





Proventa International's 6th Annual

ONCOLOGY

STRATEGY MEETING EAST COAST USA 2022

Exploring Promising Areas in Oncology Research Aimed to Expand Our Knowledge Base, Address Unmet Medical Needs and Improve Overall Clinical Success







Immunooncology



Leveraging Biomarkers



Radiopharmaceuticals



Treating AML



CAR-T



Revamping ADCs



NK cells



CID Trials G



Precision Gene-Editing



Vectors

Featuring Industry Leaders and Decision Makers



Laurent
Audoly
Founder, CEO,
Chairman of
the Board
Parthenon
Therapeutics



Geoffrey Hodge Chief Executive Officer SOTIO Biotech



Gregory Fiore Chief Executive Officer Exacis Biotherapeutics



George Tetz Chief Executive Officer CLS Therapeutics



Johanna Kaufmann Executive Vice President Oncology Codagenix Inc



Marco Gottardis
Vice President of
Oncology Innovation
Janssen
Pharmaceutical
Companies of
Johnson & Johnson



Vijay Kasturi Vice President, Medical Affairs AVEO Oncology

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STRATEGY MEETINGS!





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Proventa International's **Strategy Meetings are a** completely unique experience.



Our Vision



To be a platform for creating life-saving therapies and to facilitate the creation of a completely patient centric pharmaceutical industry.

Our Mission



By encouraging key leaders and their companies to put the patient at the very heart beat of every innovation. Sharing valuable insights and strategies to assist in the discovery, development and commercialisation of life saving therapies.

As one of the most significant areas of research in healthcare, the innovations stemming from cancer research are certainly commendable. Technologies such as the mRNA vaccine, brought to light by COVID were originally aimed at curing cancer, along with new approaches in discovering novel biomarkers, multi-omics approach to study cancer, epigenetic factors and reprogramming immune cells for enhanced efficacy, all of which are showing us a glimpse into the future. Constant advancements depict the need to keep pushing the limits of research to get to a stage where cancer is not a life-altering event, but at least a manageable, chronic condition. This strategy meeting intends to provide a stimulating environment where industry experts can discuss new discoveries, spark new collaborations and create synergy to accelerate progress in patients with cancer

Why this strategy meeting is a FOR YOU AND YOUR TEAM!

- How to Overcoming challenges for early phase I/O clinical trials and maximize opportunities for biomarker discovery
- What are the next generation ADC's?
- What is the need for a multi omics approach in cancer research?
 - How can we leverage machine learning in clinical research to transform the future of evidence generation?
- Understanding how Non-Viral Vectors Compare with Viral Vectors for Application in Gene Therapy

Our Unique Meeting Format



ROUNDTABLE DISCUSSIONS

These interactive and informal discussion groups are the hallmark of the meeting. The brightest minds in the industry are brought together in 60-minute sessions that enable participants from all over the world to share ideas, challenges and lessons learned.



PERSONALISED AGENDA

Each delegate receives a personalised agenda with the roundtable discussions that you choose. You only attend sessions and meetings that fit your challenges and interests. ensuring your time spent is focused and well-utilised.



INNOVATIVE SOLUTIONS

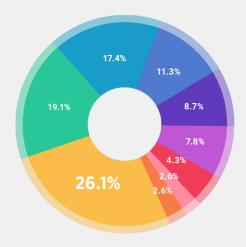
The most effective and time efficient way to assess potential partners at a strategic level. Identify key solution providers that can take your business to the next level and we will help arrange private meetings so you can connect.



STRATEGIC NETWORKING

Strategic networking opportunities form a key benefit of the meeting. Our proven format for building and strengthening alliances to make lasting connections that benefit you.

SENIORITY OF ATTENDEES



- **Director Level**
- **Scientist**
- **Department Head**
- C-Level
- Other
- President / VP
- **Team Lead**
- Manager
- Academia

Facilitator Faculty







Dr. Claudia Hesselmann, PhD Co-Founder &

CEO, Germany

Arensia



Eidus
Director Corporate
Develepment
North America, USA
Arensia



Déborah Heintze Co-founder and Chief Marketing Officer Lunaphore Technologies



Dr. Antonio Sorrentino Head of Translational Strategy Lunaphore Technologies



Andrew Jackson
U.S. Commercial Director
Sengenics



Katy McGirr
Director of Global
Marketing
Sengenics



Laurent Audoly Founder, CEO, Chairman of the Board Parthenon Therapeutics



Vivian Choi Head of Global Gene Therapy Research Takeda



Tirtha Chakraborty Chief Scientific Officer Vor Biopharma



Sashka Dimitrievska Global Therapeutic Area Head Oncology Clinical Insights Astrazeneca



Gregory Fiore
Chief Executive
Officer
Exacis
Biotherapeutics



Gottardis
Vice President of
Oncology Innovation
Janssen
Pharmaceutical
Companies of Johnson
& Johnson



Vijay Kasturi Vice President, Medical Affairs AVEO Oncology



Karuppiah Kannan Executive Director -Global Program Leader Takeda



Johanna Kaufmann Executive Vice President Oncology Codagenix Inc



Geoffrey Hodge Chief Executive Officer SOTIO Biotech



Jeffrey Humphrey Member of Board of Directors Cyteir Therapeutics



Chet Metcalf
Vice President,
Head of Chemistry
Curie
Therapeutics



Oscar Segurado Chief Medical Officer ASC Therapeutics



David Sherris
Biotechnology /
Pharmaceutical
Executive
Penrose
Therapeutics



Singh
Director,
Translational
Medicine
Takeda



George Tetz
Chief Executive
Officer
CLS Therapeutics

Intimate format events where senior leadership discuss the biggest challenges facing the industry.



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ARENSIA EXPLORATORY MEDICINE is a German operator of proprietary research clinics in Eastern Europe, dedicated to performing complex Phase Ib, IIa and PROOF-CONCEPT clinical trials with novel molecules, involving patients across numerous disease areas. Given the outstanding patient recruitment rates achieved by ARENSIA clinics, we are able to substantially reduce the overall number of sites, countries and recruitment periods for any early phase patient trial.





Lunaphore Technologies S.A. is a Swiss company born in 2014 with the vision of enabling spatial biology in every laboratory. Lunaphore has developed a game-changing chip technology which can extract spatial proteomic and genomic data from tumors and transform any simple assay into multiplex spatial biology without complexity. Lunaphore empowers researchers to push the boundaries of research to ultimately develop the next generation personalized therapies. For further information on Lunaphore and its products, please visit www.lunaphore.com.





ProTrials Research, Inc., is a full-service clinical research organization (CRO) and woman-owned business headquartered in Los Gatos, CA, with clinical operations personnel located throughout North America and across the world. ProTrials works with sponsors in the pharmaceutical, biotechnology and medical device industries to deliver high-quality clinical development services in a wide range of therapeutic areas. Founded in 1996, ProTrials has been successfully supporting clinical research for over 26 years.





Base Pair Biotechnologies is based just 10 miles south of the Houston, Texas Medical Center. Our team of 11 scientists has a combined 50+ person-years of experience in developing aptamers. Base Pair owns the sole worldwide rights to patents for multiplexed aptamer discovery. We have the capability to select aptamers to up to 30 protein, peptide, or small molecule targets in true competitive, multiplexed fashion. The resulting aptamers are therefore more specific for their particular targets. Using this technology, we have completed contracts from the National Cancer Institute, the CDC, and many large pharma and biotech firms.





Sengenics is a precision medicine company working to improve patient outcomes through physiologically relevant, data-guided decision making. Our solutions enable the discovery and validation of autoantibody biomarker signatures for patient stratification, therapeutic response prediction and development of companion diagnostics.





Inato is a global marketplace that matches community-based research sites with the right clinical trials. The marketplace allows sponsors to confidently partner with a broader range of sites, making clinical trials more accessible, inclusive and effective. Since launching in June 2020, over 1400 sites have joined the marketplace along with several top-20 sponsors. Learn more at www.inato.com





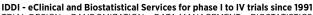
Navinci (formerly Olink AB) develops, produce and market high-performance reagent kits to serve scientists in academia, drug development and pathology research to accurately study proteins and their interplay directly in cells and tissue samples at a level not possible with other methods. Scientists use our in situ proximity ligation assay technology to better understand biological mechanisms and reveal the responses to disease and drug treatment. In particular, the effects of Immune Checkpoint Inhibitors can be studied in the tissue microenvironment giving higher chance of identifying strong drug candidates and to stratify patients that will benefit from treatment.





Haystack Oncology uses the next generation of circulating tumor DNA (ctDNA) detection technology to see through the noise and uncover the lowest levels of ctDNA in blood to report residual, recurrent, or resistant disease earlier than any other minimal residual disease (MRD) test. Designed by cancer genomics pioneers and liquid biopsy experts, Haystack's tumor-informed MRD test is best positioned to deliver the right treatment to the right person at the right time.





TRIAL DESIGN - RANDOMIZATION - DATA MANAGEMENT - BIOSTATISTICS

IDDI is an oncology expert company led by a highly-skilled biostatistical and clinical team of thought-leaders in oncology research. We combine biostatistics, regulatory & medical expertise with integrated eClinical solutions to ensure clinical data readiness for submission. SERVICES:



- Trial design
- Statistical analyses IDMCs & Interim analyses
- IRT integrated with EDC system
- Data management
- 700+ oncology trials, 12 oncology FDA-EMA approvals
- Strategic consulting services
- Biomarker and diagnostics/IVD validation Support in Contacts with Regulatory Authorities
- Drug supply
- Medical coding CDISC Data Standardization

VISIT WEBSITE

How it Works?



Real insights are shared in conversations, not lectures. That's why our unique **roundtable** format guarantees real learning and ensures you only attend discussions that deliver value to you.

Join as many as four roundtable discussions, each with up to 20 other subject experts, networking with top-level peers in an informal, relaxed and sociable environment.

Simply follow the steps below to select your roundtable discussions and we'll create your own personalised agenda for the day:



EXPLORE THE FULL AGENDA

Select which roundtable discussions you would like to join. Our sessions are divided among 6 themed tracks for easy selection - you can choose to join any session you like.



COMPLETE YOUR SCHEDULING FORM

Select your preferred roundtable discussion for each time slot. We will send this form out via email a few weeks before the event - be sure to get your first choice by completing the form quickly.



ENJOY YOUR PERSONALISED EXPERIENCE

Join your selected roundtable sessions on the day. We will give you your personalised agenda containing the time and room assignments of your chosen roundtable discussions so you won't miss it.

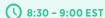
Agenda at a Glance

Oncology Strategy Overview 8th of November, 2022 - Cambridge, Boston - Le Meridien Hotel

	TRACK 1	TRACK 2	TRACK 3	TRACK 4	TRACK 5	TRACK 6
TIME EST	BIOMARKER DISCOVERY	PRECISION MEDICINE/ COMPANION DIAGNOSTICS	ANTIBODY/ADC DISCOVERY & DEVELOPMENT	IMMUNO-ONCOLOGY	CLINICAL RESEARCH & DEVELOPMENT	CELL & GENE THERAPY
ROOM ►	Jerome C. Hunsaker C	Jerome C. Hunsaker	Jerome C. Hunsaker A	Lan Jen Chu	Margaret L.A. Macvicar	Edward Pennell Brooks
08:00 - 08:30			REGISTRATION	AND WELCOME		
08:30 - 09:00	OPENING KEYNOTE PRESENTATION Drugging the Tumor Microenvironment to achieve durable response in patients					
09:00 - 10:00	Overcoming challenges for early phase I/O clinical trials and maximizing opportunities for biomarker discovery	Translating cancer 'omics' to improved outcomes	What is the next generation of ADC's?	Radioactivity - A particularly sensitive "antenna" with excellent tissue penetration	Role of machine learning in clinical research: transforming the future of evidence generation	Cell and Gene Therapy landscape in Hematological malignancies: Next- generation innovations beyond traditional Lentiviral CAR-T
10:00 - 10:10	REFRESHMENT BREAK					
10:10 - 10:30	NETWORKING / 1-1 MEETINGS					
10:30 - 10:50	NETWORKING / 1-1 MEETINGS					
10:50 - 11:10	NETWORKING / 1-1 MEETINGS					
11:10 - 12:10 SOLUTION	Auto antibody biomarkers for patient stratification: An approach to trial enrichment & drug repurposing SENGENICS	Funding early biotech companies in the Biden administration era with an emphasis on oncology		Tissue Biomarkers Innovations in Immuno- Oncology LUNAPHORE LUNAPHORE		
12:10 - 13:10	LUNCH BREAK					
13:10 - 13:40	KEYNOTE PRESENTATION Accelerated performance of complex exploratory studies in oncology: Practical insights from investigational site KEYNOTE PRESENTATION Accelerated performance of complex exploratory studies in oncology: Practical insights from investigational site					
13:40 - 14:40	Innovative approaches towards implementing Predictive Biomarkers into Immuno-Oncology	Precision Gene-Editing Tools in Personalized Cancer Medicine and the challenges they pose		Leveraging the untapped potential of engineered T-cell based therapeutics to treat solid tumors	Incorporating real-time clinical insights in clinical development process for faster submissions and accelerated approvals	How do Non-Viral Vectors Compare with Viral Vectors for Application in Gene Therapy
14:40 - 14:50	REFRESHMENT BREAK					
14:50 - 15:10	NETWORKING / 1-1 MEETINGS					
15:10 - 15:30	NETWORKING / 1-1 MEETINGS					
15:30 - 15:50	NETWORKING / 1-1 MEETINGS					
15:50 - 16:50		The use of NK therapies for treatment of solid tumors – what are the problems and how do we move forward?		Challenges and Innovative Methods in Tumor-Targeted Approaches to Treat Solid Tumor Cancers	The use of Complex Innovative Design (CID) trials in Oncology	Next Generation Allogenic Cell Therapy
16:50 - 17:20	PANEL DISCUSSION Lost in translation: from preclinical to clinical – identifying problems and overcoming obstacles					
17:20 - 18:20	DRINKS & CANAPES RECEPTION					



OPENING KEYNOTE PRESENTATION



Drugging the Tumor Microenvironment to achieve durable response in patients



Laurent Audoly Founder, CEO, Chairman of the Board **Parthenon Therapeutics**

KEYNOTE PRESENTATION

(1) 13:10 - 14:40 EST

Accelerated performance of complex exploratory studies in oncology: Practical insights from investigational site

- Unique model of dedicated research clinics in Eastern Europe
- Strategies for fast patient enrollment and retention
- Operational tips for flawless study conduct
- Takeaways after operating under COVID-19 pandemic



Dr. Claudia Hesselmann, PhD Co-founder & CEO, Germany **ARENSIA**





Tatiana Eidus Director Corporate Development North America, USA, ARENSIA

PANEL DISCUSSION

(1) 16:50 - 17:20 EST

Lost in translation: from preclinical to clinical - identifying problems and overcoming obstacles



[CHAIR] Reshma Singh Director, Translational Medicine Takeda



George Tetz CLS Therapeutics



Johanna Kaufmann **Executive Vice President Oncology** Codagenix Inc



Jeffrey Humphrey Member of Board of Directors **Cyteir Therapeutics**



TRACK 1: **BIOMARKER DISCOVERY**

Genetic, epigenetic, proteomic, glycolic, and imaging biomarkers can be used for cancer prognosis, diagnosis, and epidemiology. While some can be used to predict how aggressively your cancer will grow, and are therefore useful for assessing your prognosis, the most promising use of biomarkers today is to identify which therapies a patient's cancer may or may not respond to.

() 09:00 - 10:00 EST

ROUNDTABLE 1:

Overcoming challenges for early phase I/O clinical trials and maximizing opportunities for biomarker discovery

- What are appropriate patient populations for early phase clinical trials using I/O drugs both as monotherapy and in combination?
- How do we leverage novel trial designs for iterative testing of combinations and dose regimens and what are appropriate endpoints?
- How can we improve the identification of rational combination partners and optimally leverage biomarker programs?
- How much personalization is desirable or necessary in I/O drug development?



Johanna Kaufmann **Executive Vice President Oncology** Codagenix Inc.

(1) 11:10 - 12:10 EST



SOLUTION FOCUS ROUNDTABLE 2: Autoantibody biomarkers for patient stratification: An approach to trial enrichment & drug repurposing

- What clues is the immune system giving us about the patient population?
- How can biomarkers with high predictive value be found in heterogenous populations?
- How can novel technology overcome past limitations?



Andrew Jackson U.S. Commercial Director Sengenics





Katy McGirr Director of Global Marketing Sengenics



(1) 13:40 - 14:40 EST

ROUNDTABLE 3:

Innovative approaches towards implementing Predictive **Biomarkers into Immuno-Oncology**

- Reviewing common issues encountered when translating a biomarker to the clinic: Determining eligibility for clinical study
- Practicalities that need to be considered when planning to use a biomarker to balance arms of a study
- Delineating a roadmap towards effective implementation of predictive patient-segmentation marker into an early clinical trial to facilitate subsequent development of effective CDx





TRACK 2: PRECISION MEDICINE/COMPANION DIAGNOSTICS

Precision medicine has rapidly altered the oncology diagnostic and treatment spectrum by allowing tailored treatment strategies to precisely target the molecular alteration underlying individual patients' disease. With genomic-data sequencing new disease pathways are being discovered, new therapeutic targets are being revealed and ideal treatments populations are being identified which shall be discussed further in this track.

() 09:00 - 10:00 EST

ROUNDTABLE 1: Translating cancer 'omics' to improved outcomes

- If a striking heterogeneity exists even at the very genomic level between histologically similar tumors, can that be leveraged in detecting important molecular alterations & identifying novel targets
- Understanding as target discovery further deepens, how important a multi omics approach which combines proteomics and genomic, is in cancer to considerably improve patient outcomes
- Focus on the unique advantages of a multidimensional approach and discuss the implications of the changing cancer demographic to translational omics research



11:10 - 12:10 EST

ROUNDTABLE 2: Funding early biotech companies in the Biden administration era with an emphasis on oncology

- As the Biden administration leans into the Cancer Moonshot 2.0, would this
 program provide opportunity for the life sciences industry for precision
 oncology compared to Cancer Moonshot 1.0
- Will government controlled drug pricing be problematic for biotech/pharma R&D thereby negatively affecting the death rate from cancer
- What is the level of biotech funding for oncology compared to other diseases indications
- How does cancer therapeutic modalities affect decision to fund R&D



David Sherris
Biotechnology / Pharmaceutical Executive
Penrose Therapeutics



(1) 13:40 - 14:40 EST

ROUNDTABLE 3:

Precision Gene-Editing Tools in Personalized Cancer Medicine and the challenges they pose

- Gene-editing tools CRISPR, Zinc Finger, TALENs and emerging tools hold promise to allow for engineering required to create potent, safe allogeneic cell therapies
- Discussing technical problems and that need to be addressed to incorporate innovative tools that drive next generation of molecular precision medicine
- Discuss strengths and limitations of each of the major approaches



Gregory Fiore CEO Exacis Biotherapeutics

(15:50 - 16:50 EST

ROUNDTABLE 4:

The use of NK therapies for treatment of solid tumors – what are the problems and how do we move forward?

- There seems to be a barrier to NK therapies entering the solid tumor space
- Is the continued focus on hematologic malignancies due to limitations of NK cell therapies or due to lack of imagination?
- When hematologic malignancies make up 10% of cancer, Why is the field ignoring the other 90%?



Reshma Singh
Director, Translational Medicine
Takeda







TRACK 3: ANTIBODY/ANTIBODY DRUG CONJUGATE **DISCOVERY & DEVELOPMENT**

Traditional immunoglobulin antibodies have been further engineered for better efficacy and safety to specifically bind to a target antigen and release the cytotoxic drug into the cancer cell. This represents the newer generation of therapeutic antibody formats- bispecific antibodies and antibody-drug conjugates. This track explores the latest findings on antibody-based therapeutics and how to push the envelope in ADC manufacturing so they can be more effective.

() 09:00 - 10:00 EST

ROUNDTABLE 1: What is the next generation of ADC's?

- New technologies expanding the scope of ADCs
 - Novel Targets is cellular internalization of the target necessary?
 - Novel Toxins what are the best or new targets?
 - Novel Linkers cleavable and non-cleavable
 - Radioactive payloads
- Why understanding cell biology is critical to appropriately select the recommended phase II dose
- Novel approaches using "in vivo" chemistry to release linker-payload





Jeffrey Humphrey Member of Board of Directors **Cyteir Therapeutics**





TRACK 4: IMMUNO-ONCOLOGY

Immuno-oncology is a form of cancer treatment that uses the power of the body's own immune system to prevent, control, and eliminate cancer. Of the many types of immunotherapy, checkpoint inhibitors form one arm of the treatment that works by blocking proteins that stop the immune system from behind their resistance and explore the potential of an upcoming checkpoint

() 09:00 - 10:00 EST

ROUNDTABLE 1: Radioactivity - A particularly sensitive "antenna" with excellent tissue penetration

- Can Radio pharmaceuticals deliver radiation to cancer cells, independent of their tumor type and location in the body?
- The advantages of nuclear imaging via SPECT and PET over other imaging modalities regarding sensitivity
- A major obstacle to development and effective application of novel therapeutic regimens is optimized patient selection and response assessment
 - Noninvasive imaging using novel immunoconjugate radiopharmaceuticals can assess for expression of cell surface immune markers, such as programmed cell death protein ligand-1 (PD-L1), akin to a virtual biopsy
- Targeted radiopharmaceuticals provide very effective systemic treatment options due to the specific delivery of lethal radioactive doses to cancer tissues, while sparing healthy tissues
- How to leverage Radio-theranostics for diagnosis, staging, patients' stratification, and treatment, thus reinforcing precision medicine in oncology



Chet Metcalf Vice President, Head of Chemistry **Curie Therapeutics**

(1) 11:10 - 12:10 EST







- Immunotherapy Predictive Spatial Biomarkers: opportunities and challenges in the path to the clinic
- The Digital Pathology ecosystem and the role of pathologists, digital tools and techniques

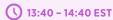


Déborah Heintze Co-founder and Chief Marketing Officer Lunaphore Technologies

Lunaphore Technologies







ROUNDTABLE 3:

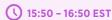
Leveraging the untapped potential of engineered T-cell based therapeutics to treat solid tumors

- Improving manufacturing to produce more potent T cells
- Next generation CAR-Ts—novel transgenes to improve T cell performance
- Combination therapies with CAR-T



Geoffrey Hodge CEO SOTIO Biotech





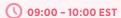
ROUNDTABLE 4:

Challenges and Innovative Methods in Tumor-Targeted Approaches to Treat Solid Tumor Cancers

- How Immunotherapy falls ineffective when the tumor microenvironment (TME) is dominated by immunosuppressive cell types that impair immune response
- Discuss an approach that depletes such cells and increases the number of cancer-fighting immune cells
- Address key limitations to enable more effective treatment of solid tumor



TRACK 5: **CLINICAL RESEARCH AND DEVELOPMENT**



ROUNDTABLE 1:

Role of machine learning in clinical research: transforming the future of evidence generation

- ML has a role to play in maximizing the success and efficiency of trials during the planning phase through application of simulation techniques to large amounts of data
 - Study simulation may optimize the choice of treatment regimens for testing in non-small cell lung cancer
- An appealing application of ML is to automate data collection into case report forms, decreasing the time, expense, and potential for error associated with human data extraction
 - Though this use requires overcoming variable data structures and provenances, it has shown early promise in cancer



Karuppiah Kannan Executive Director - Global Program Leader



(13:40 - 14:40 EST

ROUNDTABLE 3:

Incorporating real-time clinical insights in clinical development process for faster submissions and accelerated approvals

- Realizing the value of DS&AI driving oncology trial data for existing products label extension
- Can we apply clinical data insights from early to late stage trial designs to enable smarter trial designs
- Technology and analytics enabling biomarkers data to be gathered in real time how do we apply this data to drive clinical trial endpoints



Sashka Dimitrievska Global Therapeutic Area Head Oncology Clinical Insights Astrazeneca

(15:50 - 16:50 EST

ROUNDTABLE 4:

The use of Complex Innovative Design (CID) trials in Oncology

- The traditional cancer drug development pathway is increasingly being superseded by Complex Innovative Design (CID) trials that address multiple
- Advantages of CID in assessing safety and toxicity of novel anticancer medicines and efficacy in biomarker-selected patients, specific cancer cohorts or in combination with other agents



Vijay Kasturi Vice President, Medical Affairs AVEO Oncology





TRACK 6: **CELL & GENE THERAPY**

() 09:00 - 10:00 EST

ROUNDTABLE 1:

Cell and Gene Therapy landscape in Hematological malignancies: Next-generation innovations beyond traditional Lentiviral CAR-T

- Cell Therapy Innovations for AML treatment: Regulatable and universal CARs, including T, NK and Macrophage
- Manufacturing challenges for viral and non-viral components
- Geno-Toxicity in cell and gene therapy and the expectation of the regulators



(13:40 - 14:40 EST

ROUNDTABLE 3:

How do Non-Viral Vectors Compare with Viral Vectors for Application in Gene Therapy

- Limitations posed by current vectors such as immunogenicity, manufacturability. and payload
- AAV & beyond: progression in novel viral vectors
- Are there any particular disease areas/cell types that would benefit from a specific delivery modality?
- What are the current limitations of non-viral vectors, and how can they be overcome?



Vivian Choi Head of Global Gene Therapy Research

15:50 - 16:50 EST

ROUNDTABLE 4: Next Generation Allogeneic Cell Therapy

- What are the new cell types being considered by allogeneic cell therapy? Advantages & Disadvantages
- What strategies are being used to achieve hypo immunity?
- What hurdles are there to develop allogeneic cell therapies from IPSCs



Marco Gottardis

Vice President of Oncology Innovation

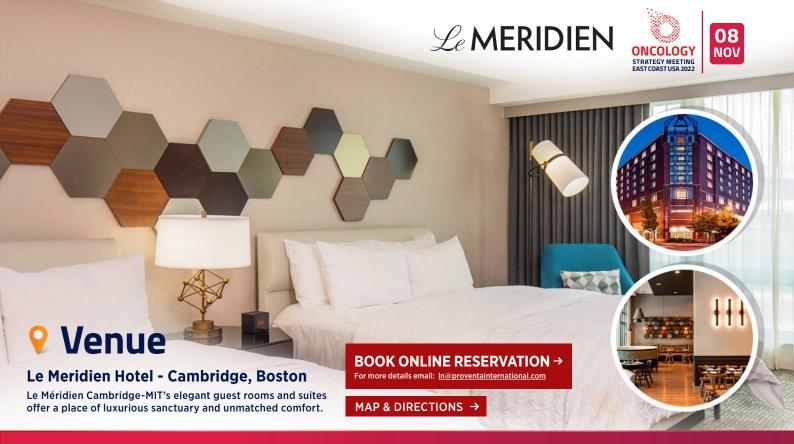
Janssen Pharmaceutical Companies of Johnson & Johnson

DRINKS & CANAPES RECEPTION









OUR FACE-TO-FACE MEETING IN OCTOBER & NOVEMBER 2022

Strategy Meeting Zurich and Boston

Zurich - Europe

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Clinical Operations Strategy Meeting 2022

Radisson Blu Hotel Zurich Airport

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BIOMANUFACTURING

Biomanufacturing Strategy Meeting 2022

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Clinical Trial Supply Chain Strategy Meeting 2022

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Cell & Gene Therapy Strategy Meeting 2022

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Exploring Promising Areas in ONCOLOGY RESEARCH Aimed to Expand Our Knowledge Base, Address Unmet Medical Needs and Improve Overall Clinical Success