

ALCOSCRUB HAND CLEANER- alcohol liquid
Erie Scientific, LLC.

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.

Alcoscrub

Ethyl Alcohol 62 % v/v

For handwashing to decrease bacteria on skin.

Keep Out of Reach of Children

If swallowed, contact a physician or poison center

Indications & Usage Section

For hand cleaning to decrease bacteria on skin

Flammable. Keep away from heat of flames. For external use only. Do not use in the eyes.

Discontinue use if redness and irritation develop. If condition persists for more than 72 hours, consult a physician. If swallowed, contact a physician or poison control center. Keep out of the reach of children.

Dosage & Administration Section

Apply a small amount to palm. Briskly rub, covering hands with product until dry.

Inactive ingredients

Water, Glycerin, Carbomer, Propylene glycol, Isopropyl Myristate, Triethanolamine, Aloe Barbadensis Leaf, Tocopheryl acetate



ALCOSCRUB HAND CLEANER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55863-700
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	1 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Isopropyl Myristate (UNII: 0RE8K4LNJS)	
Aloe Vera Flower (UNII: 575DY8C1ER)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
Trolamine (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55863-700-04	118 mL in 1 BOTTLE		
2	NDC:55863-700-16	472 mL in 1 BOTTLE		
3	NDC:55863-700-27	800 mL in 1 POUCH		
4	NDC:55863-700-32	944 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/03/2009	

Labeler - Erie Scientific, LLC. (361605223)

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Erie Scientific, LLC.