MAXSANI HAND SANITIZING AND CLEANING WIPES- benzalkonium chloride cloth DYNAMIC WIPE INDUSTRIES LTD.

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.

MaxSani Hand Sanitizing & Cleaning Wipes

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Use:

Hand sanitizer to help reduce bacteria on the skin. For use when soap and water not available.

Warnings

For external use only

When using this product

- Avoid getting into the eyes.
- In case of eye contact immediately flush eyes thoroughly with water.

Stop use and ask a doctor if

- Irritation or redness develops
- Condition persist for more than 72 hours
- Redness is present

Keep out of reach of children.

In case of accidental ingestion, contact a doctor or Poison Control Center immediately.

Directions:

- Throughly wipe hands with cloth
- Rub hands together until dry
- No rinsing required

Other information

• Dispose of properly, do not flush.

Inactive Ingredients

Water, Polysorbate 20, Glycerin, Potassium Sorbate, Aloe (Aloe Barbadensis) Leaf Extract, Dehydroacetic Acid, Phenoxyethanol, Citric Acid

Package Labeling:20wipes



Package Labeling:50wipes





Package Labeling:400wipes



MAXSANI HAND SANITIZING AND CLEANING WIPES benzalkonium chloride cloth Product Information Product Type HUMAN OTC DRUG Route of Administration HUMAN OTC DRUG TOPICAL

Active Ingredient/Active Moiety						
Ingredient Name	Basis of Strength	Strength				
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL				

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
POLYSORBATE 20 (UNII: 7T1F30V5YH)			
GLYCERIN (UNII: PDC6A3C0OX)			
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
DEHYDRO ACETIC ACID (UNII: 2KAG279R6R)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)			

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:77645- 003-01	20 in 1 PACKAGE	09/15/2020			
1		3 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)				
2	NDC:77645- 003-02	50 in 1 PACKAGE	09/15/2020			
2		3 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)				
3	NDC:77645- 003-03	80 in 1 CANISTER	09/15/2020			
3		3 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)				
4	NDC:77645- 003-04	400 in 1 CANISTER	09/15/2020			
4		3 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	09/15/2020		

Labeler - DYNAMIC WIPE INDUSTRIES LTD. (532208402)