HAND SANITIZER- ethyl alcohol liquid
Everbrands, Inc. dba EverPure, 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.

Hand Sanitizer

Drug Facts

Active ingredients

Alcohol 80%

Purpose

Antiseptic

Uses Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap & water are not available.

Warnings
For external use only. Flammable. Keep away from heat or flame.

Do not use
• in children less than 2 months of age
• on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Other Information • Store between 15-30C (59-86F) • Avoid freezing & excessive heat above 40C (104F)

Directions •Place enough product on hands to cover all surfaces. •Rub hands together until dry

Inactive ingredients glycerin, hydrogen peroxide, purifed water USP

Product Label

H&™ by EverPure™

unscented

Hand

Sanitizer

99.9% effective

against most germs

antiseptic

4 FL. OZ.

visit everpurecare.com for more information

EverPure, Inc.
401 N. Oak St.
Inglewood, CA
90302

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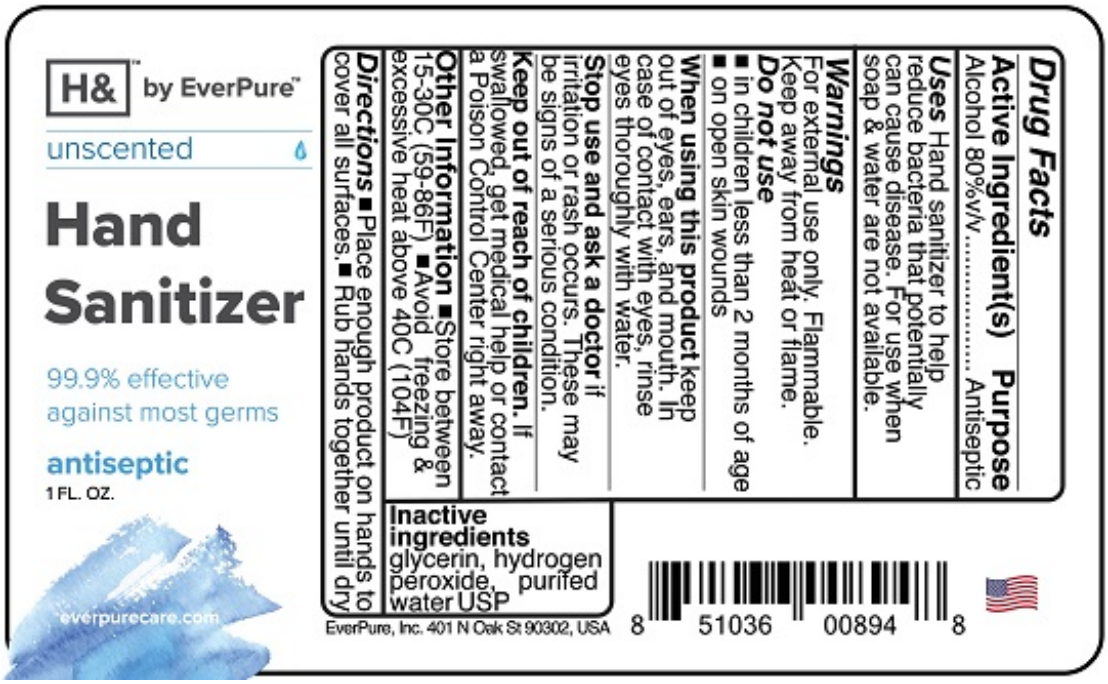
1 OZ



2 OZ

4OZ

res

HAND SANITIZER

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72655-021
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72655-021-04	29.5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/09/2020	
2	NDC:72655-021-05	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/09/2020	
3	NDC:72655-021-06	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/09/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/09/2020	

Labeler - Everbrands, Inc. dba EverPure, Inc. (080314845)

Registrant - Everbrands, Inc. dba EverPure, Inc. (080314845)

Establishment

Name	Address	ID/FEI	Business Operations
Everbrands, Inc. dba EverPure, Inc.		080314845	manufacture(72655-021)

Revised: 4/2020 Everbrands, Inc. dba EverPure, Inc.