	ADDITIONAL REQUIREMENTS	SIGNIFICANCE
Reserpine Tablets, USP, 0.25 mg, 1000s	Classification of Defects	See Explanatory Notes.
Isoniazid Tablets, USP, 0.3 Gram, 100s	Classification of Defects	See Explanatory Notes.
Isoniazid Tablets, USP, 100 mg, 100s	White Tablets	To limit deterioration which is evidenced by discoloration.
Pseudoephedrine Hydrochloride Syrup, NF, 30 mg per 5 cc, 1 pt (473 cc)	Taste/Palatability For Active Ingredient: Tighter Melting Range Tighter Specific Rotation	See Explanatory Notes. To reduce incidence of impurity isomers.
Pseudoephedrine Hydrochloride Tablets, NF, 30 mg, 100s	Classification of Defects For Active Ingredient: Tighter Melting Range Tighter Specific Rotation	See Explanatory Notes. To reduce incidence of impurity isomers.
Pyrimethamine Tablets, USP, 25 mg, 100s	Classification of Defects	See Explanatory Notes.
Promazine Hydrochloride Injection, NF, 50 mg per cc, 2 cc, 25s	Classification of Defects	See Explanatory Notes.
Prochlorperazine Suppositories, NF, 2.5 mg, 6s Prochlorperazine Suppositories, NF, 25 mg, 6s	Melting Point	To assure that the suppositories are made with a base which will melt within a specified time frame to release its medicament. No such requirement exists in the compendia.
	Accelerated Aging Test	See Explanatory Notes.
	Limits for Foreign Substances (Color, Unchlorinated Compound for Active Ingredient and 4-Chlorophenothiazine for Intermediate)	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.