JOLO ACTIVE- alcohol gel
FUSION ACCELERATE, 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

Alcohol 62% v/v

Purpose

Antiseptic

Use

To sanitize hands without requiring water or a rinse – Kills 99.9% of most common bacteria in 15 seconds

Warnings

For external use only

Flammable

Do not use near heat or flame

Do not use

In 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 throughly with water.

Stop use and ask a doctor

if irritation or rash occurs.

They 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 atv 1-800-222-1222

Directions

Apply gel liberally to unsoiled hands and rub in thoroughly

Allow to dry without wiping or rinsing

Other information

- Store at room temperature.

Inactive ingredients

Water, Aloe barbadensis Leaf Juice, Carbomer, Diisopropylamine, Glycerin, Isopropyl Myristate, Fragrance, Phenoxyethanol, Tocopheryl Acetate, Yellow 10, Yellow 5, Blue 1.

Product label

JOLO ACTIVE

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79989-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
DIISOPROPYLAMINE (UNII: BR9JLI40NO)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79989-001-01	150 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/15/2020	

Labeler - FUSION ACCELERATE, LLC (081331914)

Revised: 9/2020 FUSION ACCELERATE, LLC