Clinical Trial Supply East Asia

Clinical Trial Supply East Asia 2023

Seoul, South Korea, JW Marriot Seoul

5th-6th December 2023

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The leading clinical trial supply conference in East Asia is returning on 5th-6th December 2023. With a focus on providing delegates with practical take-aways and solutions to their most current operational and outsourcing challenges in clinical trials, this is an event not to be missed.

Focusing on the regional pharmaceutical business in Asia, highlighting the industry's current difficulties as well as its exciting innovations and growth prospects

2023 Speakers

Moon Hwan Kim, Chief Technology Officer, Veraverse
Jaehyun Park, Managing Director, Animuscure
Tina Sun, Study & Site Operations Country Head (Taiwan)- Novartis
Sangho Ma, Chief Development Officer, Bilix
Guoqing Yang, Executive Director, Clinical Quality Assurance,(China)Shanghai Henlius Biotech
Sol Han, Chief Medical Officer, Cyrus Therapeutics
Nari Yun, Executive Director, Clinical Development, GI Innovation
Taegyun Park, Head of R&D, Qratis
Jessica Thongcharen, Director Global Patient Safety, Clinical Operations(Singapore), Takeda
YoungSu Noh, Clinical Science Manager, Hanmi Pharma
Sueun Song, Division Head of Clinical Development, Celltrion
KangPyo Lee, Vice President, Hanul TL
Seoyeon Hyun, Clinical Development Director, ILIAS Biologics
Jungwon Jung, Manufacturing Manager, HK inno-N
DaeMan Moon, Director, Team Leader, Global Clinical Supply Management Team, Celltrion
Juhee JEON, PV Manager, PV Team Leader, HANA pharm
HyeJung Yang, Clinical Project Manager, MedPacto
Yohan Bae, Executive Director, Clinical Development Division, Kangstem Biotech
Eunsun Lee, Director, TiumBio
SoYoung Chun, Medical Director, Trial Informatics
Shin-Il Kim, Chief Scientific Officer, THERABEST
Baek-Jae KIM, Country Manager Korea, IATA
Hyesung Shin, Managing Director, Division Head Clinical Research Development Division, TiumBio
A Young Kim, R&D division Team leader, Jeil pharm
Yooni Kim, Vice President Clinical Services, Novotech
Marine du Jardin, APAC Regional Manager & Partnership Lead, N-SIDE
Amaury Jeandrain, Vice President of Strategy, N-SIDE

Join our new CTS community on LinkedIn

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Clinical Trial Supply East Asia

DAY 1 – 5th December 2023

8:00	**Registration and refreshments**
8:50	**Chairperson's opening remarks**
9:00	**KEY NOTE** **Choosing the right CRO: Essential considerations for a successful partnership** • Recognizing common misconceptions when selecting a CRO that may not align with your values and trial requirements • Determining the essential resources and capabilities that the CRO must possess to ensure the success of your trial • Managing that all important budget to avoiding overspending **Sueun Song, Division Head of Clinical Development, Celltrion**
9:30	**Reducing drug waste - how Technology makes clinical supply more efficient** • Drug waste in clinical trials: why is it a big deal? • Identifying the main drivers of waste • How technology enables more accurate and sustainable supply planning, leading to costs and drug waste reduction • Illustration through case studies **Marine du Jardin, APAC Regional Manager & Partnership Lead, N-SIDE** **Amaury Jeandrain, Vice President of Strategy, N-SIDE**
10:00	**Master protocols in biotech: Design and implementation** • Concept of master protocols • FDA perspectives on mater protocols • Operation of complex study design • Examples of mater protocols **Nari Yun, Executive Director, Clinical Development, GI Innovation**
10:30	**The Future of Clinical Trials: Clinical Innovation Solutions Utilizing of Artificial Intelligence (AI)** • The key trends in the clinical trials landscape • How to do CROs use AI in clinical trials • Real-world cases of AI and CRO **Yooni Kim, Vice President Clinical Services, Novotech**
11:00	**Morning refreshments and networking**
	STREAM B: Clinical Trial Supply

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11:30	**Cold chain management and optimal shipping practices** • Exploring how organisations can improve cold chain management, focusing on temperature-sensitive product handling, storage • Providing insights and strategies for effectively managing cold chain operations, such as risk assessment • Discussing the challenges of shipping temperature-sensitive products and outlining best practices **KangPyo Lee, Vice President, Hanul TL**
12:00	**Reserved for the Event Partner**
12:30	**Clinical supply strategies: tailoring approaches for successful trials** • Addressing the significance of tailoring clinical supply strategies to match the unique requirements of each trial • Examining proactive steps to manage risks, such as regulatory changes, supply chain disruptions, and unexpected patient enrollment • Discussing the ability to adapt supply strategies in response to unforeseen challenges, minimizing disruptions • **DaeMan Moon, Team Leader Development Supply Chain, Celltrion**
13:00	**Lunch and networking**
14:00	**IATA Temperature Control Regulations (TCR) on Clinical rials supply chain** • Influence of TCR (Time, Temperature, and Condition) on Pharmaceutical Transportation • Effective Application of Time and Temperature Sensitive Labels • Ensuring Compliance with TCR Regulations in Clinical Trials **Baek-Jae KIM, Country Manager Korea, IATA**
14:30	**Reserved for the Event Partner**
15:00	**Disclosing approaches to managing temperature effectively in IP/IMP management during transfer** • Addressing typical occurrences of temperature fluctuations during the distribution process • Highlighting the significance of real-time online tracking tool to identify, trace, track, locate and protect perishable drugs on their journey through supply chain to provide immediate support to achieve high quality **SoYoung Chun, Medical Director, Trial Informatics**
15:30	**Afternoon refreshments and networking**

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16:00	**Efficient strategies for IRTs and Just-in-Time packaging, labelling, and shipment in clinical trials** • Exploring strategies to improve the efficiency of Interactive Response Technology (IRT) systems in clinical trials • Discussing effective risk management strategies to anticipate and mitigate potential challenges associated with just-in-time operations, ensuring uninterrupted supply and minimizing delays • Investigating cutting-edge technological solutions that facilitate real-time tracking, monitoring, and coordination **HyeJung Yang, Clinical Project Manager, MedPacto**
16:30	**PANEL DISCUSSION: Mitigating unnecessary delays in clinical supply logistics: Huddle and Solutions** • Strategies to Minimize Delays in Clinical Supply Logistics • Implementing Effective Communication through Huddles • Practical Solutions for Streamlining Supply Chain Processes **Jaehyun Park, Managing Director, Animuscure** **Baek-Jae KIM, Country Manager Korea, IATA** **HyeJung Yang, Clinical Project Manager, MedPacto**
17:15	**Chairperson's closing remarks**
END OF DAY 1 AND NETWORKING DRINKS	

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DAY 2 – 6th December 2023

8:15	**Registration and refreshments**
8:50	**Chairperson's opening remarks**
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9:00	**Selecting the optimal CMO Partner for product manufacturing: Key considerations and best practices** • Understanding the importance of aligning with a CMO that shares your values and vision to foster a strong partnership • Evaluating the criteria to assess CMO capabilities • Emphasizing the importance of clear communication channels and effective collaboration between your team and the CMO **Jungwon Jung, QA Director, HK inno-N**
9:30	**Reserved for the Event Partner**
10:00	**Start-up Pharma & Biotech's; Establishing a relationship, engaging with your supply chain and selecting the right partners for you** • Deconstructing how to create a clinical supply chain as a small start-up pharmaceutical company • Key steps to have a progressive supply chain strategy with the potential for long-term company growth • Hear insight on how to manage supply chain activities through effective partnership management **Yohan Bae, Executive Director, Clinical Development Division, Kangstem Biotech**
10:30	**How to navigate sourcing for clinical trials in China** • Are you confused about where to start sourcing in China? • Do you have concerns about quality and transparency when investing into a trial in China? • Have time, language, and cultural differences affected the efficiency of your project? **Guoqing Yang, Executive Director, Clinical Quality Assurance, Shanghai Henlius Biotech**
11:00	**Morning refreshments and networking**
11:30	**Streamlining the End-to-End pharmaceutical supply Chain: Enhancing efficiency and quality** • Understanding the complexities and challenges of the end-to-end pharmaceutical supply chain, from manufacturing to patient delivery • Addressing regulatory requirements and compliance • How did we implement effective strategies to enhance the end-to-end pharmaceutical supply chain **Hyesung Shin, Managing Director**, **Division Head Clinical Research Development Division, TiumBio**
12:00	**Emphasizing the collaboration between clinical supplies and quality assurance teams to ensure compliant trials, mitigating potential delays and costs** • Aligning early phase vs late phase quality assurance considerations with supply chain protocols to minimize disruptions and backlogs in supply chain • Understanding importance of role and responsibility of a Qualified Person and timelines required for QP release • Improving communication channels between clinical supplies and QA teams **A Young Kim, R&D division Team leader, Jeil pharm**

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12:45	**Lunch and networking**
13:45	**Prize Draw** **Join us for a chance to win one of our prizes, including Apple products**
14:00	**Speaker Hosted Roundtables** Interactive roundtable sessions offer a unique opportunity to come together with your peers to share best practice and develop solutions to critical challenges facing the industry as a whole. Hosted by industry experts and each focused on a single issue, roundtables are an exciting, interactive way to build your personal network and learn from the experience and expertise of others. Roundtable session lasts for 45 minutes, and we'll have a 45min wrap up session
	ROUNDTABLE 1: Establishing strategies for vendor management and oversight to strengthen collaboration **Seoyeon Hyun, Clinical Development Director, ILIAS Biologics**
	ROUNDTABLE 2: Navigating global supply chain hurdles when shipping cold chain supplies oversea **Guoqing Yang, Executive Director, Clinical Quality Assurance, Shanghai Henlius Biotech**
	ROUNDTABLE 3: Decentralized clinical trials and technology **Moon Hwan Kim, Chief Technology Officer, Veraverse**
13:30	**Chairperson's closing remarks**
END OF CONFERENCE	

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