HAND SANITIZER GEL- ethyl alcohol gel
NV Labs

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.

Amber and Black HAND SANITIZER GEL (4 Fl. Oz & 16 Fl Oz.)

Active Ingredient

Active Ingredient Purpose

Ethyl Alcohol 70% v/v Antimicrobial

USES

- Hand Sanitizer to help reduce bacteria on the skin that could cause disease
- Recommended for repeated use

WARNING

Warnings

Flammable Keep away from Flame

For External Use only

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appeara and lasts

Keep out of reach of children, If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Directions

- Place enough product in your palm to thoroughly cover your hands
- Rub hands together briskly until dry
- Children under 6 years of age should be supervised when using this product

Other information

Other Information

- Store below 110F (43C) May discolor certain fabrics or surfaces

In active Ingredients

Water (Aqua) , Glycerin, Aloe Barbadensis Leaf Juice, Isopropyl Myristate, Tocopheryl Acetate, Carbomer, Triethanolamine, Fragrance (Parfum)

Manufactured By:
Questions or Comments

Manufactured By: Reforma Group, Southfield, Michigan 48033

Questions or comments? Call 1-248-358-9022

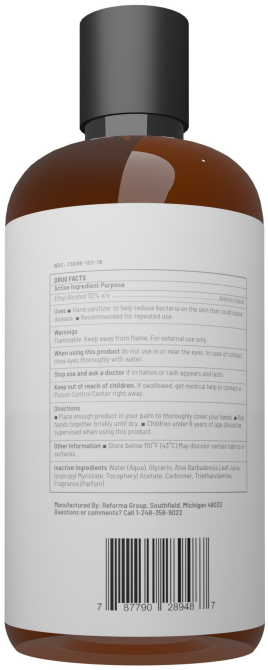
Keep out of Reach of Children

Keep out of Reach of Children. If swallowed , get medical help or contact a Poison Control Center right away.

Purpose

Antimicrobial

PRINCIPLE DISPLAY PANEL



Amber and Black

HAND SANITIZER GEL

MADE IN USA

NDC 73696-102-04

KILLS 99.99% of Germs

4 fl. oz (120 mL)

Amber and Black

HAND SANITIZER GEL

MADE IN USA

NDC 73696-102-16

KILLS 99.99% of Germs

16 fl.oz (473 mL)

HAND SANITIZER GEL

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73696-102
Route of Administration	TOPICAL		

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

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	29.278 mL in 100 mL
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	0.001 mL in 100 mL
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	0.35 mL in 100 mL
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	0.25 mL in 100 mL
TROLAMINE (UNII: 9O3K93S3TK)	0.12 mL in 100 mL
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.001 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73696-102-04	120 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/31/2020	
2	NDC:73696-102-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/31/2020	

Labeler - NV Labs (019662814)

Registrant - NV Labs (019662814)

Establishment

Name	Address	ID/FEI	Business Operations
NV Labs		019662814	manufacture(73696-102)

Revised: 1/2022 NV Labs