

HAND SANITIZER- ethanol liquid

Tazza Brands East 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.

TAZZA INSTANT HAND SANITIZER FOAM

Drug Facts

Active ingredient

Ethanol 70%

Purpose

Antiseptic

Uses

- for hand sanitizing to decrease bacteria on the skin
- recommended for repeated use

Warnings

For external use only

Flammable. Keep away from fire or flame.

When using this product avoid contact with the eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation or redness develop

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

Directions

- apply liberally into palm of hand
- rub hands together until dry
- no need to rinse

Inactive ingredients Purified Water, Diethanolamine, Propylene Glycol, Glycerin, Aloe Barbadensis Leaf Juice

Manufactured for Terraboost Media, LLC, 3109 Grand Avenue #300, Miami, FL 33133

A Product of Terraboost® • Made in USA [Made in Taiwan] • www.tazzabrands.com

Tazza Instant Hand Sanitizer Foam

alcohol formula

Kills 99.99% of Germs

fragrance free

6,000 PUMPS

81.15 FL.OZ. (2,400mL)



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Directions • apply liberally into palm of hand

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Other information • do not store above 100°F (38°C)

• non-staining • may discolor some fabrics

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HAND SANITIZER

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76370-0003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
DIETHANOLAMINE (UNII: AZ E05TDV2V)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76370-0003-5	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/27/2015	
2	NDC:76370-0003-6	2400 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/27/2015	
3	NDC:76370-0003-7	300 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/23/2020	
4	NDC:76370-0003-8	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/23/2020	
5	NDC:76370-0003-9	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/23/2020	
6	NDC:76370-0003-1	10000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/02/2020	
7	NDC:76370-0003-3	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/02/2020	
8	NDC:76370-0003-2	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/02/2020	
9	NDC:76370-0003-4	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/02/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/27/2015	

Labeler - Tazza Brands East Inc. (117842371)

Revised: 5/2022

Tazza Brands East Inc.