

DETOX PERFECT HAND GEL- alcohol gel
HUMEX,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
Active Ingredients

DETOX

Perfect Hand Gel

Waterless Hand Cleaner
Pure & Clean Hand Wash

Drug Facts

Active Ingredient

Ethyl Alcohol 65%

Purpose

Antimicrobial

Use Hand sanitizer to help reduce bacteria on the skin

Warnings

Flammable. Keep away from fire or flame.

This product is intended for external use only

When using this product, do not use in or near the eyes. In case of contact, rinse with water to remove.

Discontinue use if irritation or redness develops. If condition persists for more than 72 hours, consult a doctor.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

▪ Place enough product in your palm thoroughly cover your hands

▪ Rub hands together and wet until absorbed

- Rub hands together and pat until absorbed
- Children under 6 years of age should be supervised when using this product.

Other Information

- Store below 110°F (43°C)
- May discolor certain fabrics or surfaces

Inactive Ingredients

Water, Propylene Glycol, Dipropylene Glycol, Glycerin, Butylene Glycol, Sodium Hyaluronate, Hyaluronic Acid, Centella Asiatica Extract, Aloe Vera Leaf, Kale Leaf Extract, Avocado Fruit Extract, Celery Extract, Homopolymer, Hexacosyl Glycol, Trolamine

Questions or Comments?

Call 1-714-741-0088
www.humexlab.com

Distributed by Humex, Inc.
Brea, CA 92821-4824
Made in Korea

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USE

Hand sanitizer to help reduce bacteria on the skin

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70ml NDC: 76344-8004-1

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DETOX PERFECT HAND GEL

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76344-8004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	0.5 mL in 100 mL
CELERY (UNII: 44IDY6DTKX)	0.5 mL in 100 mL
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	0.01 mL in 100 mL
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.5 mL in 100 mL
HEXACOSYL GLYCOL (UNII: ULV0C76D5H)	0.01 mL in 100 mL
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	4 mL in 100 mL
TROLAMINE (UNII: 9O3K93S3TK)	0.01 mL in 100 mL
AVOCADO (UNII: SDS87L369F)	0.5 mL in 100 mL
CENTELLA ASIATICA (UNII: 7M867G6T1U)	0.5 mL in 100 mL
KALE (UNII: 0Y3L4J38H1)	0.5 mL in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	2 mL in 100 mL
DIPROPYLENE GLYCOL (UNII: E107L85C40)	4 mL in 100 mL
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	0.5 mL in 100 mL
HYALURONIC ACID (UNII: S270N0TRQY)	0.5 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76344-8004-2	70 mL in 1 TUBE; Type 0: Not a Combination Product	04/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/01/2020	

Labeler - HUMEX,INC. (017945311)**Establishment**

Name	Address	ID/FEI	Business Operations
MESO PHARM CO., LTD		695708527	manufacture(76344-8004)

Revised: 5/2020

HUMEX,INC.