Digital Disruption in Life Sciences - a Look at NextGen CDM

21-22 April 2017 | Vivanta by Taj, M G Road, Bengaluru

DIA

PROGRAM CHAIR

Shobhit Shrotriya
Vice President
Life Science R&D Operations
&
Capability Lead – Clinical Data Management
Accenture

PROGRAM VICE CHAIR

Raghavendra Kalmadi
Senior Director
Clinical Data Management
QuintilesIMS

PROGRAM COMMITTEE

Archana Subramanya
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Research & Development
GlaxoSmithKline

Nara Iyer
Group Head – Oncology
Novartis Healthcare Pvt. Ltd.

Jayathirtha Gopalakrishnan
Principal Sales Consultant,
Oracle Inc.

Sanjeev Sachdeva
Chief Technical Officer
Tata Consultancy Services

The pivot of the Life Sciences industry is a shift from cost optimization to technology driven innovations which is an enabler for Research & Development. With advent of digital disruption, this shift is more prominent in the rather conservative industry like ours. This conference aims to bring case studies and thought leadership on how NextGen Clinical Data Services can deliver better business outcomes which impact quintessential indicator of 'speed to market' aimed towards improved quality of outcome and enhanced regulatory compliance.

Learning Objectives

At the end of the conference, the participants should be able to:

- The role of technology in the evolution of clinical data management
- Technological innovations in clinical data management
- Understanding data standards

WHO SHOULD ATTEND

Clinical research professionals involved in management and analysis of Clinical Trial Data including:

- Clinical Data Managers
- Clinical Database Programmers
- SDTM Programmers
- Biostatisticians
- Statistical/Clinical Programmers
- Clinical Project Managers

MEETING MANAGER

Manoj Trivedi
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AGENDA

DAY 1 | Friday, 21st April

8:30 – 9:30 **REGISTRATION**

9:30 – 10:00 **OPENING CEREMONY**

10:00 – 10:30 **KEYNOTE PRESENTATION**

Are we ready for the digital world in life sciences?
Suresh Ramu
Co-Founder & CEO
Cytecare Hospitals

10:30 – 11:00 **Tea / Coffee Break**

11:00 – 13:30 **Session 1: Data Standard & data analytics**

Session Chair
Nara Iyer
Group Head – Oncology
Novartis Healthcare Pvt. Ltd.

11:00 – 11:45 | **Clinical Data Standards key to efficient Data Integration and Analytics**
Arunava Chakravarthy
Sr. Group Head, Biostatistics
Novartis Healthcare Private Limited

11:45 – 12:15 | **Clinical Metadata Repository (MDR)**
Tushar Sakpal
Senior Domain Consultant - Biostatistics & Statistical Programming
TCS

12:15 – 12:45 | **Data Analytics in Clinical Research**
Deven Babre
Associate Vice President
Clinical Data Management
TCS

12:45 – 13:30 | **Data Quality and Regulatory Compliance management for next generation CDM**
Savitha Hanuman
VP, Accenture Life Sciences Operations
Accenture

13:30 – 14:30 **LUNCH**

14:30 – 16:00 **SESSION 2: Digital Clinical Trials**

14:30 – 15:00 | **Digital Transformation of Clinical Trials – Where we are and what's in it for Subject, Investigator and Pharma**
K K Naveen
Senior Director, Data Sciences
QuintilesIMS

15:00 – 16:00 | **Panel discussion – Perspective on Digital Clinical Trials: The Road Ahead**

Moderator
Sanjeev Sachdeva
Chief Technical Officer
Tata Consultancy Services

Panelists
Sowmya Srinivasan
Managing Director - Life Sciences Technology
Accenture

K K Naveen
Senior Director - Data Sciences
QuintilesIMS

16:00 – 16:30 **Tea / Coffee Break**

16:30 – 18:00 **SESSION 3: Automation adoption using NLP, Machine Learning & Robotics**

Session Chair
Shobhit Shrotriya
Vice President
Life Science R&D Operations &
Capability Lead – Clinical Data Management
Accenture

Speakers

16:30 – 17:15 | **Robotic Process Automation and Artificial Intelligence in Drug Development**
Mandar Ghatnekar
Managing Director
Accenture Life Sciences Technology

17:15 – 18:00 | **Machine Learning as used in the Healthcare Industry**
Shobhit Shrotriya
Vice President
Life Science R&D Operations &
Capability Lead – Clinical Data Management
Accenture

18:00 **DAY END**

AGENDA

DAY 2 | Saturday, 22nd April

9:30 – 10:00 **Key note Presentation**

Denis Xavier
Professor and Head
Dept. of Pharmacology
St. John's Medical College

10:00 – 10:30 **TEA / COFFEE Break**

10:30 – 12:00 **Session 4: Big Data explosion & its impact on future of Clinical Trials**

Session Chair
Raghavendra Kalmadi
Senior Director
QuintilesIMS

Speakers

10:30 – 11:15 **Redefining Clinical data services leveraging emerging technologies**

Sowmya Srinivasan
Managing Director
Accenture Life Sciences Technology

11:15 – 12:00 **Real-world data and analytics for Life Sciences**

Jeyaseelan Jeyaraj
APAC Director – Solutions Consulting Health Sciences Global Business Unit
Oracle

12:00 – 13:30 **Session 5: eSource Industry View and Road ahead**

Session Chair
Sanjeev Sachdeva
Chief Technical Officer
Tata Consultancy Services

Speakers
TBC

13:30 – 14:30 **LUNCH**

14:30 – 16:00 **SESSION 6: Pushing boundaries in Technology**

Session Chair
Jaythirtha Gopalakrishna
Principal Sales Consultant
Oracle Inc.

Speakers
T R Anil Kumar
Senior Director - Biostatistics
QuintilesIMS

16:00 – 16:30 **Conference Wrap up**

DIA India 2017 Events

USFDA–EMA– CDSCO-DIA Multicentre GCP Workshop

12th & 13th May
St Regis, Mumbai
15th & 16th May
Taj Krishna, Hyderabad

DIA and US FDA will host a 2-day workshop on guidance and polices from the US Food and Drug Administration (FDA), European Medicines Agency (EMA) and Central Drugs Standard Control Organization (CDSCO). GCP inspections related to the quality of clinical trials in Mumbai and Hyderabad. Additional topics discussed at the workshop will include, Data Integrity and BE Study. Interact with experts from USFDA, CDSCO, EMA and industry, who will share experiences and views of the global GCP environment.

Exploring the Scientific Art of Medical Writing: Blending Complexity with Simplicity

June 16-17 | Mumbai

As the world becomes more complex, writers struggle to play a fine balancing act between dealing with more complex challenges such as authoring complex protocols, drafting risk management plans, publication planning, authoring aggregate reports, drafting comprehensive economic and outcomes literature reviews, to stepping down the complexity a notch, trashing the jargon and developing the art of authoring lay language summaries, understanding the value and nuances of QC'g safety narratives and balancing the freedom of expression with the importance of compliance. It is a tight rope walk, with the need to stay abreast with the latest tools and technologies, evolving regulations and yet not lose the fine art if writing.

Deciphering the Why, When and How of Risk Based Monitoring

September 22-23 | Mumbai

The latest ICHE6 R2 guidance has driven even the nay-sayers to focus on developing RBM strategies. Without experience and expertise to support the same, one may actually end up adding risk to a study. The industry has an urgent need to understand the strategies involved in RBM, including the development of the IQRMP, the de-velopment of a risk scoring model, the identification of the KRIs, the SDV strategy, site-tiering strategy, the road-blocks and the pre-emptive measures, the identification of the proof points and the latest regulations and the tools and technologies involved. In addition, this conference will touch on the role of the risk based monitor and the criticality of a well outlined change management strategy.

Digital Disruption in Life Sciences - a Look at NextGen CDM

Event I.D. 17660 | 21-22 April 2017 | Bengaluru, India

VENUE: Vivanta by Taj – M G Road, Bangalore
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Bangalore 560 001, Karnataka, India
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RESERVATIONS CONTACT

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CANCELLATION POLICY: ON OR BEFORE APRIL 7, 2017

- Cancellations must be in writing and received by APRIL 7, 2017. Registrants who do not cancel in writing by that date and do not attend the event will be responsible for the full registration fee paid. Registrants are responsible for cancelling their own hotel and airline reservations.
- DIA reserves the right to alter the venue, if necessary. If an event is cancelled, DIA is not responsible for any airfare, hotel or other costs incurred by registrants.
- UPON CANCELLATION, the administrative fee that will be withheld from refund amount is 25 % of the delegate fee

FULL MEETING CANCELLATION

- All refunds will be issued in the currency of the original payment

For more details, please visit www.DIAglobal.org

REGISTRATION FEES FOR TWO DAYS CONFERENCE (*Registration fee includes refreshment breaks and luncheons.*)

Early Bird on or before 24th March 2017 (Early Bird Confirmation - Subject to Payment Realization)

	BASIC RATE (INR)	SERVICE TAX 15 %(INR)	TOTAL INR
INDUSTRY MEMBER	9000	1350	10350 ❑
INDUSTRY NON- MEMBER	11000	1650	12650 ❑
ACADEMIA / GOVERNMENT	8000	1200	9200 ❑
STUDENT	5000	750	5750 ❑

After 25th March 2017

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INDUSTRY MEMBER	10000	1500	11500 ❑
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STUDENT	5000	750	5750 ❑

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STUDENT REGISTRATIONS

A student is an undergraduate/graduate who can document enrollment in a signature accredited, degree granting, academic program. Please send completed registration form, payment and copy of student identification.

DRUG INFORMATION ASSOCIATION

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Address

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