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By Val Theisz

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enabled implementation of good review practices. Journal of Medical Device Regulation Val Theisz, MSc RAC, is the regulatory affairs and quality assurance
http://certificationbody.com.au/documents/JMDRAug2012_RPS.pdf

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http://ca.linkedin.com/pub/dir/%2B/Theisz/

Medical devices regulations. of international best practices is a priority for the the regulations of medical devices as one of the medical
http://www.who.int/medical_devices/safety/en/

Medical device software traceability, Medical device software effective and regulatory compliant traceability practices. allow digital signatures to be
http://www.academia.edu/1863158/Medical_Device_Software_Traceability

Risk Management for Medical Devices Val Theisz, MSc, RAC, is the regulatory & QA manager for Certification pliance with applicable state radiation regulations.
http://www.feemedicare.org/tag/radiation/

and practical recommendations that bridge the gap between regulatory theory and practice medical device regulatory Val Theisz is a regulatory
http://www.bokus.com/bok/9789814669108/medical-device-regulatory-practices/

Overview of regulations for medical devices: CDRH regulates radiation-emitting electronic products (medical and non Electronic Products; Medical devices are
http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/Overview/

An International Perspective by Val Theisz for professionals in the medical device the gap between regulatory theory and practice.
http://gkopu.com/books/2015-8-6/medical

Mar 24, 2013 This guide is intended to introduce entrepreneurs to FDA regulation around medical devices medical app? Electronic digital health device.
http://www.slideshare.net/RockHealth/fda-101-a-guide-to-the-fda-for-digital-health-entrepreneurs

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http://www.amazon.com.au/gp/new-releases/digital-text/2537024051

Risk Management for Medical Devices Welcome to Regulatory Reconnaissance, Posted 22 June 2012 By Val Theisz.
http://www.raps.org/Regulatory-Focus/Features/Transfered-Features/2012/06/22/7003/Risk-Management-for-Medical-Devices---A-Practical-Approach/

Medical Device Regulatory Practices: An International Perspective - Biomedical Engineering - Books on Diseases - Valuable medical/health info related to diseases,
http://www.medical-books.medindia.com/3-14116-9814669105-Medical_Device_Regulatory_Practices_An_International_Perspective

Medical Instruments and Devices Principles and Medical Instruments and Devices: Principles and Practices originates from the medical instruments and devices
http://www.tandf.net/books/subjects/SCEC02/

510(K) SUMMARY . In accordance with the Food and Drug Administration Rule to implement provisions of the Safe Medical Devices Act of 1990 and in conformance with 21
http://www.accessdata.fda.gov/cdrh_docs/pdf14/K142585.pdf

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English summary of the August 2012 issue of the Journal of Medical Device Regulation. medical devices, Val Theisz devices; Good Manufacturing Practice:
http://www.globalregulatorypress.com/features/browse_issues_aug12en.shtml

Research has shown that medical device regulations cannot be completely development practices within the medical device 2008, Val D Isere
http://limerick.academia.edu/ValentineValCasey/Papers

Traceability is central to medical device software development and essential for regulatory approval. For compliance to be achieved, an effective traceability process
http://www.academia.edu/1863262/A_lightweight_traceability_assessment_method_for_medical_device_software

These requirements are being made pursuant to section 32 of the Medical Devices Regulations, Guidance for Industry - Device Licence Applications for Val
http://www.hc-sc.gc.ca/dhp-mps/md-im/activit/announce-annonce/md_notice_im_avis_CIII_CIV_format_mise_en_forme-eng.php

In addition to traditional medical devices, The Corporate Practice of Cooley has digital health regulatory attorneys experienced in structuring business
http://www.jdsupra.com/legalnews/alert-key-regulatory-considerations-for-73235/

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