

**BOOK BEFORE AUGUST 23
TO SAVE UP TO \$600!**

November 18 - 20, 2019 | Boston, MA



3rd Annual AFDD

Anti-Fibrotic Drug Development

Translating Therapeutic Success Across Clinical Phases and Fibrotic Conditions

The roadmap to anti-fibrotic drug development advancements

37 expert speakers including:



Matthew Breyer
Distinguished
Scientist
Janssen



Lisa Hazwelwood
Principal Scientist
Research, Liver
Disease and Fibrosis
AbbVie



Glenn Rosen
SVP Preclinical
Translational
Sciences Small
Molecules
Coherus BioSciences



Tim Johnson
Executive Director
Non-Haemophilia
Commercial
Assessment
UCB



**Rosza Schlenker -
Herceg**
Executive Director
- Clinical Research,
Medical Affairs &
Respiratory
Boehringer-Ingelheim



Weilin Xie
Senior Principal
Scientist
Biotherapeutics
Department
Celgene

Partners:



14
in-depth case
studies



11
hours
networking



42
expert
speakers



100+
like-minded
peers

Welcome to the 3rd Annual Anti-Fibrotic Drug Development Summit (AFDD) 2019

The industry's definitive guide to turning fibrotic mechanisms into clinically effective therapeutics across disease areas



Join over 100 fellow drug developers to disseminate the latest findings, build a comprehensive network of like-minded peers who share the same challenges and search for drug repurposing opportunities.



Hear from pharma and biotech companies of all sizes, as well as clinicians and leading academics for a complete picture of the current challenges and opportunities.



Find commonalities across disease areas that can be applied back at the lab with presentations from specialists across all fibrotic conditions including **NASH, IPF, CKD, cardiovascular, gastro-intestinal, Scleroderma, rare disease** and **ocular fibrosis**.

The next generation of anti-fibrotic treatments will be born at the 3rd AFDD Summit.

Hear what previous attendees have to say

“This conference looks at things from a different angle than others in the field, with the emphasis on industry developments and direction rather than academic research. It was nice to hear from companies developing technologies and solutions to aid research in fibrosis as well as those looking for partners. I enjoyed listening to how different companies tackled problems”

Tim Johnson, Director Fibrotic Remodeling, UCB

“Excellent panel of speakers, thought provoking discussions”

Lynn Williams, Group Leader, University of Oxford

Five Unmissable Highlights



Compelling Case Studies

The AFDD Summit provides a forum for leaders to share the newest advancements in drug development for fibrotic conditions



Diverse Discussions

Our panels bring together multi-specialty, cross background professionals to discuss the biggest challenges faced by the industry and discover the solution together



Academic Showcase

Look to the future with the latest discoveries from the most prestigious academic institutions



Productive Networking

From constructive conversations over coffee to dexterous discussions over drinks, our varied networking program has something for everyone



Hands-on Workshops

We believe in practical learning and nothing champions this better than our 3 hour workshops

An Unrivaled Speaker Line-Up



Zoe Johnson
Chief Scientific
Officer
iOnctura



Lisa Hazelwood
Principal Scientist
Liver Disease and
Fibrosis Discovery
Abbvie



John Atkinson
Senior Research
Scientist Tissue
Remodelling
UCB



Richard Stratton
Consultant Physician
and Honorary Senior
Lecturer
**University College
London**



Glenn Rosen
Senior Vice
President Preclinical
Translational
Sciences Small
Molecule
Coherus Biosciences



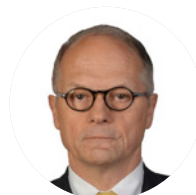
Matthew Breyer
Distinguished
Scientist CVM
Janssen



Tim Johnson
Director Immunology
Therapeutic Area
UCB



Ravi Kumar
Senior Vice President
and Chief Scientific
Officer
Acceleron Pharma



Pieter Muntendam
Chief Executive
Officer and Founder
G3 Pharmaceuticals



Aftab Taiyab
Research Scientist
McMaster University



Kirby von Kessler
Associate Director
Clinical Development
Lung Biotechnology



Masha Poyurovsky
Vice President
Discover Biology
**Kadmon
Corporation LLC**



Alexey Lugovskoy
Chief Development
Officer
**Morphic
Therapeutics**



Sten R. Sørensen
Chief Executive
Officer
Cereno Scientific



Shervin Assassi
Professor of Medicine
and Co-Director
of the UTHealth
Scleroderma Program
**University of Texas
Health Science
Center at Houston**



Robert Lafyatis
Adjunct Professor
Medicine
Boston University



Anie Philip
Professor Department
of Surgery
**Research Institute
of the McGill
University Health
Center**



Yaron Ilan
Director Department
of Medicine
**Hebrew University
Hadassah Medical
Center**



Sydney Montesi
Instructor in Medicine
Harvard University



Ganesh Raghu
Professor Medicine
in the Division
of Pulmonary
and Critical Care
Medicine
**University of
Washington**



Min Lu
Morphic Therapeutics
Head of Fibrosis



Rozsa Schlenker - Herceg
Executive Director
Clinical Research,
Medical Affairs and
Respiratory
Boehringer-Ingelheim



Rick Keenan Ph.D.
Chief Scientific
Officer
OptiKira



Weilin Xie
Senior Principal
Scientist
**Celgene
Corporation**



Eliezer Zomer Ph.D.
Vice President
Drug Discovery and
Manufacturing
**Galectin
Therapeutics**



Lee Borthwick
Fibrosis Biology
Lecturer and Chief
Operating Officer
FibroFind
**Newcastle
University**



**Jean-Francois
Thibodeau**
Research Scientist
**Prometic
Biosciences**



Gilles Tremblay
Director Preclinical
Development
Forbuis



Scott Macdonnell
Senior Staff Scientist
Regeneron



**Mehran
Moghaddam**
Chief Executive
Officer
OROX Biosciences



Scott Turner
Vice President
Translational
Sciences
Pliant Therapeutics



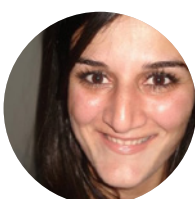
Emily Offer
Principal Scientist
and Team Leader
Redx Immunology



Komathi Stem
Chief Executive
Officer
MonArc Bionetworks



Bertram Pitt
Professor of Medicine
Emeritus
**University of
Michigan School of
Medicine**



Rana Herro
Junior Faculty
Instructor
**La Jolla Institute for
Immunology**



Resat Cinar
Staff Scientist
**National Institute
on Alcohol Abuse
and Alcoholism**



Robert Edwards
Director and CVM
Clinical Project
Specialist
Janssen



**Moustapha El-
Amine**
Director Global
Search and Evaluate
AstraZeneca



Gail Walkinshaw
Vice President
Research
FibroGen



Darcey Black PhD
Co-founder and
Director, Translational
TherapeutAix



Simon Cruwys PhD
Co-founder
and Director,
Pharmacology
TherapeutAix



Dr John O'Neill
Chief Scientific
Officer
XYLYX Bio

Pre-Conference Workshops

Monday November 18 2019

Registration and Welcome Coffee

8:30 – 9:00

Workshop A

9:00 – 12:00

Biomarkers 101

- Novel biomarkers: how to create them and how to apply them
- Hands-on demonstration of biomarker application to preclinical and clinical models
- Medical imaging and blood tests: the pros and cons of the latest technology
- Translating *in vivo* and *in vitro* models to clinical trials through consistent biomarker application

Workshop Leader



Sydney Montesi
Instructor in Medicine
Harvard University

Lunch Networking Break

12:00 – 13:00

Workshop B

13:00 – 16:00

How to cure multifactorial fibrotic diseases “Kill Two Birds With One Stone: MRI-1867, Hybrid Inhibitor of Peripheral Cannabinoid Receptor 1 (CB1R) & Inducible Nitric Oxide Synthase (iNOS), for the Treatments of NASH & Liver, Lung and Skin fibrosis”

- Outlining rationale for multi-target approach by polypharmacology to develop effective therapies in complex progressive diseases such as NASH and fibrotic diseases
- Identifying dual targeting of peripheral cannabinoid receptor 1 (CB1R) and inducible nitric oxide synthase (iNOS) as an effective therapeutic strategy in obesity, diabetes, AFLD, NAFLD, NASH, liver, lung and skin fibrosis
- Introducing the concept of third-generation cannabinoid receptor 1 (CB1R) antagonists for metabolic and fibrotic disorders
- Demonstrating preclinical efficacy of MRI-1867, an orally bioavailable small molecule antagonist of peripheral CB1R/iNOS, in fibrotic disorders: liver, lung and skin fibrosis

Workshop Leader



Resat Cinar
Staff Scientist
National Institute on Alcohol Abuse and Alcoholism

Workshop C

13:00 – 16:00

Mechanistic workshop: Inhibitors

- Discover fibrotic inducers to create the perfect mechanistic antidote
- Decipher the biological elements lacking in a fibrotic patient but found in a healthy patient
- The creation of inhibiting treatments: from R&D to clinical trials

Workshop Leaders



Richard Stratton
Consultant Physician and Honorary Senior Lecturer
University College London



Rana Herro
Junior Faculty Instructor
La Jolla Institute for Immunology

Networking Coffee to Stay or Go

16:00 – 16:30

Conference Day One

Tuesday November 19 2019

8.00 Registration and Welcome Coffee

Alexey Lugovskoy
Chief Development
Officer
**Morphic
Therapeutics**

8.20 Chair's Opening Remarks

Keynote Panel Sessions

8.30 Panel: One Vision for the Future of Fibrosis

- Understand the commonalities between varying fibrotic conditions to better prevent and reduce inflammation
- Develop a systematic approach to target fibrosis across organs
- Share data to learn from previous failures and streamline the drug development process
- Decide whether treatments should be targeting fibrosis as a disease itself or as a response to the disease within that organ

Moderator:



Ganesh Raghu
Professor Medicine
in the Division
of Pulmonary
and Critical Care
Medicine
**University of
Washington**

Panellists:



Matthew Breyer
Distinguished
Scientist CVM
Janssen



Alexey Lugovskoy
Chief Development
Officer
**Morphic
Therapeutics**



Lisa Hazelwood
Principal Scientist
Liver Disease and
Fibrosis Discovery
Abbvie

9.10 Panel: The Trials and Tribulations of Animal Models

- Decide whether animal models are necessary or whether they can be bypassed
- Weigh the relevancy of precision-cut organ slices against the longevity of animal models
- Consider testing multiple animal models for a smoother transition to clinical trial

Moderator:



Alexey Lugovskoy
Chief
Development
Officer
**Morphic
Therapeutics**

Panellists:



John Atkinson
Senior
Research
Scientist Tissue
Remodelling
UCB



Lee Borthwick
Fibrosis
Biology
Lecturer
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Corporation
LLC**



Glenn Rosen
Senior Vice
President
Preclinical
Translational
Sciences Small
Molecule
**Coherus
Biosciences**



9.50 Speed Networking

Establish meaningful business connections at a rapid rate. Efficiency at its finest.

10.50 Morning Networking Break

Research and Discovery Sessions: Novel Targets, Molecules and Pathways

Mehran Moghaddam
Chief Executive
Officer
OROX Biosciences

11.20 Case Study: Novel Two Pronged Target Approach

- Develop unique molecules that attack two targets simultaneously to stop the progression of fibrosis across the body
- Ensure secondary targets are relevant and do not enhance toxicity of treatment

Eliezer Zomer Ph.D. Vice President Drug Discovery and Manufacturing Galectin Therapeutics	11.40 Therapeutics for Chronic Fibrotic Diseases - The next generation of Galectin-3 inhibitors: from R&D through to phase III clinical trials - Discovery of functional allosteric inhibitors
Dr John O'Neill Chief Scientific Officer XYLYX Bio	12.00 Disease-Specific Extracellular Matrix Cell Culture Substrates to Improve Predictive in Vitro Models of Fibrosis - Introducing a standardized, fully humanized commercial 3D cell culture platform for fibrosis research - Describing the extracellular matrix (ECM) composition of the substrates that recapitulates the human disease environment in vitro - Discussing how this platform can reduce the dependence on animal models, and enable more relevant scientific results leading to improved drug discovery process
Scott Turner Vice President Translational Sciences Pliant Therapeutics	12.15 Case Study: Therapeutic Integrin Inhibition - Selective integrin inhibitors block TGF-β activation in a cell and tissue specific manner for the treatment of NASH and IPF - PK and PD evaluation of dual $\alpha\beta6/\alpha\beta1$ inhibitors in humans and primates
Pieter Muntendam Chief Executive Officer and Founder G3 Pharmaceuticals	12.35 Case Study: Galectin-3 as a Potential Treatment Target - Identify patients with galectin-3 mediated fibrogenesis by plasma galectin-3 levels - Using the galectin-3 carbohydrate recognition domain as an on-off switch for treatment - The promise of galectin-3 inhibition in cardiac and other conditions
Darcey Black PhD Co-founder and Director, Translational TherapeutAix Simon Cruwys PhD Co-founder and Director, Pharmacology TherapeutAix	12.55 Building a platform of evidence that will lead to clinical and partnering success in fibrotic disease - A view of the future clinical landscape in fibrotic disease – exemplified by IPF and NASH, and the ability of currently available assays to address this - The opportunity to improve translation via the use of biomarkers - Data sets required by industry partners, VCs, PIs and regulators
Emily Offer Principal Scientist and Team Leader Redx Immunology	13.10 Case Study: Selective ROCK-2 Inhibitor Programme - ROCK2 as a kinase central to the signalling processes involved in aberrant wound healing and fibrosis. - Up-regulation in acute and chronic inflammation - Inhibition of ROCK2 reduces the pro-fibrotic and pro-inflammatory response disease relevant in vitro and in vivo models
	13.30 Q&A Panel: Novel Targets, Molecules and Pathways
	13.45 Lunch Seminar Maximize productivity by pairing food with food-for-thought at the optional lunchtime seminar.
	13.45 Lunch Networking Break
Preclinical Sessions: Translating Biology from Animal to Patient	
John Atkinson Senior Research Scientist Tissue Remodelling UCB	14.45 Case Study: Successful Clinical Trials Start with Carefully Curated Animal Models - Specify readouts based on the biology of the target to reflect the chosen mechanism of action - Adopt a poly-pharmacy approach for increased effectiveness - Move away from inbred strains in animal models and diversify environment and age ranges

Lee Borthwick Fibrosis Biology Lecturer and Chief Operating Officer FibroFind Newcastle University	15.05 Case Study: Next Generation <i>In Vitro</i> Models • Ensure a smoother transition to clinical trial with precision cut slice technology that mimics the function of the organ • Recreate the organ culture by replicating the biology and cells that would be found in a person • Stimulate the culture with a cause relevant to the chosen clinical disease for authentic progression of fibrosis • Use of precision cut slices for the modelling of fibrosis
	15.25 Afternoon Networking Break
Preclinical Sessions: Translating Biology from Animal to Patient	
Scott Macdonnell Senior Staff Scientist Regeneron	15.55 Case Study: Defining “Activated” Fibroblasts using Single Cell Sequencing • Understand the characteristics of the underlying fibroblast cell population beyond the current recognition as the central mediators of extra-cellular matrix deposition in IPF • Use of an unbiased single cell RNA-sequencing analysis of a bleomycin-induced pulmonary fibrosis model to characterize molecular responses to fibrotic injury at the single cell level • Analysis of gene expression as analyzed by 3 complementary techniques which together generated a 49-gene signature that defined an activated subpopulation of fibroblasts
Komathi Stem Chief Executive Officer MonArc Bionetworks	16.15 Case Study: Develop Patient Relationships at the Preclinical Stage to Identify the Best Patient Population for your Treatment • Generate evidence for clinical trial stage from early-on to make the transitional period seamless and ensure success • Application of RWE in IPF clinical trials – translating success from the real world and fibrotic patient experience • Identify patients despite low diagnosis rate by correlating data and maintaining a registry with specific disease indicators
Tim Johnson Director Immunology Therapeutic Area UCB	16.30 Case Study: Use of Technology in the Development of TG2 Inhibitory Therapeutics • Development of a glomerular sclerosis organoid model • Responses in organoid models which recapitulate rodent <i>in vivo</i> data • Creation of CKD primate models for antibody therapies • Use of surrogate biomarkers for <i>in vivo</i> studies
	16.50 Q&A Panel: Translating Biology from Animal to Patient
	17.10 Speed Learning Roundtable discussions like you’ve never seen them before. Uncover the story behind four fibrotic therapeutic developments in one quick-fire session. Each table will be hosted by leading specialist who will share the secrets of their most high-impact strategy; you then get the opportunity to question the host before moving on to your next table.
	18.10 Poster Session and Drinks Reception Constructive conversations over cocktails and canapés to the backdrop of posters.


 The only meeting to attend if you want to learn
 about state of the art developments in fibrosis 
 Vincenza di Modugno, Director, Roche

Conference Day Two

Wednesday November 20 2019

	8.00 Welcome Coffee
Alexey Lugovskoy Chief Development Officer Morphic Therapeutics	8.20 Chair's Opening Remarks
Spotlight on Fibrotic Disease Areas	
Jean-Francois Thibodeau Research Scientist Prometic Biosciences	8.30 Case Study: Novel Compounds to Target Alstrom - Develop a drug that works through binding activation and inhibition of receptors found in macrophages and immune cells - Target the initial inflammatory process to prevent the progression of fibrotic disease
Sten R. Sørensen Chief Executive Officer Cereno Scientific Bertram Pitt Professor of Medicine Emeritus University of Michigan School of Medicine	8.50 Fireside Chat: Advancements in Cardiovascular Fibrosis - Discover new therapeutic targets outside of diabetes and blood-pressure which contribute to cardiovascular disease - Detect myocardial fibrosis with collagen and IL11 biomarkers - Selective small molecule inhibitors within cardiac fibrosis

9.10 Panel: Advancements in Metabolic Fibrosis

- Combination therapies for NASH and Chronic Kidney Disease
- The relationship between diabetes and CKD – can common therapies be effectively used to treat both conditions?
- Tissue targeting in CKD – increasing efficacy, reducing toxicity
- Novel molecular targets including Galectin-3 inhibitors and HSP47

Moderator:



Matthew Breyer
Distinguished Scientist CVM
Janssen

Panellists:



Min Lu
Morphic Therapeutics
Head of Fibrosis



Tim Johnson
Director Immunology Therapeutic Area
UCB



Lisa Hazelwood
Principal Scientist Liver Disease and Fibrosis Discovery
Abbvie



Resat Cinar
Staff Scientist
National Institute on Alcohol Abuse and Alcoholism

9.50 Panel: Advancements in Pulmonary Fibrosis

- The relationship between Pulmonary Arterial Hypertension and IPF – common mechanisms to target and combination therapies
- The transference of inhalant technology from COPD to IPF
- Small molecule therapies that target novel mechanisms

Moderator:



Scott Macdonnell
Senior Staff Scientist
Regeneron

Panellists:



Kirby von Kessler
Associate Director Clinical Development
Lung Biotechnology



Rozsa Schlenker - Herceg
Executive Director Clinical Research, Medical Affairs and Respiratory
Boehringer-Ingelheim



Weilin Xie
Senior Principal Scientist
Celgene Corporation



Rick Keenan
Chief Scientific Officer
OptiKira



Moustapha El-Amine
Director Global Search and Evaluate
AstraZeneca

10.30 Morning Networking Break

Spotlight on the Relationship between Fibrosis and Cancer

Gilles Tremblay
Director Preclinical
Development
Forbius

11.00 Case Study: Novel Inhibitors of TGF Beta for Fibrotic Cancer

- Tackle the barrier created by fibroblasts in the tumour microenvironment which prevents immune cells from targeting cancerous cells
- Understand the role of TGF Beta within fibrotic cancers and create a novel inhibitor of TGF Beta Ligands
- Consider the role of stromal components and the secretion of cytokines within cancer fibroblast to find a combined immune checkpoint and TGF Beta inhibitors treatment

11.20 Panel: You Can't Teach an Old Pathway New Tricks; the Case for and Against TGF Beta as a Primary Target for Fibrosis and Fibrotic Cancers

- Can inhibiting TGF Beta alone stop the progression of fibrosis?
- Does the reduction of TGF Beta in animal models result in strong histological end points at clinical trial?
- Is there a common TGF Beta pathway to target which will attack fibrotic cancers?

Moderator:

 **Gilles Tremblay**
Director Preclinical
Development
Forbius

Panellists:

 **Ravi Kumar**
Senior Vice President
and Chief Scientific
Officer
Acceleron Pharma

 **Zoe Johnson**
Chief Scientific
Officer
iOnctura

 **Gail Walkinshaw**
Vice President
Research
FibroGen

Clinical Sessions: Overcoming Failures

Gail Walkinshaw
Vice President
Research
FibroGen

12.00 Tactics to move a clinical trial to phase III with minimal risk of failure

- Understand the main causes of AFDD clinical trial failures and discover solutions that prevent future breakdowns
- Clinical trial design which is failure-proof
- Learn to pre-empt and adapt in the face of problems to ensure success

Robert Edwards
Director and CVM
Clinical Project
Specialist
Janssen

12.20 A Blueprint for Success: J&J's CREDENCE Clinical Trial

- Patient recruitment: a global multidisciplinary effort required perseverance
- Adjudication: a concerted cross-functional effort required diligence and determination
- Patient retention: a top priority from Day 1 left no stone unturned

12.40 Q&A Panel: Overcoming Clinical Trial Failures

13.00 Lunch Break

14.00 Wrap Up Roundtables

10 tables, 10 problems, 10 solutions. Ready, set, resolve!

- Select a pathway outside of TGF Beta to target in the treatment of fibrosis - define the drug development plan for this pathway
- Inhibitors vs inducers: create a plan to develop either an inhibiting or inducing anti-fibrotic treatment. Explain why you selected the inhibitor/ inducer.
- Design a novel biomarker and explain its application to both the preclinical and clinical stage
- Define a common mechanism to target in order to create an anti-fibrotic drug which will treat all fibrotic disease areas
- Design a preclinical plan with animal models that will effectively translate to the clinic
- Design a preclinical plan with only in vitro models
- Incorporate the use of technology into the development of an anti-fibrotic drug
- Develop a treatment for a patient with fibrotic cancer
- Translate J&J's CREDENCE trial into a clinical trial design for an anti-fibrotic treatment
- Design a patient recruitment programme for an anti-fibrotic clinical trial



Look to the Future: Academic Showcase

Aftab Taiyab
Research Scientist
McMaster University

15.00 Ocular Fibrosis: See Common Mechanisms in a New Light

- The non-canonical signalling pathway indicative of ocular fibrosis which is also found in IPF and renal
- Targeting the transformation to myofibroblasts, from epithelial to mesenchymal cells
- Learnings from ocular fibrosis mechanisms that share commonalities with other fibrotic conditions and discovery of shared biomarkers

Yaron Ilan
Director Department of Medicine
Hebrew University Hadassah Medical Center

15.20 The Gut as a Target for Anti-Fibrotic Therapy in NASH

- Using the gut for systemic immune modulation without immune suppression
- Generating an immune signal in the gut for re-directing the immune system in an anti-inflammatory way
- Targeting the gut microbiome for alleviation of fibrosis

15.40 Panel: Spotlight on Scleroderma

- *In vitro* skin based assays and how they can be applied to other conditions
- Genetic skin models including rare disease RDEB
- Blood test biomarkers indicative of scleroderma



Shervin Assassi
Professor of Medicine and
Co-Director of the UTHealth
Scleroderma Program
**University of Texas Health
Science Center at Houston**



Robert Lafyatis
Adjunct Professor Medicine
Boston University



Anie Philip
Professor Department of
Surgery
**Research Institute of the
McGill University Health
Center**

Alexey Lugovskoy
Chief Development
Officer
**Morphic
Therapeutics**

16.20 Chair's Closing Remarks



16.30 Networking Coffee to Stay or Go

Event Themes:

Gain insights into the latest **novel molecules and pathways** to identify the **best fibrotic mechanism** to target

Explore opportunities for **combination therapies** in **fibrosis** and **oncology**

Ensure **successful translation** from *in vivo* and *in vitro* models to clinical trial through the consistent application of **biomarkers** and **clinical endpoints**

Maximize your ROI at the only conference to bring together industry and leaders across fibrotic disciplines



Spotlight Partner:

monARC Bionetworks is a healthcare technology company on a mission to

modernize the clinical research industry by empowering patients to share their data and participate anytime, anywhere. By allowing patients to directly donate their data, monARC is creating a new data-sharing economy that will transform the highly sequential and siloed drug development model into an integrated and a collaborative model with patients.

www.monarcbio.com



Spotlight Partner:

The development of better medicine begins with better disease models. Unfortunately,

current in vitro models poorly mimic the pathophysiology of fibrosis, leaving a major gap in compound screening and testing efforts in anti-fibrosis drug discovery.

Xylyx Bio's native, disease-specific extracellular matrix substrates allow drug candidates to be investigated in vitro in a disease-relevant environment, enabling cell models that are significantly more predictive of human pathophysiology. Xylyx Bio products comprise both the mechanical properties and complex ratios of ECM components specific and unique to each organ type and disease state and are available in a variety of 2D and 3D product formats to support high-throughput assays

www.xylyxbio.com



TherapeutAix

Spotlight Partner:

TherapeutAix is a life sciences consultancy company based in Aachen, Germany, combining

drug discovery and development expertise with a fully integrated R&D network, to deliver practical, cost and time effective solutions to drug discovery projects. It supports investors and project teams focus on the next step, progress the right assets through robust decision-making steps, with a clear line of sight to the next value-inflection point and the clinic. Since its formation in 2018, TherapeutAix has successfully conducted project reviews, developed strategies and operationalised programs for biotech, pharma and investors in fibrosis, NASH and other therapeutic areas

www.therapeutaix.com



Exhibition Partner:

Biomodels LLC, a preclinical CRO, conducts predictive and translational studies

for biotechnology and pharmaceutical companies in the areas of fibrosis, pulmonary and inflammatory diseases, in addition to several other clinical indications. Biomodels specializes in (non-GLP) customized efficacy studies that optimize dose, schedule, and define mechanism of action. Additionally, Biomodels' state of the art Germ-Free and Gnotobiotic animal facility allows for the analysis of the role of the microbiota both in disease pathogenesis and in therapeutic efficacy.

www.biomodels.com

Why the AFDD Summit?

The **3rd AFDD Summit** is the fastest route to in-depth discussions with organizations prioritizing anti-fibrotic drug development. AFDD provides an unrivaled opportunity for your brand, your message and your reputation to be showcased in front of the leading minds of the growing 'Fibrosis Industry'.

Who do I get to meet?

Gathering stakeholders and key opinion leaders, the AFDD Summit is the ultimate opportunity to position yourself as an expert in front of 100+ drug developers. Elevate your company's standing and influence the future of anti-fibrotic drug development.

What can the AFDD Summit do for you?

Boost your brand; **engage** industry decision makers; **demonstrate** thought leadership

We understand each business is different so we'll work with you to build a bespoke partnership opportunity to fulfill your 2019/2020 business objectives.

How is AFDD different?

At this year's Anti-Fibrotic Drug Development Summit, you can expect:



A HIGHER CALIBER OF CONVERSATIONS: with over 10 hours of networking you'll have more opportunities than ever for significant discussions with your key prospects



DEDICATED ICEBREAKER SESSIONS FROM THE START: starting a conversation is never easy, so let us start them for you



INTERACTIVE SESSIONS AS STANDARD: engage your audience in solution-focused exchanges at this year's panel discussions, speed learning roundtables and poster session

Get Involved



Jakub Nunuk

Partnership Manager

Tel: +1617 455 4188

Email: sponsor@hansonwade.com

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3 Easy Ways To Book



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Jack Bainbridge
Delegate Experience Manager
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Email: info@hansonwade.com

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Please note that discounts are only valid when three or more delegates from one company book and pay at the same time.

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Contact: register@hansonwade.com

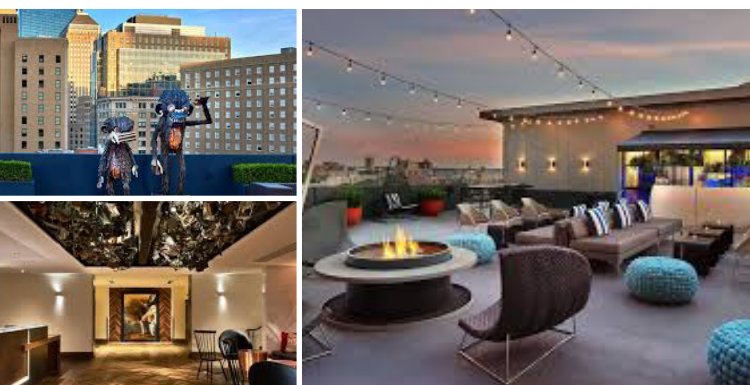
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Industry Pricing	Register & Pay By Friday, August 23	Standard Rate
Full Access: Conference + Workshop Day (A in AM) & (B or C in PM)	\$3697 (save \$600)	\$3997 (save \$300)
Conference Only	\$2499 (save \$400)	\$2899
Workshop Day Only (A in AM) & (B or C in PM)	\$1299	

*workshops can be purchased individually contact info@hansonwade.com

** Start up and academics are entitled to 40% discount off the industry price.

*** Past attendees (NASH, CKD3 and IPF) and event speakers are entitled to 10% discount off the industry price.



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