

Advancing Global Convergence of Medical Technology Regulation

IMDRF Stakeholder Session September 17, 2019

Philippe Auclair



Outline

- Principles for Advancing Global Convergence
- Impact of Differing Regulations
- Global Harmonization Movement
- Key Elements to Promote Convergence
- Global Convergence Benefits
- Recommendations to Support Convergence



Principles for Advancing Global Convergence

Health & Safety

 Right to protect health and safety and to establish regulations to determine safety and quality

A Proportionate Approach

 Application of the principle of proportionality to regulation (use the least burdensome measure)

Transparency, Efficiency and Predictability

Transparent, efficient and predictable regulatory controls



Impact of Differing Regulations

- Market entry hurdles
- Increased costs
- Technical barriers that impede trade
- MOST IMPORTANTLY, delay in patient access to medical technologies



Global Harmonization Movement

- Thanks to GHTF and now IMDRF, international regulatory convergence has been underway for nearly 20 years
- Training needed to ensure consistent application of IMDRF and GHTF guidance documents
 - APEC Medical Device Priority Work Area established
 - GMTA pilot training proposal



Key Elements to Promote Convergence

- Implementation of GHTF & IMDRF guidance documents into national laws
- Regulator training programs to ensure consistency in implementation
- Adoption of international consensus standards
- Reliance on regulatory decisions (MDSAP)



Global Convergence Benefits Promotes Public Health

- Supporting innovation/development of new therapies
- Creating opportunities for regulator to regulator collaboration
- Fosters transparent engagement from industry
- Strengthening industry compliance with regulatory requirements

Provides opportunity for regulatory reliance

Leveraging resources by regulators

- MOST IMPORTANTLY FOR PATIENTS

Improve patient timely access to life-saving and life-sustaining technologies around the world



Principles of Labeling for Medical Devices and IVD Medical Devices

Aligned labelling interpretation has a significant impact on market access and efficiency for the regulators and for the MedTech industry.

Recognition and/or use of ISO symbols and E-IFU adoption are key to enhance patient access.





Recommendations to Support Convergence

- Interactions with other harmonization groups
- Inclusion of industry experts in relevant IMDRF working groups
- Inclusion of experts from non-IMDRF regulators on working groups
- Support training programs on IMDRF documents
- Measure progress on implementation of guidance documents