PURCOL MOISTURIZING HAND SANITIZER- alcohol gel
Purocol 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.

Purcol Moisturizing Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl alcohol 70% v/v

Purpose

Antimicrobial

Uses

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

Warnings

Flammable. Keep away from fire or 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 appears and lasts.

Keep out of reach of children.

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

Directions

- Place enough product in your palm to thoroughly cover your hands
- Rub hands together until dry
- No rinsing required
- No towels needed

Other Information

- Store below 104°F (40°C)
- May discolor certain fabrics or surfaces

Inactive Ingredients

Water, Glycerin, Aloe Barbadensis Leaf Extract, Hydroxypropyl cellulose, Lauryl lactate, Myristyl

lactate, Cetyl lactate, Fragrance.

Package Labeling:60ml

Package Labeling:251.3ml

Package Labeling:473.1ml

PURCOL MOISTURIZING HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75266-685
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
LAURYL LACTATE (UNII: G5SU0BFK7O)	
MYRISTYL LACTATE (UNII: 1D822OC34X)	
CETYL LACTATE (UNII: A7EVH2RK4O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75266-685-60	60 mL in 1 TUBE; Type 0: Not a Combination Product	04/29/2020	
2	NDC:75266-685-61	251.3 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/29/2020	
3	NDC:75266-685-62	473.1 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/29/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/29/2020	

Labeler - Purocol LLC (117479056)

Revised: 5/2020 Purocol LLC