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DIA 2014

9th Annual India Conference

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The Future of Indian Healthcare: Patients, Access and Innovation

16-18 October
Palladium Hotel
Mumbai, India
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All attendees will receive
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KEYNOTE SPEAKERS



DAY 1 | October 16

M K Bhan

Former Secretary
Department of Biotechnology (DBT)
Govt. of India



DAY 2 | October 17

G N Singh

Drug Controller
General of India
(DCGI)



DAY 3 | October 18

Kiran Mazumdar Shaw

Chairperson and
Managing Director
Biocon

PROGRAM CO-CHAIRS



Padmashree Prof. Ranjit Roy Chaudhury
Chairman, Task Force for Research
Apollo Hospitals Group



Alexandra Pearce
Senior Vice President and Head
Global Regulatory Affairs
Glenmark Pharmaceuticals

PROGRAM COMMITTEE



Abby Abraham
Vice President & Co-Founder,
Clinical Solutions,
Algorithm Inc.



Vivek Ahuja
Director
Research & Development
PATH India



Alberto Grignolo
Corporate Vice President
PAREXEL, United States



Madhur Gupta
Technical Officer
Pharmaceuticals
WHO Country Office for India



Jaideep Kaneria
AVP Lifesciences
TCS Services



Sowmyanarayan Srinivasan
Head, R&D Solutions and Consulting
Cognizant Technology Solutions



Arun Mishra
Director and Head, Regulatory Affairs
Abbott India Ltd.



Goutam Pujari
Vice President
Head Global Clinical
Research Operations
Glenmark Pharmaceuticals



Santosh Savarkar
Associate Vice President
Regulatory Affairs
Alembic

PROGRAM OVERVIEW

Good health is a fundamental human right. But despite rapid improvements in the recent past, there are unprecedented healthcare challenges in India today that demand immediate attention and action. The most critical aspect among these is ensuring safety, efficacy, quality, affordability and accessibility of medicines and treatment to patients at all times. The key imperatives to overcome these continual challenges and catalyze the transformation include putting patients first, fostering innovation, and enhancing access. The **DIA 2014 9th Annual India Conference** will bring together leading local and global experts in healthcare, pharmaceutical industry members, academia, researchers, regulators, payers and patients under one roof to deliberate on the emerging trends, opportunities and challenges.

PROGRAM OBJECTIVES

- Focusing on 'Patient First' approach and various patient centric paradigms covering innovation and access of Indian healthcare
- Hear the views of leading experts from the academia, government, industry, researchers, patient representatives and regulators on current local and global trends and best practices
- Bringing all healthcare stakeholders together to discuss current issues/topics
- Covering important issues/topics such as, patient health and safety, pharmacovigilance, clinical trial, quality and manufacturing, health economics, healthcare delivery, public-private partnership in innovation and healthcare access, case studies on Indian innovation and **many more**

WHO SHOULD ATTEND

- Drug development and clinical research managers and associates
- Pharmaceutical physicians and medical directors
- Drug safety and drug surveillance personnel
- Professionals engaged in discovery research
- Clinical pharmacology scientists
- Pharmacologists
- Regulatory affairs managers
- Quality professionals
- Academic scientists
- Biostatisticians
- Data managers
- Medical writers
- Outsourcing and marketing managers
- IT professionals
- Patient Engagement Stakeholders

Support / Exhibiting Companies as of 2 September 2014



DIA Global Center
21 Dupont Circle NW, Suite 300
Washington, DC 20036

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DAY 1 | THURSDAY, OCTOBER 16**8:30-10:00 AM** REGISTRATION**10:00-10:30 AM** OPENING CEREMONY**10:30-11:00 AM** KEYNOTE PRESENTATION**M K Bhan**

Former Secretary
Department of Biotechnology, (DBT)
Government of India

11:00 AM-12:00 PM PLENARY SESSION 1**Patients**

The opening session sets forth the tone and content of the prevailing thinking and practice in the clinical research and patient care space which is undergoing a paradigm shift. The patients' voice and perspective is coming to the fore as a definite patient centric approach. The topics will help provide holistic insight to the current issues and orient future thinking.

SESSION CHAIR

**Vikas Sharma**

Senior Director
Medical Ethics & Research Office of Chief Medical and
Scientific Officer
Quintiles

PANELISTS

**Urmila Thatte**

Professor and Head
Seth G S Medical College and
KEM Hospital

**Satish Chandra**

Senior Specialist and Head Clinical Research
TAWAM Hospital (associate of John Hopkins)
UAE

**Y K Gupta**

Professor and Head
Department of Pharmacology
AIIMS

**Pooja Sharma**

Leader (Hon)
CANKIDS Patient Advocacy Group and
Senior Scientist, Clinical Research
Medanta the Medicity

**Harl Menon**

Professor
Department of Medical Oncology
Tata Memorial Hospital

12:00-13:30 LUNCH AND EXHIBIT VISIT**12:30-13:30** WHITE PAPER PRESENTATION - TCS**JOIN A DIA COMMUNITY!**

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Communities in India:

Clinical Data Management

Clinical Safety and Pharmacovigilance

Global Outsourcing

Medical Writing

Regulatory Affairs

CONCURRENT SESSIONS

13:30-15:00 PM

SESSION 1A

Patient Health Safety and Compliance for Prescription Medicine

This session will address issues surrounding the use of medicines by patients in India, including compliance with prescribed medicines and how to ensure the safe use of medicines.

SESSION CHAIR

Padmashree Prof. Ranjit Roy Chaudhury

Chairman, Task Force

Apollo Hospitals Educational and Research Foundation (AHERF)

Regulatory Initiatives to Enhance Compliance

V G Somani

Joint Drug Controller

CDSO

Preventing Errors in Using Medicines

Pramil Tiwari

Head and Professor

Department of Pharmacy Practice

NIPER

Increasing Medication Compliance in Patients

Prem Pais

Professor of Medicine

Head of Division of Clinical Research Training

St. John's Institute

13:30-15:00 PM

SESSION 1B

Pharmacovigilance in India

The speakers in this session will describe the status of pharmacovigilance regulation and practices in India, with a focus on monitoring the safe use of markets drugs and ways to take prompt remedial action when and if necessary; Clinical trial safety regulations to protect safety of clinical trial participant in India, recent changes in regulations, pitfalls and global perspective; PV outsourcing industry in India contributing to patient safety globally, also increasing PV expertise and awareness in the country.

SESSION CHAIR

Ajoy Roy

Senior Medical Director and Head Medical Affairs

PAREXEL, India

Postmarketing Pharmacovigilance Regulations and Practices in India

Y K Gupta

Professor and Head

Dept. of Pharmacology

AIIMS

A SWOT Analysis of Growing Pharmacovigilance Industry in India

Mangesh Kulkarni

Head Safety Operations

Drug Safety & Epidemiology

Novartis Healthcare Pvt. Ltd.

15:00-15:30

TEA / COFFEE BREAK & EXHIBIT VISIT

CONCURRENT SESSIONS

15:30-17:00 PM

SESSION 2A

Status of Patient Advocacy Groups in India—Panel Discussion

This session will provide current information on the status and influence of patient advocacy groups in India, their goals and aspirations, their success and ongoing concerns, and the issues that remain to be addressed to protect patient safety and promote patient health.

SESSION CHAIR

Mira Shiva

Founder Member & Advisory Committee Member -

People's Health Movement,

Former Chairperson - Health Action International Asia-Pacific

Y K Sapru

Chairman & CEO

Cancer Patients Aid Association India

Vikas Ahuja

President

The Delhi Network of Positive People, India

Blessina Kumar

Health Activist & Public Health Consultant

Chair, Global Coalition of TB Activists

15:30-17:00 PM

SESSION 2B

Patient Participation in Clinical Trials

This session will focus on the important ethical issues and practical logistics (for example, travel from rural areas) affecting the participation of patients in clinical trials in India, the stricter applicable regulations (for example, videotaped informed consents), the protection of the rights on trial participants and of the validity of the data, and the oversight of clinical investigators.

SESSION CHAIR

Barbara E. Bierer

Professor of Medicine, Harvard Medical School

Brigham and Women's Hospital Faculty Co-Director

Multi-Regional Clinical Trials Center, Harvard University

Participant Understanding and Informed Consent

Kimberly Hensle Lowrance

Managing Director

Public Responsibility in Medicine and Research (PRIM&R)

Return of Summary Results to Participants:

Regulation, Content and Process

Barbara E. Bierer

Professor of Medicine, Harvard Medical School

Brigham and Women's Hospital Faculty Co-Director

Multi-Regional Clinical Trials Center, Harvard University

International Clinical Trials Data Sharing: Principles and Mandate

Mark Barnes

Partner, Ropes & Gray LLP, Lecturer on Law

Harvard Law School, Faculty Co-Director

Multi-Regional Clinical Trials Center, Harvard University

17:00 PM

MEETING ADJOURNED

DAY 2 | FRIDAY, OCTOBER 17

10:30-11:00 AM KEYNOTE PRESENTATION

G N Singh

Drug Controller General of India

11:00 AM-12:30 PM PLENARY SESSION 2

Access

The plenary session intends to generate directional thought leadership to arrive at a unique and integrated healthcare model. The session will help to analyse and suggest future model which leverages principle of accessibility, affordability and accountability by adopting a patient-centric approach towards delivery of quality healthcare to the masses. The session will also invite opinions from the panel members and participants to understand possible impact and adoption of Health Care Economic Outcomes Research (Pharmacoeconomics), Health Insurance, Public Private Partnerships, Diagnostics & Pharma Industry – covering the Healthcare Value Chain.

SESSION CHAIR



Nata Menabde

WHO Representative to India

PANELISTS



Swati Pirmal

Vice Chairman
Piramal Enterprise Ltd.



N K Ganguly

Former Director General
Indian Council of Medical Research



Devi Shetty

Chairman
Narayana Hrudayalaya Group of Hospitals



Rajesh Balkrishnan

Associate Professor Health Outcomes
University of Michigan



Ferzaan Engineer

Co-Founder and Chairman
Cytespace

12:30-13:30 PM LUNCH AND EXHIBIT VISIT



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- ☑ **Global Forum** - your member digital newsmagazine featuring practical tips, regulatory and global updates. (6 issues)
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- ☑ **Communities** - DIA's Communities—discipline specific groups that network regionally and globally, share best practices, knowledge resources and the latest information related to your field—all online and available 24-7. Current Regional India Communities Include:
 - Clinical Safety and Pharmacology | Clinical Data Management
 - Global Outsourcing | Medical Writing | Regulatory Affairs
 - Risk-based Monitoring Scientific Working Group (SWG)
- ☑ **DIA Online Member Directory** - Network locally and globally via DIA ConneX—your platform for sharing documents and discussions, connecting with colleagues and developing your own network of colleagues and industry thought leaders.
- ☑ **Knowledge resources at www.diahome.org** featuring videos, podcasts, presentation downloads, publication archives, and robust searching and sorting tools to get the information you need, when you need it.

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- ☑ **DIA Daily** - This daily newsletter features summaries of breaking news in the pharmaceutical, biotechnology, and medical device fields gathered from thousands of global news sources.
- ☑ **SmartBrief** - Two times per week, receive the latest news and information related to diagnostic and therapeutic product innovations.
- ☑ **Career Center** - Search for employment opportunities and submit your resume, all confidentially and online.

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CONCURRENT SESSIONS

13:30-15:00 PM

SESSION 3A

Universal Health Coverage and Health Insurance

The goal of this session is to take stock of the current state and future prospects for universal health coverage and health insurance for patients in India, with particular reference to patient access to registered safe and effective medicines.

SESSION CHAIR

Indrani Gupta

Professor & Head, Health Policy Research Unit
Institute of Economic Growth, India

Access to Medicines and Health Technologies in the Context of Universal Healthcare Coverage in India: Opportunities and Challenges

Madhur Gupta

Technical Officer, Pharmaceuticals
WHO Country Office for India

Universal Health Coverage: Concerns and Options

Indrani Gupta

Professor & Head, Health Policy Research Unit
Institute of Economic Growth, India

Universal Health Coverage - Scope for Policy and Systems Reform

Priya Balasubramaniam

Senior Public Health Specialist
Public Health Foundation of India

13:30-15:00 PM

SESSION 3B

Medicine Pricing and Reimbursement

Speakers in this session will illustrate the various existing public and private mechanisms for the pricing and reimbursement of medicines in India, with a focus on how patients at all income levels can have access to the medicines they need, on what obstacles may exist to such access, and on possible ways to remove such obstacles.

SESSION CHAIR

Manu Sehgal

Vice President
PAREXEL, India

Insight into India's Pharmaceutical Pricing Policy: Will the Revised Policy Improve Access to Medicines?

Anita Kotwani

Associate Professor, Department of Pharmacology
Vallabhai Patel Chest Institute
University of Delhi, India

Medicine Reimbursement in India Present and Future

Anuj Kapoor

Senior Manager Operations
Health Economics & Outcome Research Practice
Optum Global Solutions

India's medicine reimbursement and market access policy in light of experience from Europe and Asia

Manu Sehgal

Vice President
PAREXEL, India

15:00-15:30 PM

TEA / COFFEE BREAK AND EXHIBIT VISIT

CONCURRENT SESSIONS

15:30-17:00 PM

SESSION 4A

Good Manufacturing Practices—Panel Discussion

Pharmaceutical GMP in modern times is not just an assessment through testing the finished product but is an ongoing discipline that one has to imbibe from product development to product distributed to the patients. GMP poses a significant challenges in terms of bridging - What Agency Wants and What Industry Understands. The GMP session will open with a brief presentation by USFDA Regulator and would lead into a panel discussion covering aspects of GMP Challenges for Pharmaceutical/Healthcare industry and the path to be traversed moving forward.

PRESENTATION

Thomas Arista

Investigator/National Expert-Biotech
FDA

SESSION CHAIR

Ganadhish Kamat

Executive Vice President, Quality
Lupin

MODERATOR

Vikas Dandekar

India Bureau Chief
PharmAsia News

Ajaz Hussain

President & CEO
Insight Advice & Solutions

Nick Cappuccino

Chief Technical Officer & Scientific Advisor
Dr. Reddy's Laboratories

15:30-17:00 PM

SESSION 4B

Compulsory Licensing—Increasing Tool for Easy Access of Medicines—Panel Discussion

This session will examine and debate the important and delicate issue of compulsory licensing of medicines and both its intended (affordable access to medicines) and unintended (obstacle to the entry of innovation into India) consequences.

SESSION CHAIR

Aliasgar Dholkawala

Senior Associate, Legal Services
Wadia & Gandhi & Co.

Milind Sathe

Deputy General Manager, IPM & Projects
Unichem

Alka Mehta

Head IP
Cipla

Tabrez Ahmad

Secretary General
Organization for Pharmaceutical Products of India (OPPI)

Smita Karve

Professor of Law
Jitendra Chauhan College of Law

17:00 PM

MEETING ADJOURNED

DAY 3 | SATURDAY, OCTOBER 18

10:00-10:30 AM KEYNOTE PRESENTATION

Kiran Shaw Mazumdar

Chairperson and Managing Director
BIOCON

10:30-11:00 AM TEA / COFFEE BREAK
AND EXHIBIT VISIT

11:00-12:00 noon PLENARY SESSION 3

Innovations

A country must nurture a multidisciplinary ecosystem to conceive, support and reward innovation for its people. Increasingly innovation is seen as the currency of the century. Knowledge based economies are now depending on its ability to generate new ideas, processes and solutions, and converting into social good and economic growth. Innovations could be a way forward delivering affordable and quality healthcare to India's billion-plus people. The session will explore challenges and opportunities for India centric healthcare innovation including drug discovery and development.

SESSION CHAIR

Arun Mishra

Director and Head, Regulatory Affairs
Abbott India Ltd.

PANELISTS



Padmashree Prof. Ranjit Roy Chaudhury

Chairman, Task Force
Apollo Hospitals Educational and Research Foundation
(AHERF)



Mohan D. Nair

Consultant to Pharmaceutical Industry



Romi Singh

Executive Director
Amgen



Manoj Malhotra

CEO
Pyramid Healthcare
Salient Business Solutions, Avantha Group

12:00 noon-13:30 PM

SESSION 5

Innovation in Drug Discovery and Development

Speakers in this session will illustrate the current status of pharmaceutical innovation in India - from scientific discovery to the development of innovative drugs that change the practice of medicine. Can Indian R&D improve its competitiveness on world markets?

SESSION CHAIR

Neelima Khairatkar Joshi

Senior Vice President and Head
NCE Drug Discovery & Research
Glenmark

The Need for New Innovative Approaches to New Drug Discovery

Mohan D. Nair

Consultant to Pharmaceutical Industry

Clinical Trials in 2020

Rajesh Jain

Director, Life Sciences Business Process Services
Cognizant Technology Solutions India Pvt. Ltd.

From Innovation to Cure: India Perspective on Today & Tomorrow...

Neelima Khairatkar Joshi

Senior VP and Head,
NCE Drug Discovery & Research
Glenmark

13:30-14:30 PM

LUNCH AND EXHIBIT VISIT

14:30-16:30 PM

SESSION 6

Innovations in Healthcare Solutions

This session will address examples of how certain Indian healthcare solutions have been highly innovative and have delivered concrete benefits to millions of patients. Speakers will address what additional actions need to be taken to expand such innovation to the entire Indian patient population.

SESSION CHAIR

Sudharshan Jain

Managing Director
Healthcare Solutions
Abbott India Ltd.

Indigenous Innovation in Medical Technology: An Emerging Paradigm

Siraj Dhanani

Founder and CEO
Innaccel Pvt. Ltd.

Empowering Frontline Health Workers with Technology

Kanav Kahol

Division Head
Affordable Health Technologies Division
Public Health Foundation of India

Big Data Analytics for Developing Nations: How Secondary use of EHR Data can Help India

Gunjan Jain

Principal Sales Consulting Head
South Asia
Oracle HSGBU

16:30-17:00

CONFERENCE WRAP UP

<p>VENUE DETAILS Palladium Hotel, 462 Senapati Bapat Marg, Lower Parel Mumbai, Maharashtra, 400013, India, +1.888.734.8503</p> <p>CONTACT Tejarani Pednekar, Reservations, Palladium Hotel tel: + 91 22.6162.8000 fax: + 91 22.6162.8888 Tejarani.Pednekar@palladiumhotel.in</p>	<p>MEETING MANAGER</p> <p>Manoj Trivedi Senior Manager Marketing and Program Development DIA (India) Private Limited cell: +91 98.1977.7493 Manoj.Trivedi@diaindia.org</p> <p>Rajnish Kumar cell: +91 98.1909.8457 Rajnish@cimglobal.net</p>
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<p>Signatory</p>		