

10086 COMPETITIVE PROBLEMS IN THE DRUG INDUSTRY

NAME	ADDITIONAL REQUIREMENTS	SIGNIFICANCE
Aspirin, Phenacetin and Caffeine Tablets, NF	Moisture Content	To assure the stability of the product, in that excessive moisture may cause deterioration. See Explanatory Notes.
Aspirin Tablets, USP, 0.324 Gram, 100s	Classification of Defects	See Explanatory Notes.
Aspirin Tablets, USP, 0.324 Gram, 1000s	Accelerated Aging Test Hardness Moisture Content	See Explanatory Notes. See Explanatory Notes. To assure the stability of the product, in that excessive moisture may cause deterioration. See Explanatory Notes.
Calcium Lactate Tablets, NF, 0.65 Gram, 100s	Classification of Defects	See Explanatory Notes.
Chlorthalidone Tablets, USP, 100 mg, 100s	Classification of Defects Hardness limit	See Explanatory Notes. See Explanatory Notes.
Chlorpromazine Hydrochloride Tablets, USP, 25 mg, 1000s Chlorpromazine Hydrochloride Tablets, USP, 50 mg, 1000s Chlorpromazine Hydrochloride Tablets, USP, 100 mg, 1000s	Classification of Defects Limits for foreign substances (Color, pH, unchlorinated compound for active ingredient and 4-Chlorophenothiazine for intermediate)	See Explanatory Notes. 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.
Chlorotrianisene Capsules, NF, 72 mg, 48s	Classification of Defects	See Explanatory Notes.
Caffeine and Sodium Benzoate Injection, USP, 0.25 Gram per cc, 2 cc, 12s	Color Limits Leakage Test for Ampuls Classification of Defects	See Explanatory Notes. See Explanatory Notes. See Explanatory Notes.