ANTIMICROBIAL HAND SANITIZER - alcohol spray Hanover Pen Corp dba HPC Global

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.

Antimicrobial Hand Sanitizer

Active Ingredient

Ethyl Alcohol 66.5 percent

Purpose

Antiseptic

Uses

To decrease bacteria on the skin. Recommended for repeated use. For external use only. Flammable, keep away from fire or flame. Do not use in the eyes, if this happens rinse thoroughly with water. Stop use and ask a doctor if irritation of redness develops and persists for more than 72 hours.

Keep out of reach of children.

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

Directions

spray hands throughly with product allow to dry without wiping children under six should be supervised while using this product.

Inactive Ingredients

aloe vera, fragrance, purified water, triethanolamine Drug Facts

Drug Facts Active Ingredient Purpose	tizer Dz.				
Ethyl Alcohol 66.5%. Antiseptic Artiseptic	I Sani 30 fl. c Slobal 17331 SA				
<i>Warnings</i> For external use only. Flammable, keep away from fire or flame.	Hand Itrus HPC (HPC (r PA, 1 e in U;				
Do not use in the eyes, if this happens rinse thoroughly with water. Stop use and ask a doctor if irritation and redness develop & persists for more than 72 hours.					
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ANTIMICROBIAL HAN	D SANITIZER				
alcohol spray					
Product Information					
Product Type	HUMAN OTC DRUG	Ite m Cod	e (Source)		NDC:51811-455
Route of Administration	TOPICAL				
Active Ingredient/Active Moi	ety				
Ingredient Name Basis of		Basis of Streng	th	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL		5.985 mL in 9 mL
Inactive Ingredients					
	Ingredient Name				Strength
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 2	X)				
WATER (UNII: 059QF0KO0R)					
TROLAMINE (UNII: 903K93S3TK)					
Packaging					
			Marketing Star	÷	Marketing End

# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1 NDC:51811-455- 20	9 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/13/2010				
Marketing Information						
Marketing Categ	ory Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not	final part333E	09/13/2010				

Labeler - Hanover Pen Corp dba HPC Global (003022670)

Registrant - Robert Houghland (003022670)

Establishment

Name	Address	ID/FEI	Business Operations
Hanover Pen Corp dba HPC Global		003022670	repack(51811-455), label(51811-455)

Revised: 1/2020

Hanover Pen Corp dba HPC Global