INSTANT HAND SANITIZER- hand sanitizer solution
American Screening 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.

Active Ingredients

Ethyl Alcohol 70%

Purpose

Antibacterial

Use

For rinse-free hand sanitizing to remove 99.9% bacteria on skin.

Warnings

Flammable. Keep away from fire or flame. For external use only.

When using this product

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

Stop use and ask a doctor if

irritation and redness develop and persist.

Keep out of reach of children.

If swallowed, get medical health promptly

Directions

Wet hands thoroughly with product and allow to dry without wiping.

Other Information

Store under 105°F

Inactive Ingredients

Aloe Barbadenisis Leaf, Glycerin, Hydroxypropylcellulose, Vitamin E, Water

4 ounces (118 mL) - NDC 00000-000-00

8 ounces (236 mL) - NDC 00000-000-00

16 ounces (437 mL) - NDC 00000-000-00

INSTANT HAND SANITIZER

hand sanitizer solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73817-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 L in 100 L

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73817-100-04	0.118 L in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/04/2020	
2	NDC:73817-100-08	0.236 L in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/04/2020	
3	NDC:73817-100-16	0.473 L in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/04/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/04/2020	

Labeler - American Screening LLC (169065054)

Establishment

Name	Address	ID/FEI	Business Operations
American Screening LLC		169065054	manufacture(73817-100)

Revised: 12/2021 American Screening LLC