EPOCH - alcohol gel
NSE Products, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Epoch® Antiseptic Hand Sanitizer

Active Ingredient

SD Alcohol 40-B (62%)

Purpose

Antimicrobial.

Usage

Disinfects and provides protection by killing 99.99% of the most common harmful germs and bacteria.

Warnings

For external use only.

Keep out of reach of children.

Keep out of eyes.

Discontinue use if irritation develops. If condition persists for more than 72 hours consult a physician.

Flammable, keep away from fire or flame.

Directions

Place a teaspoon of product in one hand. Spread on both hands, applying liberally. Rub gently until dry.

Inactive Ingredients:

Water (Aqua), Lavandula Angustifolia (Lavender) Oil, Lonicera Japonica (Honeysuckle) Flower Extract, PEG-40 Hydrogenated Castor Oil, Carbomer, Aminomethyl Propanol, Fragrance (Parfum), Chlorphenesin.

Questions?

1-888-742-7626

Mfd. in the U.S.A. exclusively for
NSE Products, Inc., Provo, UT 84601

PRINCIPAL DISPLAY PANEL - 100 ml Bottle Label

epoch®

with Lavender,

Honeysuckle Extract,
and Chlorphenesin

Antiseptic

Hand Sanitizer

100 ml e (3.4 fl. oz.)

EPOCH

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62839-1083
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	61.9937 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

LAVENDER OIL (UNII: ZBP1YXW0H8)	
LONICERA JAPONICA FLOWER (UNII: 4465L2WS4Y)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CHLORPHENESIN (UNII: I670DAL4SZ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62839-1083-3	100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2009	
2	NDC:62839-1083-4	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/27/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	06/01/2009	

Labeler - NSE Products, Inc. (803486393)

Revised: 4/2020 NSE Products, Inc.