

HAND SANITIZER- hand sanitizer gel
TRI-COASTAL DESIGN GROUP 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.

TricsdlFrtressIvryOrngYllwlHndSntzrAlch62

Active ingredient

Alcohol 62%

Purpose

Antiseptic

Use

- For handwashing to decrease bacteria on the skin ■ Recommended for repeated use

Warnings

- **For external use only**
- **Flammable**, keep away from open flame and sources of heat
- Does not contain grain alcohol; do not drink. If taken internally will produce gastric disturbances.

When using this product

- Avoid the eyes and mucous membranes
- In the case of eyes or mucous membranes contact, rinse area thoroughly with water

Stop use and ask a doctor if

- Condition worsens
- Redness or irritation develops
- Condition persists for more than 3 days

Keep out of reach of children

- If swallowed, contact a doctor or Poison Control Center immediately

Directions

- Rub dime sized amount between hands until dry.
- Supervise children in the use of this product.
- In the case of eye contact, rinse eyes thoroughly with water.

Other information



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PEEL
 CPSIA#R830561-20220/20220706
 Batch# R82000692207060404 EXP:07/2022
 This product has been granted the Good Housekeeping Seal.
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Drug Facts

Active Ingredient Purpose
 Alcohol 62% Antiseptic

Use ■ For handwashing to decrease bacteria on the skin ■ Recommended for repeated use ▶

Drug Facts (continued)

Warnings
 ■ For external use only
 ■ Flammable, keep away from fire and flame
 ■ Does not contain grain alcohol; do not drink. If taken internally will produce serious gastric disturbances.

When using this product
 ■ Avoid the eyes and mucous membranes ▶

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Drug Facts (continued)

■ In the case of eyes or mucous membranes contact, rinse area thoroughly with water

Stop use and ask a doctor if
 ■ Condition worsens
 ■ Redness or irritation develops
 ■ Condition persists for more than 3 days

Keep out of reach of children ▶

Drug Facts (continued)

■ If swallowed contact a doctor or Poison Control Center immediately.

Directions
 ■ Rub dime sized amount between hands until dry.
 ■ Supervise children in the use of this product.
 ■ In the case of eye contact, rinse eyes thoroughly with water.

Other information
 ■ Store below 105 °F ▶

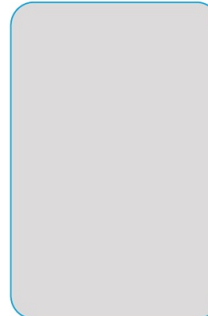
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Drug Facts (continued)

■ May discolor some fabrics.

Inactive ingredients
 Water, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Triethanolamine, Fragrance, PEG-40 Hydrogenated Castor Oil, FD&C YELLOW NO.5, D&C RED NO.33.

Questions?
 ■ 1-800-278-9218



Package Label



Drug Facts
(continued)

Warnings

- For external use only
- Keep this and all other products away from fire and flame
- Do not use on large areas of skin
- If used excessively, it may produce minor, painful, redness, dryness, and irritation

When using this product

- Avoid the eyes and mucous membranes

Drug Facts
(continued)

Directions

- Apply small amount between hands until dry
- Do not rub into skin of the face
- In the case of eye contact, rinse eyes thoroughly with water

Other information

- Store below 105 °F

Drug Facts
(continued)

Directions

- In the case of eye or mucous membranes contact, rinse area thoroughly with water

Stop use and ask a doctor if

- Condition worsens
- Redness or irritation develops
- Condition persists for more than 2 weeks

Keep out of reach of children

Drug Facts
(continued)

Warnings

- May irritate some fabrics

Inactive Ingredients

Hydroxyethyl Cellulose, Propylene Glycol, Carbomer, Triethyl Citrate, Sodium Hydroxide, Water, Ethanol, Glycerin, Fragrance, Benzyl Alcohol, Methylparaben, Propylparaben, Potassium Sorbate, Sodium Benzoate, Sodium Chloride, Sodium Citrate, Sodium Hydroxide, Sodium Lauryl Sulfate, Sodium Phosphate, Sodium Sulfate, Sodium Tartrate, Sodium Xanthate, Water, Xanthan Gum

Questions?

- 1-800-278-9218

Drug Facts
(continued)

Warnings

- May irritate some fabrics

Inactive Ingredients

Hydroxyethyl Cellulose, Propylene Glycol, Carbomer, Triethyl Citrate, Sodium Hydroxide, Water, Ethanol, Glycerin, Fragrance, Benzyl Alcohol, Methylparaben, Propylparaben, Potassium Sorbate, Sodium Benzoate, Sodium Chloride, Sodium Hydroxide, Sodium Lauryl Sulfate, Sodium Phosphate, Sodium Sulfate, Sodium Tartrate, Sodium Xanthate, Water, Xanthan Gum

Questions?

- 1-800-278-9218

Drug Facts
(continued)

Warnings

- May irritate some fabrics

Inactive Ingredients

Hydroxyethyl Cellulose, Propylene Glycol, Carbomer, Triethyl Citrate, Sodium Hydroxide, Water, Ethanol, Glycerin, Fragrance, Benzyl Alcohol, Methylparaben, Propylparaben, Potassium Sorbate, Sodium Benzoate, Sodium Chloride, Sodium Hydroxide, Sodium Lauryl Sulfate, Sodium Phosphate, Sodium Sulfate, Sodium Tartrate, Sodium Xanthate, Water, Xanthan Gum

Questions?

- 1-800-278-9218

HAND SANITIZER

hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49852-702
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
WATER (UNII: 059QF0K00R)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49852-702-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/30/2020	

Labeler - TRI-COASTAL DESIGN GROUP INC. (609734900)

Revised: 8/2020

TRI-COASTAL DESIGN GROUP INC.