

Managing the Device Supply Chain

Best Practices

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Appendices (<i>can be found on the CD in the back of this report</i>)	
A. 21 CFR Part 820 – Quality System Regulation	
B. Quality Management System – Medical Devices – Guidance on the Control of Products and Services Obtained from Suppliers	
C. 21 CFR Part 820.50 – Purchasing Controls	
D. Quality Systems Approach to Pharmaceutical cGMP Regulations	
E. Contract Manufacturing Arrangements for Drugs: Quality Agreements	
F. SG4-N84: Guidelines for Regulatory Auditing of Quality Management Systems of Medical Device Manufacturers – Part 5: Audits of Manufacturer Control of Suppliers	