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Internal Audit Report Audit Report Date: Last Audit Report ... ISO 9001 ISO 14001 System Audited: Quality Management System Authorised: Marcus Maxwell Authorised Date: 2.7.18 Report Number(s): 1 Start Time: 12.30pm Finish Time: 4.30pm Objectives Of Audit To Confirm That The Management System Conforms To May 21th, 2024 GMP GMP And GLP DATABASE - Version 34 ( May 14 ) GLPUS FDA : 21 CFR Part 210 & 211, Process Validation And OOS Draft Guidance 1997, 21 CFR 58.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, And 902 (GMP, Sterile, Biolo Apr 3th, 2024 Toxicology Studies- GMP Or Non-GMP? The Limits Allowed For Any New Impurities In The Clinical Trial Materials, That Were Not Present In The GLP Batches, Is Addressed By ICH Q3A (Drug Substance) And ICH Q3B (Drug Product) To Be No More Than 0.15% In Drug And 0 Jun 22th, 2024.

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