15-18 November 2021
GS1 Healthcare 3rd Online Summit
Global Standards for Global Health

Agenda
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On Monday, 15 November we will host special GS1 standards introduction sessions to global standards. For more information click here

Day 1
Tuesday, 16 November
First session
How do digital technology and global standards improve clinical outcomes?
Second session - Part 1
Medical Device Regulation Spotlight
Second session - Part 2
Supplier Experiences

Day 2
Wednesday, 17 November
First session
Leading implementation case studies - progress continues
Second session
Driving healthcare change across the world

Day 3
Thursday, 18 November
First session
Pharmaceutical Regulation Spotlight
Second session
Helping Healthcare emerge smarter: What does the future of healthcare need to consider?

Replays with a live host are scheduled every day. For more information, click here
First session
AEDT 4pm-6pm | PST 11pm-1am | EST 2am-4am | CET 8am-10am

Summit Opening

- Miguel Lopera, President & CEO, GS1 Global Office, Belgium – confirmed

How do digital technology and global standards improve clinical outcomes?

In a rapidly changing healthcare environment, global standards bring the relevant data into focus helping healthcare leaders make informed decisions and support the care teams around them to deliver the best health outcomes and positive experiences.

- Andy Crosbie, Manager of Post Market Surveillance Strategy, Medicines and Healthcare Products Regulatory Agency (MHRA), UK – confirmed
- Roger Dukers, Expert Application Manager SAP, Zuyderland Medical Centre, the Netherlands – confirmed
- Peter O’Halloran, Chief Information Officer & Executive Group Manager, ACT Health Directorate, ACT Health, Australia – invited
- Dr. Alberto Sanna, Center for Advanced Technology in Health and Well-Being, Director, San Raffaele Hospital, Italy – confirmed

Second session - Part 1
AEDT 12am-1am | PST 7am-8am | EST 10am-11am | CET 4pm-5pm

Medical Device Regulation Spotlight

Hear regulators from around the world share developments relating to unique medical device identification (UDI).

- Dr. Azzam O. Al-Othman, Director of Surveillance - Medical Devices, Saudi Food and Drug Authority (SFDA), Saudi Arabia – confirmed
- Leandro Rodrigues Pereira, General Manager - General Management of Health Products Technology, GGMON / DIRE5 / ANVISA, Brazil – confirmed

Second session - Part 2
AEDT 1am-2am | PST 8am-9am | EST 11am-12pm | CET 5pm-6pm

Supplier Experiences

Suppliers have been working to implement unique medical device identification (UDI) – learn from their implementation experiences.

- Dennis Black, Director, e-Business, Becton, Dickinson and Company (BD), USA – confirmed
- Mr. Sixtus Yv, Technical Director, Foosin Medical Supplies Inc., Ltd., China – confirmed
- Liliana Zuluaga Idárraga, Technical Director, Industrias Medicas Sampedro S.A.S., Colombia – confirmed
First session
AEDT 4pm-6pm | PST 11pm-1am | EST 2am-4am | CET 8am-10am

Leading implementation case studies – progress continues
Growing implementations: Learn about exciting progress made in standards implementations
- Chair – Feargal McGroarty, National Haemophilia System Project Manager, St James’s Hospital, Ireland – invited
- Pierre Fernandez-Barbereau, R&D Clinical Supply Chain Operations, Industrial Development, Sanofi, France – confirmed
- Sinead Moran, Special Feeds Unit Manager, Children’s Health Ireland (CHI), Ireland – confirmed

Second session
AEDT 12am-2am | PST 7am-9am | EST 10am-12pm | CET 4pm-6pm

Driving healthcare change across the world
Building on the Call to Action: Africa Strategy for Pharmaceutical Supply Chain Traceability, hear from healthcare humanitarian and donor organisations, as well as national regulators, about the role of global standards in helping everyone across the globe access authentic, safe medical products.
- Further details for this session, including confirmed speakers, will be provided.
First session
AEDT 4pm-6pm | PST 11pm-1am | EST 2am-4am | CET 8am-10am

Pharmaceutical Regulation Spotlight
The use of globally unique identification and traceability are critical to help fighting falsified and sub-standard medicinal products and to support the healthcare supply chain resilience. During this session regulators, will share updates about this important work.

• Georgios Ampartzidis, Logistics Manager Humanitarian Aid Department, World Federation of Hemophilia (WFH), Canada – confirmed
• Jeff Denton, Vice President, Global Secure Supply Chain, AmerisourceBergen Corporation, USA – invited
• Daniela Marreco, Specialist, GGMON / DIRE5 / ANVISA (Brazilian Health Regulatory Agency, Anvisa), Brazil – confirmed
• Speaker to be confirmed, Pfizer, France – confirmed

Second session
AEDT 12am-2am | PST 7am-9am | EST 10am-12pm | CET 4pm-6pm

Helping Healthcare emerge smarter: What does the future of healthcare need to consider?
Those who work in health and care fill essential roles in our society. But the health and care systems are facing enormous change. It must reinvent itself in the face of new and increasing demands and limited resources. We will discuss why global standards combined with advancing technology will drive impactful change not only for the patient but the clinician and healthcare system too.

• Chair - William Smart, former National CIO, NHS England, current Global Director External Relations, Dedalus Group, Italy – confirmed
• Paul Coplan, VP and Head, Medical Device Epidemiology & Real World Data Sciences, Johnson & Johnson, Germany – confirmed
• Dr. Eric Hans Eddes, Senior Advisor Benchmarking, International Consortium for Health Outcomes Measurement (ICHOM), the Netherlands – confirmed
• Andrew Raynes, Chief Information Officer, Royal Papworth Hospital NHS Foundation Trust, Cambridge, UK – confirmed
• Alex Zimmerman, System Director Supply Chain Technology, Baylor Scott & White Health, USA – confirmed

Summit Closing
• GS1 Healthcare Tri-chair - Gerry Collins, Global Platform Leader Parenterals, Janssen, Pharmaceutical Companies of Johnson & Johnson – confirmed