

**TAYLOR AND FINCH INSTANT HAND SANITIZER ALOE VERA- alcohol gel
COGNESCENTI PTY. LTD.**

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.

hand sanitizer

Drug Facts

Active ingredient

Ethyl Alcohol 75%

Purpose

Antiseptic

USE

- hand sanitizer to help reduce bacteria on the skin.

Warnings

For external use only.

Flammable. Keep away from fire and flame.

When using this product

avoid contact with eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if

- irritation and redness develops.

Keep out of reach of children.

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

Directions

- Put enough on you palm to cover hands. Rub hands together briskly until dry.
- Children under 6 years of age should be supervised when using this product.

Other information:

- Store at 20° to 25° (68° to 77°).
- May discolor some fabrics or surface.

Inactive ingredients

Purified Water (Aqua), Cross Polymer, Triethanolamine, Parfum (Fragrance), Citronellol, Linalool, Lillial, Hexyl Cinnamic Aldehyde, Benzyl Salicylate, Hydroxycitronellal, D-Limonene, Geraniol, Cinnamyl Alcohol, Lyral.

package



TAYLOR AND FINCH INSTANT HAND SANITIZER ALOE VERA

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79978-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0K00R)	
TROLAMINE (UNII: 9O3K93S3TK)	
.ALPHA.-AMYL CINNAMALDEHYDE (UNII: WC51CA3418)	
GERANIOL (UNII: L837108USY)	
.ALPHA.-HEXYL CINNAMALDEHYDE (UNII: 7X6O37OK2I)	
HYDROXYCITRONELLAL (UNII: 8SQ0VA4YUR)	
CINNAMYL ALCOHOL (UNII: SS8YOP444F)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79978-006-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/31/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/31/2020	

Labeler - COGNESCENTIPTY. LTD. (745278551)

Establishment

Name	Address	ID/FEI	Business Operations
COGNESCENTIPTY. LTD.		745278551	manufacture(79978-006)

Revised: 7/2020

COGNESCENTIPTY. LTD.