³Division of General Pediatrics and Adolescent Medicine, Johns Hopkins University School of Medicine, Baltimore, MD.

POSTERS – Thursday, June 6, 2024 – continued

APPLICATION OF IN VIVO ORGANELLAR IMMUNOPURIFICATION TO REVEAL MECHANISMS UNDER-LYING GENETIC INTERSTITIAL LUNG DISEASE. **David Ziehr**^{1,2,3}*, Jason Yang¹, Jack Bush¹, Raghu Chivukula^{1,2,3}, ¹Center for Genomic Medicine, Massachusetts General Hospital; ²Division of Pulmonary and Critical Care Medicine, Massachusetts General Hospital; ³Harvard Medical School, Boston, MA.

THE IPF-ASSOCIATED GENETIC VARIANT rs12417955 RESIDES IN A NOVEL 3' REGULATORY REGION FOR MUC5B THAT RESPONDS TO AIRWAY EPITHELIAL DIFFERENTIATION SIGNALS AND ENVIRON-MENTAL CUES. Sarah K. Sasse^{1*}, Arnav Gupta^{1,2}, Anna Peljto², Rachel Blumhagen¹, Fabienne Gally¹, Evgenia Dobrinskikh², Michael R. Weaver¹, Robin D. Dowell^{3,4,5}, Ivana V. Yang², David A. Schwartz² and Anthony N. Gerber^{1,2}, ¹Department of Medicine, National Jewish Health, Denver, CO; ²Department of Medicine, University of Colorado, Aurora, CO; ³BioFrontiers Institute, ⁴Molecular, Cellular and Developmental Biology, ⁵Computer Science, University of Colorado, Boulder, CO.

AUTOMATED CLASSIFICATION OF USUAL INTERSTITIAL PNEUMONIA ON COMPUTED TOMOGRA-PHY IS ASSOCIATED WITH PROGRESSION OF INTERSTITIAL LUNG ABNORMALITIES IN THE COPDGENE STUDY. **Stephen Humphries**^{1*}, Devlin Thieke¹, Samuel Ash², Hiroto Hatabu³, Gary M. Hunninghake⁴, David Lynch¹, ¹Department of Radiology, National Jewish Health, Denver, CO; ²Critical Care, South Shore Hospital, South Weymouth, MA; ³Department of Radiology and 4. Department of Pulmonary and Critical Care Medicine, Brigham and Women's Hospital, Harvard Medical School, Boston, MA.

CLINICAL AND GENETIC BIOMARKERS OF PULMONARY FIBROSIS AMONG GULLAH SYSTEMIC LU-PUS ERYTHEMATOSUS PATIENTS. Robert Campbell, Jr.* — NHLBI PRIDE AGOLD Scholar, Julius Nyalwidhe, - Leroy T. Canoles Jr. Cancer Research Center, Eastern Virginia Medical School, Norfolk, VA; Christopher Gignoux, - Colorado Center for Personalized Medicine, University of Colorado - Anschutz Medical Campus, Aurora, CO.

FUNCTIONALIZING HUMAN DECM FOR INCORPORATION INTO 3D PULMONARY FIBROSIS MODELS. Haley Noelle^{1*}, Mikala C. Mueller¹, Dema H. Essmaeil², Rachel Blomberg¹, Chelsea M. Magin^{1,3,4}, ¹Department of Bioengineering, University of Colorado, Denver | Anschutz Medical Campus; ²Department of Biology, University of Colorado, Denver; ³Department of Pediatrics, University of Colorado Anschutz Medical Campus; ⁴Division of Pulmonary Sciences & Critical Care Medicine, Department of Medicine, University of Colorado, Anschutz Medical Campus, Aurora, CO.