SPAROOM HAND SANITIZER- alcohol liquid Natural Essentials, 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.

Drug Facts

Active Ingredients

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

• For handwashing to decrease bacteria on the 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 or redness develop and persist.

Keep out of reach of children.

If swallowed, get medical help promptly.

Directions

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

Other Information

Store under 105°F

Inactive Ingredients

Aloe Barbadensis Leaf, Dimethicone, Water. May contain fragrance.

Principal Display Panel – Bottle Label

sparoom®

HAND SANITIZER

70% Alcohol

Kills 99.99% of Common Germs

8.0 fl oz (236.6 mL)

Inactive Ingredients Aloe Barbadensis Leaf, Dimethicone, May contain fragrance.	Other Information Store under 105°F	Directions Wet hands thoroughly with product dry without wiping.	Stop use and ask a doctor if irritation or redness develop and p Keep out of reach of children. If swallowed, get medical help pro	mings mable. Keep away from xternal use only. n using this product of use in eyes. In case of rinse with water.	Uses •For handwashing to decre on the skin.	Drug Facts Active Ingredients Ethyl Alcohol 70%	Asparoom HAND SANITIZER	Ltd. / SpaRoom siand OH 44128 oom.com h U.S.& foreign components 8 50015 33395 8
thicone, Water.		oduct and allow to	n persist. en. p promptly.	n fire or flame. f contact with	decrease bacteria	2	Kills 99.99% of Common Germs 8.0 fl oz (236.6 mL)	Distributed by Unitrex Ltd. / SpaRoom 5060 Taylor Rd, Cleveland OH 44128 © 2020 Unitrex Questions? www.sparoom.com Made in the U.S.A with U.S.& foreign components

SPAROOM HAND SANI	ΓIZER				
alcohol liquid					
Product Information					
Product T ype	HUMAN OTC DRUG	Item Code (Source)		NDC:66902-645	
Route of Administration	TOPICAL				
Active Ingredient/Active Moi	ety				
Ingredi	ent Name		Basis of Strength		Strength
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)			Alcohol		$0.70\ mL$ in $1\ mL$
Inactive Ingredients					
	Ingredient Name				Strength
ALOE VERA LEAF (UNII: ZY81Z83H0)					
DIMETHICO NE (UNII: 92RU3N3Y1O)					
TRIISOPROPANOLAMINE (UNII: W91					
WATER (UNII: 059QF0KO0R)					
Packaging					

# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1 NDC:66902-645- 08	236.6 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/03/2020				
Marketing Information						
Marketing Catego	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not fin	al part333A	06/03/2020				

Labeler - Natural Essentials, Inc. (947484713)

Establishment

Name	Address	ID/FEI	Business Operations
NATURAL ESSENTIALS, INC.		947484713	MANUFACTURE(66902-645)

Revised: 6/2020

Natural Essentials, Inc.