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The Validation Master Plan for Product Distribution (VMP) describes the policies and strategies of the qualification program for product shipment qualification and defines the requirements for the storage and transport of Drug Substance (DS), Bulk Drug Product (BDP), and finished Drug Product (DP) manufactured at company-operated sites or approved contracted manufacturing sites.

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It covers the planning of validation activities related to the manufacturing and control of the registered stages of Drug Product or Active Pharmaceutical Ingredient (API) for clinical use, validation or sale. All manufacturing activities concerned with: - The receipt and

establishment of new Drug Products or API□s.

The Preparation of Validation Master Plan - Gmpsop
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A Validation Master Plan, also referred to as "VMP", outlines the principles involved in the qualification of a facility, defining the areas and systems to be validated, and provides a written program for achieving and maintaining a qualified facility. GMP Validation for a pharmaceutical/biotech product or process

High Quality Validation Master Plans (VMP) for FDA | EU ...
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