MEDI WIPER HAND SANITIZING WIPES- benzalkonium chloride liquid Timk Inc.
Active Ingredient
Benzalkonium Chloride 0.1%
Inactive ingredients
Water, Phenoxyethanol, Glycerin, Sodium Benzoate, Polysorbate 20, Disodium EDTA, Tocopheryl Acetate, Citric Acid
Purpose
Antiseptic
Warnings
For external use only
Stop use and ask a doctor if ■ hypersensitivity symptoms such as erythema, itching and dermatitis happen ■ skin irritation happens
Do not use ■ in combination with soap or antibacterial cleansing agents ■ the product for a long time in the same area as swelling, inflammation or sickness may occur due to absorption through the skin
When using this product ■ avoid using repeatedly in the same area, skin irritation may occur ■ avoid getting into the eyes (if contact occurs, wash well with clean water) ■ if following abnormal symptoms persist, discontinue use: Irritation around the eyes, ears, mucous membranes, including the mouth, skin irritation and rashes
Keep out of reach of children

■ If swallowed, get medical help or contact a Poison Control Center right away

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Use

■ Instant hand antiseptic to decrease bacteria on the skin

Directions

- Clean with wipes and let dry
- Do not flush

Other information

- Read the directions and warnings before use
- Avoid freezing and excessive heat above 40°C (104°F)



MEDI WIPER HAND SANITIZING WIPES benzalkonium chloride liquid Product Information Product Type HUMAN OTC DRUG Item Code (Source) Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g	

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
POLYSORBATE 20 (UNII: 7T1F30V5YH)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
GLYCERIN (UNII: PDC6A3C0OX)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)				
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83930- 010-02	108 g in 1 CONTAINER; Type 0: Not a Combination Product	12/01/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	12/01/2024	

Labeler - Timk Inc. (121537031)

Registrant - Timk Inc. (121537031)

Establishment				
Name	Address	ID/FEI	Business Operations	
NAICO		694725335	manufacture(83930-010)	

Revised: 9/2024 Timk Inc.