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Manufacturing Practices (GMP), And The Term Of
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- Version 34 (May 14) GLPUS 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211, Process Validation And OOS Draft Guidance 1997,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,

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Materials, That Were Not Present In TheGLP Batches, Is

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(Drug Product) To Be No More Than0.15% In Drug And

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12.1 Validation Policy 12.2 Validation Documentation ... Current Step 4 Version, Dated 10 November 2000. Increasing GMP Requirements
TRS957.indd 135 21.04.10 11:03. 136 2. Mar 9th, 2024
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