HANDS ON HYDRATING HAND SANITIZER- alcohol liquid
D-Time Limited Liability Company

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 Ingredient

Alcohol 79% v/v.

Purpose

Antiseptic skin cleanser

Use

For personal hand hygiene to help prevent the spread of bacteria

Warning

For external use only
When using this product avoid contact with eyes.
Flammable. Keep away from heat and flame.

Ask doctor

Stop use and consult a healthcare professional if irritation develops.

Keep out of reach of children.

If swallowed, contact a Poison Control Center or , get medical help right away.

Directions

Adults and children over 2 years: · For occasional and personal domestic use · Supervise children when they use this product · Spray onto hands and rub thoroughly for at least 30 seconds. Allow to dry.

Inactive ingredients

Water (Aqua), Glycerin, Hydrogen peroxide, Isopropyl alcohol, Bergamot leaf oil, Hyaluronic acid

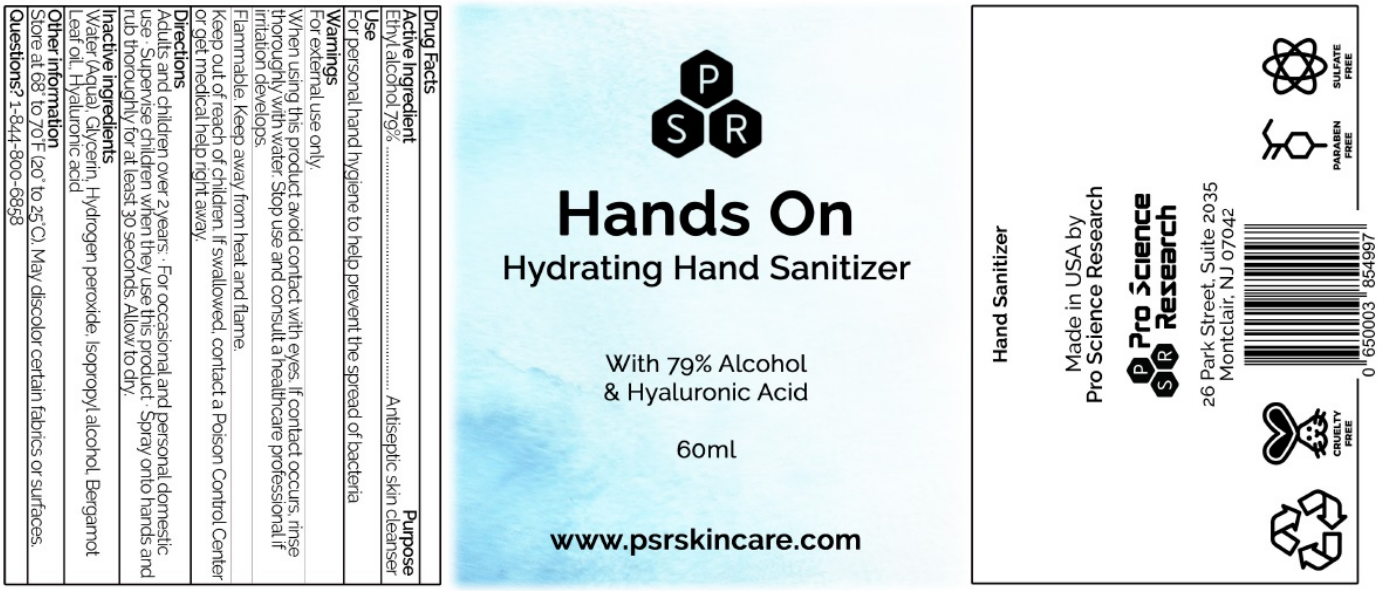
Other information

Store at 68 to 70 F (20 to 25 C). May discolor certain fabrics on surfaces

Questions?

1-844-800-6858

Package Label



HANDS ON HYDRATING HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75306-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	79 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
BERGAMOT OIL (UNII: 39W1PKE3JI)	
HYALURONIC ACID (UNII: S270N0TRQY)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:75306-001-01	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020	
2	NDC:75306-001-02	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020	
3	NDC:75306-001-	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination	04/01/2020	

3	03	Product	04/01/2020	
4	NDC:75306-001-04	750 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020	
5	NDC:75306-001-05	1000 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020	
6	NDC:75306-001-06	250 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/01/2020	

Labeler - D-Time Limited Liability Company (081728006)

Establishment

Name	Address	ID/FEI	Business Operations
D-Time Limited Liability Company		081728006	manufacture(75306-001)

Revised: 5/2020 D-Time Limited Liability Company