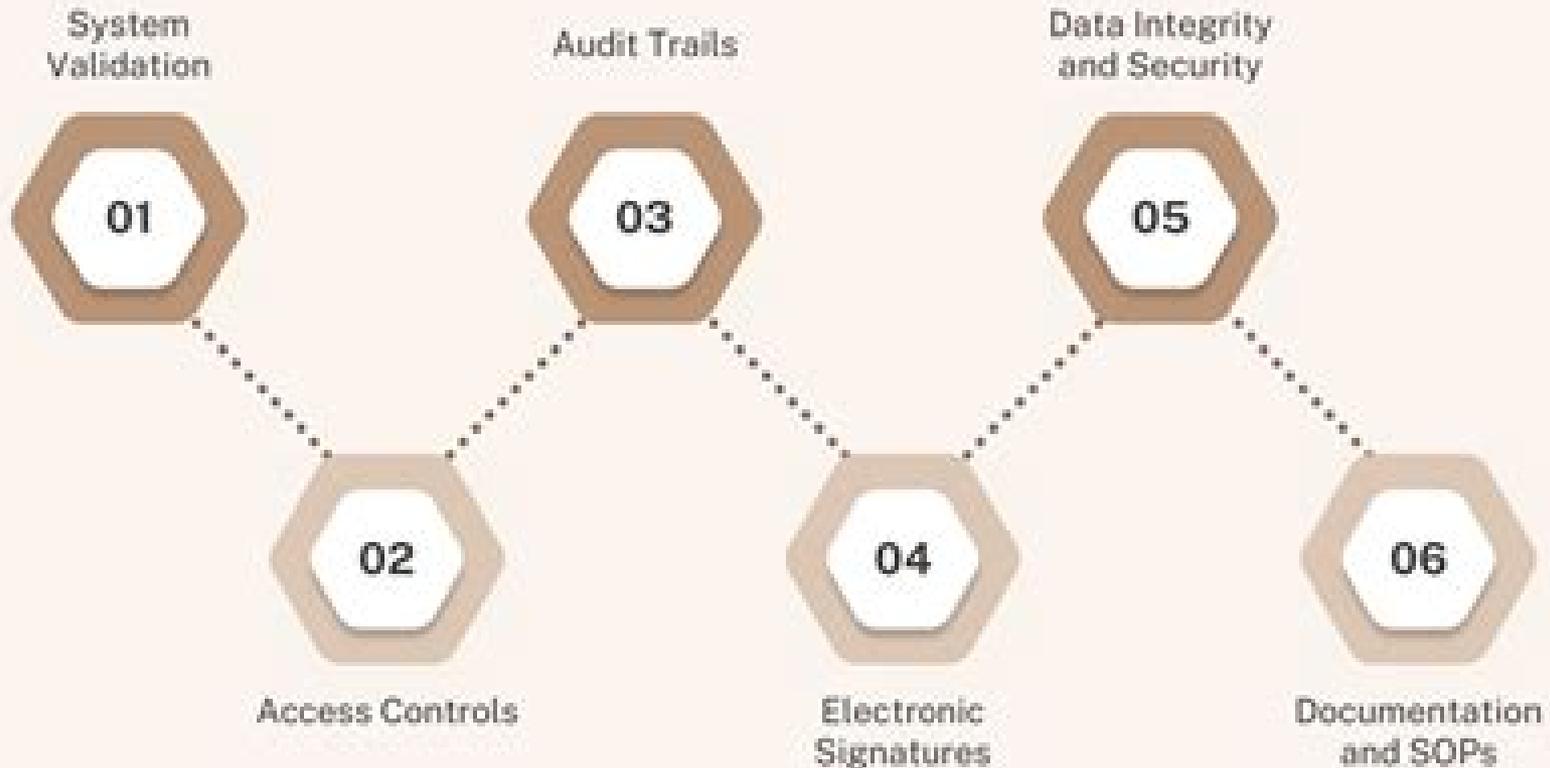


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Handbook of Computer and Computerized System Validation for the Pharmaceutical Industry Stephen Robert Goldman, 2003 This handbook details methods for sustainable compliance with GxPs and 21 CFR Part 11 validation requirements regarding computerized systems in the pharmaceutical biotechnology and medical device industry The

handbook follows FDA guidelines and best industry practices in defining roles responsib **Guidance for industry** ,2001

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