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Proventa International's 11th Annual

CLINICAL OPERATIONS & CLINICAL TRIAL SUPPLY CHAIN

STRATEGY MEETING WEST COAST USA 2024

Thursday, 23rd May 2024 **Hard Rock Hotel San Diego**

Jumpstart Clinical Operations on Patient Centricity, Innovation and Efficiency

Featuring Industry Leaders and Decision Makers:

Amish Patel
Senior Vice President, Technical Operations
Calidi Biotherapeutics

Tanya Russell Kirkpatrick
Vice President, Clinical Operations Head - Oncology and Rare Disease - Global Product Development
Pfizer

Joseph Shan
Vice President, Clinical Operations
Adcentrx Therapeutics

Janal Urich
Vice President, Global Portfolio and Program Management
Atara Biotherapeutics

Moya Daniels
Chief Quality Officer
Diakonos Oncology

Tara Lehner
Vice President, Clinical Operations
INmune Bio Inc.

Catherynne Cruz-Scheckner
Executive Director, Clinical Supplies
Ionis Pharmaceuticals, Inc.

24 ROUNDTABLE DISCUSSIONS

6 TRACKS

2 KEYNOTE PRESENTATIONS

1 PANEL DISCUSSION

1 LOCATION

What Makes Our Strategy Meetings So Unique?

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

CLINICAL OPERATIONS & CLINICAL TRIAL SUPPLY CHAIN

STRATEGY MEETING WEST COAST USA 2024

Thursday 23rd May 2024 | Hard Rock Hotel San Diego

We're committed to delivering long-term value across our extensive life science network. Through our carefully crafted meetings, collaborative experiences and services Proventa International can offer you the perfect opportunity to meet your business goals, whatever they may be.

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.

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

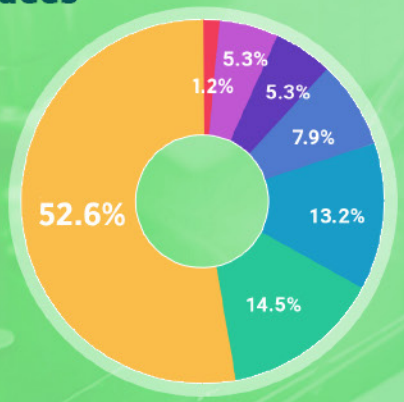
The most effective and time efficient way to assess potential partners at a strategic level. Identify key solution providers that can take your business to the next level and we will help arrange private meetings so you can connect.

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
- President / VP
- Department Head
- Other
- Team Lead
- C-Level
- Scientist
- Academia
- Manager
- Biology Specialist



Clinical Operations

- Clinical Development
- Clinical Operations
- Clinical IT
- Clinical Data
- Clinical Innovation
- Early Phase Trials
- Late Phase Trials
- Risk Based Monitoring
- Biostatistics
- Study Feasibility / Site Head / Site Operations
- Patient Recruitment

Clinical Trial Supply Chain

- CEO
- VP
- Director
- Global Clinical Program Head
- Clinical Trial Logistics
- Supply Chain
- Clinical Distribution
- Clinical Supplies Management
- Warehousing
- Cold Chain
- Quality Inspection

Meet Investors

- Venture Capital
- Private Equity
- Large Pharma/Biotech
- Corporate Venture Capital
- Institutional
- High Net Worth
- Family Office/Private Wealth
- Government Organisation/
- Sovereign Wealth Fund
- Angel

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PROVENTA INTERNATIONAL

Facilitator Faculty

Amish Patel
Senior Vice President, Technical Operations
Calidi Biotherapeutics

Catherynne Cruz-Scheckner
Executive Director, Clinical Supplies
Ionis Pharmaceuticals, Inc.

Janal Urich
Vice President, Global Portfolio and Program Management
Atara Biotherapeutics

Joseph Shan
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Adcentrx Therapeutics

Tanya Russell Kirkpatrick
Vice President, Clinical Operations Head - Oncology and Rare Disease - Global Product Development
Pfizer

Tara Lehner
Vice President, Clinical Operations
INmune Bio Inc.

Moya Daniels
Chief Quality Officer
Diakonos Oncology

How Has Our **STRATEGY MEETINGS** Benefit The Life Science Industry

This was a very important insightful event and I have few takeaways messages/practices which I will be implementing on my current and future projects."

Jean-Pierre Metabanzoulou -
Senior Director, CMC & RA, Acasti Pharma

This was a very informative, significantly interactive and highly educational event. "

Domenico Merante -
VP Clinical Development, Sosei Heptares

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CRIO, the leader in eSource technology, is an enterprise grade eClinical solution transforming clinical research with the latest in cloud technology. CRIO eSource enables remote monitoring and immediate data review with a powerfully integrated eSource/EDC solution that protects data integrity. With CRIO, users enter data into a simple, interactive application, capturing data only once - at the moment of immediacy - without re-entering data into the EDC. Alerts, automatic calculations, and custom controls make data entry more efficient, timely and accurate. And, because everything is available online, CRAs can monitor site operations remotely, and data managers can view data immediately.

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Proventa's end-to-end consulting division gather real-time business intelligence on the industry's **needs, challenges, budgets** and **investment areas**. We combine this information with your specific needs to enhance your business development strategy. With the wealth of intel we provide, Proventa guarantees tangible results for your business within twelve months of the event.

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Agenda at a Glance

CLINICAL OPERATIONS & CLINICAL TRIAL SUPPLY CHAIN
STRATEGY MEETING WEST COAST USA 2024
Thursday, 23rd May 2024 | Hard Rock Hotel San Diego

TIME (PT)	TRACK 1: PATIENT RECRUITMENT / DECENTRALIZED & HYBRID TRIALS AND EMERGING BIOPHARMA / OUTSOURCING	TRACK 2: CLINICAL TECHNOLOGIES & INNOVATION	TRACK 3: GLOBAL SITE SELECTION AND FEASIBILITY STUDY	TRACK 4: DIVERSITY, EQUITY, AND INCLUSION IN PATIENT RECRUITMENT & ENGAGEMENT	TRACK 5: CLINICAL SUPPLIES / STORAGE & GLOBAL NETWORKS
ROOM ►					
08:00 - 08:30	BREAKFAST & REGISTRATION				
08:30 - 09:00	OPENING KEYNOTE PRESENTATION				
09:00 - 10:00 PHARMA/ BIOTECH	**Patient Recruitment 2.0: Strategies And Approaches for Success in Decentralized/ Hybrid Trials**	**The Latest Trends in Electronic Data Capture (EDC) Systems and eClinical Solutions for Digital Transformation in Clinical Operations** Joseph Shan, *Vice President, Clinical Operations,* Adcentrx Therapeutics	**Embarking on the Pinnacle of Clinical Trial Success in 2024: Unveiling Global Site Selection Strategies, Where Patient Population Precision Aligns with Study Criteria, and Ethical and Cultural Considerations Illuminate the Path to Ethical and Culturally Sensitive Trials**	**Implementing Diversity, Equity, and Inclusion (DEI) Initiatives: Insights from Company Case Studies and Exemplary Practices**	**A Strategic Discussion on Clinical Supplies, Storage Innovation, and Global Networks Reshaping the Landscape of CTM, Storage Facilities, Packaging, and Labeling in Clinical Trials** Amish Patel, *Senior Vice President, Technical Operations,* Calidi Biotherapeutics
10:00 - 10:05	REFRESHMENT BREAK				
10:05 - 10:25	NETWORKING / 1-1 MEETINGS				
10:25 - 10:45	NETWORKING / 1-1 MEETINGS				
10:45 - 11:05	NETWORKING / 1-1 MEETINGS				
11:10 - 12:10 SOLUTION	**The Role of AI/ML in Enhancing Outsourcing Efficiency for Emerging Biopharma** SPONSOR	**AI and ML in 2024: Innovations Reshaping Clinical Trial Supply Chains** SPONSOR	**How Can Global Site Selection Evolve to Optimize Assessments of Site Infrastructure, Ensuring Facilities, Equipment, and Resources Meet the Demands of Modern Clinical Trials?** SPONSOR	**Navigating The Future Involves Pioneering Strategies For Diversity, Equity, And Inclusion In Patient Recruitment And Engagement Through Comprehensive Demographic Data Collection** SPONSOR	**Interrogating the Precision of Carrier Selection, Route Optimization, and Timely Delivery of Clinical Supplies and Storage with Global Networks and Distribution Logistics** PELI BIOTHERMAL
12:15 - 13:15 PHARMA/ BIOTECH	**How are Emerging Biopharma Firms Tackling Infrastructure Challenges to Expedite Drug Development?** Janal Urich, *Vice President, Global Portfolio and Program Management,* Atara Biotherapeutics	**Pros and Cons of Randomization and Trial Supply Management Systems vs. Fully Integrated IRTs** Catherynne Cruz-Scheckner, *Executive Director, Clinical Supplies,* Ionis Pharmaceuticals, Inc.	**How Can Sponsors and Researchers Harness the Power of Global Site Selection and Feasibility Studies to Anticipate Challenges and Optimize Conditions for Successful Global-Scale Trials?** Tara Lehner, *Vice President, Clinical Operations,* INmune Bio Inc.	**Eliminating Health Inequities: How Can We Improve The Diverse Representation Of Clinical Trial Programs To Drive Health Equity? (Topic TBC)** Tanya Russell Kirkpatrick, *Vice President, Clinical Operations Head - Oncology and Rare Disease - Global Product Development,* Pfizer	**Mastery of Clinical Logistics: Unleashing the Power of Quality Control and Assurance in Clinical Supplies/Storage**
13:15 - 14:00	NETWORKING LUNCH				
14:00 - 14:20	NETWORKING / 1-1 MEETINGS				
14:20 - 14:40	NETWORKING / 1-1 MEETINGS				
14:40 - 15:10	AFTERNOON KEYNOTE PRESENTATION				
15:10 - 16:10 SOLUTION	**Managing the Complications of Collaborations: Obstacles and Resolutions in Outsourcing for Developing Biotech/Pharma Companies**	**Harnessing the Power of Digital Platforms for Enhanced Monitoring (EMC), Communication, and Decision-Making Across the Clinical Trial Ecosystem In Global Clinical Trials**	**An In-Depth Exploration of Emerging Trends and Cutting-Edge Best Practices in Elevating Clinical Trial Awareness and Revolutionizing Recruitment Strategies**	**Involving Patients in Decision-Making: Exploring a Scope of Strategies, Statements, and Transformations for Engaging Underrepresented Communities and Patient Engagement** Moya Daniels, *Chief Quality Officer,* Diakonos Oncology	**Strategies and Approaches for Seamless Integration of Clinical Supplies, Storage, Global Networks, and Regulatory Compliance through Dynamic Partnerships and Supplier Networks**
16:10 - 16:30	AFTERNOON REFRESHMENT BREAK				
16:30 - 17:00	PANEL DISCUSSION: Investing in Health Technology Companies with a Clinical Focus in Pharma and Biotech				
17:00 - 18:00	DRINKS & CANAPES RECEPTION				

Event Day | Keynote Presentations

A great way to open the roundtable discussions is through a timely presentation from a top-tier biotech/ pharmaceutical company. Listen as we hear this 30-minute exposition on this meeting's pressing topic.

08:30 - 09:00 PT **OPENING KEYNOTE PRESENTATION**

Topic TBC

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

14:40 - 15:10 PT **AFTERNOON KEYNOTE PRESENTATION**

Topic TBC

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

16:30 - 17:00 PT **AFTERNOON KEYNOTE PRESENTATION**

Investing in Health Technology Companies with a Clinical Focus in Pharma and Biotech

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

Event Day

CLINICAL OPERATIONS & CLINICAL TRIAL SUPPLY CHAIN
STRATEGY MEETING WEST COAST USA 2024
Thursday, 23rd May 2024 | Hard Rock Hotel San Diego

TRACK 1

Patient Recruitment / Decentralized & Hybrid Trials And Emerging Biopharma / Outsourcing

A mix of risk assessments, analytical capabilities, and cutting-edge technology is required to provide an effective risk-based monitoring platform. Various discussions will focus on how we can include successful risk-mitigation methods and how we can fast improve patient safety to guarantee we are moving in clinical trials

08:00 - 08:30 PT — **BREAKFAST & REGISTRATION**

08:30 - 09:00 PT — **OPENING KEYNOTE PRESENTATION**
See Page 6

09:00 - 10:00 PT — **ROUNDTABLE 1**

Patient Recruitment 2.0: Strategies And Approaches for Success in Decentralized/Hybrid Trials

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

10:00 - 11:10 PT — **REFRESHMENT BREAK & NETWORKING / 1-1 MEETINGS**

11:10 - 12:10 PT — **SOLUTION FOCUS ROUNDTABLE 2**

The Role of AI/ML in Enhancing Outsourcing Efficiency for Emerging Biopharma

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

12:15 - 13:15 PT — **ROUNDTABLE 3**

How are Emerging Biopharma Firms Tackling Infrastructure Challenges to Expedite Drug Development?

Janal Urich
Vice President, Global Portfolio and Program Management
Atara Biotherapeutics

ABOUT THE SPEAKER
Speaker TBC

13:15 - 14:40 PT — **REFRESHMENT BREAK & NETWORKING / 1-1 MEETINGS**

14:40 - 15:10 PT — **AFTERNOON KEYNOTE PRESENTATION**
See Page 6

15:10 - 16:10 PT — **ROUNDTABLE 4**

Managing the Complications of Collaborations: Obstacles and Resolutions in Outsourcing for Developing Biotech/ Pharma Companies

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

16:10 - 16:30 PT — **AFTERNOON REFRESHMENT BREAK**

16:30 - 17:00 PT — **PANEL DISCUSSION**
See Page 6

17:00 - 18:00 PT — **DRINKS & CANAPES RECEPTION**

Event Day

TRACK 2 Clinical Technologies & Innovation

This track will highlight important discussions to help you stay ahead of the curve and retain a competitive advantage in the industry by examining major trends that will emerge and impact the future of Clinical Operations.

08:00 - 08:30 PT **BREAKFAST & REGISTRATION**

08:30 - 09:00 PT **OPENING KEYNOTE PRESENTATION**
See Page 6

09:00 - 10:00 PT **ROUNDTABLE 1**

The Latest Trends in Electronic Data Capture (EDC) Systems and eClinical Solutions for Digital Transformation in Clinical Operations

- EDC Systems in Clinical Research: What's working and what's not?
- How will clinical data collection evolve over the next 10 years?
- What are best practices for implementing novel eClinical solutions in Clinical Operations?

Joseph Shan
Vice President, Clinical Operations
Adcentrx Therapeutics

ABOUT THE SPEAKER
Joseph (Joe) Shan, M.P.H., VP, Clinical Operations at Adcentrx Therapeutics, has nearly 30 years of clinical experience in biopharma and medical device companies. Nearly 25 years were focused on clinical development of oncology therapeutics, including radiopharmaceuticals, small molecules, targeted therapy, immunotherapy and cellular therapy. Prior to Adcentrx, Joe held leadership roles in Clinical Development Operations at Mosaic ImmunoEngineering, MEI Pharma, Elevar Therapeutics, Kiadis Pharma (formerly CytoSen Therapeutics) and was VP, Clinical & Regulatory Affairs and a corporate officer of Peregrine Pharmaceuticals for nearly a decade. Mr. Shan received his M.P.H. from George Washington University and B.S. from UCLA

10:00 - 11:05 PT **REFRESHMENT BREAK & NETWORKING / 1-1 MEETINGS**

11:10 - 12:10 PT **SOLUTION FOCUS ROUNDTABLE 2**

AI and ML in 2024: Innovations Reshaping Clinical Trial Supply Chains

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

12:15 - 13:15 PT **ROUNDTABLE 3**

Pros and Cons of Randomization and Trial Supply Management Systems vs. Fully Integrated IRTs

- IRT full capabilities
- RTSMs role in clinical trials
- Tailoring the system for maximum flexibility vs maximum compliance

Catherynne Cruz-Scheckner
Executive Director, Clinical Supplies
Ionis Pharmaceuticals, Inc.

ABOUT THE SPEAKER
Catherynne is an Executive Director at Ionis Pharmaceuticals, heading up the clinical supplies department. She started her career as a pharmacist in a retail and hospital setting and has worked for large pharmaceutical companies such as Warner Lambert, Parke Davis, Pfizer, and Valeant Pharmaceuticals, as well as smaller biotech companies like Ascenta Therapeutics and Phenomix. Catherynne is a clinical supplies management professional with 28 years of pharmaceutical experience and a proven track record of building cross-functional collaboration resulting in on-time delivery of project goals. Background encompasses pre-formulation/formulation, manufacturing, clinical supplies, IRT development, supply chain, and CMC project management.

13:15 - 14:40 PT **REFRESHMENT BREAK & NETWORKING / 1-1 MEETINGS**

14:40 - 15:10 PT **AFTERNOON KEYNOTE PRESENTATION**
See Page 6

15:10 - 16:10 PT **ROUNDTABLE 4**

Harnessing the Power of Digital Platforms for Enhanced Monitoring (EMC), Communication, and Decision-Making Across the Clinical Trial Ecosystem In Global Clinical Trials

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

16:10 - 16:30 PT **AFTERNOON REFRESHMENT BREAK**

16:30 - 17:00 PT **PANEL DISCUSSION**
See Page 6

17:00 - 18:00 PT **DRINKS & CANAPES RECEPTION**

Event Day

CLINICAL OPERATIONS & CLINICAL TRIAL SUPPLY CHAIN
STRATEGY MEETING WEST COAST USA 2024
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TRACK 3

Global Site Selection and Feasibility Study

Due to the COVID outbreak, many companies are struggling to enrol patients in the appropriate sites that are closer to them. They're up against the task of finding the correct user-friendly data innovation in order to identify matches with different sites that will allow the study to go more quickly, as well as access to diverse locations that will offer trials to patients of various ethnicities.

08:00 - 08:30 PT BREAKFAST & REGISTRATION

08:30 - 09:00 PT OPENING KEYNOTE PRESENTATION
See Page 6

09:00 - 10:00 PT ROUNDTABLE 1

Embarking on the Pinnacle of Clinical Trial Success in 2024: Unveiling Global Site Selection Strategies, Where Patient Population Precision Aligns with Study Criteria, and Ethical and Cultural Considerations Illuminate the Path to Ethical and Culturally Sensitive Trials

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

10:00 - 11:05 PT REFRESHMENT BREAK & NETWORKING / 1-1 MEETINGS

11:10 - 12:10 PT SOLUTION FOCUS ROUNDTABLE 2

How Can Global Site Selection Evolve to Optimize Assessments of Site Infrastructure, Ensuring Facilities, Equipment, and Resources Meet the Demands of Modern Clinical Trials?

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

12:15 - 13:15 PT ROUNDTABLE 3

How Can Sponsors and Researchers Harness the Power of Global Site Selection and Feasibility Studies to Anticipate Challenges and Optimize Conditions for Successful Global-Scale Trials?

- How to scope out and assess what is required for trial success
- How to partner for optimal feasibility results
- Navigating assumptions that arise through different cultural lenses
- Managing internal stakeholders when planned trial designs change because of feasibility results

Tara Lehner
Vice President, Clinical Operations
INmune Bio Inc.

ABOUT THE SPEAKER
Ms. Lehner is Vice President of Clinical Operations at INmune Bio. She has spent over 24 years in the pharmaceutical industry as an employee and consultant working across pharma, biotech, and CROs in Phases I–IV, with a focus on oncology and CNS. She has led long-term strategic partnerships between CROs and pharma and emphasizes the humanness of business relationships. Ms. Lehner holds an MS from the Temple University School of Pharmacy. Tara lives on a ranch in Southern California with her partner and two daughters, ages 7 and 8.

13:15 - 14:40 PT NETWORKING LUNCH & NETWORKING / 1-1 MEETINGS

14:40 - 15:10 PT AFTERNOON KEYNOTE PRESENTATION
See Page 6

15:10 - 16:10 PT ROUNDTABLE 4

An In-Depth Exploration of Emerging Trends and Cutting-Edge Best Practices in Elevating Clinical Trial Awareness and Revolutionizing Recruitment Strategies

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

16:10 - 16:30 PT AFTERNOON REFRESHMENT BREAK

16:30 - 17:00 PT PANEL DISCUSSION
See Page 6

17:00 - 18:00 PT DRINKS & CANAPES RECEPTION

Event Day

CLINICAL OPERATIONS & CLINICAL TRIAL SUPPLY CHAIN
STRATEGY MEETING WEST COAST USA 2024
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TRACK 4

Diversity, Equity, And Inclusion In Patient Recruitment & Engagement

Delays in patient recruitment and engagement activities can cause major setbacks and become remarkably costly. This track is aimed at identifying key pitfalls, and exploring effective patient recruitment plans that leverage the latest tools, technologies and analytical approaches to deploy into having a successful clinical trial.

08:00 - 08:30 PT BREAKFAST & REGISTRATION

08:30 - 09:00 PT OPENING KEYNOTE PRESENTATION
See Page 6

09:00 - 10:00 PT ROUNDTABLE 1

Eliminating Health Inequities: How Can We Improve The Diverse Representation Of Clinical Trial Programs To Drive Health Equity? (Topic TBC)

Tanya Russell Kirkpatrick
Vice President, Clinical Operations
Head - Oncology and Rare Disease - Global Product Development
Pfizer

ABOUT THE SPEAKER
Speaker TBC

10:00 - 11:05 PT REFRESHMENT BREAK & NETWORKING / 1-1 MEETINGS

11:10 - 12:10 PT SOLUTION FOCUS ROUNDTABLE 2

Navigating The Future Involves Pioneering Strategies For Diversity, Equity, And Inclusion In Patient Recruitment And Engagement Through Comprehensive Demographic Data Collection

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

12:15 - 13:15 PT ROUNDTABLE 3

Eliminating Health Inequities: How Can We Improve The Diverse Representation Of Clinical Trial Programs To Drive Health Equity?

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC

13:15 - 14:40 PT NETWORKING LUNCH & NETWORKING / 1-1 MEETINGS

14:40 - 15:10 PT AFTERNOON KEYNOTE PRESENTATION
See Page 6

16:20 - 17:20 PT ROUNDTABLE 4

Involving Patients in Decision-Making: Exploring a Scope of Strategies, Statements, and Transformations for Engaging Underrepresented Communities and Patient Engagement

Moya Daniels
Chief Quality Officer
Diakonos Oncology

ABOUT THE SPEAKER
Speaker TBC

16:10 - 16:30 PT AFTERNOON REFRESHMENT BREAK

16:30 - 17:00 PT PANEL DISCUSSION
See Page 6

17:00 - 18:00 PT DRINKS & CANAPES RECEPTION

new for 2024

Event Day

TRACK 5

Strategic Partnerships, Investment & Collaborations

Bringing together Managing Partners & C-level executives from investment firms supporting innovation in the clinical trial process, strategies from pharma companies and corporate arms with a vested interest in driving this innovation. It comes with the purpose of identifying and collaborating with potential drug development partners whilst addressing industry challenges in this area

09:00 - 10:00 PT ROUNDTABLE 1

A Strategic Discussion on Clinical Supplies, Storage Innovation, and Global Networks Reshaping the Landscape of CTM, Storage Facilities, Packaging, and Labeling in Clinical Trials

- The importance of global networks for clinical supply delivery of clinical trial materials to study sites on time, considering factors such as transportation modes, customs regulations, and local regulations
- Explore the challenges and best practices for managing and storing clinical trial materials. Consider the importance of temperature-controlled storage, proper labeling, documentation, and inventory management to ensure accurate tracking and efficient supply chain management
- Discuss the importance of appropriate temperature-controlled packaging materials and strategies to ensure the safe and efficient distribution of clinical supplies

Amish Patel
Senior Vice President Technical Operations
Calidi Biotherapeutics

ABOUT THE SPEAKER

Dr. Amish Patel has over 20 years of leadership experience in the global biotech industry, providing strategic guidance throughout the entire biologic product life-cycle and been involved with fifteen+ INDs and commercial product launch. Dr. Patel is responsible for oversight of Global Manufacturing, Quality Control, Quality Assurance and Supply Chain to produce Clinical and Commercial cancer therapies utilizing Calidi's innovative oncolytic viral therapies with stem cell-based delivery platforms to treat cancers with significant unmet needs. Dr Patel has managed the CDMO network and supplied clinical products for phase 1 and phase 3 trials globally. Dr Patel has extensive experience in licensing and out licensing several technologies from Academia and non profit organizations and even commercialized such assets. He serves as Technical Activities Committee of the National Institute for Innovation in Manufacturing Biopharmaceuticals (NIIMBL) part of Manufacturing USA® and funded through the U.S. Department of Commerce with significant additional support from its members. He was most recently, the Senior Director of Product Development at Emergent BioSolutions (formerly PaxVax, Inc., prior to acquisition), where he successfully built and directed cross-functional teams working on numerous biologic Clinical and commercial products.

11:10 - 12:10 PT SOLUTION FOCUS ROUNDTABLE 2

Interrogating the Precision of Carrier Selection, Route Optimization, and Timely Delivery of Clinical Supplies and Storage with Global Networks and DIstribution Logistics

Speaker TBC

PELI BIOTHERMAL

ABOUT THE SPEAKER
Speaker TBC

12:15 - 13:15 PT ROUNDTABLE 3

Mastery of Clinical Logistics: Unleashing the Power of Quality Control and Assurance in Clinical Supplies/Storage

Speaker TBC

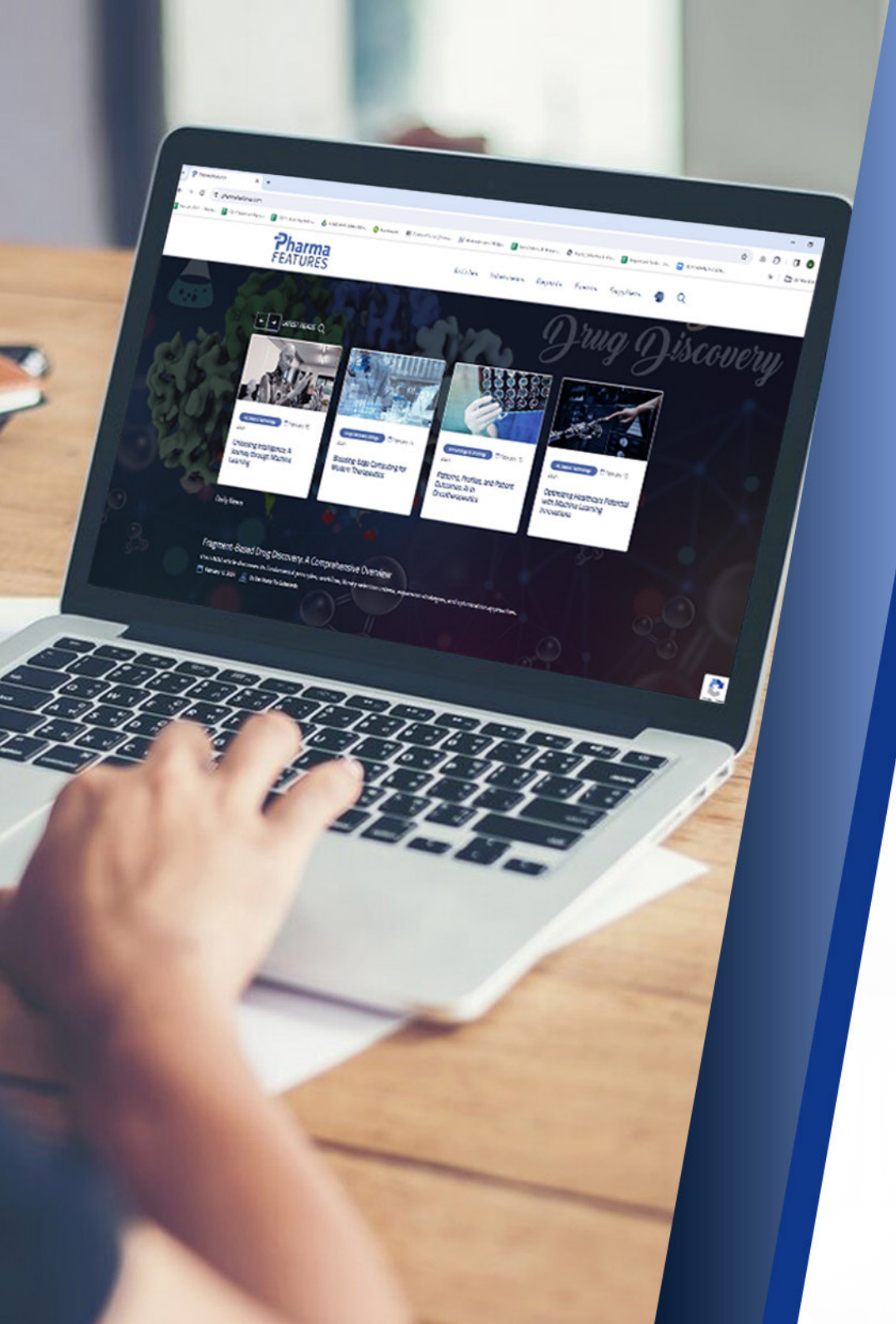
ABOUT THE SPEAKER
Speaker TBC

15:10 - 16:10 PT ROUNDTABLE 4

Strategies and Approaches for Seamless Integration of Clinical Supplies, Storage, Global Networks, and Regulatory Compliance through Dynamic Partnerships and Supplier Networks

Speaker TBC

ABOUT THE SPEAKER
Speaker TBC



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CLINICAL OPERATIONS & CLINICAL TRIAL SUPPLY CHAIN

STRATEGY MEETING WEST COAST USA 2024

Thursday 23rd May 2024 | Hard Rock Hotel San Diego

Delegate Price *Pharma/Biotech/ Academia — $699

Vendor/ Service Provider — $2999

OUR FACE TO FACE MEETING IN MAY 2024

STRATEGY MEETING EAST COAST USA
Hyatt Regency Princeton

- MAY 14 — DRUG DISCOVERY BIOLOGY & BIOINFORMATICS — STRATEGY MEETING EAST COAST USA 2024
- MAY 15 — MEDICINAL CHEMISTRY — STRATEGY MEETING EAST COAST USA 2024
- MAY 16 — CLINICAL OPERATIONS & CLINICAL TRIAL SUPPLY CHAIN — STRATEGY MEETING EAST COAST USA 2024

STRATEGY MEETING WEST COAST USA
Hard Rock Hotel San Diego

- MAY 21 — DRUG DISCOVERY BIOLOGY & BIOINFORMATICS — STRATEGY MEETING WEST COAST USA 2024
- MAY 22 — MEDICINAL CHEMISTRY — STRATEGY MEETING WEST COAST USA 2024
- MAY 23 — CLINICAL OPERATIONS & CLINICAL TRIAL SUPPLY CHAIN — STRATEGY MEETING WEST COAST USA 2024

Hotel & Venue

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