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

Touchpoint Wipes Fragrance Free

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,

1500 Wipe Pouch Label

Touch

Point™

Sanitizing Wipes

1500 Wipes • 8"x 6"

Fragrance Free

Kills 99.99%

of most common germs that may cause illness

DO NOT REMOVE ROLL FROM BAG

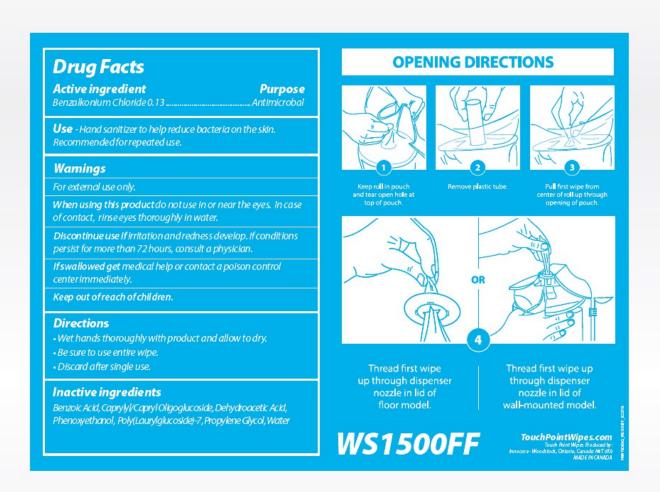
FRAGRANCE FREE



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TOUCHPOINT WIPES FRAGRANCE FREE SANITIZING WIPES benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70924-002
Route of Administration	CUTANEOUS		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
	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)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:70924-002- 02	2 in 1 BOX	09/09/2016		
1	NDC:70924-002- 01	1500 in 1 POUCH; Type 0: Not a Combination Product			
2	NDC:70924-002- 04	2 in 1 BOX	12/11/2018		
2	NDC:70924-002- 03	1800 in 1 POUCH; Type 0: Not a Combination Product			
3	NDC:70924-002- 06	2 in 1 BOX	12/11/2018		
3	NDC:70924-002- 05	2400 in 1 POUCH; Type 0: Not a Combination Product			
4	NDC:70924-002- 09	2 in 1 BOX	12/11/2018		
4	NDC:70924-002- 07	400 in 1 POUCH; Type 0: Not a Combination Product			
5	NDC:70924-002- 11	6 in 1 BOX	12/11/2018		
5	NDC:70924-002- 10	320 in 1 CANISTER; Type 0: Not a Combination Product			
6	NDC:70924-002- 13	6 in 1 BOX	10/16/2019		
6	NDC:70924-002- 12	150 in 1 CANISTER; Type 0: Not a Combination Product			
7	NDC:70924-002- 15	2 in 1 BOX	11/11/2019		
7	NDC:70924-002- 14	2200 in 1 POUCH; Type 0: Not a Combination Product			
8	NDC:70924-002- 17	2 in 1 BOX	09/21/2022		
8	NDC:70924-002- 16	650 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