

ADVANCE HAND SANITIZER MOISTURIZING FORMULA WITH ALOE AND VITAMIN

E- alcohol gel

All Pharma 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.

Active ingredient

Ethyl Alcohol 70.0%

Purpose

Antimicrobial

Uses

To help decrease bacteria on the skin. Recommended for repeat use.

Warnings

For external use only.

Flammable. Keep away from heat and flame.

When using this product

- Keep out of eyes. In case of contact, rinse eyes thoroughly with water
- Avoid contact with broken skin
- Do not inhale or ingest

Stop use and ask a doctor if

If irritation or redness develops.

Keep out of reach of children.

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

Directions

- Wet hands thoroughly with product and allow to dry without wiping
- For children under 6, use only under adult supervision
- Not recommended for infants

Inactive ingredients

water, glycerin, propylene glycol, carbomer, fragrance, triethanolamine, isopropyl myristate, aloe barbadensis powder, tocopheryl acetate, FD&C Blue 1



bWell Products



ENRICHED WITH MOISTURIZERS
FOR SOFT HANDS

advanced hand sanitizer

NDC 53149-1120-8

TOPICAL ANTISEPTIC



Drug Facts

Active ingredient	Purpose
Ethyl Alcohol 70% w/w	Antiseptic

Use For use when soap and water are not available to help decrease bacteria on the skin. Recommended for repeat use.

Warnings For external use only

Flammable, keep away from fire or flame.

Do not use • On children less than 2 months of age • On open skin wounds

When using this product • Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water • Avoid contact with broken skin • Do not inhale or ingest

Stop use and consult a doctor • If irritation or rash occurs. These may be signs of a serious condition.

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

Directions • Place enough product on hands to cover all surfaces. Rub hands together until dry. • For children under 6, use only under adult supervision • Not recommended for infants

Other information • Store at 68-77°F (20-25°C) • May discolor some fabrics • Harmful to wood finishes and plastics

Inactive ingredients water, glycerin, propylene glycol, carbomer, fragrance, triethanolamine, isopropyl myristate, aloe barbadensis powder, tocopheryl acetate, FD&C Blue 1

Manufactured by:
All Pharma, LLC
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Hialeah, FL 33016

Questions or comments?
800-491-7908



8 FL OZ (236 mL)

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ADVANCE HAND SANITIZER MOISTURIZING FORMULA WITH ALOE AND VITAMIN E

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53149-1120
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)	
CARBOMER 980 (UNII: 4Q93RCW27E)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TROLAMINE (UNII: 9O3K93S3TK)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53149-1120-3	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	
2	NDC:53149-1120-1	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	
3	NDC:53149-1120-2	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	
4	NDC:53149-1120-4	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	
5	NDC:53149-1120-8	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	
6	NDC:53149-1120-5	3790 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/01/2020	

Labeler - All Pharma LLC (078572520)**Registrant** - All Pharma LLC (078572520)**Establishment**

Name	Address	ID/FEI	Business Operations
All Pharma LLC		078572520	manufacture(53149-1120)

