GIANT EAGLE ANTIBACTERIAL HAND WIPES- benzalkonium chloride cloth Giant Eagle

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.

Giant Eagle Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

- for hand sanitizing to decrease bacteria on the skin.
- recommended for repeated use

Warnings

For external use only.

When using this product

avoid contact with eyes. In case of 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 the reach of children.

If swallowed, get medical help or contact Poison Control Center right away (1-800-222-1222)

Directions

- adults and children 2 years and over: apply to hands; allow to dry without wiping.
- children under 2 years: ask a doctor before use.

Other information

- store at room temperature
- keep closed tightly.
- may discolor certain fabrics or surfaces

Inactive ingredients

Purified Waterr, SD Alcohol 40, Phenoxyethanol, Decyl Glucoside, Potassium Sorbate, Sodium Benzoate, Disodium EDTA, Citric Acid, Aloe Barbadensis (Aloe) Leaf Extract, Fragrance.

Principal Display Panel

giant eagle

Antibacterial

HAND

WIPFS

FRESH SCENT

40 WIPES



Denzalkonium chloride cloth Product Information Product Type HUMAN OTC DRUG Item Code (Source) Route of Administration Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength

BENZALKONIUM 0.13 a

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -

UNII:7N6JUD5X6Y) CHL	.ORIDE	in 100
----------------------	--------	--------

Inactive Ingredients			
Ingredient Name	Strength		
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)			
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)			
WATER (UNII: 059QF0KO0R)			
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
ALCOHOL (UNII: 3K9958V90M)			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:56194-575- 40	40 in 1 PACKAGE; Type 0: Not a Combination Product	06/22/2022	

Marketing In	Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	06/22/2022		

Labeler - Giant Eagle (929989007)

Revised: 11/2022 Giant Eagle