WHY? SUMMITS

Updated: 20 October, 2024
for the latest programe update, please download agenda on conference website

October 22 - 23
2024

Mumbai
India
Goldfinch Hotel - Mumbai

2024
GLOBAL DRUG SAFETY & PV OUTSOURCING SUMMIT

World's greatest minds in drug safety, pharmacovigilance, data analysis, reporting, government policy, and innovative technology.

Silver Sponsors

ETHICARE Clinical Trial Services Science | Quality | Integrity

COGNIFAI Automate | Innovate | Accelerate

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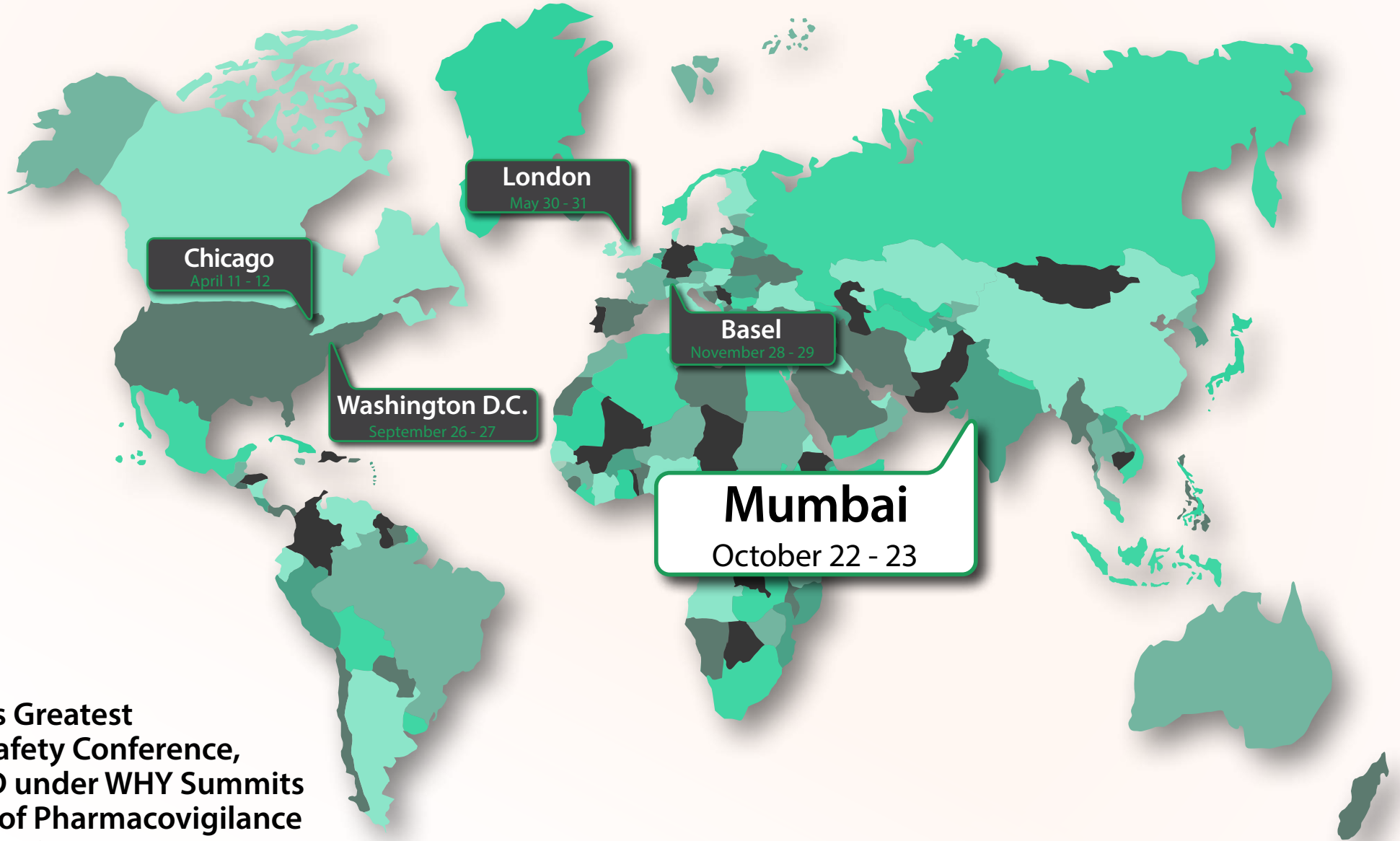
ALWIS

2024 Pharmacovigilance Summits Worldwide



World´s Greatest Drug Safety Conference, UNITED under WHY Summits Future of Pharmacovigilance WORLD TOUR

Why attend:

- ✓ Stay updated on global pharmacovigilance regulations for compliance assurance.
- ✓ Learn about the latest technology implementations in pharmacovigilance.
- ✓ Explore future career prospects in modern pharmacovigilance.
- ✓ Enhance skills through ieractive workshops.
- ✓ Get regulatory updates for compliance assurance.
- ✓ Explore innovative solutions in the technology showcase.
- ✓ Learn from real-world case studies and best practices.

Only in 2023, WhySummits brought together over 2,000 senior pharmaceutical professionals from around the globe, fostering engaging discussions and networking opportunities through our dynamic conferences. Now, we're excited to introduce our newest annual event, dedicated to addressing the increasingly crucial topic of drug safety, as we continue our mission to facilitate meaningful connections and knowledge sharing within the industry.

Main topics:

Updates on Global PV Market Regulation

Strategic Position of CROs & BPOs in Local PV Market

Intertwining Global and Local Collaboration in Expanding PV Market

Latest Technology in PV - A.I. & M.L implementation

Updates on Global Signal Detection & Risk Management Strategies

Future of Communication in safety Services & Career in world of modern pharmacovigilance

With 25 industry leading speakers and a number of carefully selected and dedicated delegates from pharma industry, this conference offers a unique opportunity for meaningful interaction and in-depth discussions.

From thought-provoking keynote presentations, real-world problem-solving case studies to interactive workshops and roundtable discussions, each participant plays a pivotal role in shaping the future of pharmacovigilance and drug safety.

Professionals attending:

- Senior-level decision-makers
- Regulatory experts
- Life-science technology and pharma thought leaders
- Drug safety and PV professionals in original pharma
- Biotechnology firms
- Contract research organizations (CROs)
- Regulatory agencies and academia.
- Senior professionals from generic companies
- Pharmacovigilance Executives
- Regulatory Affairs Professionals
- Quality Assurance Specialists
- Drug Safety Officers
- Clinical Research Leaders
- Healthcare Compliance Experts

"Unlike some other similar events, I found relevance in every session within your conference. The content was neither too simplistic nor too advanced. The participants and presenters provided a diverse view of the issues that are present for most in this industry."

Christine Clearwater, Manager, Safety Operations and Vendor Management, Baxter, Global Patient Safety

"Great dialogue on key issues. Everyone shared truthful insight and did not hold back, even on negative experiences."

Sameer Thapar, Assistant Professor & Advisor, Drug Safety and Pharmacovigilance, Rutgers University

"Loved it. It was very practical and provided valuable insight into practical methods that are actionable for patient safety."

Daniel Naranjo, Global Safety Lead, Global Patient Safety Evaluation, Takeda

Key features

- Cutting-edge Insights from Industry Leaders
- Interactive Workshops for Skills Enhancement
- Regulatory Updates for Compliance Assurance
- Technology Showcase of Innovative Solutions
- Case Studies and Best Practices Sharing
- Roundtable Discussions for Collaborative Problem-solving

Summit in line with 7 (i)s:

- Intelligence – Senior stakeholders with strong expertise in Drug Safety
- Ideas – Exploring new trends
- Interaction – Benchmarking best of breed strategies
- Inspiration – What are new strategies and how to capitalize from them
- Innovation – Driving the effectiveness of PV organization
- Implementation – Moving forward from plans on paper to real impacts
- Improvement – Achieving excellence

Our Sponsors

Ethicare – headquartered in Ahmedabad, India – is a leading full-spectrum PV & CTM service provider. Ethicare provides quality-driven Service to Pharmaceutical, Biotech, Medical Devices and Nutraceutical Companies across the Globe. We provide end-to-end Pharmacovigilance Services in various stages of the product cycle which includes Clinical Trial Safety Management and Post Marketing Safety Management.

Ethicare team has expertise in Medical Information Call Centre, Global Literature Management, ICSR (Adverse Event) Processing & Submissions, Aggregate Reports (DSUR, PADER, PSUR, PBRER, ASR) preparation, Signal Management, Risk Management Plan (RMP), Risk Evaluation Mitigation Strategy (REMS), Safety Management Plan (SMP), Global / Country Specific Pharmacovigilance System Master File (PSMF/ PvMF), QPPV/RPPV/PvOI Service, Compliance Management, Audi/Inspection Readiness & Support, PV System Set-up for MAH. Our tailor-made and cost-effective business models help to achieve maximum client satisfaction along with meeting technical compliance.

CognifAI Solutions Pvt Ltd is a pioneering AI startup dedicated to harnessing the power of artificial intelligence to drive transformative change across various industries. While we specialize in delivering innovative AI solutions for the Pharma and Life Sciences sectors, our expertise extends across multiple domains.

Our mission is to enhance operational efficiency, optimize decision-making, and elevate customer engagement through advanced AI technologies. Some of our key solutions include:

1. Medical Literature Monitoring for Pharmacovigilance: Cutting-edge tools for real-time tracking and analysis of medical literature to ensure drug safety and compliance.
2. CliniOps: An advanced platform for medical document processing, including medical narration, writing, and translation.
3. Image Validation: Precise technology for validating pharmaceutical drug labels, ensuring accuracy and regulatory adherence.

At CognifAI Solutions Pvt Ltd, we are committed to delivering customized AI solutions that address the unique challenges of each industry we serve, driving innovation and excellence across the board.

At **Alwis Group,** we're pioneering AI-driven solutions for the healthcare and pharmaceutical industries. Our innovative tools are transforming pharmacovigilance, enhancing drug safety monitoring, and streamlining regulatory compliance. Our suite of products includes:

- ReTrans Enterprise: AI-powered literature surveillance
- ReTrans Extension: Browser plugin for real-time safety data analysis
- ScoMed: Intelligent medical literature scoring
- ZiNex: Advanced medical document processing
- ZiQuel: AI-driven quality complaint management
- ZiTrack: Smart inbound receipt management

We're committed to improving patient safety and operational efficiency through cutting-edge technology. Our solutions reduce costs, minimize errors, and accelerate processes, allowing healthcare professionals to focus on what matters most – patient care.

Clinevo Technologies is an IT firm that provides regulatory compliant and user-friendly cloud-based Software Solutions for Drug Safety / Pharmacovigilance and Clinical Trials to over 200+ Pharma / CROs.

Clinevo Technologies help companies perform Pharmacovigilance in a highly efficient and cost-effective manner through their Integrated Drug Safety Platform with Database, AS2 gateway, Signal detection, Case Intake, Literature management and SDEA agreements capabilities.

Clinevo Cloud-based software / databases have been validated by several international authorities, including the USFDA, EMA, MHRA, Health Canada, DCGI, and others, and meet regulatory requirements such as HIPAA, ICH, GxP, 21 CFR Part 11, Annex 11, GDPR, and others.

Speakers and Panelists

Dr. Shubhadeep Debabrata Sinha
Senior Vice President and Medical Director
Hetero

Syed Zafeeruddin
Global Pharmacovigilance Manager
Julphar

Rohan Shinde
Assistant General Manager
Jamp pharmaceuticals

Chanbasha Shaik
Global Pharmacovigilance Head
Bridgewest Group

Chaitanya Kulkarni
General Manager Pharmacovigilance
Marksans Pharma Ltd

Dr Mukesh Gori
Director - PS & PV External Engagements
Novartis Pharma

Geeta Narendra Shanbhag
Vice President - Pharmacovigilance & Medico-regulatory Affairs
Ipca Laboratories Ltd.

Dr. Abhijit Surwade
Manager Global Pharmacovigilance
Colgate-Palmolive

Dr. Anuja Jawale
Associate Director – R&D Procurement and Supplier Management
Organon

Dr. Anju Agarwal
Global Director, Global patient safety
ADVANZ PHARMA

Dr Devang Patel
Head of Global Pharmacovigilance
Zydus Lifesciences Limited

Chairman

Gurpreet Singh
Vice President, Managing Director Integrated Safety
IQVIA

Deepak Shankarappa
Director, Pharmacovigilance
Teva Pharmaceuticals Ltd

Vivek Gupta
Associate Director, R&D Supplier Management & Procuremen
Organon

Speakers and Panelists

Venkata Kishore Kumar Darisi
SERM Associate Scientific Director
GSK

Dr Jamal Baig
Director & MCSH- South Asia & IndoChina Region
Sanofi

Dr Arunima Sen
Associate Vice President, Product Safety and Risk Management
Viatris

Joydeep Sengupta
Global Pharmacovigilance - Site Head, Sun Pharma

Dr. Vaibhav Salvi
Director and Head – Clinical Study Unit, India and Southeast Asia, Sanofi R & D – Clinical Sciences and Operations
Sanofi

Sunil Nighot
Manager, Medical Safety Performance & Compliance
Kenvue

Vynie Ann Rao
Manager-DSRM (SOPs Lead and QA)
Lupin Limited

Dr Santanu Kumar Tripathi
Principal of JIMSH & Chief Technical Advisor
JIMSH & IHS

Dr Santosh Hulawale
Senior Manager, PvOI (India), SERM
Glenmark Pharmaceuticals

Dr. Sharad Patel
Manager, Global Pharmacovigilance
Glenmark Pharmaceuticals

Dinesh Wagh
Manager Pharmacovigilance
Precision for Medicine

Dr Ranjit Chavan
Global Head Pharmacovigilance, Clinexel Life Sciences

Day1

8:00AM REGISTRATION OPENING REMARKS & ICEBREAKER

Start the conference on a positive note with warm welcoming remarks from our distinguished hosts. Dive into an engaging icebreaker activity designed to foster connections, spark conversations, and set the tone for a collaborative and productive event.

Get ready to break the ice and kick off our conference with energy and enthusiasm.

9:15AM OFFICIAL WELCOME FROM WHYSUMMITS

9:20AM OFFICIAL WELCOME AND OPENING REMARKS

9:30AM **OPENING KEYNOTE:** PHARMACOVIGILANCE: STRATEGIC DEVELOPMENT, EMERGING OPPORTUNITIES, AND OVERCOMING CHALLENGES

- Evaluation of existing processes, technologies, and compliance measures.
- Emphasis on the integration of automation and artificial intelligence (AI) in PV processes.
- Transformation of vendors and Contract Research Organizations (CROs) into strategic partners.
- Focus on employee training and development to build a skilled and knowledgeable PV team.
- Setting up robust systems and processes to manage the entire lifecycle of PV activities on a global scale.

Chanbasha Shaik, *Global Pharmacovigilance Head,* **Bridgewest Group**

10:00AM COFFEE & NETWORKING

Updates on Global PV Market Regulation

10:30AM **PRESENTATION:** RECENT CHANGES IN PV REGULATIONS OF INDIAN CENTRAL DRUGS STANDARD CONTROL ORGANIZATION (CDSCO)

- Introduction to recent regulatory updates in Indian pharmacovigilance landscape.
- Overview of key changes and amendments in pharmacovigilance regulations.
- Implications of these regulatory changes on pharmaceutical companies operating in India.
- Strategies for ensuring compliance and adapting to the evolving regulatory environment.
- Drawing parallels between Indian regulations and those of prominent regulatory bodies such as the European Medicines Agency (EMA) and the U.S. Food and Drug Administration (FDA).
- Identifying similarities and differences in pharmacovigilance requirements and compliance standards across regions.

Geeta Narendra Shanbhag, *Vice President - Pharmacovigilance & Medico-regulatory Affairs,* **Ipca Laboratories Ltd.**

11:00AM **PANEL DISCUSSION:** ENSURING COMPLIANCE WITH EVOLVING REGULATORY REQUIREMENTS

- Critical importance of adapting to evolving regulatory requirements in pharmacovigilance.
- Exploration of the multifaceted challenges faced by industry stakeholders in maintaining compliance.
- Insights from diverse panelists representing regulatory bodies, pharmaceutical companies, and industry experts.
- Strategies and best practices shared by panelists for effectively navigating and complying with regulatory changes in pharmacovigilance.

Moderator:

Joydeep Sengupta, *Global Pharmacovigilance - Site Head,* **Sun Pharma**

Panelists:

Dr Devang Patel, *Head of Global Pharmacovigilance,* **Zydus Lifesciences Limited**

Gurpreet Singh, *Vice President, Managing Director Integrated Safety,* **IQVIA**

11:30AM **CASE STUDY:** FROM DIVERGENCE TO CONVERGENCE: A PHARMACOVIGILANCE ODYSSEY

- Current Pharmacovigilance Scenario
- Can A Convergence be achieved in Pharmacovigilance regulations
- What are the challenges in achieving convergence
- If convergence, is achieved what are the benefits of the same
- What efforts we can take towards harmonization and convergence

Dr. Anuja Jawale, *Associate Director – R&D Procurement and Supplier Management,* **Organon**

12:00PM NETWORKING LUNCH

Trends and Analytics of Technology Innovation in PV Divisions

1:00PM **PRESENTATION:** PV CURRENT TRENDS, CHALLENGES AND OPPORTUNITIES KEY POINTS

- Increased Focus on Quality, Compliance and Quality Management System Requirements of Audit and Inspection readiness
- Process Enhancements, Changes, Improvements
- Further adoption of Technology and Tools, Database migrations
- Focus on Data Analytics and Trends
- Organisational Culture Enhancement –Focus on People Development, Training and Retention
- Change Management – Mergers / Acquisitions and Integrations

Gurpreet Singh, *Vice President, Managing Director Integrated Safety,* **IQVIA**

Day1

1:30PM **PANEL DISCUSSION:** STRATEGIC PARTNERSHIPS IN OUTSOURCING PHARMACOVIGILANCE ACTIVITIES

- How has the role of outsourcing evolved in pharmacovigilance over the years?
- What role do outsourcing partners play in the integration and implementation of new technologies?
- What are the critical factors for building and maintaining effective partnerships with pharmacovigilance service providers?
- Improving collaboration and transparency: need for strong sponsor- vendor relationship

Moderator:
Gurpreet Singh, *Vice President, Managing Director Integrated Safety,* **IQVIA**
Panelists:
Dr Mukesh Gori, *Director - PS & PV External Engagements,* **Novartis Pharma**
Chanbasha Shaik, *Global Pharmacovigilance Head,* **Bridgewest Group**
Vivek Gupta, *Associate Director, R&D Supplier Management & Procurement,* **Organon**
Dr Arunima Sen, *Associate Vice President, Product Safety and Risk Management,* **Viatris**

2:00PM **CASE STUDY:** CASE STUDY: SUCCESSFUL OUTSOURCING MODELS IN INDIAN PHARMACOVIGILANCE

- Exclusive case study presentation showcasing successful outsourcing models in Indian pharmacovigilance, tailored for potential sponsors or solution providers.
- Detailed examination of proven strategies and best practices adopted by pharmaceutical companies in outsourcing pharmacovigilance activities.
- Showcase of real-world examples and case studies highlighting the effectiveness and benefits of different outsourcing models.
- Opportunities presented for sponsors or solution providers to leverage their expertise and solutions in supporting successful outsourcing initiatives in the Indian pharmacovigilance landscape.

Chaitanya Kulkarni, *General Manager Pharmacovigilance,* **Marksans Pharma Ltd**

2:30PM **MATCHMAKING SESSION AND NETWORKING BREAK:** "CONNECTING FOR SAFER MEDICINES: LET'S DISCUSS TOGETHER!"

- Join fellow professionals for interactive discussions, sharing experiences, and exploring opportunities to enhance drug safety together.
- Discover potential collaborators, whether you're seeking partners for innovative projects, regulatory guidance, or technological solutions to strengthen drug safety practices.

Intertwining Global and Local Collaboration in Expanding PV Market

3:00PM **PRESENTATION:** BUILDING BRIDGES: ENHANCING GLOBAL AND LOCAL COLLABORATION IN PHARMACOVIGILANCE

- Enhancing collaboration between global and local stakeholders in pharmacovigilance.
- Examination of the importance of fostering collaboration across borders for effective pharmacovigilance practices.
- Adapting to Global Budget Cuts: Discuss strategies for maintaining effective collaboration despite global budget constraints.
- Discussion on the mutual benefits, challenges, and best practices associated with bridging global and local efforts in pharmacovigilance.
- Ways to leverage emerging opportunities for collaboration, including technology and partnerships.

Dr. Anju Agarwal, *Global Director,* **Global patient safety at ADVANZ PHARMA**

3:30PM **PANEL DISCUSSION:** END-TO-END PHARMACOVIGILANCE SERVICES AND SETTING UP THE PV ORGANIZATION TO A NEW MORE EFFECTIVE MODEL

- Advantages and challenges of globalizing pharmacovigilance operations versus maintaining localized approaches.
- Diverse perspectives shared by panelists representing global and local pharmaceutical companies, regulatory bodies, and industry experts providing diverse viewpoints on the globalization-localization spectrum.
- Insights into end-to-end Pharmacovigilance services and setting up the PV Organization to a new more effective model

Moderator:
Dr. Abhijit Surwade, *Manager Global Pharmacovigilance,* **Colgate-Palmolive**
Panelists:
Venkata Kishore Kumar Darisi, *SERM Associate Scientific Director,* **GSK**
Dr. Shubhadeep Debabrata Sinha, *Senior Vice President and Medical Director,* **Hetero**
Rohan Shinde, *Assistant General Manager,* **Jamp pharmaceuticals**

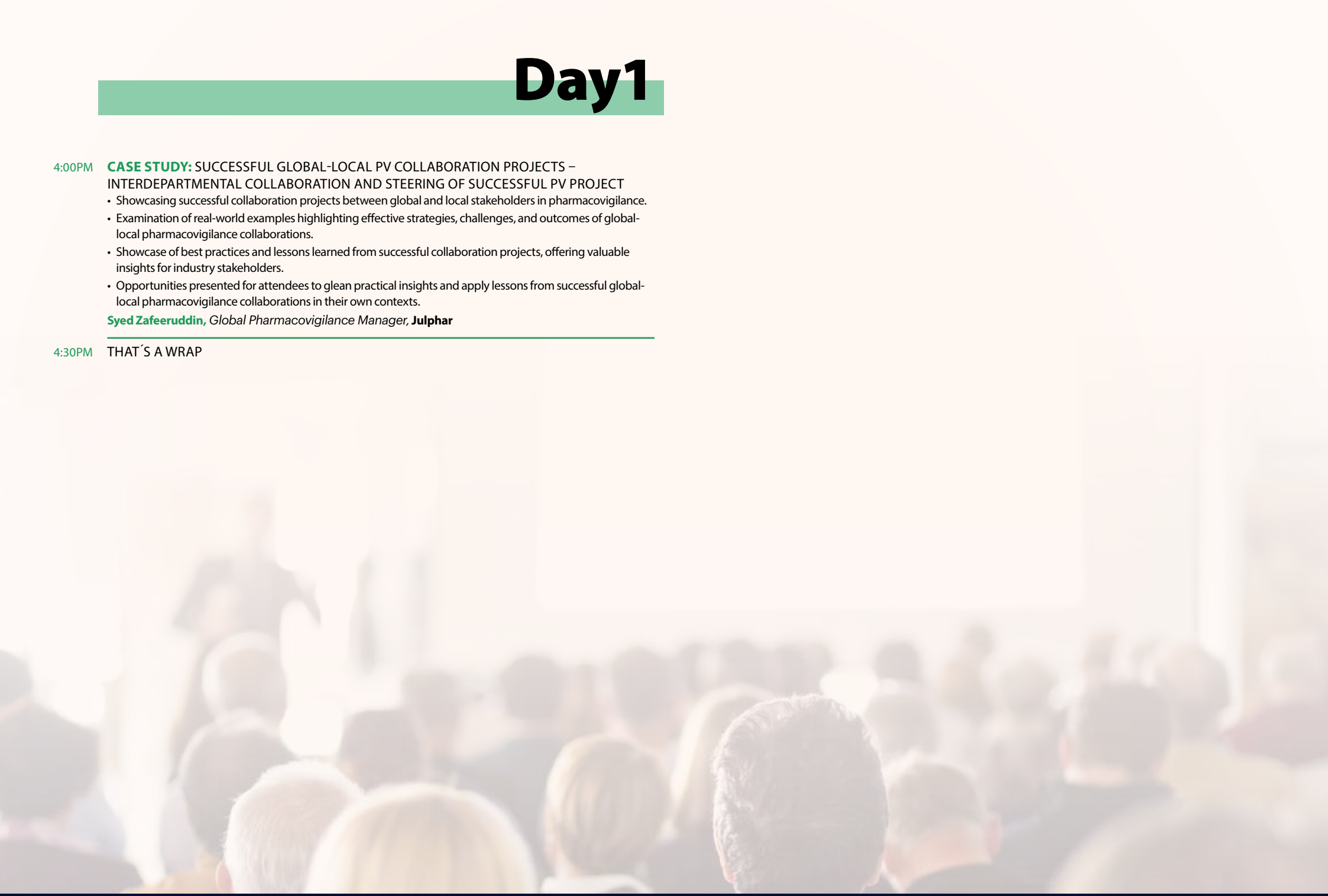
Day1

4:00PM **CASE STUDY:** SUCCESSFUL GLOBAL-LOCAL PV COLLABORATION PROJECTS – INTERDEPARTMENTAL COLLABORATION AND STEERING OF SUCCESSFUL PV PROJECT

- Showcasing successful collaboration projects between global and local stakeholders in pharmacovigilance.
- Examination of real-world examples highlighting effective strategies, challenges, and outcomes of global-local pharmacovigilance collaborations.
- Showcase of best practices and lessons learned from successful collaboration projects, offering valuable insights for industry stakeholders.
- Opportunities presented for attendees to glean practical insights and apply lessons from successful global-local pharmacovigilance collaborations in their own contexts.

Syed Zafeeruddin, *Global Pharmacovigilance Manager,* **Julphar**

4:30PM THAT´S A WRAP



Day2

Latest Technology in PV - A.I. & M.L implementation

9:00AM **KEYNOTE:** PIONEERING PHARMACOVIGILANCE SOLUTIONS FOR NICHE THERAPIES: FOCUS ON BIOSIMILARS

- Principles of Pharmacovigilance in Biosimilars – Variability, Traceability, Immunogenicity, Regulatory
- Operational Requirements for Pharmacovigilance in Biosimilars
- Case processing, Aggregate safety and Risk Management
- Effective Management of Global PV system in Biosimilars BU
- PV integration in Biosimilar Strategy, Development and Marketing

Dr. Shubhadeep Debabrata Sinha, *Senior Vice President and Medical Director,* **Hetero**

9:30AM **PANEL DISCUSSION:** ADDRESSING INCONSISTENCIES IN DATA REPORTING AND LEVERAGING TECHNOLOGY SOLUTIONS, HOW TECHNOLOGY CAN HELP ?

- Identifying and resolving inconsistencies in data reporting, leveraging diverse expertise for comprehensive solutions.
- Discuss strategies for fostering collaboration between stakeholders and vendors to effectively address pharmacovigilance technology challenges.
- Big Data: approaches and tools for handling large volumes of data in pharmacovigilance, ensuring efficient processing and analysis.
- Best practices for identifying and resolving inconsistencies in data reporting, including data standardization, quality control measures, and cross-functional collaboration.

Moderator:
Dr Mukesh Gori, *Director - PS & PV External Engagements,* **Novartis Pharma**
Panelists:
Chaitanya Kulkarni, *General Manager Pharmacovigilance,* **Marksans Pharma Ltd**
Dr. Vaibhav Salvi, *Director and Head – Clinical Study Unit, India and Southeast Asia, Sanofi R & D – Clinical Sciences and Operations,* **Sanofi**

10:00AM **ENHANCING PATIENT SAFETY THROUGH A LIFE CYCLE APPROACH**

- Life Cycle Approach to Drug Monitoring: Emphasize the importance of patient safety throughout the drug monitoring process, focusing on continuous oversight from drug development to post-market surveillance.
- Ethical Aspects and Adverse Event Detection: Discuss the combination of ethical patient care, timely detection of adverse events, and the role of accurate and quality reporting in ensuring better patient outcomes.

Dr Santanu Kumar Tripathi, *Principal of JIMSH & Chief Technical Advisor,* **JIMSH & IHS**

10:30AM COFFEE & NETWORKING

Updates on Global Signal Detection & Risk Management Strategies

11:00AM **KEYNOTE:** ADVANCES IN SIGNAL DETECTION AND RISK MANAGEMENT

- In-depth presentation unveiling the latest advancements in global signal detection within pharmacovigilance.
- Exploration of cutting-edge methodologies, technologies, and approaches enhancing signal detection on a global scale.
- Insightful analysis of emerging trends, challenges, and opportunities shaping the future of signal detection in pharmacovigilance.
- Practical insights and strategies shared to empower attendees with the knowledge and tools needed to optimize signal detection in their organizations.

Dinesh Wagh, *Manager Pharmacovigilance,* **Precision for Medicine**

11:30AM **PANEL DISCUSSION:** ADDRESSING CHALLENGES IN SIGNAL DETECTION AND RISK MANAGEMENT

- Addressing the multifaceted challenges encountered in signal detection and risk management within pharmacovigilance.
- Comprehensive exploration of the key challenges faced by industry stakeholders, including regulatory complexities, data integration issues, and emerging safety concerns.
- Diverse perspectives shared by panelists representing regulatory agencies, pharmaceutical companies, academia, and technology providers.
- Collaborative examination of innovative strategies, best practices, and collaborative approaches to overcome challenges and enhance signal detection and risk management effectiveness in pharmacovigilance.

Moderator:
Deepak Shankarappa, *Director, Pharmacovigilance,* **Teva Pharmaceuticals Ltd**
Panelists:
Venkata Kishore Kumar Darisi, *SERM Associate Scientific Director,* **GSK**
Dr Santanu Kumar Tripathi, *Principal of JIMSH & Chief Technical Advisor,* **JIMSH & IHS**
Dr Santosh Hulawale, *Senior Manager, PvOI (India), SERM,* **Glenmark Pharmaceuticals**
Dr Jamal Baig, *Director & MCSH- South Asia & IndoChina Region,* **Sanofi**
Gurpreet Singh, *Vice President, Managing Director Integrated Safety,* **IQVIA**

12:30PM LUNCH BREAK

Day2

Future of Operational effectiveness and Risk Management

1:30PM **OPERATIONAL EFFECTIVENESS WITHIN END-TO-END PV SERVICES**

- Managing end to end global Pharmacovigilance operations
- PV/SDEA Agreements: Lifecycle management of PV Agreements
- People Management and Effective Recruitment and Training
- Innovation in communication with local PV personnel to fulfill the local regulatory requirements for approved products, social media monitoring, reconciliation and regulatory intelligence updates.

Dr. Sharad Patel, *Manager, Global Pharmacovigilance,* **Glenmark Pharmaceuticals**

2:00PM **PANEL DISCUSSION:** CHANGE MANAGEMENT SYSTEM IN PV AND DEVELOPMENT OPPORTUNITIES IN PHARMACOVIGILANCE

- Dynamic panel discussion delving into Change Management System and development opportunities within the field of pharmacovigilance.
- Exploration of diverse Change Management trajectories, including roles in drug safety, regulatory affairs, medical affairs, and data analysis.
- Insights into the evolving skill sets, competencies, and qualifications sought after by employers in the pharmacovigilance industry.
- Interactive dialogue among panelists representing academia, industry, and recruitment agencies, offering valuable guidance and advice for aspiring professionals seeking to advance their careers in pharmacovigilance.

Moderator:

Dr Jamal Baig, *Director & MCSH- South Asia & IndoChina Region,* **Sanofi**

Panelist:

Syed Zafeeruddin, *Global Pharmacovigilance Manager,* **Julphar**

Dr. Vaibhav Salvi, *Director and Head – Clinical Study Unit, India and Southeast Asia, Sanofi R & D – Clinical Sciences and Operations,* **Sanofi**

Sunil Nighot, *Manager, Medical Safety Performance & Compliance,* **Kenvue**

2:30PM **PRESENTATION:** FUTURE OF PHARMACOVIGILANCE: NAVIGATING OPERATIONAL RISKS AND DRIVING CONTINUOUS IMPROVEMENT

- Discovering key insights on Operational risks in PV and the need for continuous improvement to navigate these challenges that can impact compliance and patient safety.
- Exploring strategies for mitigating risks through process optimization and proactive monitoring.
- Learning about the importance of continuous improvement in adapting to evolving regulatory requirements and industry standards.

Vynieann Rao, *Manager-DSRM (SOPs Lead and QA),* **Lupin Limited**

3:00PM COFFEE & NETWORKING

3:30PM **PRESENTATION:** ROLE OF CROS & BPOS IN SHAPING THE INDIAN PHARMACOVIGILANCE LANDSCAPE

- Exclusive presentation highlighting the pivotal role of Contract Research Organizations (CROs) and Business Process Outsourcing (BPOs) in shaping pharmacovigilance in India.
- In-depth exploration of how CROs and BPOs contribute to the efficiency, innovation, and compliance of pharmacovigilance processes.
- Showcase of case studies and success stories illustrating the significant contributions of CROs and BPOs in driving quality outcomes and regulatory adherence.
- Insights into potential partnership opportunities for solution providers to collaborate with CROs and BPOs, delivering value-added pharmacovigilance solutions.

Dr Ranjit Chavan, *Global Head Pharmacovigilance,* **Clinexel Life Sciences**

4:00PM **ROUNDTABLE DISCUSSION:** CHARTERING THE COURSE FORWARD: ADVANCING PV INNOVATION

- Come together to reflect on the insights gained throughout the conference
- Chart a course forward for advancing Change Management System in PV
- Engage in collaborative problem-solving and share actionable strategies for fostering career growth and development in the ever-evolving field of pharmacovigilance.

Roundtable leaders:

Gurpreet Singh, *Vice President, Managing Director Integrated Safety,* **IQVIA**

Syed Zafeeruddin, *Global Pharmacovigilance Manager,* **Julphar**

5:00PM THAT´S A WRAP!

2024 World tour at a glance

1. April 11 - 12, 2024 Chicago, IL, United States
   American Medical Device Safety & Compliance Conference
2. May 30 - 31 London, United Kingdom
   9th Global Drug Safety & RWE Forum
3. September 26 - 27 Washington D.C., United States
   American Drug Safety Summit
4. October 22 - 23 Mumbai, India
   Global Drug Safety & PV Outsourcing Summit
5. November 28 - 29 Basel, Switzerland
   World Drug Safety Summit Basel

Sponsorship

Additional sponsorship opportunities are available for those who wish to further customize their involvement.

Exhibiting

With a large and senior audience and decision makers, thoroughly selected, exhibiting at any Summit at 2024 FoP SUMMIT WORLD is a popular sponsorship option with great value for solution providers.

Sponsorship includes

- Selected Summit Three Access Passes
- Exhibition space
- Helping to prearrange face to face meetings with selected participants

Dinner Sponsorship

2024 FoP SUMMIT WORLD TOUR will host a series of dinners These dinners bring together thoughtfully selected groups of 15-20 peers from established pharma, biotech, healthcare, and medtech companies. The dines start with a 30-minute networking reception followed by a 60-minute seated dinner, with the option for participants to remain afterward to continue networking.

- Selected Summit Three Access Passes
- 30-minute reception, and 60 minute seated dinner

Speaking

Limited speaking opportunities are available for our sponsoring partners to demonstrate the expertise of their organization. Be sure to ask about these early so we can ensure your presentation flows seamlessly with the overall content. Speaking sponsorships has several options – keynote presentations, case study presentations, expert presentations, panel discussions, workshops, or roundtable leadership. Speaking opportunities are available for experts in the field of Drug safety specialists, QPPVs, Safety Heads, C-level pharmaceutical and biotech executives, hospital management, clinicians, epidemiologists, pharmacologists, Project and Portfolio Management, Contract Management, Consultancy, CROs, Data Management, Artificial Intelligence, Robotics and Digital Innovation experts

Contact us

Updated: 20 October, 2024
for the latest programe update, please download agenda on conference website

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