Virtue Insight

PRESENTS

#VIct

Clinical TRIALS

Series 2025

25th September 2025

Sterlings Mac Hotel, Bengaluru, India

www.virtueinsight.com

KEY SPEAKERS INCLUDE

KHALID AHMED KHAN
Deputy Drugs Controller
Food Safety and Drug Administration

MILIND ANTANI
Leader, Pharma and Healthcare
Nishith Desai Associates

TAPANKUMAR SHAH
Senior Director
AstraZeneca

MAYUR PARMAR
Drugs Inspector (Deputy Collector, Gujarat Government), FDA

KRISHNENDU SENGUPTA
Director - Centralized Monitoring
AstraZeneca

RAHUL RATHOD
Director Medical Affairs
Abbvie

SUBHASH THULUVA
Sr. Vice President & Head - Clinical Development, Biological E

MONISHA SHARMA
Senior Director, APAC. Global Clinical Trial Operations, MSD

KHALEEL AHMAD
Associate Director, Line Manager, Global Trial Management
Bristol Myers Squibb

KAMALA RAI
Vice President, Senior Global Program Head, Novartis

VIJAY PARTHASARATHY
Senior Director, Clinical Operations
Novo Nordisk

ASHISH PAWAR
Associate Director, Clinical Data Science, Eli Lilly

ANIL KUKREJA
Executive VP - Clinical Research & Medical Affairs, Emcure Pharmaceuticals

DEEPAK NEMA
Director, Clinical Data Sciences
Pfizer

PRITAM GUPTA
Associate Director, Biostatistics
Bristol Myers Squibb

SHUBHADEEP SINHA
Senior Vice-President & Medical Director
Hetero

FRANCIS MICHEAL
Director
Viatris

ANUP CHOUDHURY
Medical Director, Global Clinical Development, Novartis

LATEEFUDDIN SHARIFF
Senior Director - Global Lead CDMO
AstraZeneca

RAJESH PICHAPILLAI
Associate Director, Clinical Programming, AstraZeneca

ANIL BHOJWANI
Associate Director, Medical Writing Asset Lead, GSK

RAVI GAWARE
Head - Diabetes and Obesity, Clinical Research, Novo Nordisk

VINAYKUMAR NANDIMATH
Head Global Clinical Monitoring
Dr. Reddy's Laboratories

PRIYA RAJAGOPAL
Director Study Delivery, Vx GCD, GCO
GSK

RAJESH KUMAWAT
Head- Medical Services & Clinical Development, Himalaya Wellness

ANURADHA NAIR
Head Clinical Quality and Training
Dr. Reddy's Laboratories

SAJJAD PATIL
Therapeutic Area Head - Clinical Operations, Biocon Biologics

DHARMAPAL SHAROFF
General Manager & Head - Global Regulatory Medical Writing, Viatris

PIYUSH AGARWAL
Head Clinical Development
Dr. Reddy's Laboratories

UME ASMA
Global Medical Safety Scientist
Pfizer

AVIK GHOSH
GM & Head Clinical Operations
Sun Pharma

PRITI GUPTA
Centralized Monitoring Lead
GSK

AAKASH DHRUVA
Global Clinical Strategy Lead
Viatris

Plus More Coming Soon...

EXHIBITOR

MEDCLINICA
Accelerating Clinical Trials

CONFERENCE INTRODUCTION

The global clinical trial outsourcing market, valued at **USD 52.39 billion in 2023**, is projected to reach **USD 111.01 billion by 2032**, growing at a **CAGR of 8.7%** from 2024 to 2032. This remarkable growth is fuelled by the globalization of trials, rapid advancements in medical treatments, evolving technologies, and a rising reliance on Contract Research Organizations (CROs) to execute complex, multicentre clinical studies.

In this rapidly transforming landscape, the **Clinical Trials Series 2025** emerges as a timely and essential platform to explore the future of clinical research and innovation. Bringing together leaders from across the clinical trials ecosystem—including sponsors, CROs, regulatory bodies, academic researchers, technology providers, and patient advocates—this summit offers a collaborative space to examine the latest developments, share strategic insights, and drive forward-thinking discussions.

This year's multidisciplinary program will delve into key areas such as:

Clinical Trials – Market Analysis - Opportunities & Challenges
Decentralised Clinical Trials (DCTs) / Patients Prioritised Trials
Work Together - Sponsor / Site / CRO / Patients / Regulators
Clinical Trial Supply - What's the new way forward?
Clinical Trial Design / Clinical Trail Management
Outsourcing / Partnerships / RBQM
Inspection Readiness / Site Management / Risk Monitoring
Technology & Innovation / RWE & RWD
Key Regulatory Changes & Developments
What's the way forward? / Opportunity - Learn & Network

As clinical trials become more complex and globally integrated, now is the time to **embrace innovation, foster cross-sector collaboration**, and shape strategies that ensure **quality, efficiency, and patient-first outcomes**.

We are honoured to welcome you to the **Clinical Trials Series 2025**, and we trust that your participation will contribute significantly to both **industry progress and public health advancement**. Together, let us spark new ideas, build lasting partnerships, and lead the way toward a more resilient and responsive clinical research environment. We wish and pray that all our efforts will be beneficial to our industries and to our country at large.

WHO SHOULD ATTEND AND WHO YOU'LL MEET

CIOs, CEOs, CDOs, Vice Presidents, Presidents, Heads, Directors and Team Leaders from the following areas:

- Clinical Research & Development
- Clinical Research Services
- Clinical Operations
- Clinical Data Management
- Clinical IT
- Clinical Trials
- Medical Affairs
- Regulatory Affairs
- Compliance
- Quality control / Assurance/GCP
- Clinical Study Design
- Safety Surveillance
- Subject Recruitment
- E-Clinical System

WHY SHOULD ATTEND ?

Get more from the event, with a **broader scope bringing the whole communications value chain together.** Enjoy and make the best out of our **dedicated networking time, meet the leading international vendors** showcasing the products of tomorrow in the co-located exhibition. **Expand your knowledge** of the latest business models and strategies in the high-level conference.

08:30 – Coffee & Registration – An opportunity to meet and to network with your conference colleagues.

09:20 – Welcome Address

RBQM

09.30 – "Smarter Trials, Safer Outcomes: Driving Success with Risk-Based Quality Management (RBQM)"

- Understanding the core principles and components of RBQM in clinical trials.
- Identifying critical data and processes to focus quality efforts where they matter most.
- Shifting from reactive to proactive risk detection and mitigation strategies.
- Overcoming challenges in RBQM adoption, including culture, technology, and training gaps.
- Utilizing advanced analytics and AI for real-time risk assessment and decision-making.
- Collaborating effectively with CROs and sites to implement risk-based approaches.
- Measuring the impact of RBQM on trial timelines, cost-efficiency, and quality outcomes

KRISHNENDU SENGUPTA
Director - Centralized Monitoring
AstraZeneca

(DCTs)

10:00 – KEYNOTE PANEL DISCUSSION: Rise of Clinical Trials: Smarter, Decentralized & Patient-Centric

- Learn how DCTs improve patient engagement, flexibility, and cost efficiency.
- Hybrid Trial Approaches – Explore how combining digital and traditional methods broadens reach and data collection.
- Identify essential factors for effective DCT implementation.
- Evaluate the long-term viability and environmental impact of DCTs.
- Platform Trials – Unpack benefits, challenges, and best practices for maximizing impact.
- Personalized Patient Care – Examine how these innovations push healthcare toward more tailored treatments.
- Future Outlook – Are DCTs truly more sustainable in future? Where are we heading?

Moderator

SUBHASH THULUVA
Sr. Vice President & Head - Clinical Development
Biological E

Panelists

KAMALA RAI
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Novartis

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Senior Director, APAC. Global Clinical Trial Operations
MSD

VINAYKUMAR NANDIMATH
Head Global Clinical Monitoring
Dr. Reddy's Laboratories

RAVI GAWARE
Head - Diabetes and Obesity, Clinical Research
Novo Nordisk

ANURADHA NAIR
Head Clinical Quality and Training
Dr. Reddy's Laboratories

10:50 – Morning Coffee/Tea & Discussion

WORKING TOGETHER

11:20 – DISCUSSION WITH EXPERTS: "SPONSOR – SITE – CRO – PATIENT: Building Stronger Clinical Trial Partnerships"

- Evolving sponsor, site, CRO, and patient collaboration to meet rising clinical trial demands.
- Embedding real, measurable transparency across all stakeholder relationships.
- Implementing structural changes sponsors need to build stronger, more agile partnerships.
- Enabling CROs to drive performance through continuous improvement and benchmarking.

- Moving beyond intentions to make transparency a tangible, operational standard.
- Enhancing site preparedness and collaboration for more effective trial execution.
- Applying key learnings from past disruptions to strengthen future clinical trial success.

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PATIENT CENTRICITY

12:10 – From Sponsor-Centric to Patient-Centric: Transforming Recruitment & Engagement

- Leveraging innovative tools to boost patient recruitment and involvement.
- Addressing key barriers and challenges to patient participation.
- Enabling secure data sharing while safeguarding patient privacy.
- Navigating the evolving space with limited regulatory guidance and emerging models.

12:40 – Networking luncheon

REGULATORY

13:30 – DISCUSSION WITH EXPERTS: Outsourcing in Clinical Trials: Crafting Partnerships for Success

- Decentralized Trial Regulations – Adapting to evolving global and regional guidelines.
- AI & Compliance, Leveraging AI in regulatory submissions, safety, and inspections.
- Inspection Readiness – Best practices for FDA and global regulatory preparedness.
- GCP & Ethics – Upholding human rights, data integrity, and ethical standards in the digital era.
- Using real-world evidence to support regulatory outcomes.
- Managing Global Trials – Aligning protocols, GMP/GDP, and documentation across sites.
- Ensuring uniform compliance in multi-country trials. Maintaining consistency across borders.
- India's Regulatory Progress – Where are we heading and where should we be?

Moderator

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Leader, Pharma and Healthcare
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Pfizer

CLINICAL RESEARCH

14:20 - The Evolving Landscape of Clinical Research: Challenges to Opportunities

- Balancing data privacy, global regulations, and the rise of decentralized trials.
- Tackling data complexity while advancing trial innovation and adaptive designs.
- Breaking down barriers to diversity, inclusion, and digital accessibility in participation.
- Navigating high costs, long timelines, and funding pressures impacting trial efficiency.
- Leveraging India's growing potential as a clinical research hub.
- Strengthening vendor partnerships and optimizing patient engagement through tech-forward strategies.

14:50 – Afternoon Tea/Coffee

CHALLENGES & OPPORTUNITIES

15:10 – DISCUSSION WITH EXPERTS: Challenges & Opportunities - Clinical Trials at the Crossroads: Tackling Today, Transforming Tomorrow

- Managing trial delays, recruitment bottlenecks, and high dropout rates.
- Addressing diversity, equity, and inclusion gaps in trial participation.
- Innovative trial designs to save time and cost without compromising on the efficiency
- Ensuring patient-centricity through better engagement and personalized trial designs.
- Building agile, risk-resilient frameworks to future -proof trial operations.
- Navigating trial delays due to supply chain disruptions and geopolitical factors
- Building flexible, risk-based quality management systems

Moderator

VIJAY PARTHASARATHY
Senior Director, Clinical Operations
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Himalaya Wellness

AAKASH DHRUVA
Global Clinical Strategy Lead
Viatris

PROTOCOL DESIGN EXCELLENCE

15.50 - Master Clinical Trial Protocol

ANIL BHOJWANI
Associate Director, Medical Writing Asset Lead
GSK

IMPACT OF TECHNOLOGY

16:20 – DISCUSSION WITH EXPERTS: Revolutionizing Clinical Trials Through Technology - Innovation, Efficiency & Access

- Tech-Powered Trials: Enhancing Speed, Safety, and Patient Engagement
- The Smart Shift: How Technology is Reshaping Clinical Trials?

- Smarter Trials with AI – Discover how artificial intelligence is transforming trial design and control
- Blockchain in Data Management, Understand its role in secure, decentralized data handling
- For Smarter Trials, Smarter Tech – How digital innovation is redefining clinical research?
- Overcoming barriers to tech in trials by cracking digital adoption
- Understanding how (RWE) and (RWD) is transforming clinical trials.
- From Glitches to Gains: Navigating Tech in Trials

Moderator

LATEEFUDDIN SHARIFF
Senior Director - Global Lead CDMO
AstraZeneca

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Associate Director, Clinical Programming
AstraZeneca

PRITI GUPTA
Centralized Monitoring Lead
GSK

17:00 – Closing Remarks & End of conference

FOR SPONSORSHIP OPPORTUNITIES

Sponsorship or exhibition offers an excellent opportunity to network directly with decision makers. Our conferences, featuring world renowned speakers, attract a niche audience from across the globe, providing a unique platform to connect with the right people. This is a cost effective way to achieve targeted outreach, eliminating the need for less efficient advertising methods. To enhance your visibility, we continuously engage industry pioneers through emails, newsletters, and various media channels, ensuring the event and its sponsors receive maximum exposure.

Sponsorship Enquires -
info@virtueinsight.com

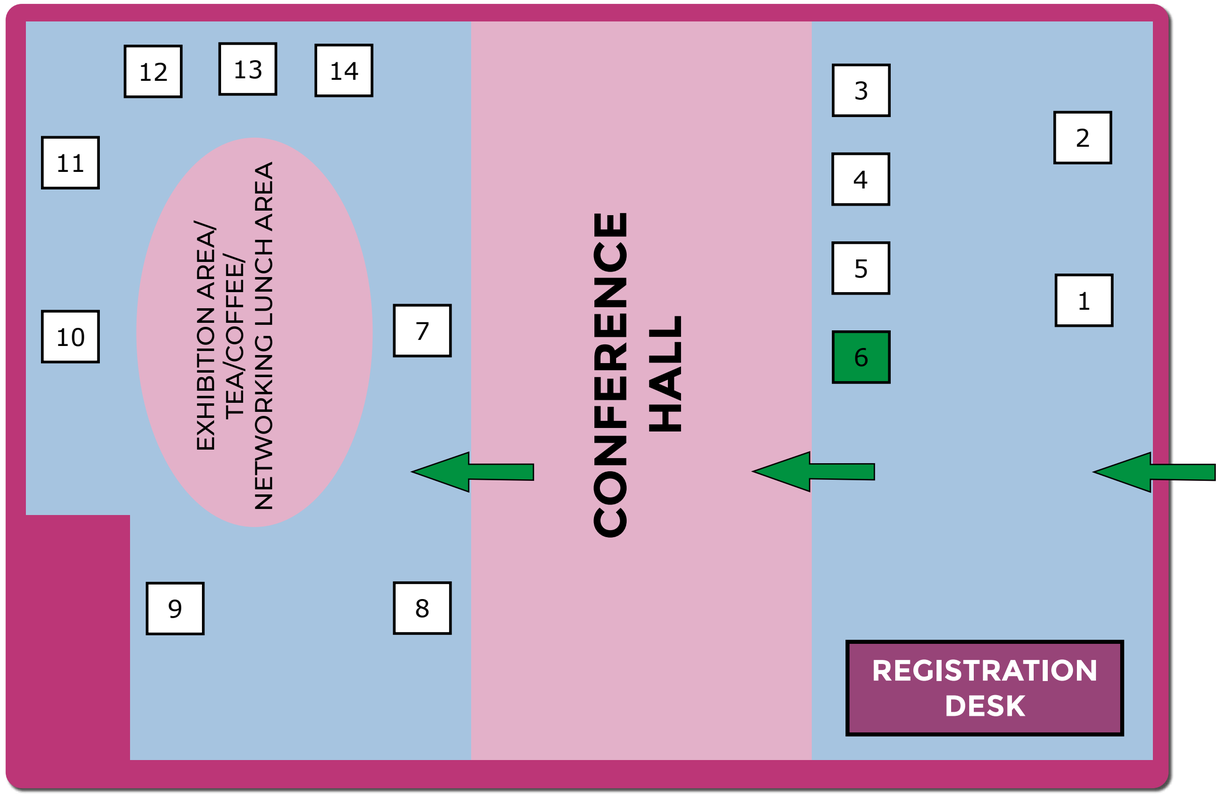
FOR DELEGATE REGISTRATIONS

Our comprehensive conference agenda, featuring the latest insights and world-class leaders as speakers, attracts delegates from across the globe. Our goal is to equip attendees with up-to-date knowledge of industry developments while providing them with valuable networking opportunities with key industry professionals.

Delegate Registration -
bookings@virtueinsight.com

FLOOR PLAN - Book your stalls now before they run out !!!

Sold Blocked Vacant



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Note :- The floorplan is subject to change at the discretion of the organisers.

Clinical Trials Series 2025

REGISTER ONLINE :
Link : https://konfhub.com/clinical-trials-series-2025
For Multiple Bookings - Photocopy this form and send it to info@virtueinsight.com

REGISTRATION FORM

RESERVATION PRICING

Exclusive Deal: 3 delegates for the Price of 2!

SUPER EARLY BIRD PRICE (Valid Until 8th August 2025)

Cost per delegate - Fee: INR 10,000 + GST (18%)

STANDARD PRICE

Cost per delegate - Fee: INR 15,000 + GST (18%)

Please email us at bookings@virtueinsight.com

Registration Form Details:

Forename .....Surename.....

Job Title .....

Company .....

GST No (If Applicable) .....

Official Contact Number.....

Address .....

Country.....Pincode.....

Phone ..... Fax .....

Email .....

I confirm that I have read & agree to the terms and conditions of booking.....(Please Tick) ☐

Signature.....

Methods of Payments:

By Cheque - complete and return the above registration form via post or email, together with your cheque payable to Virtue Insight.

By Bank Transfer:

Account Name	–	Virtue Insight
Account Type	–	Current
Account Number	–	915020031763553
Bank Name	–	Axis Bank
Bank Address	–	2/8 LAMBERT NAGAR, 1st Cross Street, Virugambakkam , Chennai - 600092
Swift Code	–	AXISINBB211
NEFT / IFSC Code	–	UTIB0000211
Micro Code	–	600211010

Queries:

Should you have any questions on bookings,

Please feel free to contact us.

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General Information Venue:

Sterling's Mac Hotel
134, old, HAL Old Airport Rd, opposite to Leela Palace Road,
Kodihalli, Karnataka - 560017
Tel: 91 80 4249 4949

TERMS AND CONDITIONS:

Payment terms: Virtue Insight requires the full amount to be paid before the conference. We may refuse entry to delegates who have not paid their invoice in full.

Cancellations: Delegates and vendors are subject to the following charges and refunds upon withdrawal or cancellation between 2-3 month's prior 75% cancellation fee/ 25% refund. Less than 2 months prior to the event Full cancellation fee / No refund.

Administration Fee: If you cancel your participation (once confirmed) and haven't paid the attendance fee you will be liable to pay an administration fee of £200

Substitutions/Name Change: If you are unable to attend you may nominate, in writing, another delegate to take your place at any time prior to the start of the event. This can be done at no extra cost.

Presentation: If you cannot attend the conference, you can still purchase the presentations (Subject to availability) - Please email to bookings@virtueinsight.com

Indemnity: Virtue Insight reserves the right to make alterations to the conference/executive briefing content, timing, speakers or venue without notice. The event may be postponed or cancelled due to unforeseen events beyond the control of Virtue Insight. If such a situation arises, we will reschedule the event.

Fee: The conference fee includes lunch, refreshments and conference papers provided on the day. This fee does not include travel or hotel accommodation.

Payment Charges: We are a UK registered company and use Barclaycard payment gateway. Some card providers do charge a small foreign transaction fees for international payment (this is not charged from our end). If not sure please contact your card provider before making payment.

Covid Situation, Natural Calamity, National Security: In the case where the government decide to go into a lockdown or restrictions on social and business gatherings during the dates of the conference, the event will be postponed to a new date. Registered clients can choose to join the conference on the new date or decide to take a credit note for their payment so that they can decide to participate for any of our future events within the time frame of next one year.