

10106 COMPETITIVE PROBLEMS IN THE DRUG INDUSTRY

NAME	ADDITIONAL REQUIREMENTS	SIGNIFICANCE
Streptomycin Sulfate, USP, Equivalent to 1 Gram of Streptomycin Base	Classification of Defects Solubility Time Limit Weight Variation	See Explanatory Notes See Explanatory Notes The USP and Code of Federal Regulations do not specifically require compliance with weight variation limits. Our require- ment assures a uniform quantity within limits, and thus, proper and uniform dosage.
Tartar Hydrate and Cocaine Elixir, USP, 4 fl. oz.	Free from sediment	To assure that impurities are detected that may arise from production procedures, or from changes in sources of materials, or in the processing of the item. The presence of these impurities is inconsistent with good manufacturing practices. See Explanatory Notes
Potassium Penicillin G, Sterile, USP, 20,000,000 Units	Classification of Defects Color Limits Weight Variation	See Explanatory Notes See Explanatory Notes The USP monograph and CFR does not specifically require compliance with content uniformity for this item. Our requirement assures uniform quantity within limits and the proper and uniform dosage. See Explanatory Notes
Sodium Methicillin for Injection, USP, 1 Gram	Classification of Defects Solubility Time Limit Weight Variation	See Explanatory Notes See Explanatory Notes The USP monograph and CFR does not specifically require compliance with content uniformity for this item. Our requirement assures uniform quantity within limits and the proper and uniform dosage.