PULLIO HAND SANITIZING WIPES PEPPERMINT- benzalkonium chloride liquid
NAICO

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.

Drug Facts

ACTIVE INGREDIENT

Benzalkonium Chloride 0.1%

INACTIVE INGREDIENTS

Water, Glycerin, Sodium Benzoate, Ethylhexylglycerin, Polysorbate20, Disodium EDTA, Citric Acid, Tocopheryl Acetate, Fragrance

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

■ aviod 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, under the skin irritation and rashes

KEEP OUT OF REACH OF CHILDREN

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

Uses

■ 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 40C (104F)

PACKAGE LABEL: Pullio hand sanitizing wipes (peppermint) / 20ct

PACKAGE LABEL: Pullio hand sanitizing wipes (peppermint) / 60ct

PULLIO HAND SANITIZING WIPES PEPPERMINT

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75536-0007
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)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
Ethylhexylglycerin (UNII: 147D247K3P)	
Polysorbate 20 (UNII: 7T1F30V5YH)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75536-0007-1	125 g in 1 CONTAINER; Type 0: Not a Combination Product	06/01/2020	
2	NDC:75536-0007-2	390 g in 1 CONTAINER; Type 0: Not a Combination Product	06/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/01/2020	

Labeler - NAICO (694725335)

Registrant - NAICO (694725335)

Establishment

Name	Address	ID/FEI	Business Operations
NAICO		694725335	manufacture(75536-0007)

Revised: 6/2020 NAICO