SANIRUS HAND SANITIZER- ethyl alcohol gel
6A Holding, 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.

Sanirus Hand Sanitizer 65

Drug Facts

Active Ingredients

Ethyl Alcohol 65% v/v

Purpose

Antiseptic

Uses: Hand Sanitizer to help reduce bacteria on skin

***Warnings*: For external use only.**

Flammable. keep away from heat or flame.

When using this product • Avoid contact with eyes. In case of contact rinse with water. • Do not inhale or ingest. • Avoid contact with broken skin.

Stop use and ask a doctor if skin irritation develops.

Keep out of reach of children. In case of accidental ingestion, get medical help or contact a Poison Control Center immediately

Directions • Wet hands thoroughly and rub together until dry. • Adult supervision for children under 6. • Not for infants.

Other information • Store between 15°-30°C (59°-86°F). • May discolor fabrics

Inactive Ingredients: Water (Aqua), Butylene Glycol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Juice, Triethanolamine, Citric Acid.

KILLS MOST OF THE GERMS

65% ALCOHOL

ALOE VERA

2X SANITIZING STRENGTH

REFRESHING GEL

GMP Quality

***DISTRIBUTED BY:* 6A HOLDING LLC, MIAMI, FLORIDA**

MADE IN CHINA

Packaging

SANIRUS HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74691-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TROLAMINE (UNII: 9O3K93S3TK)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74691-002-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/23/2020	

Labeler - 6A Holding, LLC (117474000)

Revised: 6/2020 6A Holding, LLC