

HAND SANITIZER- ethyl alcohol gel
DIVERSIFIED MANUFACTURING CORP

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.

PURE FORMULA 1 HAND SANITIZER

ACTIVE INGREDIENT

ETHYL ALCOHOL 62%

PURPOSE

SANITIZER

USES

HAND SANITIZER

ALOE VERA

WARNINGS

FOR EXTERNAL USE ONLY.

*DO NOT GET INTO EYES.

FLAMMABLE - DO NOT USE NEAR HEAT, FIRE, FLAME, OR WHILE SMOKING.

IN CASE OF EYE CONTACT, FLUSH IMMEDIATELY AND THOROUGHLY WITH WATER.

*DO NOT INGEST. IF SWALLOWED, SEEK MEDICAL HELP OR CALL POISON CONTROL IMMEDIATELY.

*STOP USE AND ASK A DOCTOR IF REDNESS AND/OR IRRITATION OCCURS AND STAYS LONGER THAN 7 DAYS.

KEEP OUT OF REACH OF CHILDREN.

DIRECTIONS

PUT SMALL AMOUNT INTO PALM OF HAND. RUB TOGETHER UNTIL DRY.

INACTIVE INGREDIENTS

PURIFIED WATER, GLYCERIN, PROPYLENE GLYCOL, ISOPROPYL MYRISTATE, TOCOPHERYL ACETATE, ALOE VERA

Pure™

FORMULA 1

HAND SANITIZER

Aloe

*Kills 99.9% of
most common germs*

4 Fl. Oz. (118ml)
Product of the USA

Hand Sanitizer, Aloe Vera

Drug Facts

ACTIVE INGREDIENTS:
Ethyl Alcohol (52%).....Sanitizer

OTHER INGREDIENTS: Purified water, Glycerin, Propylene Glycol, Isopropyl Myristate, Tocopherol Acetate, Aloe Vera



DIRECTIONS: Put small amount into palm of hand. Rub together until dry.

WARNINGS: For External use only. ***Do not get into eyes.**
FLAMMABLE - Do not use near heat, fire, flame or while smoking. In case of eye contact, flush immediately and thoroughly with water.
***Keep out of reach of children.** *Do not ingest. If swallowed, seek medical help or call poison control immediately. *Stop use and ask a doctor if redness and/or irritation occurs and stays longer than 7 days.

HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62172-112
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62172-112-18	236.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/17/2020	
2	NDC:62172-112-14	118 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package	03/17/2020	
3	NDC:62172-112-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/17/2020	
4	NDC:62172-112-11	3785 mL in 1 JUG; Type 0: Not a Combination Product	03/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/17/2020	

Labeler - DIVERSIFIED MANUFACTURING CORP (185073996)

Revised: 3/2020

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