GOOD MANUFACTURING PRACTICES FOR PHARMACEUTICAL PRODUCTS – MAIN PRINCIPLES (WHO TRS 986 – Annex 2)

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Overview

The linked document, **WHO Technical Report Series No. 986, Annex 2: Good Manufacturing Practices for Pharmaceutical Products: Main Principles**, outlines the fundamental principles of good manufacturing practices (GMP) for ensuring the consistent quality and safety of pharmaceutical products. It provides essential guidelines covering all stages of the production process, from raw materials to final product distribution, ensuring compliance with safety standards and regulatory requirements.

Suppliers are responsible for the proper implementation of GMP across all stages of manufacturing. Additionally, it is crucial to conduct thorough risk assessments to identify and mitigate potential hazards throughout the production process. These risk assessments play a vital role in maintaining product quality and ensuring compliance with GMP standards.

Link to Document

TRS 986 - ANNEX 2: WHO GOOD MANUFACTURING PRACTICES FOR PHARMACEUTICAL PRODUCTS - MAIN PRINCIPLES