

ACCELERATING ACCESS TO INNOVATIVE PRODUCTS - IMPROVING RESPONSES TO PANDEMIC DISEASES

January 29-30, 2018 | Mandarin Oriental Singapore

PROGRAM CHAIR

Dorothee GRIMALD

Director, Global Regulatory Policy MSD International GmbH (Singapore Branch)

PROGRAM CO-CHAIR

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Assistant Professor Head of Education Centre of Regulatory Excellence (CoRE) Duke-NUS Medical School

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Thean Soo (TS) LO

Director – Asia Pacific egulatory Policy Lead Global Regulatory Affairs Janssen Asia Pacifi J & J Singapore

Silke VOGEL

Associate Professor, Deputy Director Centre of Regulatory Excellence (CoRE) Duke-NUS Medical School





DIA and the Duke-NUS Centre of Regulatory Excellence (CoRE) will host a 2-day conference in Singapore on accelerating access to health products to improve the response to pandemic diseases. Over these 2 days, we will discuss the global impact of infectious diseases and approaches to expedite research and product development (including vaccines and medical devices). We will also explore the pathways to accelerate patient access to innovative products and how to effectively manage a pandemic situation in Asia. Speakers will include experts from Health Authorities, NGOs, as well as the industry and academia.

Program Highlights

- · Global impact of infectious diseases
- · Increasing antimicrobial resistance and challenges to treatment management
- Experiences from recent and current pandemics in Asia
- Accelerating product research and development for infectious and tropical diseases
- Emerging roles of Real World Evidence and Precision Medicine in pandemics
- Initiatives to enhance timely accessibility to necessary health products
- · Regulatory pathways and initiatives in Asia to expedite availability of health products
- Panel discussion Developing a cross-cutting framework for pandemic responses

Who should attend?

- Industry professionals involved in Research & Development, Regulatory Affairs and Medical Affairs in Pharmaceuticals and Medical Technology
- Regulators and personnel from Health Authorities and Ministries
- · Academia and Researchers

DIA COMMITTEE MEMBERS

Youngshin LEE

SVP/Managing Director ASEAN, India & Korea

Kanchan PATEL

Associate Director India Operations DIA (India) Private Limited

MEETING MANAGERS

Kanchan Patel

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Exhibit Opportunities Available. For more information, contact: Youngshin Lee

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DIA Global Center: Washington, DC, USA | Basel, Switzerland | Beijing/Shanghai, China | Horsham, PA, USA | Mumbai, India | Tokyo, Japan

Day 1 | Monday, January 29, 2018

8:00- 8:45 Registration/Welcome coffee
8:45- 9:00 Welcome Message

Joseph Scheeren

Chair-Elect DIA Senior Advisor R&D

Bayer AG

John Lim

Professor, Executive Director

Centre of Regulatory Excellence (CoRE)

Duke-NUS Medical School

9:00- 9:30 Opening Remarks

Jin Shun

Head - Regulatory Affairs APAC Abbott Laboratories (S) Pte. Ltd.

(Chair -Advisory Committee - DIA Singapore)

Dorothee Grimald

Director, Global Regulatory Policy

MSD International GmbH

(Singapore Branch)

(Chair - Programme Committee)

	Keynote Speech
9:30-10:15	Antimicrobial Resistance and Pandemic
	Infectious Diseases - Impact on Global
	Healthcare Systems and Patients

Mark Pearson

Deputy Director of Employment Labour and Social Affairs

OECD

10:15-10:30	Q&A
10:30-11:00	Tea / Coffee Break
11:00-12:30	Session 1: Addressing the Global Impact of Infectious Diseases

Session Co-Chair

Silke Vogel

Associate Professor, Deputy Director Centre of Regulatory Excellence (CoRE)

Duke-NUS Medical School

11:00 | Evolving Tropical Infectious Diseases - Challenges to Treatment Management

Lee Tau Hong

Associate Consultant

Tan Tock Seng Hospital

11:30 | Role of the industry – Current Challenges in Meeting needs of Infectious Diseases/ Sustaining a Market for Niche Diseases

Hiromichi Shirasawa

Vice President and Head of Japan Development MSD KK

12:00-12:30 | Q&A

12:30 – 13:30 Lunch

13:30–15:00 Session 2-A: A Focus on Current and Recent Pandemics in Asia

Session Co-Chair

Michelle CHENG

Head of Regulatory Affairs

Global Drug Development

Singapore & Asian Emerging Markets (AEM)

Novartis

13:30 | Expediting Research and Product Development – Lessons learnt

Jenny Low Guek Hong

Co-Director, Viral Research and Experimental Medicines Centre,

Senior Consultant, Department of Infectious Diseases Singapore General Hospital

Kathy Tai

Medigen Vaccine Biologics Corporation (MVC)

Damian Foo

Programme Manager

Veredus Laboratories Pte Ltd.

15:00-15:30	Tea / Coffee Break
15:30-16:30	Session 2-B: Ensuring Accessibility of Products During Pandemic Management

Session Co-Chair

James Leong

Assistant Professor

Head of Education

Centre of Regulatory Excellence (CoRE)

Duke-NUS Medical School

15:30 | Navigating the Current Development and Regulatory Landscape in a Pandemic Setting

Melvin Sanicas

Regional Medical Expert

Sanofi Pasteur, Asia & JPAC

16:10 | Health Systems Strengthening - Role of Funders

Douglas Ball

Regulatory Consultant

Asian Development Bank

16:30 -17:30	Panel Discussion	
Moderator		
James Leong		
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Panelists		

Jenny Low Kathy Tai Damian Foo

Melvin Sanicas Douglas Ball

17:30	End of Day 1
17:30- 19:00	Networking Cocktail (By invitation only)

Day 2 | Tuesday January 30, 2018

9:00-10:30

Session 3-A: Enablers for Successful Management of Health Products During Pandemics

Session Co-Chair

Thean Soo (TS) LO

Director – Asia Pacific egulatory Policy Lead Global Regulatory Affairs Janssen Asia Pacifi J & J Singapore

9:00 | Good Reliance Practices as Key to Regulatory Efficiency

Samvel Azatyan

Group Lead - Capacity Building Regulatory Systems Strengthening Essential Medicines and Health Products World Health Organisation (WHO)

9:30 | Emerging Role of Precision Medicines in Pandemics

Jorge Villacian

Chief Medical Officer J&J

9:50 | Using Real World Evidence to Complement Early Access to Pandemic Health Products

Isabelle Grau

Director of Operations

Real World Evidence Asia Pacific

ICON

10:10 | Q&A

10:30-11:00	Tea / Coffee Break
11:00-12:00	Session 3-B

Session Co-Chair

Finny Liu

APAC Regional Regulatory Policy Lead

PDR

Roche, Singapore Pte Ltd.

11:00 | Supply Chain and Logisticoា©iderations During a Pandemic

Swapnil Deshpande

Head of APAC Packaging Hub

Roche Singapore Technical Operations Pte Lte

11:20 | Bringing Timely and Affordable Treatment to the Patients – Is Collaboration a Solution for Health Security?

Geoff Clark

Senior Strategic Health Advisor (Access)
APLMA

11:40 | Q&A

12:00–13:00 Lunch

13:00–15:00 Session 4-A: Accelerating Access to Health Products

Session Co-Chair

Jin Shun

Head, Regulatory Affairs APAC Abbott Laboratories (S) Pte. Ltd

13:00 | Overview of Early Access Regulatory

Pathways for Innovation

Yoshikazu Hayashi

Associate Centre Director

New Drug Review

PMDA

13:30 | Overview of Early Access Regulatory Pathways for Innovation

Chen Wan-yu

Taiwan FDA

14:00 | Perspective of Dengue Vaccine Registration Process

Anuradha Poonepalli

Senior Regulatory Specialist

HSA

14:30-15:00	Tea / Coffee Break
15:00-16:00	Session 4-B

Session Co-Chair

Dorothee Grimald

Director, Global Regulatory Policy

MSD International GmBH

(Singapore Branch)

15:00 | Bringing Innovation to Patients – Industry Perspective on Expediting Access

Lorenz Scheppler

Director

Global Regulatory Affairs

Janssen Pharmaceutical of J&J

Petra Kaars-Wiele

Senior Director

International Regulatory Affairs and Global Labelling

Abbott

16:00-17:00 **Grand Panel Discussion and Closing Remarks**

Developing a cross-cutting framework for pandemic responses – Critical elements

Moderator

Jenny Low Guek Hong

Co-Director, Viral Research and Experimental Medicines Centre, Duke-NUS

Senior Consultant, Department of Infectious Diseases Singapore General Hospital

Panelists

Yoshikazu Hayashi, PMDA Chen Wan-yu, Taiwan FDA Anuradha Poonepalli, HSA Hiromichi Shirasawa, MSD

Petra Kaars-Wiele, Abbott

17:00 End of the conference

Accelerating Access To Innovative Products – Improving Responses To Pandemic Diseases Event I.D. 83317 | January 29-30, 2018 | Singapore

VENUE: Mandarin Oriental Singapore

5 Raffles Ave, Marina Square, Singapore 039797 Tel.: + 65 6885 3522 Fax: + 65 6338 5739

RESERVATIONS CONTACT

Conference rate for delegates: Deluxe room at \$385++ per room per night (inclusive of daily buffet for 01 and internet access)

Kindly note that the rates are valid for booking till 07 January, 2018 and subject to availbility of rooms. Rates will be subjected to changes for any bookings after 07Jan. 2018

Contact: ms eunice tan: eunicet@mohg.Com | mr jan lim: janl@mohg.Com

All rates are subject to 10% service charge and 7% goods and services tax (gst). Please note that the gst is subject to change in accordance with the government policy. Rates are quoted in singapore dollars on a per room per night basis.

MEETING MANAGER

Kanchan Patel, Associate Director Operations, DIA (India) Private Limited cell: +91 90.2909.8844 | kanchan.patel@diaglobal.org

REGISTRATION FEES FOR TWO DAYS CONFERENCE

CANCELLATION POLICY: ON OR BEFORE JANUARY 10. 2018

- Cancellations must be in writing and received by JANUARY 10, 2018.
 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 fee includes refreshment breaks and luncheons.) Early Bird on or before 17th January, 2018 (Subject to Payment Realization) Registration Fee (SGD) Member Industry Early-bird 900 Standard 1,100 On-site 1.300 Academia. Early-bird 400 \Box Non-Profit Standard 500 On-site 600 Non-Member Industry Early-bird 1,100 Standard 1,300 On-site 1.500 Early-bird 600 Academia, Non-Profit 700 Standard On-site 800 Government Early-bird 400 500 Standard Student 200

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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 identific tion.

DRUG INFORMATION ASSOCIATION

800 Enterprise Road Suite 200 Horsham, PA 19044-3595 tel: 1.215.442.6100

email: youngshin.lee@diaglobal.org

Membership		
DIA Membership	200	
2-Year Membersh	360	
Table Top/Booth	s Rate -SGD	
Early Bird	4,000	
Standard	5,000	
Sponsorship	Contact: Kanchan Patel Kanchan.Patel@DIAglobal.org	
Group Discount	A group of 3-4, 15% A group of 5 and more, 20% Contact: Kanchan Patel Kanchan.Patel@DIAglobal.org	

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PAYMENT DETAILS

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