

DECON HAND INSTANT HAND SANITIZER- alcohol liquid
Veltek Associates, Inc.

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.

Drug Facts

Purpose

Antiseptic Handwash

Active Ingredients

Ethanol 64.5 (w/w)

Uses

- For hand washing to decrease bacteria on the skin.
- Recommended for repeated use.

Warnings

Flammable, keep away from fire or flame when using this product

For External Use Only

- Do not use in eyes.
- In case of eye contact, flush with water for 15 minutes.
- Discontinue use if irritation and redness develop. If condition persists for more than 72 hours consult a doctor.
- Avoid contact with broken skin.

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.

Other Information

For Spill/Exposure Emergency Response Service in English, French and Spanish (and 23 other languages), call CARECHEM24 at 866-928-0789.

Inactive Ingredients

Acrylates/C10/30 Alkyl Acrylate Crosspolymer, EDTA, Fragrance, Glycerin, Purified Water and Tetrahydroxypropyl Ethylenediamine.

Questions?

Call 610-644-8335

Place holder text

DECON HAND INSTANT HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64307-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	64.5 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
EDETIC ACID (UNII: 9G34HU7RV0)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
EDETOL (UNII: Q4R969U9FR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64307-001-09	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/1999	
2	NDC:64307-001-10	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/1999	
3	NDC:64307-001-04	472 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/1999	
4	NDC:64307-001-06	472 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/1999	
5	NDC:64307-001-02	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/1999	
6	NDC:64307-001-05	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/1999	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/01/1999	

Labeler - Veltek Associates, Inc. (108844119)

Establishment

Name	Address	ID/FEI	Business Operations
Central Solutions Inc.		007118524	manufacture(64307-001) , label(64307-001) , pack(64307-001)

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Name	Address	ID/FEI	Business Operations
Veltek Associates, Inc.		108844119	manufacture(64307-001) , pack(64307-001) , label(64307-001)

Revised: 11/2018

Veltek Associates, Inc.