LITTLE ONES BABY FRESH SANITIZER- ethyl alcohol liquid KMART CORPORATION

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.

DRUG FACTS

ACTIVE INGREDIENT

ETHYL ALCOHOL 65%

PURPOSE

ANTISEPTIC

USES

TO HELP DECREASE BACTERIA ON THE SKIN

WARNINGS

FOR EXTERNAL USE ONLY

- FLAMMABLE
- KEEP AWAY FROM SOURCE OF HEAT OR FIRE

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE THOROUGHLY WITH WATER

STOP USE AND ASK A DOCTOR IF

IRRITATION OR REDNESS DEVELOPS OR LASTS

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY. 1-800-222-1222

DIRECTIONS

- PUT ENOUGH PRODUCT IN YOU PALM TO COVER HANDS AND RUB HANDS TOGETHER UNTIL DRY
- CHILDREN UNDER 6 YEARS SHOULD BE SUPERVISED WHEN USING THIS PRODUCT

OTHER INFORMATION

- STORE AT A TEMPERATURE BELOW 110°F (43°C)
- MAY DISCOLOR CERTAIN FABRICS OR SURFACES

INACTIVE INGREDIENTS

WATER (AQUA), PROPYLENE GLYCOL, ISOPROPYL ALCOHOL, CARBOMER, AMINOMETHYL PROPANOL, BENZOPHENONE-4, GLYCERIN, FRAGRANCE (PARFUM), RED 33 (CI 17200)

QUESTIONS OR COMMENTS?

1-800-842-7886

LABEL COPY



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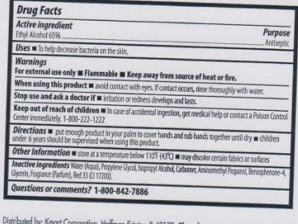


Little Ones

Hand Sanitizer

baby fresh

For adult use only. Not for use on infants



Distributed by: Kmart Corporation, Hoffman Estates, IL 60179, **Shop kmart.com** • Made in Conada Effective at eliminating over 99.89% of many common hermful germs and bacteria in as little as 15 seconds.

SATISFACTION GUARANTEE

If you are unsatisfied for any reason, return the unused portion to the store for a full refund or call 1-800-842-7886. 06-20465 8



LITTLE ONES BABY FRESH SANITIZER

ethyl alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:49738-413

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient NameBasis of StrengthStrengthALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)ALCOHOL650 mg in 1 mL

Inactive Ingredients Ingredient Name Strength WATER (UNII: 059QF0KO0R) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) ISOPROPYL ALCOHOL (UNII: ND2M416302) CARBOMER 934 (UNII: Z135WT9208)

AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
SULISOBENZONE (UNII: 1W6L629B4K)	
GLYCERIN (UNII: PDC6A3C0OX)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:49738-413- 02	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product					
2	NDC:49738-413- 08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product					
3	NDC:49738-413- 32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product					

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not final	part333E	04/27/2015					

Labeler - KMART CORPORATION (008965873)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment					
Name	Address	ID/FEI	Business Operations		
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture (49738-413)		

Revised: 4/2015 KMART CORPORATION