HAND SANITIZER- alcohol liquid True Value Company, L.L.C.

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

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

Directions: Place enough product on hands to cover all surfaces. Rub hands together until dry. Supervise children under 6 years of age when using this product to avoid swallowing.

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.

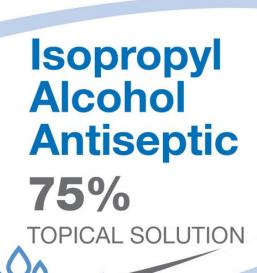
Inactive ingredients: glycerin, hydrogen peroxide, purified water USP

Use[s]: Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Active ingredient[s]: Isopropyl alcohol 75% v/v

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

Purpose: Antiseptic



Non-Sterile Solution
HAND SANITIZER

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1 GALLON (3.78L)

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HAND SANITIZER

alcohol liquid

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:73998-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL -	ISOPROPYL	0.75 mL
UNII:ND2M416302)	ALCOHOL	in 1 mL

Inactive Ingredients Ingredient Name Strength HYDRO GEN PERO XIDE (UNII: BBX060AN9V) WATER (UNII: 059QF0K00R) GLYCERIN (UNII: PDC6A3C0OX)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73998-001- 02	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2020	
2	NDC:73998-001- 04	1892 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2020	
3	NDC:73998-001- 03	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2020	
4	NDC:73998-001- 05	18927 mL in 1 PAIL; Type 0: Not a Combination Product	05/12/2020	
5	NDC:73998-001- 06	208198 mL in 1 DRUM; Type 0: Not a Combination Product	05/12/2020	
6	NDC:73998-001- 07	1040988 mL in 1 CONTAINER; Type 0: Not a Combination Product	05/12/2020	
7	NDC:73998-001- 08	1249 185 mL in 1 CONTAINER; Type 0: Not a Combination Product	05/12/2020	

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not final	part333A	03/26/2020					

Labeler - True Value Company, L.L.C. (006929681)

Registrant - True Value Company, L.L.C. (006929681)

Establishment						
Name	Address	ID/FEI	Business Operations			
True Value Company, LLC		006929681	manufacture(73998-001)			

Revised: 5/2020 True Value Company, L.L.C.