TOUCHPOINT WIPES SANITIZING WIPES- benzalkonium chloride cloth
Innocore Sales & Marketing Inc

Touchpoint Wipes
Sanitizing Wipes

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13 %

Purpose

Antimicrobal

Use

Hand sanitizer to help reduce bacteria on the skin.

Recommended for repeated use.

Warnings

For external use only.

When using this product*do not use in or near the eyes. In case of contact, rinse eyes thoroughly in water.*

Discontinue use if*irritation and redness develop. If conditions persist for more than 72 hours, consult a physician.*

If swallowed get*medical help or contact a poison control center immediately.*

Keep out of reach of children.

Directions

- *Wet hands thoroughly with product and allow to dry.*
- ***Children under 6 years of age should be supervised when using this product.***
- *Be sure to use entire wipe.*
- *Discard after single use.*

Inactive ingredients

Benzoic Acid, Caprylyl/Capryl Oligoglucoside, Dehydroacetic Acid, Phenoxyethanol, Poly(Laurylglucoside)-7, Propylene Glycol, Water

1500 Wipe Pouch Label

Touch
Point™

Sanitizing Wipes

1500 Wipes • 8"x 6"

Kills 99.99%
of most common germs that may cause illness

DO NOT REMOVE ROLL FROM BAG

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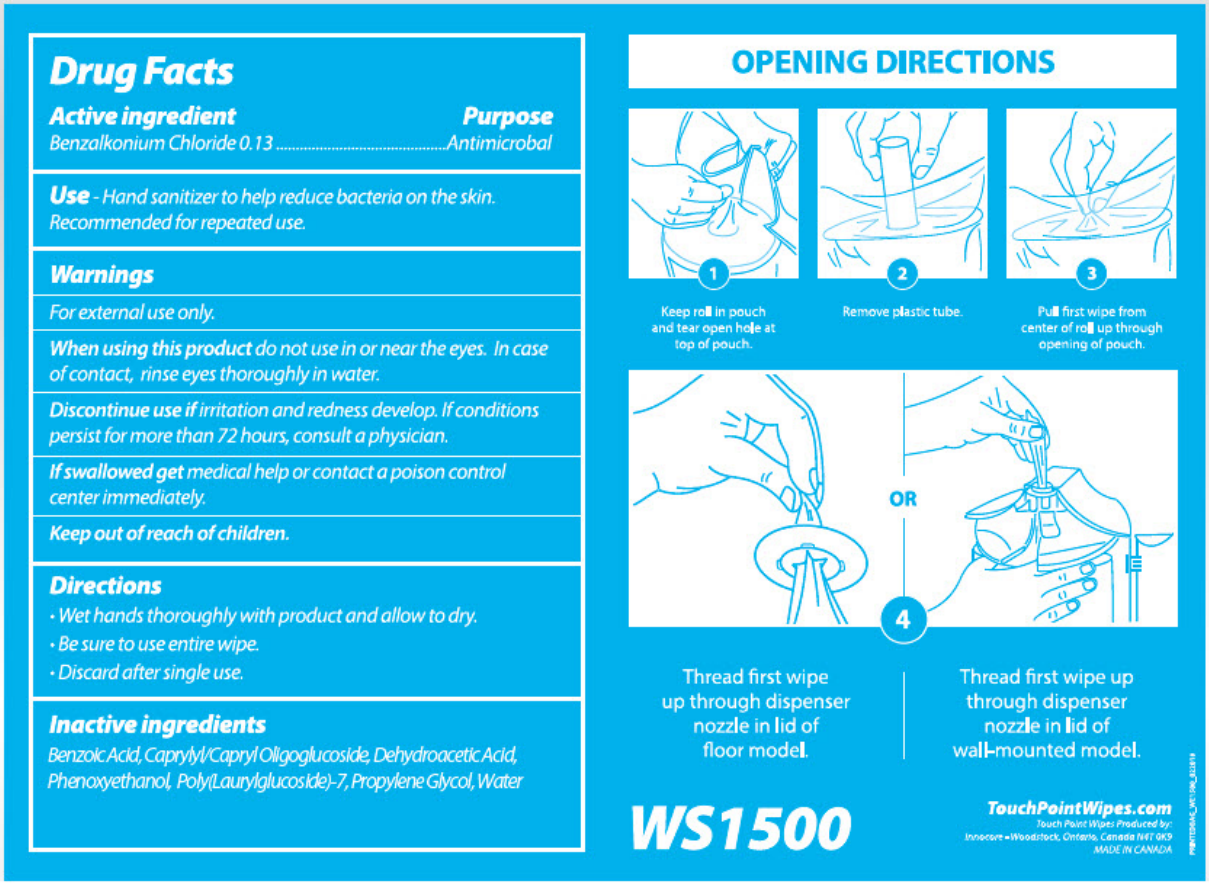

TOUCHPOINT WIPES SANITIZING WIPES

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70924-001
Route of Administration	CUTANEOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)	
POLY(LAURYLGLUCOSIDE)-7 (UNII: VB00RDE21R)	

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70924-001-02	2 in 1 BOX	09/09/2016	
1	NDC:70924-001-01	1500 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/09/2016	

Labeler - Innocore Sales & Marketing Inc (201152597)

Revised: 12/2023 Innocore Sales & Marketing Inc