HAND SANITIZER- alcohol gel
FARMACIA SAN ARCANGEL, S.A. DE C.V.

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

Ethyl alcohol v/v- 60-80%

Purposes

Antimicrobial

Uses

Hand sanitizer to decrease bacteria on the skin that could cause disease

Recommended for repetead use

Warnings

For external use only: hands

Flammable, keep away form fire and flame

When using this product

keep out of ayes, do not use in or near the eyes

In case of contact with eyes, flush thorougly with water

Avoid contact with broken skin

Do not inhale or ingest

When using this product

keep out of ayes, do not use in or near the eyes

In case of contact with eyes, flush thorougly with water

Avoid contact with broken skin

Do not inhale or ingest

Stop use and ask a doctor if

Irritation and redness develop

Condition persists for more than 72 hours

Keep out of reach of children

If swallowed, get medical help or contact a poison center right away

Directions

Put enough product in your palm to cover hands and rub hands together briskly until dry without wiping

for children under 6 years of age should be supervised when using

not recommended for infants

Other information

Store below 105 F 40C

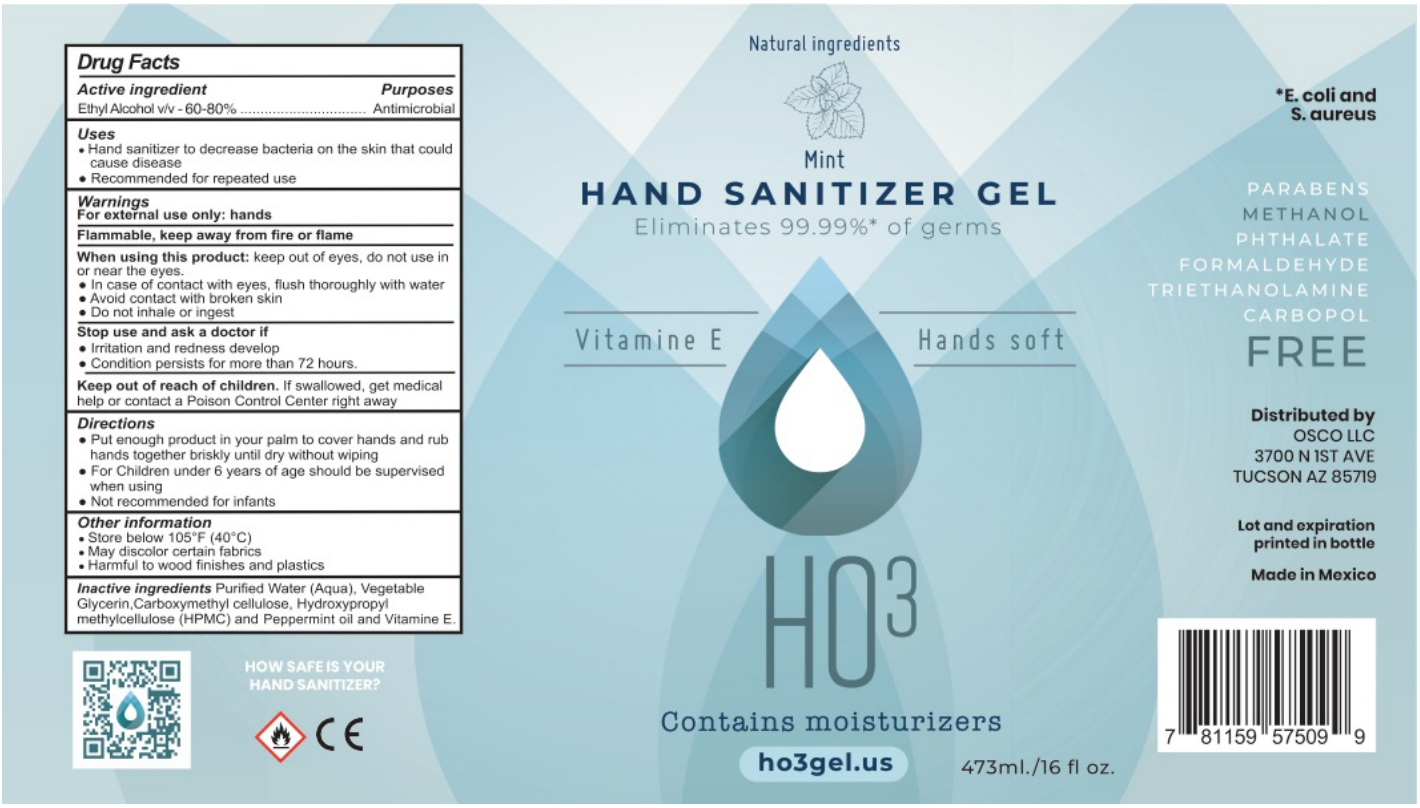
Mary discolour certain fabrics

harmful to wood finishes and plastics

Inactive ingredients

Purified water (aqua), vegetable glycerin, carboxymethyl cellulose, hydroxypropyl methylcellulose HPMC, and pepper mint oild and vitamine E

Principal display



Drug Facts

Active ingredient	Purposes
Ethyl Alcohol v/v - 60-80%	Antimicrobial

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

Warnings

For external use only: hands

Flammable, keep away from fire or flame

When using this product: keep out of eyes, do not use in or near the eyes.
- In case of contact with eyes, flush thoroughly with water
- Avoid contact with broken skin
- Do not inhale or ingest

Stop use and ask a doctor if
- Irritation and redness develop
- Condition persists for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Directions
- Put enough product in your palm to cover hands and rub hands together briskly until dry without wiping
- For Children under 6 years of age should be supervised when using
- Not recommended for infants

Other information
- Store below 105°F (40°C)
- May discolor certain fabrics
- Harmful to wood finishes and plastics

Inactive ingredients Purified Water (Aqua), Vegetable Glycerin,Carboxymethyl cellulose, Hydroxypropyl methylcellulose (HPMC) and Peppermint oil and Vitamine E.

HOW SAFE IS YOUR HAND SANITIZER?

Distributed by
OSCO LLC
3700 N 1ST AVE
TUCSON AZ 85719

Lot and expiration printed in bottle

Made in Mexico

Natural ingredients

Mint

HAND SANITIZER GEL

Eliminates 99.99%* of germs

Vitamine E

Hands soft

Contains moisturizers

*E. coli and S. aureus

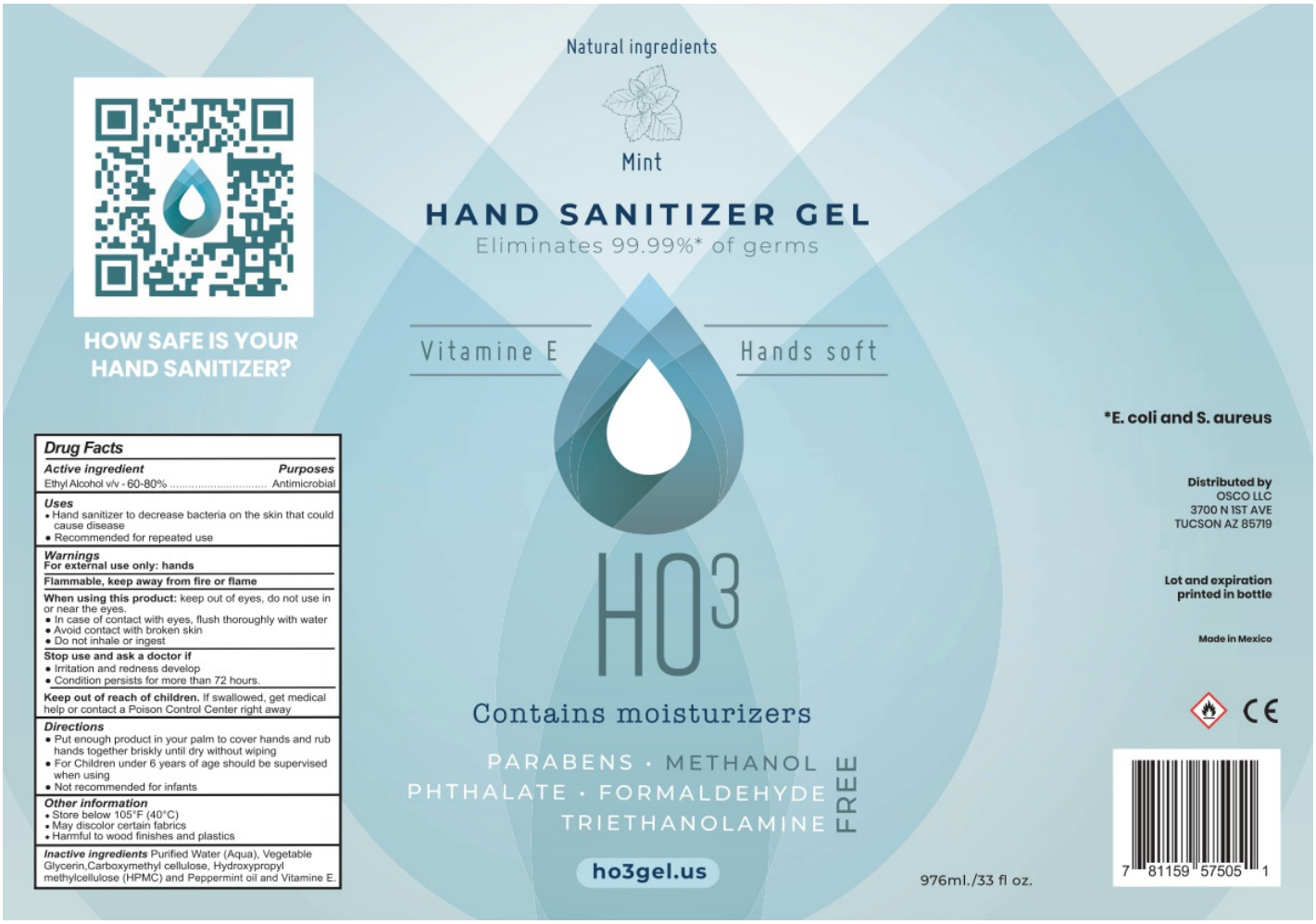
HO3

ho3gel.us

236ml./8 fl oz.

PARABENS
METHANOL
PHTHALATE
FORMALDEHYDE
TRIETHANOLAMINE
CARBOPOL
FREE

7 81159 57508 2



Natural ingredients

Mint

Hand sanitizer gel

eliminates 99.9% of germs

vitamine E hands soft

contain mosturizers

*e coli and S. areus

HO3

parabens

methanol

formaldehyde

triethanolamine

carbopol

free

HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80535-0001
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
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19X)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
HYPROMELLOSE 2208 (60000 MPA.S) (UNII: 2F7T07H9ZD)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PEPPERMINT OIL (UNII: AV092KU4JH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80535-0001-1	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/07/2020	
2	NDC:80535-0001-2	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/07/2020	
3	NDC:80535-0001-3	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/07/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/07/2020	

Labeler - FARMACIA SAN ARCANGEL, S.A. DE C.V. (951583068)

Registrant - FARMACIA SAN ARCANGEL, S.A. DE C.V. (951583068)

Establishment

Name	Address	ID/FEI	Business Operations
FARMACIA SAN ARCANGEL S.A. DE C.V.		951583068	label(80535-0001) , pack(80535-0001) , manufacture(80535-0001)

Revised: 9/2020 FARMACIA SAN ARCANGEL, S.A. DE C.V.