

10118 COMPETITIVE PROBLEMS IN THE DRUG INDUSTRY

ADDITIONAL REQUIREMENTS	SIGNIFICANCE
Hydrogen Peroxide Solution, USP, 1.1% (15.16 grams)	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
Free from sediment	
Color limits	
pH limits	The USP monograph has no specific requirement for pH. This requirement is to assure greater stability over the shelf life of the items.
Preservative shall be present.	USP monograph states that a preservative "may be present". By our P. D. requiring the presence of a preservative, we are assuring greater stability of the item.
Classification of Defects	See Explanatory Notes
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Color limits	See Explanatory Notes
Leakage Tests for Ampuls	See Explanatory Notes
Classification of Defects	See Explanatory Notes
Aluminum Iodinate Injection, USP, 50%, 30 cc, 25s	
Potassium Chloride Injection, USP, 2.16%, 10 cc, 10s	
Rescain Tablets, USP, Oral, 500,000 Units, 100s	