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#### **TECHNICAL ISO/TR REPORT 14969**

ISO 13485 Specifies The Quality Management System Requirements For Medical Devices For Regulatory Purposes (see Annex A). ISO 13485 Accommodates The Previous ISO 13488 By Permissible Exclusion As Specified In ISO 13485:2003, 1.2. When Judging The Applicability Of The Guidance In Thi 15th, 2024

# ISO 14001, ISO 50001, ISO 26000, ISO 10002, ISO 16949

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ISO 12207 9 3. History (2) ISO/IEC 12207 Sponsor: • Joint Technical Committe 1 (JTC1) (Information Technology) Of International Organization For Standardization (ISO) And International Electrotechnical Commission 7 (IEC). • Developer: Subcommittee 7 (SC7) (Software Engineering) Proposed In June 1988 Published 1 August 1995File Size: 292KBPage Count: 49 12th, 2024

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