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Review on Cleaning Validation

Salgar J. B., Bais. S. K., Sultana Amir Sayyad
Fabtech College of Pharmacy, Sangola, Solapur, Maharashtra, India

Abstract: *Each pharmaceutical industry's objective is to reliably and affordably produce goods with the necessary qualities and attributes. Method development is crucial for drug discovery, development, and evaluation in pharmaceutical formulations drug discovery, development, and evaluation in pharmaceutical formulations, method development is crucial. This review article's main goal was to examine how pharmaceutical manufacturing procedures are developed and validated from the beginning of formulation to the final commercial batch of product. The results must be trustworthy when analytical procedures are used to get high-quality results for pharmaceutical samples.A verification policy in the pharmaceutical business specifies how verification is carried out, and both the type of verification and the verification policy adhere to Good Manufacturing Practice (GMP) laws. The efficient running of pharmaceutical enterprises depends on validation. From raw ingredients to finished goods, stability and validation are performed at every stage. Accuracy, specificity, precision, limit of detection (LOD), limit of quantitation (LOQ), robustness, robustness, and system were the validation parameters for the method. In terms of conformity testing, it is explained. Both routine and stability assessments make use of all validation parameters.[1].*

Keywords: Validation, Method development, Limit of quantization, Limit of detection, Linearity, Analytical

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