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A new draft technical information report (TIR) developed establish a common understanding of postmarket tools and activities for medical devices, AAMI/ISO DTIR20416:2020 Medical devices—Post-market surveillance for manufacturers, has been approved by the AAMI Standards Board and registered by the American National Standards Institute (ANSI).

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AAMI TIR17: 2017 Compatibility of materials subject to sterilization. his technical information report provides guidance for health care product manufacturers in thequalification of polymeric materials, ceramics, and metals for use in health care products that are sterilized by the following modalities: a) radiation (gamma, electron beam, or x-ray); b) ethylene oxide; c) moist heat (steam); d ...

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of documents, it may be helpful to address the objectives of AAMI's technical document development program and see why these publications are developed in the first place.

Differentiating Standards, Recommended Practices, and ...

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This Technical Information Report (TIR) provides perspectives on the application of AGILE during medical device software development.

AAMI TIR45:2012 (AAMI TIR 45:2012) - Guidance on the use ...

AAMI TIR45 is a document released by the Association for the Advancement of Medical Instrumentation in 2012. It is important to note that this Technical Information Report is not a standard or even a recommended practice.

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