HYGENIX AF WIPE- benzalkonium chloride cloth
GM Industrial, Inc.

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

Antiseptic

For hand sanitizing to decrease bacteria on the skin

Recommended for repeated use

For external use only.

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation or redness develops, or if condition persists for more than 72 hours.

Keep out of react of children

In case of accidental ingestion, seek medical attention or contact a Poison Control Center immediately.

Directions

Wet hands thoroughly with product.

Discard wipe in trash receptacle after use. Do not flush.

Rub hands together briskly until dry.

Other information

Store in a cool dry place below 104 °F

Water, C9-11 Pareth-6, Cocamidopropyl PG-dimonium chloride phosphate, Dihydroxyethyl cocamine oxide, Acetamidoethoxyethanol, Citric acid.

90 CT 74146-243-93

74146-243-85

HYGENIX AF WIPE

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74146-243
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		

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

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	
ACETAMIDOETHOXYETHANOL (UNII: LVX2APC4XR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74146-243-96	6 in 1 BOX	06/17/2021	
1	NDC:74146-243-93	647.3 mL in 1 CANISTER; Type 0: Not a Combination Product		
2	NDC:74146-243-86	4 in 1 BOX	06/17/2021	
2	NDC:74146-243-85	810.1 mL in 1 CANISTER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/17/2021	

Labeler - GM Industrial, Inc. (025827197)

Establishment

Name	Address	ID/FEI	Business Operations
GM Industrial, Inc.		025827197	manufacture(74146-243) , label(74146-243) , pack(74146-243) , relabel(74146-243) , repack(74146-243)

Revised: 6/2021 GM Industrial, Inc.