

PRODUCT SPECIFICATIONS SHEET

Product Name Catalog # Ethyl Alcohol 50% Grain Derived Ethanol 111000100

TEST	MONO- GRAPH	SPECIFICATION	TYPICAL RESULT
Assay (v/v)	Internal	49.50% - 50.50%	50.01 %
Proof	27CFR 30	Lot Analysis	100.0
Specific Gravity	Internal	0.9332 -0.9352 @ 15.56°C	0.9342
Specific Gravity	Internal	0.9306-0.9326 @ 20.00°C	0.9316
Acidity (as acetic acid)	FCC	NMT 0.5 mL of 0.02N sodium hydroxide is required to restore the pink color. (NMT 0.003%)	Pass
Alkalinity (as NH3)	FCC	NMT 0.2 mL of 0.02N sulfuric acid is required to restore the red color. (NMT 3 mg/kg)	Pass
Organic Impurities - Fusel Oil	FCC	No foreign odor is perceptible when the last traces of alcohol leave the paper.	Pass
Identification by Infrared Absorption	FCC	Conforms to IR Spectra	Pass
Organic Impurities - Ketones, Isopropyl Alcohol	FCC	No precipitate forms within 3 min.	Pass
Inorganic Impurities - Lead	FCC	NMT 0.5 mg/kg	LT 0.5 mg/kg
Organic Impurities - Methanol	FCC	200 ppm max.	1 ppm
Nonvolatile Residue	FCC	NMT 0.003%	0.000 %
Organic Impurities - Any other single impurity	FCC	1000 ppm max.	1 ppm
Solubility in Water	FCC	No haze or turbidity develops	Pass
Organic Impurities - Substances Darkened by Sulfuric Acid	FCC	The mixture is colorless or has no more color than either the acid or the sample before mixing.	Pass

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Form: Ethanol, 100, USP/FCC, Rev 1.6, 08/23, PJM



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Organic Impurities - Substances Reducing Permanganate	FCC	The pink color does not entirely disappear.	Pass
Organic Impurities - Sum of all impurities	FCC	5000 ppm max.	0 ppm
Acidity or Alkalinity	USP	The solution is pink (30μg/g, expressed as acetic acid)	Pass
Clarity of Solution	USP	Sample solution A and Sample solution B show the same clarity as that of water, or their opalescence is not more pronounced than that of the Standard suspension A.	Pass
Color of Solution	USP	The Sample solution has the appearance of water or is not more intensely colored than the Standard solution	Pass
Identification Test B (Infrared Spectroscopy)	USP	Conforms to IR Spectra	Pass
Limit of Nonvolatile Residue	USP	NMT 2.5 mg	0.0 mg
Organic Impurities - Acetaldehyde and Acetal	USP	NMT 10μL/L, expressed as acetaldehyde	0 μL/L
Organic Impurities - Benzene	USP	NMT 2μL/L	1 μL/L
Organic Impurities - Methanol	USP	NMT 200μL/L	1 μL/L
Organic Impurities - Sum of all other impurities	USP	NMT 300μL/L	1 μL/L
UV Absorbance	USP	NMT 0.40 at 240 nm	0.02
UV Absorbance	USP	NMT 0.30 between 250 and 260 nm	0.01
UV Absorbance	USP	NMT 0.10 between 270 and 340 nm	0.00

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This product is for further commercial manufacturing, laboratory, or research use, and may be used as a process solvent for pharmaceutical purposes. It is not intended for use as an active ingredient in drug manufacturing nor as a medical device or disinfectant. Appropriate/legal use of this product is the responsibility of the user.

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