DR. DRIS HAND SANITIZER- ethyl alcohol liquid
Hanson Venture Enterprise 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.

Dr. Dri's Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl Alcohol 68% v/v

Purpose

Ethyl Alcohol 68% v/v................Antiseptic

Uses

Hand sanitizer to help reduce bacteria on the skin. For use when soap and water are not available.

Warnings

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

Do not use:

• on 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 the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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

Other Information

• store in a cool dry place. Do not overheat.

Inactive Ingredients

Purified Water, Hydroxyethyl Cellulose, Coconut Pulp Extract, Lavender Oil, Eucalyptus Oil, Aloe Vera Leaf Juice,

DR. DRIS HAND SANITIZER ethyl alcohol liquid			

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81383-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	2720 mL in 4000 mL

Inactive Ingredients	
Ingredient Name	**Strength**
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
COCONUT (UNII: 3RT3536DHY)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:81383-002-01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2021	
2	NDC:81383-002-02	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2021	
3	NDC:81383-002-03	475 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2021	
4	NDC:81383-002-04	4000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/26/2021	

Labeler - Hanson Venture Enterprise Inc (204263995)

Establishment

Name	Address	ID/FEI	Business Operations
Natures Formulae Health Products Ltd.		241385587	manufacture(81383-002)