
The Finished Product

specifications and control tests on the finished product - finished product must comply with the criteria defined by directive 75/318/eec as amended, i.e. $\pm 5\%$ for the assay of active substance(s) except when otherwise justified. 1.3 relationships between the specification of a finished product at the end of shelf life and at manufacture (at release) **finished product specification - polypeptide** - finished product specification analytical report page 2 of 2 product: deslorelin acetate analytical reference no. * analysis performed by contract laboratory # tests to be performed when retesting. **finished product specification - white labs** - finished product specification 3 11. genetic modification - spl does not use genetically modified materials in its products. 12. process controls & qc checks parameter method by whom frequency target comments **sop for finished product storage** - sop for finished product storage · once meat/poultry items are packaged and labeled, they will be master-packed (if appropriate), and immediately moved into either dry storage (jerky and other shelf-stable products), refrigerated storage, or frozen storage. · all coolers will be maintained to hold a temperature of 41°F or lower, with daily **standard operating procedure for finished good test ...** - composite wood products. for finished goods that consist of a laminated product in which one side is not laminated or coated, the product may be cut and tested as a panel with a surface coating on one side (see sop for composite wood panels). the key activity in the deconstruction process is the removal of the surface layer of a **specifications: test procedures and acceptance criteria ...** - current product and process understanding, that assures process performance and product quality. the controls can include parameters and attributes related to drug substance and drug product materials and components, facility and equipment operating conditions, in-process controls, finished product specifications, and the **guideline on manufacture of the finished dosage form** - consideration should be given in 3.2.p.2 to what extent the assurance of quality of the finished product is founded on the manufacturing process itself. the significance of the process description and process controls as part of the overall control strategy should be outlined based on development studies and evaluated. **manufacturers versus component part and raw material ...** - manufacturers versus component part and raw material suppliers: how to prevent liability by kenneth ross * introduction one of the more perplexing and potentially dangerous areas of product liability practice and law concerns the liability of the original equipment or finished **guidance for industry - food and drug administration** - notice (42 fr 57749), fda described conditions for using lal as a finished product test. 6. by 1983, fda indicated in guidance that an lal test could be used as a finished product test for endotoxins. **production and process controls - fda** - in-process materials and finished drug products subpart j - records and reports : §211.180(e)(2)(3) - annual product review §211.192 -production record review, deviation and investigation **material safety data sheet finished product. section 1 ...** - cover girl lip perfection lipstick page 5 of 6 while this finished product is not considered hazardous as defined by osha in 29 cfr 1900.1200 (d), this msds contains valuable information critical to the safe handling and proper **6120.1 finished product stdrds prog8/27/87 new line speed ...** - 6120.1 finished product stdrds prog8/27/87 opi: mpits/sisspd finished product standards program for the new line speed inspection system and the streamlined inspection system i. purpose this directive provides guidelines for applying the finished product standards program for the new line speed and streamlined inspection systems for poultry. ii ... **non-food safety consumer protection tasks** - non-food safety consumer protection tasks objectives . after completing this module, you will be able: 1. identify the statutes, regulations and primary directives that relate to non- ... fsis directive 6120.1, "finished product standards program for the new **q3b(r2) - ich** - impurities in new drug products 3 threshold (see attachment 1), and total degradation products observed in the relevant batches of the new drug product, should be reported with the analytical procedures indicated. below 1.0%, the results should be reported to the number of decimal places **ingredients and finished product - wifoodprotection** - • ingredients, finished product and environmental -lot management is important • ingredient testing when you do not have an internal kill step is a verification of your supplier's ccp • finished product testing is a verification of your ccp. • environmental monitoring is a verification of zoning and gmps • testing is not a control ...

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