

MEDICAL DEVICE CYBERSECURITY WORKING GROUP UPDATE

Working Group Co-chairs:

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GOALS

 To facilitate international regulatory convergence on medical device cybersecurity with open discussion and sharing best practices that are understandable and feasible for all stakeholders.

 Specifically, the WG goal is to produce a document providing medical device cybersecurity guidance for all responsible stakeholders, including manufacturers, healthcare providers, regulators, and users across the entire device lifecycle.

SCOPE

The document is intended to:

- Provide recommendations to aid in minimizing cybersecurity risks across the **total product lifecycle**;
- Recognize that cybersecurity is a shared responsibility among all stakeholders which are not only manufacturers but also healthcare providers, patients, regulators, and researchers;
- **Define terms** consistently and clarify the current understanding on medical device cybersecurity;
- Promote broad **information sharing policies** for cybersecurity incidents, threats, and vulnerabilities.



LINKAGES WITH EXISTING IMDRF DOCUMENTS

- IMDRF/GRRP WG/N47 FINAL: 2018, in sections 5.5.2 and 5.8 describes information security, IT environment and cybersecurity.
- IMDRF/SaMD WG/N12 FINAL: 2014 describes the importance of information security with respect to safety considerations in Section 9.3.
- It was the intent of this WG to further elaborate on and provide additional clarity and granularity on these topics.

ACTIVITIES TO DATE

- Kick-off meeting was in January 2019
- Meetings occurred every 2 weeks
- Final guidance document outline complete: February
 2019
- Guidance drafting and iterative review February to April
 2019
- In-person WG working meeting: June 10-13, 2019,
 Medical Imaging & Technology Alliance (MITA) office in Arlington, Virginia
- Submitted draft Guidance to IMDRF Management Committee : **August 2019**

KEY HIGHLIGHTS OF CURRENT DRAFT

- Document is structured into 4 main components:
 - Definitions
 - General Principles
 - Pre-market Considerations
 - Post-market Considerations
- All definitions add to N47 terms/definitions, generally align with internationally recognized standards, and include key terms such as: 'cybersecurity', 'compensating control', 'legacy device', 'patch', and 'privacy', among others



KEY HIGHLIGHTS CONT'D

- **General Principles** section covers:
 - Requirement for total product life cycle approach
 - Concept of shared responsibility among stakeholders
 - Global harmonization and concept of information sharing
- Pre-market section covers:
 - Recommendations for manufacturers only
 - Recommendations include: good design, risk management, security testing, labelling, and regulatory submission requirements

KEY HIGHLIGHTS CONT'D

Post-market section covers:

- Recommendations for <u>all stakeholders</u> including manufacturers, healthcare providers and patients (users), regulators, and security researchers
- Includes 'best practices' for <u>healthcare providers</u> to use medical devices in a secure manner
- Provides guidance on information sharing and vulnerability disclosure along with remediation concepts and incident response best practices
- Proposes level of regulatory oversight required for different categories of software maintenance
- Recommends best practices for Legacy devices (devices that cannot be reasonably protected against current cybersecurity threats)

FUTURE WORK PLAN AND MILESTONES

- Proposed document planned for Public Consultation:
 October and November 2019
- 2. Review and Organize Public Comments: **December 2019**
- In-person meeting to produce a final guidance document: mid/late January 2020 (tentatively Geneva, Switzerland)
- 4. Submit Final Guidance in **February 2020** for approval at Management Committee Meeting in **March 2020**

Public Consultation: October-November 2019

Review Comments: **December 2019** In-person WG meeting, Geneva, TBC: January 2020 Final Guidance to IMDRF Management Committee: February 2020



THANK YOU