

MICROBAN - alcohol gel
Noveko 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.

Active ingredient Purpose

70% V/V Ethyl alcohol.....Antiseptic

Uses

for handwashing to decrease bacteria on the skin

recommended for repeated use

Warnings

For external use only.

Flammable, keep away from fire or flame.

Do not use in the eyes. In case of contact, rinse eyes thoroughly with water.

Discontinue use if irritation and redness develop.

If condition persists for more than 72 hours, consult a doctor.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

wet hands thoroughly with product and allow to dry without wiping supervise children in the use of this product

Other Information

avoid temperature over 110F

may discolor certain fabrics

harmful to wood finishes and plastics

Inactive ingredients

Water, Aloe vera, Carbomer, Tocopheryl acetate, Triethanolamine, Glycerin, Fragrance.

Questions?

call 1 866 377-3030

weekdays, 9AM - 5PM EST

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ANTISEPTIC HAND SANITIZER

Microban
ORIGINAL

KILLS 99.9% OF GERMS
WATERLESS

Vitamin E and Aloe Vera enriched gel.
fl oz. (L/mL)

MICROBAN			
alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49955-600
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	0.7 L in 1 L
Inactive Ingredients			
Ingredient Name			Strength
WATER (UNII: 059QF0KO0R)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			

.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49955-600-10	1 L in 1 BOTTLE		
2	NDC:49955-600-13	1.3 L in 1 BOTTLE		
3	NDC:49955-600-50	0.5 L in 1 BOTTLE		
4	NDC:49955-600-25	0.25 L in 1 BOTTLE		
5	NDC:49955-600-60	0.06 L in 1 BOTTLE		
6	NDC:49955-600-12	0.125 L in 1 BOTTLE		
7	NDC:49955-600-02	.025 L in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333	11/18/2009	

Labeler - Noveko Inc (205614519)

Revised: 1/2010

Noveko Inc