

WIPE OUT ANTIBACTERIAL LIQUID HAND CUCUMBER MELON- benzalkonium chloride soap

TZUMI INNOVATIONS LLC

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.

77878-028, Style: WP0024 CUC

Active Ingredient(s)

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Use

For hand washing to decrease bacteria on the skin

Warnings

For external use only

When using this product do not get into eyes. If contact occurs, rinse eyes thoroughly with water

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

- Wet hands
- Apply palmful to hands
- Scrub thoroughly for at least 15 seconds
- Rinse thoroughly and dry

Inactive ingredients

Water, Cocamidopropyl dimethylamine oxide, Citric Acid, Fragrance, Kathon, Glycerin, Fatty acyl diethanol amine, Disodium EDTA, Hydroxyethyl Cellulose (250HHR), CI 51436, Blue 1 (CI 42090)

FILE NAME: WP0024-CUC Wipe-Out Hand Soap PKG		COLORS
BRAND: Wipe-Out	DIMENSIONS:	 4 Color Process PANTONE PANTONE PANTONE
ITEM #: WP0024-CUC	COMMENTS: • 4 COLOR PROCESS (CMYK) PRINTING • All measurements in mm	
PKG TYPE: Clear Label	MATERIALS: White Sticker	To FTY: please make adjustments as necessary
DATE: 11/30/20	DO NOT PRINT:  Die Line	REVISION DETAILS:
DESIGNER: Chloe	Metallic Foil Print:	

ACTUAL SIZE SHOWN



WIPE OUT ANTIBACTERIAL LIQUID HAND CUCUMBER MELON

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77878-028
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
DISODIUM HEDTA (UNII: KME849MC7A)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77878-028-01	221 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/23/2020	

Labeler - TZUMI INNOVATIONS LLC (117426322)**Establishment**

Name	Address	ID/FEI	Business Operations
Aogrand International Trade Corporation		421353092	manufacture(77878-028)

Revised: 12/2020

TZUMI INNOVATIONS LLC