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Good Manufacturing Practices (GMP) For Medicinal Products

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IS 16021 (2012): Good Manufacturing Practices (GMP) ...

ISO 10012 : 2003 Measurement Management Systems — Requirements For Measurement Processes Measuring Equipment ISO 14159 : 2002 Safety Of Machinery — Hygiene Requirements For The Design Of Machinery ISO Mar 3th, 2024

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1.1.2 21 CFR 210 AND 211: CURRENT GOOD MANUFACTURING PRACTICE FOR FINISHED PHARMACEUTICALS Parts 210 And 211 Of CFR Title 21 Are The Laws Defining Good Manufacturing Practices For Finished Pharmaceutical Products. All Manufacturers Must Follow These Regulation Jan 10th, 2024

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US FDA : 21 CFR Part 210 & 211, Process Validation And OOS Draft Guidance 1997, 21 CFR 58.on GLP 2. WHO TRS : 822, 823, 834, 863, And 902 (GMP, Sterile, Biolo May 11th, 2024

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2. Personnel - GMP ZThere Are Competent And Appropriately Qualified Personnel In Sufficient Numbers To Ensure Service Provision. ZThe Responsibilities Of All Staff Should Be Clearly Understood And Recorded. ZAll Personnel Receive Initial And Continuing Training Relevant To Their Needs. ZOnly Staff Who Have Appropriate Training ...File Size: 948KB Apr 23th, 2024

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