STUVZ HAND SANITIZER- alcohol liquid
MR. HAND, 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.

Stuvz Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Uses

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

Warnings

Flammable. Keep away from fire or flame.

For external use only.

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.

Keep out of reach of children.

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

Do Not Use

- In children less than 2 months of age.
- On open skin wounds

Directions

- Place enough product in your palm to thoroughly cover your hands.
- Rub hands briskly until they are fully dry.

Other Information

- Store between 15-30°C (59-86°F).
- Avoid freezing and excessive heat above 40°C (104°F).

Inactive Ingredients

Water, hydrolized jojoba esters, glycerin, PEG-10 dimethicone, sodium hyaluronate, lavandula hybrida.

Stuvz Hand Sanitizer

With Hyaluronic Acid

5.7 FL. OZ/168 ML

STUVZ HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79283-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

GLYCERIN (UNII: PDC6A3C0OX)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
WATER (UNII: 059QF0KO0R)	
POLYETHYLENE GLYCOL 500 (UNII: 761NX2Q08Y)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
JOJOBA OIL, RANDOMIZED (UNII: 7F0EV20QYL)	
LAVANDIN OIL (UNII: 9RES347CKG)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79283-002-01	168 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/27/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/27/2020	

Labeler - MR. HAND, LLC (117563109)

Establishment

Name	Address	ID/FEI	Business Operations
Pure Source, LLC		080354456	manufacture(79283-002)

Revised: 10/2020 MR. HAND, LLC