

30 September - 01 October 2025 | London, UK

Conference Brochure



Pioneering The Future Of Biomarkers & Precision Medicine



11Content Tracks
Across Two Days



10+ Hours of prearranged 1:1 Meetings



80+ Partners



1,000+ Attendees

Featuring 3 Groundbreaking Programmes!



Biomarkers



Spatial Biology for Precision Medicine



Digital Pathology & Al

Over 150 Industry-Leading Speakers Including...



🚔 KEYNOTE SPEAKER

MATTHEW BROWN, Chief Scientific Officer, Genomics England

Don't Miss His Keynote at 08:45 on Day One: Improving Rare Disease Diagnostic Performance Using Multiomics



SIR MARK CAULFIELD,

Vice Principal for Health, Faculty of Medicine and Dentistry Queen Mary University of London; Director of the NIHR Barts Biomedical Research Centre; President, British Pharmacological Society

Don't Miss Mark's Fireside Chat at 09:00 on Day Two



IVDR REGULATIONS & THEIR IMPACT

MIKE MESSENGER, Head of Regulatory Strategy, BIVDA

WANDERSON DOS SANTOS TRINDADE, Director Global Regulatory Affairs & CDx, Daiichi-Sankyo

CHRIS BRAY, Head Global Regulatory Affairs, Precision Medicine & Companion Diagnostics, Merck KGaA

VIHANGA PAHALAWATTA, Director Regulatory Affairs Companion Diagnostics, Abbvie

KARIN SCHMITT, Chief Operating Officer, Mursla Bio

SCOTT REID, Vice President & Global Head of Companion Diagnostics, Discovery Life Sciences

Don't Miss the Interactive Panel Discussion on Day One at 11.55

WELCOME TO

Biomarkers & Precision Medicine 2025

Welcome to Biomarkers & Precision Medicine 2025, Oxford Global's premier event that unites the leading minds of Precision Medicine under one roof at the QEII Centre in London on September 30 - October 1.

Join us for two immersive days of expertly curated content tackling the most critical challenges, spotlighting innovation and identifying emerging technologies. Experience 3 cutting edge programmes at an event which is celebrating its 20th Anniversary: Biomarkers 2025, Spatial Biology for Precision Medicine & Digital Pathology & Al.

Engage in high-level discussions and expertly tailored sessions that spotlight breakthroughs across precision medicine. Our carefully designed agenda explores cutting-edge advancements in biomarker-driven research, Al-powered pathology, and spatial multi-omics, shaping the future of diagnostics and therapy development. Join panels of leading scientists as they discuss the integration of spatial insights into biomarker discovery, the role of Al in transforming pathology workflows, and navigating the current regulatory landscape. Sessions will also examine how these innovations enhance clinical decision-making, patient stratification, and drug response monitoring, driving the next wave of precision medicine breakthroughs.

In addition to groundbreaking presentations and interactive discussions, this event offers invaluable networking and interactive opportunities with industry leaders and innovators. Explore our interdisciplinary collaboration hubs, Start-Up Zone and the Plenary Panel Discussions on the exhibition stage.

> Don't miss this opportunity to engage with pioneering experts and stay ahead of the next-generation advancements set to redefine precision medicine.



Director of Production & Content -Precision Medicine Brand, Oxford Global











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WHAT'S NEW

Benefits Of Attending



Stay ahead of evolving regulatory landscapes and navigate IVDR regulations with talks, panel discussions and roundtables focused on providing direction to this conundrum.



Maximise your time & register for the complementary pre-conference workshop being hosted by Labcorp on September 29th.



Interdisciplinary learning and knowledge share is key to driving Precision Medicine. Our Collaboration Hub sessions will enable experts and innovators from diverse complementary fields to foster collaboration, hear fresh perspectives, encourage innovation, break down silos and address specific challenges within the field.



Gain exclusive insights into cutting-edge biomarker research - explore the latest in biomarker discovery, validation, and clinical implementation across oncology, neuroscience, immunology, rare diseases etc.



Unlock the potential of Al - discover how Al-powered biomarker analysis, spatial biology, and digital pathology are revolutionising drug development, patient stratification and image analysis.



Engage with thought leaders in translational research - gain firsthand insights from leading scientists, clinicians, and regulatory experts on biomarker adoption, clinical trial design, and strategies for precision medicine implementation.



Discover groundbreaking biomarker technologies shaping the future - deep dive into proteomics, metabolomics, mass spectrometry, flow cytometry, multiplex imaging, and single-cell sequencing techniques advancing biomarker analysis.



Delve into biomarker innovations for neurodegenerative diseases & neuroscience

- hear from experts on novel imaging biomarkers, liquid biopsy approaches, and cuttingedge digital biomarker strategies for CNS and neuropsychiatric disorders.



Interactive Panels

BIOMARKERS

IVDR Regulations & Their Impact

Biomarker Strategies: Lessons From The Last Decade And The Road Ahead For Pharma & Academia

Translating Biomarkers From Bench To Bedside

Applying Single Cell And Omic Technologies In Biomarker Discovery And Development

Biomarker Strategies In Neurodegenerative Therapeutics

PRECISION MEDICINE

Al-Powered Precision Medicine

Building The Next Era Of Precision Medicine: Stakeholder Collaboration Across Biomarkers, Diagnostics, Omics & Therapeutic Innovation

DIGITAL PATHOLOGY

Building A Business Case For Digital Pathology

Digital Pathology in Pharma -Overcoming Integration Strategies & **Future Proofing**

SPATIAL BIOLOGY

Addressing The Future Needs Of Spatial Multi-Omics

Utilising Spatial Biology In Pharma R&D Exploring The Application Of Spatial Technologies In Immunology & Oncology



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Celebrate Your Industry, Acknowledge Your Impact & Accelerate Better Patient Outcomes At This Night of Celebration At The Stunning Church House Assembly Rooms Just A 3-min Walk Away. The Awards Begin Immediately After The Close of Day One.





SPECIAL EVENT FEATURES

Big Attendance Meets Intimate Connections

Special Feature:



The celebration event takes place on 30 September at the prestigious Church House Assembly Hall in Westminster, London, following the Biomarkers and Precision Medicine conference.

Join us to celebrate your achievements, light up our industry, and drive the next wave of transformation.

- Day Two -**Keynote Fireside Chat**

Join Professor Sir Mark Caulfield for an exclusive fireside chat reflecting on his groundbreaking work and the legacy of his contributions to genomic medicine. In this in-depth conversation, he will share key milestones from his career, the impact of large-scale genomic initiatives, and the lessons learned in advancing precision medicine. Looking ahead, he will offer his perspective on the future of precision medicine, emerging innovations, and the next steps in translating cutting-edge research into real-world clinical practice.

Day One - Opening Keynote Address

Matthew Brown, Chief Scientific Officer, Genomics England Recognised for his pioneering contributions to genomics and precision medicine, Matthew Brown will take to the stage to explore the transformative potential of multiomics in rare disease diagnostics. He will examine how cuttingedge approaches, including long-read DNA sequencing, transcriptomics, proteomics, and metabolomics, are enhancing diagnostic accuracy beyond traditional shortread whole genome sequencing. With a focus on the latest advancements at Genomics England and across the industry, this keynote will highlight the breakthroughs shaping the future of rare disease research and clinical implementation.





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Day One & Two Collaboration Hub Sessions

Interdisciplinary learning and knowledge share is key to driving Precision Medicine. Our Collaboration Hub sessions will provide a platform to enable experts and innovators from diverse complementary fields to foster collaboration, hear fresh perspectives, encourage innovation, break down silos and address specific challenges within the field.

Each invitation-only, focused session will last 50 minutes and with only 15 spaces available per hub, registration is essential to secure your place at the table!



Addressing the Evolving Regulatory Landscape for Personalised Medicine

Moderated by: Vihanga Pahalawatta, Director Regulatory Affairs Companion Diagnostics, Abbvie

Utilising AI in Precision Medicine

Moderated by: James Schofield, Founder & Chief Executive Officer, TopMD Precision Medicine

Bridging the Gap Between Research and Clinical Implementation in Precision Medicine

Moderated by: Richard Barker, Chairman & Founder, Metadvice & New Medicine Partners

Addressing Ethical and Legal Challenges in Genetic Data Use

Moderated by: Natalie Banner, Director of Ethics, Genomics England





WHY PARTNER WITH **OXFORD GLOBAL?**

At Oxford Global, our mission is to curate personalised experiences that foster community and inspire innovation.

We believe in the power of networking, connection, and knowledge to deliver quality products and services that exceed expectations. Partnering with Oxford Global means having a dedicated team committed to helping you achieve your goals and navigating the industry's ever-changing landscape.

✓ Arrange 1-1 Meetings

Benefit from quaranteed one-to-one face time with your key prospects, with detailed pre-meeting information provided to enable effective and productive conversations.

Speaking Opportunities

Showcase your company's recent work to a relevant and highly engaged audience.

Host Panel & Roundtable Discussions

Feature alongside key opinion leaders to discuss current hot topics and highlight your company's expertise.

Organise Workshops

Demonstrate best practice within the industry in front of your peers with case studies from your clients.

Exhibit your Products & Solutions

Promote your offerings and ensure delegates know where to find you with a prominent brand presence in the exhibition hall.

✓ Digital Marketing & Lead Generation

Accessing the Oxford Global database, amplify your thought leadership and branding messaging through our digital content initiatives.

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Join world-leading experts in biomarker research and development for this in-person meeting in the heart of London.

Discover groundbreaking insight into biomarker science, patient stratification, companion diagnostics, regulatory considerations, clinical development and trial design and explore cutting-edge innovations driving precision medicine forward.

VPs, Directors & Senior Managers from leading pharma & biotech companies and research institutions in the following fields and more:

- Biomarkers
- Biomarker Discovery
- Biomarker Identification
- Biomarker Validation
- Clinical Biomarkers
- Pre-Clinical
- Translational Science

- Clinical Development
- Precision Medicine
- Early Detection
- Enabling Technologies
- Multiplex Technologies
- Genomic Biomarkers
- Regulation

Formal and informal meeting opportunities offer delegates the chance to discuss key solutions with leading service providers. Formal 1:1 meeting opportunities will be available to arrange prior to the event which take place during the dedicated networking breaks covering:

- Biomarker **Technologies**
- Diagnostic Biomarkers

- Imaging Techniques
- Biomarker Analysis
- Proteomics
- Metabolomics
- Clinical Development
- Multiplex tools
- Digital Pathology

Day One

Track 1: Biomarkers Identification, Validation in Oncology, Immuno-Oncology & **Immunology**

- Biomarker discovery tools and technologies
- Biomarker strategies for patient identification
- Biomarkers to detect, predict and monitor response to treatment
- Check point inhibitors and dual therapies in immuno-oncology

Track 2: Diagnostic Biomarkers & Regulations

- IVDR regulations and their impact
- Companion diagnostic development
- Parallel diagnostic and drug approval strategies
- · Diagnostic-driven therapies: towards the clinic with optimised market access and patient adoption

Track 3: Biomarkers for Clinical Development

- Defining a robust clinical biomarker strategy
- Real-world applications of biomarkers in healthcare
- · Transforming clinical development through biomarker-driven clinical trial design and data analysis
- Exploratory/clinical endpoint biomarkers needed to support clinical trials
- Biomarker development gene therapy- patient responses and histology

Track 4: Genomic & Multi-Omic Approaches in Biomarker Discovery and Development

- Precision medicine approaches for personalised therapies
- · Precision oncology and molecular profiling
- Biofluid-based molecular biomarkers: cfDNA, ctDNA and CTCs
- · Validating and verifying genomic markers in preclinical drug development

Day Two

Track 1: Biomarkers Identification & Validation: Neuroscience & Co-Morbidity

- · Novel case studies and strategies in biomarker discovery, analysis and validation in neuroscience, CNS, neuropsychiatric disorders
- Imaging, digital and liquid biomarkers
- · Technologies and approaches to improve target engagement, early diagnostic biomarkers and how best to incorporate biomarkers into drug development, patient stratification and clinical trials in

Track 2: New & Emerging Biomarker Technologies for Biomarker Analysis

- Technology approaches, techniques and analysis flow cytometry, mass spectrometry, proteomics, metabolomics
- Biomarker assay optimisation and validation
- Single-plex and multiplex technologies
- · Integrating multi-omics data

Track 3: Biomarkers for Clinical Trials

- Translational biomarkers from preclinical to clinical phase studies
- The role of safety and efficacy biomarkers in translation
- Patient profiling and biosignatures in clinical development

Track 4: Biomarkers for Diagnostics & Precision Medicine

- · Liquid biopsy biomarkers for disease monitoring, multi-cancer screening and early detection /ultra-
- Latest technologies in detection and molecular characterisation of: CTCs; cfDNA, ctDNA, circulating extracellular RNA; exosomes and microvesicles
- Biofluid biomarkers for early disease detection
- · Delivering precision medicine for respiratory, autoimmune, immunology diseases and neuroscience



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Spatial Biology for Precision Medicine



Join leaders, experts, and researchers in our Spatial Biology for Precision Medicine programme, connecting global academic & research organisations as well as pharma representatives for high-level discussions on the latest innovations in spatial research & technologies.

Forward looking visionary leaders will discuss the current state of the industry, market trends and future growth areas aiding the application of spatial technologies in the clinic as well as the rise of digital pathology.

VPs, Directors & Senior Managers from leading pharma & biotech companies and research institutions in the following fields and more:

- Spatial Transcriptomics
- Spatial Proteomics
- Systems Biology

- Molecular Medicine
- Spatial Genomics
- Spatial Metabolomics

Formal and informal meeting opportunities offer delegates the chance to discuss key solutions with leading service providers. Formal 1:1 meeting opportunities will be available to arrange prior to the event which take place during the dedicated networking breaks covering:

- Spatial Imaging Platforms
- Spatial Data Analysis Tools
- Spatial Genomics

- Tissue Imaging & Prep
- Single Molecule Imaging
- Bioinformatics

Day One

Track 5: Spatial Multi Omics Tools, Techniques & Approaches

- Translating spatial imaging techniques and approaches into clinics
- Transcriptomics, proteomics, genomics and metabolomics breakthrough tools, techniques and approaches for precision medicine
- Single-cell transcriptome / RNA-seq
- Multi-modality processing
- Discovery of robust next-generation diagnostics and biomarkers that are key to designing precision medicine treatments

Track 6: Spatial Biology in Pharma & Translational Drug Research

- Accelerating the discovery and characterisation of biomarkers and drug targets using spatial tools
- Tailoring precision medicine approaches by considering the spatial distribution of drug
- Impacting disease research informing assessment of treatment and action mechanisms/deciphering dynamics of cell interactions
- Feasibility and scalability of various platforms

Day Two

Track 5: Al Image Analysis & Data Analytics In Spatial Biology

- Tissue imaging and analysis using advanced spatial profiling techniques and Al-guided technology
- Imaging data analysis/how to set a spatial experiment
- Relevant spatial parameters in different model systems
- Label-free imaging technologies imaging mass spec
- Algorithm design for spatial data
- Spatial multiplexed imaging for disease characterisation
- Intersection of Digital pathology and spatial biology
- · Computational methods to assist in the identification, classification and visualisation of complex multi-model datasets
- Overcoming challenges in multi-omic data acquisition and analysis



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AGENDA AT A GLANCE Digital Pathology & Al



Gather with leading innovators in digital pathology for an inperson event in the heart of London.

Dive into the latest advancements in Al-driven image analysis & computational pathology, as well as hearing from the challenges faced and lessons learnt from those who have already implemented DP & Al into their workflow so that you can join others in shaping the future of pathology.

VPs, Directors & Senior Managers from leading pharma & biotech companies and research institutions in the following fields and more:

- Digital Pathology
- Computational Pathology
- Image analysis
- Biomedical Informatics & Al
- Computational Medicine
- Pathology IT

- Immunohistochemistry
- Histopathology
- Toxicology and Safety Sciences
- NHS transformation
- Pathology
- Diagnostics

Formal and informal meeting opportunities offer delegates the chance to discuss key solutions with leading service providers. Formal 1:1 meeting opportunities will be available to arrange prior to the event which take place during the dedicated networking breaks covering:

- Digital Pathology Hardware
- Scanners
- Monitors
- Digital Pathology Software
- Image analysis
- Multi-modal image analysis

- Image Management
- Cloud computing/ storage
- End to End Solutions
- LIS
- Enterprise Imaging & Healthcare IT
- Al Algorithms

Day One

Track 7: Digital Pathology Implementation - Image Analysis & Al

- Advances, challenges, benefits and future developments of Digital Pathology, and the implications for pathology practice
- Analysing the business case for digital pathology
- User experience of converting to and integrating digital pathology
- Advances in image quality and processing
- Interoperability fully integrated DP and AI workflow
- Integrating pathology data with molecular data for accurate diagnosis and personalised
- Exploring both manual and fully automated image analysis and pattern recognition
- Pathologists' perspective using and working with Al
- Image format standardisation
- Multiplexed image analysis

Day Two

Track 6: Computational Pathology & Utilising Digital Pathology In Pharma Research

- · Implementing AI-based precision pathology
- Al-assisted diagnosis to improve the efficiency of pathologists
- Advancing machine and deep learning algorithms
- Improving WSI workflow efficiency
- Pathology PACS and informatics
- Data Storage addressing the challenge
- Computational Pathology in Precision Oncology
- Ground truth data and technology for biomarker assessment
- Digital image analysis in drug discovery
- Digitising toxicology
- AIML solutions for Biomarker discovery, detection, quantification and use in patient



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KEYNOTE SPEAKERS



MARIA ORR, Head of Precision Medicine, Biopharmaceuticals. AstraZeneca



MICHAEL CANNARILE, Head of Biomarkers, Early **Development Oncology** (EDO), Pharma Research & Development (pRED), Roche Innovation Center Munich



MIKE MESSENGER. Head of Regulatory Strategy, **BIVDA**



PAUL J VAN DIEST, Professor and Head, Department of Pathology, **University Medical Center** Utrecht



VIHANGA PAHALAWATTA, **Director Regulatory Affairs** Companion Diagnostics, **Abbvie**



BERND WOLLSCHEID, Head of Institute for Translational Medicine Professor & Group Head, ETH Zurich



VIRGINIA SAVOVA. Senior Director, Cell Targeted Precision Medicine. AstraZeneca

ALEXANDRA SEVKO

VP of Translational Science, Amphista Therapeutics

ALAIN VAN GOOL

Professor of Personalised Healthcare, Radboud University Medical Centre

AMONIDA ZADISSA

Associate Director of Informatics, UK Dementia Research Institute

ANTONINA MITROFANOVA

Associate Dean for Research & Associate Professor, Department of Biomedical and Health Informatics, Rutgers University

ARMAN RAHMAN

Assistant Professor of Anatomy, University College Dublin

ARTHUR LEWIS

Director of Pathology, Clinical Pharmacology & Safety Sciences, AstraZeneca

AZAM HAMIDINEKOO

Associate Director of Clinical Pharmacology & Safety Sciences, AstraZeneca

BELINDA NEDJAI

Associate Professor in Cancer Biomarkers & Epigenetics, Director of Molecular Epidemiology Laboratory, Queen Mary University of London

BERND WOLLSCHEID

Head of Institute for Translational Medicine. Professor & Group Head, ETH Zurich

CHRIS BRAY

Head of Global Regulatory Affairs, Precision Medicine & Companion Diagnostics, Merck KGaA

DANNY KAYE

Lead Digital Pathology Scientist, Leeds Teaching Hospitals NHS Trust, NPIC

DAVID CLARK

Consultant Haematopathologist, Nottingham University Hospitals NHS Trust

DAVID KRIGE

Head of Translational Sciences, Accession **Therapeutics**

DAVID SNEAD

Professor & Consultant Pathologist UHCW NHS Trust Coventry and Director of PathLAKE

DEBAYAN MUKHERJEE

Principal Scientist of Spatial Multiplex Imaging, In Vitro/In Vivo Translation Research, Pharma R&D, GSK

EDINA SILAJDZIC

Lecturer, King's College London

ELIZABETH HARRINGTON

Global Head of Translational Medicine, Targeted Therapy Franchise, AstraZeneca

ELENA MIRANDA

Director of Non-Clinical Histology, GSK

EMANUELA OLDONI

Scientific Lead of Personalised Medicine, EATRIS, European Infrastructure for Translational Medicine

EMMANUEL VALENTIN

Vice President of Translational Medicine, ImCheck Therapeutics

ESPEN WALKER

Global Head of Medical Diagnostics, AstraZeneca

FAN LIU

Professor & Group Head, FMP Berlin

FRANCESCA TRAPANI

Scientific Director of Molecular Pathology & Laboratory Head, Boehringer Ingelheim

GAYLE MARSHALL

Head of Biomarkers, Medicines Discovery Catapult

GIOVANNA LALLI

Director of Strategy & Operations, Life Arc

HARPREET SAINI

Senior Director of Bioinformatics, Astex **Therapeutics**

HELEN GRAVES

Principal Scientist, Alchemab Therapeutics

HENOCH HONG

Associate Director, Merck KGaA

HUW BANNISTER

Senior Director of Global Diagnostics, Digital & Computational Pathology, AstraZeneca

INÊS SEQUEIRA

Associate Professor, Group Leader, Spatial Biology Hub Lead, Queen Mary University London

IRENE DEL MOLINO DEL BARRIO

Principal Scientist, GSK

JAN-PHILIPP MALLM

Head of Single Cell Open Lab, DKFZ, German Cancer Research Center

JEAN-BAPTISTE LUGAGNE

Associate Professor, University of Oxford

IEAN-CHRISTOPHE OLIVO-MARIN

Professor & Head of the Quantitative Image Analysis Unit, Institut Pasteur

JENS KIECKBUSCH

Director of External Innovation, Precision Medicine, Research & Development, GSK

IIMMY BELL

Professor & Director, Research Centre for Optimal Health, University of Westminster

JIM EYLES

Director of Clinical Immuno-Oncology Discovery Group, AstraZeneca

JO TAYLOR (BEM)

Founder, After Breast Cancer Diagnosis & METUPUK

JOANNA JANUS

Research Programme Manager (Early Detection & Prevention), Cancer Research UK



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JOÃO PINTO

Medical Specialist, Institute of Molecular Pathology and Immunology, University of Porto (Ipatimup)

JOHAN LUTHMAN

Executive Vice President and Head of R&D, Lundbeck

JEROEN VAN DER LAAK

Professor of Computational Pathology, Radboud University

JOSHUA ATKINS

Senior Genomic Epidemiologist, Cancer Epidemiology Unit (CEU), University of Oxford **KARL SMITH-BYRNE**

Senior Molecular Epidemiologist, Cancer Epidemiology Unit (CEU), University of Oxford

KARIN SCHMITT

Chief Operating Officer, Mursla Bio

KAZUMASA KANEMARU

Postdoctoral Researcher, Teichmann Group, Cambridge Stem Cell Institute, University of Cambridge

KIRI GRANGER

Chief Scientific Officer, Monument **Therapeutics**

KRISTINA HOLMBERG

Head of Clinical Biomarkers, Lundbeck

LAURENT AUDOLY

Chief Executive Officer & Co-Founder PriveBio, Inc

MANI MUDALIAR

Director of Quantitative Biomarkers, Recursion

MANUELA CERINA

Scientific Director of Neurodegeneration, LifeArc

MANUEL SALTO-TELLEZ

Professor of Integrative Pathology & Director of Integrated Pathology Unit (IPU), Institute of Cancer Research

MARÍA LAURA GARCIA BERMEJO

Scientific Director & Co-chair, Ramon & Cajal Health Research Institute (IRYCIS) & EATRIS

Biomarkers Platform

MARISCA MARIAN

Oncology Market Access Strategy Leader, Bayer

MARIA ORR

Head of Precision Medicine, Biosamples & Early Oncology, AstraZeneca

MARIYA IVANOVSKA

Chief Assistant Professor, The Medical University of Plovdiv

MARKUS SCHULZE

Senior Scientist of Clinical Biomarkers and Technologies, Merck Healthcare KGaA

MATTHEW BROWN

Chief Scientific Officer, Genomics England

MATTHEW HUMPHRIES

Director of Research Operations, Leeds Teaching Hospitals NHS Trust

MICHAEL CANNARILE

Head of Biomarkers, Early Development Oncology (EDO), Pharma Research & Development (pRED), Roche Innovation Centre Munich

MICHAEL ZAIAC

Head of Medical Affairs & Oncology, Europe & Canada, Daiichi Sankyo

MIGUEL SOUTO MORA

Director of Business Development & Innovation, IDIBELL

MIKE MESSENGER

Head of Regulatory Strategy, BIVDA

MIRO VENTURI

Operating Partner, ARCHIMED

MUHAMMAD ASLAM

Consultant Pathologist, National Clinical Lead for Digital Pathology & Al, Wales, Betsi Cadwaladr University Health Board

NATALIE BANNER

Director of Ethics, Genomics

NEIL HUMPHRYES-KIRILOV

Associate Director of Human Genomics, C4X Discovery

NIK MATTHEWS

Head of Imperial BRC Genomics Facility, Imperial College London

NITIN JAIN

Director of Franchise Project Management, AstraZeneca

OLGA NISSAN

CEO, Protica Bio

PAUL I VAN DIEST

Professor & Head, Department of Pathology, University Medical Center Utrecht

PETER GROENEN

Head Of Translational Science, Owner & Consultant, Cerebrum DAO & Alpinuity Bio **GmbH**

RAHUL DEB

Consultant Histopathologist & Lead Breast Pathologist, University Hospitals of Derby & Burton

RICHARD FESTENSTEIN

Clinical Professor of Molecular Medicine, Department of Brain Sciences, Imperial College London

SIR MARK CAULFIELD

Vice Principal for Health, Faculty of Medicine and Dentistry, Queen Mary University of London; Director of the NIHR Barts Biomedical Research Centre: President. British Pharmacological Society

SONALI NATU

Consultant Cellular Pathologist, Tees Valley Pathology Services & Clinical Lead for Pathology, North East and North Cumbria

SVETLANA MUKHINA

Director of Global Regulatory Affairs, CDx, Merck Healthcare KGaA

STEPHANIE CRAIG

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THOMAS HACH

Global Programme Clinical Head, Novartis

THOMAS JENSEN

Chief Executive Officer, Allarity

VALERIE TALY

CNRS Research Director & Group Leader, University of Paris Descartes

VICTORIA GOSS

Head of Early Diagnosis and Translational Group, University of Southampton Clinical Trials Unit

VIHANGA PAHALAWATTA

Director of Regulatory Affairs Companion Diagnostics, Abbvie

VIRGINIA SAVOVA

Senior Director of Cell Targeted Precision Medicine, AstraZeneca

WANDERSON DOS SANTOS TRINDADE

Director of Global Regulatory Affairs & Companion Diagnostics, Daiichi-Sankyo

WOLFGANG BREITWIESER

Head of Molecular Biology, Cancer Research **UK Manchester Institute**

WOUTER DE JONGE

Professor of Neurogastroenterology, Amsterdam UMC



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Pre Event Workshop | Hosted by Labcorp - 29 September 2025

Empowering Precision Medicine with Comprehensive Insights and Global Scalability

This workshop offers a deep dive into accelerating precision medicine programmes, reducing development risks, and enhancing successful outcomes using insights across various therapeutic areas. Attendees will gain valuable insights into maximising ROI in biomarker-driven development and learn from industry experts sharing real-world success focused on:

- Balancing scientific innovation with operational efficiency
- Scientific and operational strategies to enhance global reach and adoption

Closing - LUCAS RIFKIN, MD Senior Medical Director, Medical Affairs, Labcorp

End of workshop

- Optimising study design for enhanced outcomes in critical therapeutic areas, including oncology, neurology, cardiometabolic and rare diseases, using novel biomarkers
- Leveraging the right technology platform in key areas such as flow cytometry, digital pathology, and next-generation sequencing to advance immune profiling, enable AI-generated insights and unlock new genomic targets

By attending, participants will gain a comprehensive understanding of the latest trends, challenges, and opportunities in precision medicine, equipping them with the knowledge to accelerate patient access to critical therapeutics while navigating the complexities of biomarker-driven development.



13:00	Registration & Networking
13:30	Welcome - LUCAS RIFKIN, MD Senior Medical Director, Medical Affairs, Labcorp
13:45	Therapeutic Area Application: Insights into Novel Clinical Biomarkers for MASH - SANDRA LJUBICIC, MSc, PhD Senior Scientist, Cardiometabolic Biomarkers, Biomarker Solution Center, Labcorp
14:15	Therapeutic Area Application: The Role of NGS in Clinical Trials for Next- Generation Therapies - TAYLOR JENSEN, PhD VP, Head of Science Enterprise Oncology, Labcorp
14:45	Break
15:15	Current Landscape and Future Perspectives for the Role of Flow Cytometry in Drug Development, From Biomarker Discovery to Personalized Medicine - CHRISTÈLE GONNEAU, PhD Global Scientific Director, Flow Cytometry, Labcorp
15:30	Advancing Biomarker Discovery in Clinical trial through Digital Pathology and Artificial Intelligence - PAUL MESANGE, PhD Global Director of Operations, Histology, Labcorp
15:45	Effective Personalized Medicine and Disease Monitoring with Circulating DNA Biomarkers - ALEX FORT, PhD Lead Scientist, Genomics, Labcorp
16:00	Q&A Panel
16:30	Blueprint for Companion Diagnostics: Translating biomarker Insights - TUC AHMAD, Scientific Director, Diagnostic Development Services, Labcorp



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Collaboration Hub Sessions Day 1 - 30 September 2025

Bridging the Gap Between Research and Clinical Implementation in Precision Medicine

- Discussion Points: Identifying the barriers to translating research findings into clinical practice and exploring strategies to improve collaboration and align academic research with clinical needs, regulatory requirements and healthcare systems
- Invited Specialists: Academic researchers, clinicians, healthcare administrators, technology or knowledge transfer, regulatory officials and translational medicine experts

Moderated by: RICHARD BARKER, Chairman & Founder, Metadvice & New Medicine Partners

Addressing Ethical and Legal Challenges in Genetic Data Use

- Discussion Points: How to create ethical guidelines and legal frameworks for responsible genetic data use in precision medicine
- · Invited specialists: Bioethicists, legal experts, geneticists, patient advocates, government policy makers
- Moderated by: NATALIE BANNER, Director of Ethics, Genomics England

Addressing the Evolving Regulatory Landscape for Personalised Medicine

- Discussion Points: How to prepare for and adapt to shifts in regulations related to personalised health care, whilst streamlining processes
- Invited Specialists: Regulatory officials, patient representatives, clinicians

Moderated by: VIHANGA PAHALAWATTA, Director of Regulatory Affairs & Companion Diagnostics, Abbvie

Collaboration Hub Sessions Day 2 - 01 October 2025

Overcoming Barriers to Technology Adoption/Transfer in Precision Medicine

- Discussion Points: How to align stakeholders and design strategies to overcome the challenges in adopting precision medicine technologies, such as cost, accessibility, technology transfer and integration into clinical workflows
- Invited specialists: Technology developers, implementation specialists, knowledge/ technology transfer, healthcare administrators, clinicians and regulatory officials
- Moderated by: ADAMA IBRAHIM, Vice President of Digital Transformation, Research and Early Development (R&D), Novo Nordisk

Utilising AI in Precision Medicine

- Discussion Points: How to maximise the benefit of AI in precision medicine and explore challenges around managing risk, ethical considerations, regulatory compliance and integrating into existing systems
- Invited Specialists: Al specialists, data scientists, clinicians, regulatory officials, bioethicists and IT infrastructure experts
- Moderated by: James Schofield, Founder & Chief Executive Officer, TopMD Precision Medicine

Integrating and Sharing Data in Precision Medicine

- Discussion Points: Designing strategies to improve integration and analysis of diverse data sources (genomics, proteomics, patient health records) and also the sharing of datasets internally and externally to further precision medicine research
- · Invited Specialists: Data scientists, bioinformaticians, clinicians and IT infrastructure experts
- Moderated by: TBA

12:15

15:25

15:25

16:15

12:10

13:00



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08:40 **Oxford Global's Welcome Address**

Exhibition Room - Plenary Keynote Address - Improving Rare Disease Diagnostic Performance Using Multiomics

• Short-read whole genome sequencing provides genetic diagnoses for about 30% of patients referred with rare diseases. Although many of these patients will have oligo- or polygenic diseases, or diseases of environmental origin.

• Many patients with monogenic diseases are thought to remain undiagnosed using this single technology approach.

- 08:45 New approaches, including long-read DNA sequencing, transcriptomics, proteomics and metabolomics have potential to provide diagnoses for a significant proportion of those where short-read sequencing is not diagnostic.
 - Future perspectives at England and wider industry.

	MATTHEW BROWN, Chief S	MATTHEW BROWN, Chief Scientific Officer, Genomics England											
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	Track Chair: Deepak Bhere, Assistant Professor & Principal Investigator, University of South Carolina	Track Chair: EVA RODRIGUEZ-SUAREZ, Biomarker Lead/ Biomarker Strategy Immunology, Novartis	<i>Track Chair:</i> Alex Dulovic, Group Leader, NMI	Track Chair: Jose Izarzugaza, Director, Biomarker & MoA Insights, Novo Nordisk	Track Chair: Nik Matthews, Head of Imperial BRC Genomics Facility, Imperial College London	Track Chair: Stephanie Traub, Associate Director, BSM & Biomarker Lead, UCB	Track Chair: Jean-Christophe Olivo-Marin, Professor & Head of the Quantitative Image Analysis Unit, Institut Pasteur						
	Track Keynote Address: Biomarker Strategies For Patient Identification	Track Keynote Address: CDx Regulatory Framework In The US And EU	Track Keynote Address: The Impact Of Biomarkers In Early Clinical Trial Design And Decision Making	Track Keynote Address: Precision Medicine, Proteomics And Problem Solving	Track Keynote Address: Spatial Multiplex & Transcriptomics Applications In Drug Discovery	Track Keynote Address: Cell-Cell Interaction And Tissue Architecture In Health And Disease	Conference Room 7 Track Keynote Address: Al Implementation: Who Pays The bills?						
9:15		While companion diagnostics help to identify the best treatment options for patients, their development represents a major challenge due to the complexity of regulatory requirements and demands efficient coordination between drug developers, IVD companies and Health Authorities. This presentation shares the perspective from a Pharma company on the regulatory strategy that enables successful drug/CDx co-development and co-approval in the US and EU.	Decision-making for new molecular entities in early clinical development often suffers from the risk of false positive and negative decisions driven by small heterogeneous, and often advanced stage patient populations. This talk will explore how biomarker concepts can mitigate such risks by enhancing patient enrichment, supporting dose selection, and demonstrating evidence of mechanism in Phase I clinical development.	Why genetics are not enough? Examples and limitations of proteomics in common and rare diseases How can academia, industry, patients and regulators (alphabetically listed) find pragmatic solutions?	Advances in multiplexing and computational analysis have revolutionised our comprehension of tissuespecific biology and therapeutic interactions. Spatial multiplexing facilitates drug discovery by characterising patient biopsies, elucidating model systems, and identifying novel biomarkers. It also contributes to understanding drug distribution, molecular heterogeneity, and the mechanisms underlying off-target toxicity. This presentation provides an overview of recent developments in the field, along with examples from projects.	Cell-cell interactions and tissue architecture are crucial in maintaining health, influencing cellular behavior and tissue homeostasis. Disruptions in these interactions contribute to disease pathogenesis, including cancer and autoimmune disorders. Understanding these dynamics can reveal mechanisms of disease and identify therapeutic targets, highlighting the interplay between cellular microenvironments and tissue architecture as pivotal in health and disease management.	The possibilities for clinically using AI in daily diagnostic pathology are rapidly increasing with many good algorithms that are brought to the market, often with certification. However, implementation is not taking off rapidly for two main reasons: implementation is technically still challenging, and the business case needs to be developed. In this presentation, the ins and outs of the business case for AI implementation in pathology will be discussed.						
	MICHAEL ZAIAC, Head of Medical Affairs, Oncology, Europe & Canada, Daiichi Sankyo	SVETLANA MUKHINA, Director of Global Regulatory Affairs & CDx, Merck Healthcare KGaA	MICHAEL CANNARILE, Head of Biomarkers, Early Development Oncology (EDO), Pharma Research & Development (pRED), Roche Innovation Centre Munich	THOMAS HACH, Global Programme Clinical Head, Novartis	ELENA MIRANDA, Director of Non-Clinical Histology, GSK e between conference rooms	VIRGINIA SAVOVA, Senior Director of Cell- Targeted Precision Medicine, AstraZeneca	PAUL J VAN DIEST, Professor and Head, Department of Pathology, University Medical Centre Utrecht						



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Poster Displays & 1-2-1 Meetings x4

MORNING NETWORKING BREAK & REFRESHMENTS:

10:05



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	From Concept to Clinic: The Emerging Role Of Next Generation Multi- Omics In Biomarker- Driven Drug Development	Running The CDx Gauntlet: Developing A Commercial Ready CDx, Navigating IVDR, And Preparing For Market Access	Solution Provider Presentation	Solution Provider Presentation	Scaling Spatial Biology With The Phenocode™ IO60 Panel: Real-World Experience From Propath	Maximise Insights, Maximise Success: Shaping the Future Of Biomarker Development With 10x Genomics Single Cell And Spatial Technologies	Pixels with Purpose: Advancing Precision Medicine With Digital Pathology			
	SAPIENT	6			AKOYA BIOSCIENCES*					
	How mass spectrometry offers a single platform	DISCOVERY			During this session, Akoya Biosciences (A Quanterix	10× GENOMICS	Due sision as adiaire ademande as sue the selicent terrented			
11:25	for multi-omics analyses of dynamic proteins, metabolites, and lipids across sample matrices, species, and development phases • How these approaches de-risk and accelerate translation of known and novel drug targets and biomarkers, from target identification to clinical assays • How Sapient's DynamiQ™ Insights Engine provides a clinical and molecular reference database to strengthen decision confidence by validating findings across diverse populations	 Overview of developing prototype companion diagnostics, balancing early trial performance with commercialisation, illustrated through case studies Use of LDTs to reduce costs and timelines, with compliance to IVDR, UKCA, and FDA requirements supported by our Kassel laboratory Pathways to transition from LDTs to commercial CDx solutions, accelerating patient sample access and ensuring readiness for market launch 	Personalis*	mission bio	Company) will present the PhenoCode™ IO60 panel, an ultrahigh-plex solution for deep spatial phenotyping of the tumor microenvironment. Kelly Hunter, Chief Scientific Officer at Propath, a scientifically driven CRO with extensive wet and dry lab expertise, will share his experience automating IO60 for scalability. Together, Akoya and Propath will highlight how automation and scientific partnership enable reproducibility, accelerate translational research, and generate actionable biological insights.	10x Genomics accelerates biopharma research with single cell, spatial, and in situ multiomics- advancing drug development, biomarker discovery, and translational insights. This presentation will highlight our key technologies and latest innovations.	Precision medicine demands more than just targeted therapies—it requires targeted tools. Digital pathology is emerging as a critical enabler in identifying, validating, and operationalizing biomarkers that drive patient stratification a therapeutic success. In this session, we'll share how Al-power image analysis, multiplex imaging, and robust data platforms to deepen biological understanding and accelerate drug development. Through concrete examples, we'll demonstrate how digital pathology workflows can integrate seamlessly into biomarker discovery and companion diagnostic programs.			
	MO JAIN, Founder and Chief Scientific Officer Sapient	SCOTT REID, VP & Global Head of Companion Diagnostics, Discovery Life Sciences	Senior Representative, Personalis	Senior Representative, Mission Bio	KELLY HUNTER, Chief Scientific Officer, Propath UK	Senior Representative, 10x Genomics	PAUL MESANGE, Global Director of Histology, Labcorp			



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	Transforming Cancer Patient Management Through Digital Pathology And Spatial Profiling	IVDR Performance Evaluation - A Roadmap For Innovators	Building An Early Diagnosis Trial Group: Previous, Current And Future Studies	Precision Phenotyping For Precision Medicine	Spatial Omics Technologies In A Cancer Research Facility	Approaches To Leverage Spatial Biology And Machine Learning For Clinical Biomarker Development	Transforming Cellular Pathology At Nottingham University Hospitals - Delivering 100% Digital Pathology Reporting And Integrating Al Into Routine Diagnostic Practice					
:50	This talk highlights how digital pathology and spatial profiling can potentially revolutionise care for Head and Neck Squamous Cell Carcinoma (HNSCC). To address overtreatment driven by limited biomarkers, we explore single-cell spatial multiomics for novel prognostic and predictive marker discovery. Biomarkers to segregate patients into high- and lowrisk recurrence groups can guide personalised therapy and improve outcomes.	This talk will provide a strategic roadmap for innovators navigating performance evaluation under the EU IVDR. It will introduce the core pillars of scientific validity, analytical performance, clinical performance, and human factors, outlining key frameworks for compliance. Through practical insights and case examples, we will explore how to optimise efficiency, reduce costs, and streamline timelines in regulatory submissions.	In July 2023, the Southampton Clinical Trials Unit established a dedicated Early Diagnosis and Translational Group to support the increasing number of studies in this field. This talk will provide an overview of these trials and share first-hand insights into the highs and lows of running studies in early diagnosis and translation.	By using in-depth MRI-based phenotyping we are enabling precise, non-invasive assessment of organ structure and function. This approach supports the development of novel biomarkers to stratify disease risk, guide early diagnosis, tailor treatments, and optimise clinical trial design. By integrating rich imaging-derived phenotypes into precision medicine, we advance predictive, preventive, and therapeutic strategies across diverse patient populations.	Spatial omics processing involves complex workflows requiring sophisticated skills in histology, molecular biology, as well as scientific computing and bioinformatics. In our institute, for handling spatial transcriptomics projects, multiple core facilities provide a comprehensive service that includes all aspects of sample processing including tissue QC, staining and imaging, region selection and capture, followed by library prep, sequencing and data processing using dedicated computational pipelines.	 The importance of spatial context in drug development Spatial context and antibody drug conjugates (ADCs) Derivation of context information from H&E images using machine learning: Tumor microenvironment and HRD Resolving spatial context by multiplex immunofluorescence 	Nottingham University Hospitals (NUH) cellular pathology is one of the largest acute teaching hospital trusts in the NHS. The department has implemented Indica labs HALO AP digital pathology system. 100% of cases are scanned an reported with a completely digital workflow. Pathologists work in a paperless, glassless virtual workspace. Al tools have been integrated into the routine clinical workflow including the IBEX Galen prostate and breast tools and the Indica labs Macrodissect tools. This presentation will describe the implementation journey, outcomes and lesson learned.					
	ARMAN RAHMAN,	MIKE MESSENGER,	VICTORIA GOSS, Head of Early Diagnosis and Translational Group,	JIMMY BELL, Professor & Director, Research	WOLFGANG BREITWIESER, Head	MARKUS SCHULZE, Senior Scientist of Clinical Biomarkers &						

Q&A session & transition time between conference rooms

Centre for Optimal

Westminster

Health, **University of**

of Molecular Biology,

Cancer Research UK

Manchester Institute

Merck Healthcare

DAVID CLARK, Consultant Haematopathologist, **Nottingham University Hospitals NHS Trust**

Technologies,

KGaA

Assistant Professor of

Anatomy, **University College Dublin**

Head of Regulatory

Strategy, **BIVDA**

University of

Trials Unit

Southampton Clinical



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Integrates with Swiss/ ETH projects to enhance clinical decisions

BERND WOLLSCHEID,

Translational Medicine,

Head of Institute for

My talk will explore its potential to enhance diagnostics, and accelerate translational research, using the example of application to clinical PD-L1 reflex testing. Drawing on experience across national molecular programmes and digital pathology innovation, I will reflect on key challenges and opportunities in advancing this powerful technology at

MATTHEW HUMPHRIES, Director of Research

Update On Cellular Pathology AI Deployment

An overview of the recent developments and experiences in deploying AI technologies across Wales in cellular pathology. It covers case studies, key challenges, and measurable outcomes, with a focus on digital pathology and Al integration in clinical setting. The talk aims to highlight both technical progress and the diagnostic impacts of Al deployment in the region.

MUHAMMAD ASLAM, Consultant Pathologist, National Clinical Lead for Digital Pathology and Al, Wales, Betsi Cadwaladr University Health Board

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12:40

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KIRI GRANGER, Chief Scientific Officer, **Monument Therapeutics**



signalling

• Precision Medicine:

Professor & Group Head, ETH Zurich

INTERACTIVE

Operations, **Leeds Teaching Hospitals NHS**

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	Conference Room 1 - Pand The Next Era Of Precision Collaboration Across Bior Omics & Therapeutic Inno	Medicine: Stakeholder narkers, Diagnostics,	Conference Room 2 - Panel Discussion: The Future Of Precision Medicine And Benefits For The Patient		Conference Room 5 - Par Addressing The Future No Omics • Future of spatial biology • Moving towards the practical findings	eeds Of Spatial Multi-	Conference Room 7 - Panel Discussion: Building A Business Case For Digital Pathology • Wants and needs from different stakeholders • ROI • Clinical benefits • Efficiency & quality gains	
3:05	Moderator: MIKE MESSENGER, Head of Regulatory Strategy, BIVDA Panellists: SENIOR REPRESENTATIVE, Labcorp MARISCA MARIAN, Oncology Market Access Strategy Leader, Bayer ALEXANDRA SEVKO, VP of Translational Science, Amphista Therapeutics KIRI GRANGER, Chief Scientific Officer, Monument Therapeutics		Moderator: LAURENT AUDOLY, Cheif Executive Officer & Co-Founder, PriveBio, Inc Panellists: JOHAN LUTHMAN, Executive Vice President and Head of R&D, Lundbeck MIRO VENTURI, Operating Partner, ARCHIMED MO JAIN, Founder & Chief Scientific Officer, Sapient BELINDA NEDJAI, Associate Professor & Director of Molecular Epidemiology Laboratory, Queen Mary University of London		Panellists: AZAM HAMIDINEKOO, Associate Director, Clinical Pharmacology & Safety Sciences, AstraZeneca FAN LIU, Professor & Group Head, FMP Berlin INÊS SEQUEIRA, Associate Professor, Group Leader, Spatial Biology Hub, Queen Mary University London AMONIDA ZADISSA, Associate Director of Informatics, UK Dementia Research Institute NIK MATTHEWS, Head of Imperial BRC Genomics Facility, Imperial College London Senior Representative, Ariadne.ai		Moderator: DAVID SNEAD, Professor & Consultant Pathologist UHCW NHS Trust Coventry and Director of PathLAKE Panellists: HUW BANNISTER, Senior Director of Global Diagnostics & Digital and Computational Pathology, AstraZeneca MATTHEW HUMPHRIES, Director of Research Operations, Leeds Teaching Hospitals NHS Trust PAUL J VAN DIEST, Professor and Head, Department of Pathology, University Medical Center Utrecht SONALI NATU, Consultant Cellular Pathologist, and Clinical Lead for Pathology, Northeast and North Cumbria, Tees Valley Pathology Services VANDANA MALLEMPATI, Senior Director of Product	
		INTERACTIVE		INTERACTIVE		INTERACTIVE	Management, Iron Mountain INTERACTIVE	
3:35				LUNCH BREAK:	Poster Displays & 1-2-1 M	leetings x3		
	<i>Track Chair:</i> Folake Orafidiya, Senior Scientist, Avacta	Track Chair: ALAIN VAN GOOL, Professor of Personalised Healthcare, Radboud University Medical Centre	Track Chair: Emily Adams, Associate Professor, University of Oxford	Track Chair: Harpreet Saini, Senior Director Bioinformatics, Astex Therapeutics	Track Chair: TBA	Track Chair: Stephanie Traub, Associate Director, BSM & Biomarker Lead, UCB	Track Chair: TBA	
	Solution Provider Presentation	Start-Up Zone Pitches 10-Minute Presentation: UltraBright Biotech	Designing For Success: Navigating Early- Phase Translational Medicine Studies	Proteomics To Advance Precision Medicine Seer Application of different omic	Single-Cell Identity And Heterogeneity In Its Environment: Where Innovations Push The Understanding Of Spatial Biology	Spatial Proteomic Characterisation Of Renal Cell Carcinomas Identifies Metabolic Reprogramming Of The TME Associated With	Silver Level Solution Provider Presentation	
	Quanterix Discovery Fueled by Ultra-Sensitivity	10-Minute Presentation: EndoGene.bio	Practical insights and real-world case studies from ARC's collaborations with pharmaceutical and	technologies is now feasible at population scale. This talk will present examples of how the integration of proteomics in large patient and population studies can	Leica weedstatus	Disease Progression Comparison Spatial proteomics provides valuable insights into cell characterisation within the	Attendees Are Welcome To Attend Co-Located Sessions	
4:35		endo gene .bio	diagnostic teams on early- phase translational and precision medicine trials.	help to predict disease risk, understand mechanisms, and reveal shared connections between rare and common diseases. Studies include different proteomic technologies and an investigation how these can be combined and how their complementarity can be employed for synergistic insights into human health.		tumour microenvironment. Using our Paletrra™ Aldriven solution, we explored the relationship between the immunogenic profile of clear cell renal cell carcinoma (ccRCC) and metabolic reprogramming. This end-to-end platform generates whole-tissue data with enhanced histological context, enabling unbiased spatial analysis to advance		
	Senior Representative, Quanterix		BETHANY MONTGOMERY, Technology Specialist, ARC Regulatory	CLAUDIA LANGENBERG, Director, Seer	PRZEMEK FLESZAR, Advanced Workflow Specialist, Leica Microsytems	understanding and therapeutic development. Senior Representative, Neo Genomics		

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	Why Genomic Profiling Is Important For Patients' Survival And Outcomes?	Navigating Complexity Of Biomarker Subgroups To Accelerate Drug Approval	Translating Cancer Research Into Novel Therapies	Predicting Therapeutic Response Using Genomics In Inflammatory Diseases And Beyond	Revealing Spatial Interactome Of The Cell By Cross-Linking Mass Spectrometry	IMMUcan: Large-Scale Spatial IO Profiling For Biomarker And Target Discovery	The NPIC AI FORGE: A Unique Multi-Scanner Facility For Data Acquisition For Digital Pathology Artificial Intelligence			
15:00		The precision medicine strategy gets complex for oncology studies involving multiple biomarker subgroups and diagnostics tests. For these trials, regulators expect robust evaluation of efficacy in each biomarker subgroups enrolled in the trial before deciding the final drug approval label. These perspectives will be discussed as well as to how effective biomarker/ diagnostic testing strategies can minimise delays to drug approval.	 Biomarker strategies for the identification of the right patients for targeted therapy Biomarkers for monitoring drug response Translational analysis of mechanisms of resistance to guide future combination strategies 	Many patients with inflammatory diseases do not respond to individual therapies. Genomics plays a role in this process. We have used a Bayesian approach to analyse common variants in an ulcerative colitis cohort to identify a genomic signature that can predict response to anti-integrin therapy. This will be developed for use in clinical trial design and/or treatment selection.	The specific functions of cellular organelles and sub-compartments depend on their protein content, which can be characterised by spatial proteomics approaches. Here, we develop a cross-linking assisted spatial proteomics (CLASP) to map sub-organelle proteomes and membrane protein topology and revealed spatial interactomes of organelles such as the mitochondria and even the whole cell.	 IMMUcan is an EU-funded public-private consortium profiling the TME of >2,500 cancer patients The novelty is to combine molecular and spatial cellular readouts at large scale Use cases for biomarker and drug target discovery will be discussed 	The development and use of AI within digital pathology is becoming more common and offers a myriad of potential uses. Despite the promise of AI, it is not without limitations. My talk will discuss our novel multi-scanner facility, which can support the creation of richer datasets through the replication of pathology slides across multiple scanners from different vendors, supporting the creation of more generalisable AI.			
	JO TAYLOR BEM, Founder, After Breast Cancer Diagnosis & METUPUK	NITIN JAIN, Director of Franchise Project Management, AstraZeneca	ELIZABETH HARRINGTON, Global Head of Translational, Medicine &Targeted Therapy Franchise, AstraZeneca	NEIL HUMPHRYES- KIRILOV, Associate Director of Human Genomics, C4X Discovery	FAN LIU, Professor & Group Head, FMP Berlin	HENOCH HONG, Associate Director, Merck KGaA	DANNY KAYE, Lead Digital Pathology Scientist, Leeds Teaching Hospitals NHS Trust, NPIC			
				Q&A session & transition tin	me between conference rooms					
	From Plasma To Placenta: Translational	Solution Provider Presentation	Comprehensive Genomic Profiling To Support Global	Practical Implementation of Clinical Standards	MACSima™: The Complete Toolbox For Spatial Biology	Creative Solutions To Scientific And Technical Challenges In	Silver Level Solution Provider			
	Proteomic Signatures Across Diverse Clinical	SURFIX Redefining Point-of-Care Testing.	Oncology Clinical Trials	and IVDR Compliance in Multi-Omic Data		Digital Pathology	Attendees Are Welcome To Attend Co-Located Sessions			
	Matrices		RULES BASED MEDICINE an IQVIA business	GEN SEQ	À	Concept Life Sciences a Maliver of Paracytical Load				
15:25	Synexa Life Sciences applies Olink® PEA to diverse matrices, enabling multiplex proteomic analysis across disease areas. We identified neuroinflammatory markers in PTSD serum, phenotypespecific differences in HIV plasma and CSF, and immune dysregulation in placental tissue. These studies showcase Olink's sensitivity and highlight sample selection's critical role in translational biomarker research.		Illustrating IQVIA Laboratories' approach to supporting our sponsor's global oncology clinical trials, including regional considerations - such as IVDR in Europe - and assay validation and verification approaches, with a focus on Illumina's TruSight Oncology 500 tissue and ctDNA assay portfolio of next-generation sequencing assays for mutation profiling.	Genseq, an ISO15189, CAP/ CLIA clinical diagnostics company, delivers multi- omic genomics solutions for clinics and biopharma. This talk explores applying clinical, regulatory, and sustainability standards to genomics workflows, covering validation of single-cell/nuclei workflows in NASH/NAFL tissue, ISO15189 companion diagnostic services, and IVDR-compliant whole exome sequencing development.	The emergence of spatial biology heralds a paradigm shift in our understanding of biological systems, revealing previously unseen layers of complexity. Here, we will discuss how MACSima is driving advancements in disease therapies and unraveling the complexity of the tumor microenvironment.					
	ANDREIA SOARES, Head of Scientific Strategies, Synexa Life Sciences	Senior Representative, Surfix Diagnostics	DAVID LATTO, Director Biomarker Liason, Rules Based Medicine	Senior Representative, GenSeq	LILIA DRAGANOVA, PhD, Instrument Sales Specialist - UK North & Ireland, Miltenyi Biotec	Dr Marta Czapranska Senior Scientist, Concept Life Sciences				

Q&A session & transition time between conference rooms

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	Day One 30 September 2025										
		BIOMARKERS & PR	ECISION MEDICINE		SPATIAL BIOLOGY FOR	PRECISION MEDICINE	DIGITAL PATHOLOGY & AI				
	CONF ROOM 1: WESTMINSTER - BIOMARKER IDENTIFICATION, VALIDATION, IN ONCOLOGY, IO & IMMUNOLOGY	CONF ROOM 2: ST JAMES - DIAGNOSTIC BIOMARKERS & REGULATIONS	CONF ROOM 3: MOORE - BIOMARKERS FOR CLINICAL DEVELOPMENT	CONF ROOM 4: ABBEY - GENOMIC & MULTI- OMIC APPROACHES IN BIOMARKER DISCOVERY & DEVELOPMENT	CONF ROOM 5: GIELGUD & BURTON - SPATIAL MULTI- OMICS TECHNIQUES & APPROACHES	CONF ROOM 6: OLIVIER - SPATIAL BIOLOGY IN PHARMA & TRANSLATIONAL DRUG RESEARCH	CONF ROOM 7: RUTHERFORD - DIGITAL PATHOLOGY IMPLEMENTATION - IMAGE ANALYSIS & AI				
	Use Of Mass-Spec Based Proteomics For Biomarker Discovery	Precision Medicine Beyond Oncology: Unlocking the Next	Of A Novel AI- Based Alzheimer's	Automating Sample Prep: Next-Generation Workflows for	Accelerating Spatial Proteomics: Innovations In Spectral	Spatial Biology At Scale With Orion Technology	Bronze Level Solution Provider				
	In Clinical Plasma And Tissue Samples	Frontier in Biomarker- Driven Therapies		Diagnostic. From Bior Assay Concept To	Biomarker Discovery opentrons	Multiplex Fluorescence Imaging With The Invitrogen™ EVOS™	RARECYTE	Attendees Are Welcome To Attend Co-Located Sessions			
15:50	BIOGNOSYS NEXT GENERATION PROTEOMICS	Precision medicine has significantly impacted cancer drug development and treatment. This talk will review how lessons from the oncology field are now being applied to other therapeutic areas, including cardiometabolic and neurodegenerative diseases.	And Regulatory Submission CellCarta	How automation streamlines reproducible LC-MS sample prep for proteomics and genomics New technologies to enable fully automated workflows Increase throughput without increasing footprint	Thermo Fisher S C I E N T I F I C Genseq is an ISO15189, CAP/CLIA clinical diagnostics company providing clinics and biopharma partners with multi-omic genomics solutions. The talk will summarise the experience of applying multiple clinical, regulatory and sustainability standards to genomics data generation workflows including validation of single cell/nuclei workflows in NASH/NAFL tissue, parallel development of ISO15189 companion diagnostic services for treatments and IVDR compliance to whole exome sequencing.	Orion revolutionises spatial biology with high-throughput, single-round imaging, eliminating iterative cycling and enhancing image quality. Delivering 18 plex in a single scan, Orion streamlines workflows and accelerates discovery. It enables comprehensive microenvironment profiling, uncovering disease mechanisms and advancing prognostic and predictive biomarker discovery. Case studies showcase its transformative applications.					
	DANIEL REDFERN, Senior Director of Business Development, Biognosys AG	DAVID FABRIZIO, VP, Commercial Strategy and Medical Innovation, Foundation Medicine	TODD CHERMAK, SVP & Global Business Head Immunology & Proteomics, Cellcarta	DANIEL LORD, Sales Director, Opentrons	ADYARY FALLARERO, Senior Product Manager, Thermo Fisher	MICKAEL MEYRAND, Field Application Scientist, Rarecyte					



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AFTERNOON NETWORKING BREAK & REFRESHMENTS:

Poster Displays & 1-2-1 Meetings x3

Translation In The

DEBAYAN MUKHERJEE,

Spatial Multiplex Imaging,

Research, Pharma R&D,

In Vitro/ In Vivo Translation

Principal Scientist of

Clinic





Preclinical Strategies To Define Immuno-**Oncology Biomarkers For Clinical Translation**

the Clinical Immuno-

JIM EYLES, Director of

Discovery Group,

AstraZeneca

AstraZeneca.

Oncology Discovery team at

The In Vitro Diagnostic Immuno-oncology agents are often combined Regulation and the FDA with standard-of-care LDT Rule represent a therapies or other significant paradigm shift in requirements related to immunomodulatory molecules, presenting the use of biomarker tests challenges in tracking in pharmaceutical clinical biomarkers intrinsically trials. Clinical trial sponsors associated with the need to understand and contribution of specific proactively address changes components. To address while navigating the these challenges and complexities to ensure that enhance the effectiveness the data generated through the trial is compliant with the of clinical development regulations and can support programmes, rational experimentation using subsequent regulatory suitable preclinical submissions. This session is models is invaluable. This geared towards identifying presentation will outline specific challenges and strategies employed by potential solutions.

VIHANGA PAHALAWATTA, Clinical Immuno-Oncology Director of Regulatory Affairs & Companion

Diagnostics, Abbvie

Involving Lab

Navigating The Path: ICT01, First-In-Class Considerations Anti-BTN3A mAb **For Clinical Trials Selectively Activating** y9δ2 T Cells: **Developed Tests Translational Results** From Phase I And **Dose Selection**

> ICT01 is an anti-BTN3A monoclonal antibody that selectively activates y9δ2 T cells, triggering a downstream immunological cascade through secretion of pro-inflammatory cytokines, and further augmenting the antitumour immune response. Favourable safety profile together with promising efficacy signals is observed in multiple haematological and solid tumour patients, and the 10 mg ICT01 dose is selected as RPD2 based on PK/PD modelling.

EMMANUEL VALENTIN, Vice President of Translational Medicine, **ImCheck Therapeutics**

Omics Technologies And Computational Approaches In Drug Discovery

Discovering and developing new drugs is a timeconsuming and expensive process. Use of genomewide, high-throughput omics technologies and computational approaches for integrating multi-dimensional data in drug discovery can lead to more informed drug target selection, biomarker identification and accelerated drug development. In my talk, I will present examples of how multi-omics data integration approaches can be used for identifying

biomarkers for patient

and clinical settings.

stratification in pre-clinical

HARPREET SAINI, Senior Director of Bioinformatics, **Astex Therapeutics**

Uncovering Scarless Healing: A Multiomics Atlas Of Oral & Skin Fibroblast Heterogeneity

Oral mucosa heals with minimal scarring, unlike skin, but its cellular diversity remains poorly understood. Using single-cell and spatial omics, we mapped fibroblast heterogeneity across human and mouse oral tissues and skin, revealing key regenerative populations. Our multimodal atlases uncover molecular mechanisms of scar-free healing and offer a valuable resource for advancing tissue repair research.

INÊS SEQUEIRA, Associate Professor, Group Leader, Spatial Biology Hub Lead, **Queen Mary University** London

Digital Pathology Multimodal Analysis And Complex Biomarkers In Oncology

There is a mismatch between the complexity of the biology targeted by new drugs and the simplicity of the predictive biomarkers in use; this is why DNA damage response or immuno-oncology related drugs have a small repertoire of predictive biomarkers. In this lecture, we will explore ways to create a new generation of complex biomarkers applicable in the routine setting.

MANUEL SALTO-TELLEZ, Professor of Integrative Pathology & Director of the RMH/ICR Integrative Pathology Unit, Institute of Cancer Research

17:40

NETWORKING DRINKS & END OF DAY ONE



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BREAKFAST ROUNDTABLES

Biomarkers

Improving Support For Industry-Academic Collaboration

This session will explore how industryacademic collaboration can accelerate digital pathology and biomarker discovery for cancer. We will specifically discuss how smaller industry partners can leverage academic settings as their R&D engine, utilising Al-driven and multiomics platforms to bridge the gap between research and clinical application. The goal is to personalise cancer treatment and advance the future of precision medicine.

Moderator: ARMAN RAHMAN,

College Dublin

Assistant Professor, **University**

Biomarkers

Introducing Al-Derived Digital Biomarkers For Clinical Trials And Patient Monitoring

Assessing Clinical Trials For Global IVD Regulatory And Quality Compliance

Biomarkers

Biomarkers play a pivotal role in evaluating clinical trials, offering measurable indicators of safety, efficacy, and compliance. Their integration is essential for meeting global in vitro diagnostic (IVD) regulatory standards and ensuring quality assurance. Standardised biomarker validation supports harmonisation across jurisdictions, strengthens trial credibility, and accelerates translation of innovations into practice, ultimately improving patient outcomes and regulatory confidence worldwide.

Moderator: MARIYA IVANOVSKA, Chief Assistant Professor, **The Medical University of Plovdiv**

Spatial Biology

FAIR Data Practices And Project Design For Spatial Profiling Techniques

Interactive discussion about unresolved issues regarding FAIR spatial

Topics (among others):

- Open questions / issues regarding metadata management
- Data management plan: sample experimental setup - data acquisition
- Repositories for sharing data & data linking
- Questionnaire: share your challenges

Key results will be shared at the end of the conference by Amonida Zadissa (see programme).

Moderator: JAN-PHILIPP MALLM, Head of Single Cell Open Lab **DKFZ**, **German Cancer Research Center &** AMONIDA ZADISSA, Associate Director of Informatics, **UK Dementia Research Institute**

Digital Pathology & Al

Imaging Based Standardisation In Digital Pathology: A **Necessary Step Or A Barrier To** Innovation?

Imaging based standardisation in digital pathology promises interoperability, safety and AI readiness enabling cross-site reporting and consistent quality. Strict standards may limit vendor innovation, would be associated with un-necessary transition costs and risk lowest common denominator solutions. Balancing harmonisation and flexibility is crucial. The roundtable is a discussion on how to balance harmonisation with flexibility so that it can be an enabler of progress and not be a barrier if applied to rigidly.

Moderator: SONALI NATU, Consultant Cellular Pathologist, **Tees Valley Pathology Services and Clinical Lead for** Pathology, Northeast and **North Cumbria**

Digital Pathology & Al

Real World Learning From Deploying Digital And Computational Pathology At Scale

Moderator: HUW BANNISTER, Senior Director of Global Diagnostics, Digital and Computational Pathology,

AstraZeneca

Conference Room 1: Fireside Chat with Professor Sir Mark Caufield

Moderator: RICHARD

of Molecular Medicine,

Imperial College London

FESTENSTEIN, Clinical Professor

Department of Brain Sciences,

08:40

08:05

PROFESSOR SIR MARK CAULFIELD, Vice Principal for Health, Queen Mary's Faculty of Medicine and Dentistry; Director of the NIHR Barts Biomedical Research Centre; President, **British Pharmacological Society**

Moderator: MIRO VENTURI, Operating Partner, ARCHIMED

Q&A session & transition time between conference rooms



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	CONF ROOM 1: WESTMINSTER - BIOMARKER IDENTIFICATION & VALIDATION: NEUROSCIENCE & CO- MORBIDITY	AARKER IDENTIFICATION - NEW & EMERGING BIOMARKER TECHNOLOGIES TRIALS CONF ROOM 3: MOORE - BIOMARKER FOR CLINICAL TRIALS		CONF ROOM 4: ABBEY - BIOMARKERS FOR DIAGNOSTICS & PRECISION MEDICINE	CONF ROOM 5: GIELGUD & BURTON - AI IMAGE ANALYSIS & DATA ANALYTICS IN SPATIAL BIOLOGY	CONF ROOM 6: OLIVIER - COMPUTATIONAL PATHOLOGY & UTILISING DIGITAL PATHOLOGY IN PHARMA RESEARCH
	<i>Track Chair:</i> Giovanna Lalli, Director of Strategy & Operations, Life Arc	Track Chair: James Schofield, Founder & Chief Executive Officer, TopMD Precision Medicine	Track Chair: Paul Rhyne, Senior Director, Scientific Solutions, Luminex	Track Chair: Valerie Taly, CNRS Research Director & Group Leader, University of Paris Descartes	Track Chair: Priya Narayanan, Senior Scientific Officer, Institute of Cancer Research	<i>Track Chair:</i> Vandana Mallempati, Senior Director of Product Management, Iron Mountain
	Track Keynote: Precision Medicine In Neuroscience: The Role Of Digital Biomarkers	Track Keynote: Proteomics-Based Predictive Biomarkers For Immunotherapy Response	Track Keynote: Patient Selection Approaches in AstraZeneca	Track Keynote: Multimodal Diagnostics For Precision Oncology: An Industry Perspective	Track Keynote: Advancing Biomarker Discovery Through Al-Driven Spatial Biology: A Comprehensive Pipeline Approach	Track Keynote: Next-Gen Al-Based Breast Cancer Grading For Improved Therapy Selection
9:10	Neuroscience drug development has historically suffered a low success rate, in part due to a lack of diagnostic specificity. We are entering a new era of precision medicine in psychiatry and neurology, enabled by novel digital biomarkers. Monument Tx is a clinical-stage biotech company developing stratified therapeutics for schizophrenia and neuroinflammation, exemplifying this approach.	Immune checkpoint inhibitors show limited response in Head and Neck cancer. We developed a proteomics-based classification system, analysing 128 patients' FFPE samples with 11,000+ proteins per sample. Machine learning-based classification achieved 86% sensitivity, 90% PPV. Identified resistance mechanisms offer insights for personalised immunotherapy strategies in HNSCC.	Precision Medicine is central to AstraZeneca's scientific endeavours. By deepening our understanding of response biology and advancing diagnostic innovation, we tailor treatments to individuals for optimal outcomes. This presentation showcases our groundbreaking approaches to delivering the right treatments to patients.	Precision oncology has evolved rapidly, with over 65% of solid cancer approvals targeting specific genomic alterations. Industry, associations, and clinicians need to work together to see the full potential of diagnostics. Combining IHC, NGS, ctDNA, and digital pathology will define the new era of multimodal diagnostics.	Spatial Biology, powered by AI, is advancing biomarker discovery and therapeutic innovation. We developed and utilised an agile pipeline to translate and unlock the information in spatial imaging techniques and decipher cell interactions to inform treatment dynamics. By integrating spatial data with AI-driven analysis, in our use case, the results demonstrated enhanced understanding of cellular dynamics, supporting biomarker identification and enabling personalised treatment strategies along with tailored therapeutic approaches.	We studied the use of deep learning to assess a range of breast-cancer related tissue features: presence of lymph node metastases, extent of lymphatic infiltrate within tumours, and the components of tumour grading. It was shown that DL enables reproducible, quantitative tumour feature extraction, showing a good correlation with pathologists' scores and with patient outcome. Our current research involves larger-scale validation with pathologists to study the added value of the developed algorithms in terms of efficiency and diagnostic accuracy. Al-based biomarkers to select the optimal treatment for individual patients were also studied.
	KIRI GRANGER, Chief Scientific Officer, Monument Therapeutics	OLGA NISSAN, CEO, Protica Bio	MARIA ORR, Head of Precision Medicine Biosamples & Early Oncology, AstraZeneca	ESPEN WALKER, Global Head of Medical Diagnostics, AstraZeneca	AZAM HAMIDINEKOO, Associate Director of Clinical Pharmacology & Safety Sciences, AstraZeneca	JEROEN VAN DER LAAK, Professor of Computational Pathology, Radboud University
			Q&A sessi	on & transition time between conference	e rooms	
	Diamond/Platinum Level Solution Provider Presentation	Diamond/Platinum Level Solution Provider Presentation	Multiplex Luminex Assay Development For Malaria Vaccine Immunology	Diamond/Platinum Level Solution Provider Presentation	Solution Provider Presentation	Diamond/Platinum Level Solution Provider Presentation
						Attendees Are Welcome To Attend Co-Located Sessions
9:35	Attendees Are Welcome To Attend Co-Located Sessions	Attendees Are Welcome To Attend Co-Located Sessions	Luminex	Attendees Are Welcome To Attend Co-Located Sessions	Mavinci Navinci	
			SAMUEL PROVSTGAARD- MORYS, PHD Student/ Research Assistant, Luminex		CRESCENS TIU, Clinical Scientist, Navinci	
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			Day Tu	w o October 01, 202	?5	
		BIOMARKERS & PRE	SPATIAL BIOLOGY FOR PRECISION MEDICINE	DIGITAL PATHOLOGY & AI		
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	Gold Level Solution Provider Presentation	Solution Provider Presentation	Challenges And Solutions In Receptor Occupancy Assay Development - Flow Cytometry	Planning For Success When Developing & Validating Assays For Global Clinical Trials	Mapping Biology In High Definition: Spatial Multiomics For Biomarker Discovery & Translational Breakthrough	Navigating Pathology In Translational Research & Clinical Trials
		♦ Olink	PRECISION for medicine	ALMAC	BRUKER	DIAGNE LIA
:20	Attendees Are Welcome To Attend Co-Located Sessions		Receptor occupancy (RO) assays are essential tools to assess drug-target engagement but are influenced by biological and technical variables. Changes in receptor expression, cell population dynamics, drug effects, and sample handling, among others can all skew RO calculations. Careful optimisation and validation are essential to ensure reliable results and avoid delays in clinical development.	 Key considerations for the development and analytical validation of assays for clinical stratification Implementation of clinical trial assays across global jurisdictions (EU, US and China) Quality control, surveillance of assay performance and clinical trial monitoring 	Join our presentation as we explore the future of multiomics research with Bruker Spatial Biology. Discover our fit for purpose solutions, integrating spatial transcriptomics, proteomics, genomics and bulk multiomics to unlock new biological insights.	Diagnexia Analytix acts as a lighthouse in translational research and clinical trials, guiding sponsors through storms of unclear endpoints, CRO variability, and limited pathology expertise. By embedding expert pathology leadership into workflows, we bring clarity, consistency, and regulator-ready data. This session highlights a case study showing how expert pathology support illuminates the path to trial success.
		Senior Representative, Olink	CIRO NOVAES, Principal Scientist, Cell Biology, Precision For Medicine	CHERYL MCFARLANE, Associate Director, Assay Development & Validation, Almac Diagnostic Services	JOANA CAMPOS, Senior Study Manager, ProPath UK RICHARD BUUS, Senior Field Applications Scientist, Bruker Spatial Biology	GEORGIA CRESWELL, Pathology Solutions Specialist, Diagnexia Analytix
			Q&A session	& transition time between conference room	ıs	
	Improving Early Detection Of Dementia By Multimodal Biomarker Integration	Blood And Spatial Tumour Proteomics Inform The Future Precision Prevention Of Prostate Cancer	End-To-End Automation: Moving Flow Cytometry Into The Digital Age	Methylation Based Biomarkers For Cancer Patient Follow-Up	Beyond IHC - The Added Value Of Morphometrics In Biomarker Analysis & Precision Medicine	Emerging Roles For Large Language Models And Agentic AI In Cellular Pathology
:45	Dementia is one of the biggest global health challenges. Emerging blood and digital biomarkers are opening novel, potentially transformative avenues for early detection of disease risk as well as early diagnosis. With Alzheimer's disease as the trailblazer, this talk will also discuss challenges and opportunities for multimodal data integration in promoting more accurate diagnosis across different dementia subtypes.	In EPIC's study (620 matched pairs), 5,420 plasma proteins were measured; meta-analysis with two other prospective studies (totalling 3,198 prostate cancer cases) identified significant associations: ACP3, CNTNAP2, GP2, TSPAN1, KLK15 (positive) and FLT3LG (inverse). Further spatial tumour profiling shows differing gene and protein expression by clonal lineage. These findings reveal novel mechanisms with potential for early detection.	We developed six harmonised spectral flow cytometry panels with shared backbones, standardised markers, and gating strategies to streamline reporting and enable integration with CDISC's Cell Phenotyping domain. Through modular automation spanning sample processing to analysis, we reduce manual steps and enhance efficiency, consistency, and scalability in our regulated GCP lab.	Liquid biopsy approaches offers great advantages for cancer patient follow-up. Among the several cancer specific components that can be tracked in liquid biopsy, circulating tumor DNA (ctDNA) is especially suited to fine tracking of cancer evolution. This presentation will focus on the combined use of cancer specific methylation markers and digital PCR for ctDNA quantitative detection in several cancers.	How IHC is being used clinically as a biomarker in precision medicine? How spatial transcriptomics and multiplex IHC can provide context to biomarker results? Added value of digital pathology and artificial intelligence models to reflect underlying biology in refined morphometric analysis	Examines the emerging use of large language models and Agentic AI systems in cellular pathology. The ability of these systems to present data from multiple different sources in a succinct meaningful way is relevant to cellular pathology in both research and diagnostic work-up of clinical cases. Other applications are emerging in quality management systems where important efficiencies can be realised, particularly in laboratories grappling with short staffing.

GIOVANNA LALLI, Director of Strategy & Operations, **Life Arc**

KARL SMITH-BYRNE, Senior Molecular Epidemiologist, Cancer Epidemiology Unit (CEU), University of Oxford

IRENE DEL MOLINO DEL

GSK

BARRIO, Principal Scientist,

VALERIE TALY, CNRS Research Director & Group Leader, University of Paris **Descartes**

University Belfast

STEPHANIE CRAIG, Lecturer in Precision Medicine, Queen's

DAVID SNEAD, Professor & Consultant Pathologist, UHCW NHS Trust Coventry and Director of PathLAKE



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10	From Biospecimen To Breakthrough: Powering Precision Biomarker Discovery With Disease- State Insights	Solution Provider Presentation	Dynamic Monitoring And Biomarker Discovery: ctDNA-Driven Insights Across The Cancer Continuum	Beyond Symptoms: Improving The Diagnosis Of Bipolar Disorder With Dried Blood Spot Metabolomics	Solution Provider Presentation	Optimising Pharma R&D Through Digital Pathology, AI And CRO Integration
	BLO IVE	ноговіс:	Burning Rock Dx	biocrates The future of research and health	vizgen	OracleBio
	This presentation highlights how purpose-built, regulatory-grade biospecimens are essential for reliable biomarker discovery, validation, and translation. Discover how these high-quality, disease-state samples power precision medicine programs, from initial research insights to achieving significant clinical impact and regulatory approval across every phase.		Discover how Burning Rock supports clinical and exploratory studies with ctDNA-based genomic and methylation assays, offering biomarker insights and dynamic monitoring strategies tailored to global and regional oncology development programmes.	Dried blood spot metabolomic profiling combined with digital assessment can help differentiate between bipolar disorder and major depression, enabling earlier, more objective and personalised diagnosis.		This presentation explores practical strategies in quantitative digital pathology to support Pharma R&D, focusing on workflow efficiency, Al-driven data quality, and effective CRO engagement to drive faster, more reliable decision-making across the drug development pipeline.
	BRAD EVANS, Senior Scientific Advisor for Disease State, Scientific Affairs, BiolVT	Senior Representative, Hologic Diagenode	MICHAEL CHEN, Senior Translational Medicine Scientist, Burning Rock DX	JAKUB TOMASIK, Assistant Research Professor, University of Cambridge	Senior Representative, Vizgen	LORCAN SHERRY, Chief Scientific Officer, OracleBio
			Q&A	session & transition time between confer	ence rooms	
	Silver Level Solution Provider Presentation	Solution Provider Presentation	Capture The Landscape Of Disease Using Femtogram Level Sensitivity	Solution Provider Presentation	From Sample To Report: Automated Spatial Biology Workflow With PhenoScout Al	How I Learned to Stop Worrying and Love IHC Biomarkers
:35	Attendees Are Welcome To Attend Co-Located Sessions	Attendees Are Welcome To Attend Co-Located Sessions	Upgrades in immunoassay technology are advancing biomarker research. The new FemtoQuest™ system yields femtogram/mL sensitivity in many sample types, in both dual and single plex formats. This presentation will discuss high sensitivity detection of IL-17A and IL-17F in autoimmune disease.		We present an end-to-end spatial biology workflow integrating ZEISS Axioscan 7 spatial biology, SlideStream, and Mindpeak's PhenoScout AI. Foundation AI models enable robust, reproducible analysis across diverse mIF/IHC stainings without any prior image analysis expertise, advancing scalable and robust spatial biology.	PATHOMATION From Tamoxifen to AI, this talk shows how IHC biomarkers shape targeted therapies. We'll cover HER2, PD-L1, Claudin18, training's impact on adoption, lessons from failures, and how AI can extend pathologists' expertise in precision medicine.
			DANIEL GARCIA-WEST, Technology Manager, Proteins Pathways and Analysis, Merck Life Science KGaA	Senior Representative, Hurdle	CHRISTOPER MILLS, Senior Scientist, Concept Life Sciences	RUDY HOVELINCK, CEO, Pathomation

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	Conference Room 1 - Panel Dis Medicine:		5 - Panel Discussion - Exploring ogies In Immunology & Oncolog		Conference R Pharma – Ove	oom 6 - Panel Discussion - Digital Pathology in ercoming Integration Strategies & Future Proofing					
3:00	Moderator: LAURENT AUDOLY, CEO, Co-Four Panellists: MANI MUDALIAR, Director of Qual JIMMY BELL, Professor & Director University of Westminster DAVID SNEAD, Professor & Const Coventry and Director of Pathl BELINDA NEDJAI, Associate Profe London ADAMA IBRAHIM, Vice President Early Development (R&D), Novo	antitative Biomarkers, Recursion or, Research Centre for Optimal I ultant Pathologist, UHCW NHS TOLAKE essor, Queen Mary University o f Digital Transformation, Resea Nordisk	Teaching Hospitals BERND WOLLSCHEIL Professor & Group H	TTHEW HUMPHRIES, Director of Research Operations, Leeds aching Hospitals NHS Trust RND WOLLSCHEID, Head of Institute for Translational Medicine, affessor & Group Head, ETH Zurich MUHAMMAD ASLAM, Consultant Pathologist, for Digital Pathology & AI, Wales, Betsi Cadwa Board Senior Representative, Diagnexia Analytix			ASLAM, Consultant Pathologist, National Clinical Lead nology & AI, Wales, Betsi Cadwaladr University Health entative, Diagnexia Analytix IIVYAN THAYAPARAN, Director, External Innovation &				
3:30			LUNCI	H BREAK: Poster Displays	& 1-2-1 Meetings x4						
	Track Chair: Cristina Legido Quigley,Principal Investigator in Systems Medicine, Kings College London	Track Chair: James Schofield, Founder & Chief Executive Officer, TopMD Precision Medicine	<i>Track Chair:</i> Paul Rhyne, Senior Director, Scientific Solutions, Luminex	Director, Scientific Group Leader, University of		Hamidinekoo, r, AstraZeneca	Track Chair: TBA				
	Solution Provider Presentation	Productising Your Biomarker	Key Considerations For High Quality Data Generation In Global Clinical Trials	Realising The Full Potential Of Dx Concordance Studies: From Pragmatic Study Design To Accelerating Impactful Outcome	From Tissue To R Ensuring Quality Image Analysis	For Robust	Unlocking Prognostic Biomarkers - Integrating Imaging And Omics With Foundation Models SONRAI				
	ALAMAR BIOSCIENCES	MEDTECH TO MARKET	0000	Diaceutics Better Testing, Better Treatment							
14:50		Providing an overview of the design, verification and validation stages for launching new biomarker technologies. Covering the regulatory requirements, key bench and clinical validation studies needed for a technical file submission.	The primary goal of clinical trials is to protect participant safety, with secondary goals including generating accurate, precise, and complete analytical datasets for global studies. High-quality databases enable efficient evaluation of a compound's safety and efficacy. This session highlights key factors influencing data quality in clinical trial execution.	Traditional concordance studies lack real-world relevance. At Diaceutics, we rapidly mobilise scientific studies across independent labs in our network with pragmatic designs that reflect every day clinical practice. Our method delivers robust science along with meaningful operational insights to enhance launch strategies and overall brand impact.			Pathology foundation models are redefining biomarker discovery. Learn how WSI embeddings and omics integration can unlock prognostic signals, with a practical pancreatic cancer case study built on public datasets.				
	MAURA MALPETTI, Race Against Dementia ARUK Fellow Senior Research Associate, UK DRI at Cambridge	MATT PEARCE, Director, Medtech To Market	ANDREW ROCHE, Senior Director of Scientific Affairs, ICON Laboratory Solutions	KEN RUPPEL, VP Scientific & Medical Services, Diaceutics DR CHRISTINE QUINN, Senior Director, Diaceutics	TOM ASHMORE , E Development Mar Histologix		MATT LEE, Director of Al and Medical Imaging, Sonrai Analytics				



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			Da	y Two October O	1, 2025	
		BIOMARKERS & PR	ECISION MEDICINE		SPATIAL BIOLOGY FOR PRECISION MEDICINE	DIGITAL PATHOLOGY & AI
	CONF ROOM 1: WESTMINSTER - BIOMARKER IDENTIFICATION & VALIDATION: NEUROSCIENCE & CO- MORBIDITY	CONF ROOM 2: ST JAMES - NEW & EMERGING BIOMARKER TECHNOLOGIES FOR ANALYSIS	CONF ROOM 3: MOORE - BIOMARKERS FOR CLINICAL TRIALS	CONF ROOM 4: ABBEY - BIOMARKERS FOR DIAGNOSTICS & PRECISION MEDICINE	CONF ROOM 5: GIELGUD & BURTON - AI IMAGE ANALYSIS & DATA ANALYTICS IN SPATIAL BIOLOGY	CONF ROOM 6: OLIVIER - COMPUTATIONAL PATHOLOGY & UTILISING DIGITAL PATHOLOGY IN PHARMA RESEARCH
15:15	Panel Discussion: Biomarker Strategies In Neurodegenerative Therapeutics	Panel Discussion: Application and Integration Of Proteomic Biomarkers In Precision Medicine	Panel Discussion: Translating Biomarkers From Bench To Bedside	Panel Discussion: Navigating Trends In Biomarker & Diagnostic Partnerships	IO-FAST: Initiative For FAIR Spatial Data	Scaling Computational Pathology for the Next Generation Of Cancer Therapies: Challenges and Opportunities
	 Enabling technologies and Imaging approaches to improve target engagement and early diagnostic biomarkers How best to incorporate biomarkers into drug development and clinical trials for neurodegenerative diseases Recent advancements in developing biomarkers for clinical utility 	Biomarker discovery, Diagnostics and prognostics, Patient stratification and precision medicine	Patient identification Endpoints		IO-FAST (Initiative for FAIR Spatial Data) addresses the urgent need for standardised, interoperable spatial omics frameworks. By uniting academia, industry, and data repositories, we support the research community in harmonising data formats, streamlining metadata annotation, and	This presentation explores the crucial role of the pharmaceutical industry in expanding digital pathology accessibility across global immunohistochemistry laboratories. By leveraging computational approaches, we can revolutionise biomarker detection in cancer patients. The session will highlight innovative strategies and collaborative efforts needed to overcome current challenges and drive advancements in personalised medicine, ultimately enhancing patient outcomes on a global scale.
	Moderator: KIRI GRANGER, Chief Scientific Officer, Monument Therapeutics Moderator: KARL SMITH-BYRNE, Senior Molecular Epidemiologist, Cancer Epidemiology Unit (CEU), University of Oxford Panellists: MANUELA CERINA, Scientific Director of Neurodegeneration LifeArc Moderator: EMANUELA OLDONI, Scientific Lead of Personalised Medicine, EATRIS, European Infrastructure for Translational Medicine	Moderator: SUSANNE MUNKSTEAD, CSO, Diaceutics Panellists: JENS KIECKBUSCH, Director of External Innovation, Precision Medicine, Research & Development, GSK	fostering community collaboration- empowering researchers to share and integrate complex spatial datasets.			
	NATASA GIALLOUROU, Field Metabolomics Scientist, Metabolon KRISTINA HOLMBERG, Head	Protica Bio JOSHUA ATKINS, Senior Genomic Epidemiologist, Cancer Epidemiology Unit (CEU), University of Oxford ALEXANDRA SEVKO, VP of Translational Science ATASA GIALLOUROU, Field JOSHUA ATKINS, Senior Genomic Epidemiologist, Cancer Epidemiology Unit (CEU), University of Oxford ALEXANDRA SEVKO, VP of Translational Science Panellists: MIKE MESSENGER, Head or Regulatory Strategy, BIVDA RISTINA HOLMBERG, Head ACHIEVAL TO ATKINS, Senior Genomic Epidemiologist, Cancer Epidemiology Unit (CEU), University of Oxford ALEXANDRA SEVKO, VP of Translational Science ADEVICE BIO MIKE MESSENGER, Head or Regulatory Strategy, BIVDA RISTINA HOLMBERG, Head ACHIEVAL TO ATKINS, Senior Genomic Epidemiologist, Cancer Epidemiology Unit (CEU), University of Development Oncology (EDO), Pharma Research & Development (pRED)	MIKE MESSENGER, Head of Regulatory Strategy, BIVDA MICHAEL CANNARILE, Head of Biomarkers, Early Development Oncology (EDO), Pharma Research & Development (pRED) Roche Innovation Center Munich	CHRIS BRAY, Head of Global Regulatory Affairs, Precision Medicine & Companion Diagnostics, Merck KGaA	AMONIDA ZADISSA, Associate Director of Informatics, UK Dementia Research Institute	HUW BANNISTER, Senior Director of Global Diagnostics, Digital and Computational Pathology, AstraZeneca
15 10	Lundbeck			NITIN JAIN, Director of Franchise Project	Q&A session & trans.	ition time between conference rooms
15:40				Management, AstraZeneca	Data-Driven Analysis And Spatiotemporal Control Of Bacterial Colonies	Revolutionising Image Analysis In Drug Discovery
			Translational Sciences, Accession Therapeutics LAURA HUBBARD, Operations Director,		The morphology of bacterial microcolonies influences key properties such as resource uptake and resistance to environmental stressors, including antibiotics. We present machine learning–based tools for analysing microscopy data and for spatiotemporal control of gene expression to precisely manipulate microcolony shape. These capabilities open new avenues for research and applications in biofilms, antibiotic resistance, diagnostics, and engineered biomaterials.	At AstraZeneca, we are embedding AI, image analysis and frameworks for integrating emerging multi-omic data into our digital toxicological pathology workflows. This is providing novel insights and a dynamic, integrated understanding of tissues, disease, and the impact of our therapies upon drug therapeutic indices. This enables a seamless and scalable digital revolution to support our accelerated drug discovery portfolio.
	INTERACTIVE	INTERACTIVE	INTERACTIVE	INTERACTIVE	JEAN-BAPTISTE LUGAGNE, Assosiate Professor, University of Oxford	ARTHUR LEWIS, Director of Image Analysis & Platform, Pathology, AstraZeneca



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_	CONF ROOM 1: WESTMINSTER - BIOMARKER IDENTIFICATION & VALIDATION: NEUROSCIENCE & CO- MORBIDITY	CONF ROOM 2: ST JAMES - NEW & EMERGING BIOMARKER TECHNOLOGIES FOR ANALYSIS	CONF ROOM 3: MOORE - BIOMARKERS FOR CLINICAL TRIALS	CONF ROOM 4: ABBEY - BIOMARKERS FOR DIAGNOSTICS & PRECISION MEDICINE	CONF ROOM 5: GIELGUD & BURTON - AI IMAGE ANALYSIS & DATA ANALYTICS IN SPATIAL BIOLOGY	CONF ROOM 6: OLIVIER - COMPUTATIONAL PATHOLOGY & UTILISING DIGITAL PATHOLOGY IN PHARMA RESEARCH	_
	Antibodies From Resilient Individuals: A Novel Approach For Neurodegeneration Drug Discovery	Biomarkers For Treatment Response In Prostate Cancer	Accelerating Clinical Trials In Neurodegenerative Disease Using Wearable Sensors And Al-Derived Kinematic Biomarkers	Epigenetic Detection Of HPV-Related Cancers: Validation Of The S5 Classifier In African And Global Low-Resource Settings	Spatially Resolved Multiomics Of Human Cardiac Niches	Al Quantification In H&E Stained Tissue	-
16:05	Alchemab use a patient first, target agnostic approach to drug discovery that often results in novel targets with unexplored biology and with no high quality commercial reagents available to study them. This talk will provide	 Mechanism-centric biomarker discovery Patient risk of treatment failure assessment Novel therapeutic vulnerabilities MYC+NME2 as markers and 	Our work demonstrates how data-derived wearable biomarkers can track personal disease trajectories and indicates the potential of such biomarkers for substantially reducing the duration or size of clinical trials testing	The S5 methylation classifier is a liquid biopsy-based tool that detects early cancer-related epigenetic changes across multiple HPV-associated malignancies. Applicable to self-collected or clinician-obtained	We combine single-cell and spatial transcriptomics to discover human cardiac niches. We profile cardiac conduction system cells, revealing that the pacemaker cells exhibit distinct characteristics in the adult and developing heart. In the		_
	examples of overcoming these challenges by developing novel assays and tools.	therapeutic targets for advanced prostate cancer	disease-modifying therapies and for enabling behavioural transcriptomics.	samples, it has demonstrated high accuracy in diverse global populations. Its non-invasive, scalable design supports early detection in low-resource settings, highlighting its potential for broad integration into biomarker-driven cancer screening programmes.	ventricle, we illustrate the processes of cardiomyocyte compaction and maturation. Overall, we offer a comprehensive map of the human heart, both in adults and during development.		_
			DICHARD FECTENCTEIN	curred selecting programmes.			_
	HELEN GRAVES, Principal Scientist, Alchemab Therapeutics	ANTONINA MITROFANOVA, Associate Dean for Research & Associate Professor, Rutgers University	RICHARD FESTENSTEIN, Clinical Professor of Molecular Medicine, Department of Brain Sciences, Imperial College London	BELINDA NEDJAI, Associate Professor, Queen Mary University of London	KAZUMASA KANEMARU, Postdoctoral Researcher, Cambridge Stem Cell Institute, University of Cambridge	FRANCESCA TRAPANI, Scientific Director & Molecular Pathology Laboratory Head, Boehringer Ingelheim	F
	Q&A session & transition time between conference rooms						
16:30	Mapping Neurodegeneration: Multi-Omics Insights Into Disease and Preclinical Models	A Gene Expression Based Biomarker For Predicting Response To Treatment With Stenoparib	Epigenetic Biomarkers For Inflammatory Disease	Biomarkers For Early Detection Of Cancer And/ Or Enabling Development & Delivery Of Cancer Preventative Interventions		How Al Is Shaping Prostate Cancer Diagnosis	3
	Use multi-omics platforms to better understand neurodegenerative diseases and assess how well preclinical models reflect human disease. Combined targeted and	By analysing gene expression in cancer cell lines that are sensitive and resistant to the dual PARP and Tankyrase inhibitor Stenoparib, Allarity has developed a response predictor. Validation in a clinical		Novel interventions to reduce risk and intercept cancer at its earliest stages could benefit both patients and health systems. To make precision prevention a reality, there is a critical need to	Attendees Are Welcome To Attend Co-Located Sessions	 IPATIMUP experience with Al. Clinical advantages of Al-guided prostate biopsies. Transformative potential of Al in shaping the future of prostate cancer diagnosis. 	,
	untargeted omics to analyse plasma, CSF, and brain tissue, revealing differentially expressed molecules. Multi-omics shows promise for biomarker discovery, and clinical relevance.	trial is ongoing.		develop cancer risk biomarkers and surrogate endpoints. This talk provides an overview of CRUK's cancer prevention strategy, and our perspectives on the challenges and opportunities for cancer prevention biomarkers.			
	GAYLE MARSHALL, Head of Biomarkers, Medicines Discovery Catapult	THOMAS JENSEN, Chief Executive Officer, Allarity	WOUTER DE JONGE, Professor of Neurogastroenterology, Amsterdam UMC	JOANNA JANUS Research Programme Manager, (Early Detection & Prevention), Cancer Research UK		JOÃO PINTO, Medical Specialist, Institute Molecular Pathology and Immunology, University of Porto (Ipatimup)	
16:55	END OF CONFERENCE						

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VENUE INFORMATION

Experience London

Biomarkers & Precision Medicine 2025 will be held at the Queen Elizabeth II Centre.

Since 1986, The QEII Centre has been London's premier venue for domestic and international events. Their venue offers world-class facilities for high-profile conferences, conventions, exhibitions and corporate events with a capacity of up to 2,500.

Located opposite Westminster Abbey, the QEII Centre is less than a five minute walk from the Thames, and a two-minute walk from a royal park. The Centre is served by exceptional transport links providing easy access to everything the city has to offer, from world-class restaurants and nightlife to shopping and culture. Within an hour's transfer from five international airports, you will have an effortless journey to the Centre.

DRIVING

Blue badge holders are permitted to park on the QEII Centre forecourt free of charge. Parking in this location is subject to availability so please call ahead on +44 20 7798 4000 to enquire.

Disabled passengers arriving by taxis or alternative vehicles may disembark on the forecourt.

PUBLIC TRANSPORT

The nearest step-free tube station is Westminster, which is served by the Jubilee, District and Circle lines.

Venue Website





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