BIOCENTRIS E2 HANDWASH- benzalkonium chloride liquid Biocentris Pharmaceuticals, LLC

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.

Biocentris E2 Foam Handwash

Drug Facts

Active ingredient

Benzalkonium chloride 0.13% w/w

Purpose

Antiseptic

Uses

- Handwash to help reduce bacteria on the skin that potentially can cause disease.
- Recommended for repeated use.

Warnings

For external use only.

When using this product keep away from eyes. In case of eye contact, flush eyes with water.

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

Keep out of reach of children. If swallowed, contact a physician or poison control center.

Directions

- Dispense a golf ball size amount of foam onto dry hands and rub vigorously for 15 seconds
- Rinse with water and towel dry.

Inactive ingredients Water, Ethanol, Cocamidopropyl PG-Dimonium Chloride Phosphate, Phenoxyethanol, Polyquatemium 11, Decyl Glucoside, Cetrimonium Chloride, Polyquatemium-10, Chlorhexidine Gluconate

Manufactured for:

biocentris PHARMACEUTICALS

24 Executive Parkway, Ringwood, NJ 07456

888-497-4375

info@biocentris.com • www.biocentris.com

E2 FOAM HANDWASH FOR FOOD INDUSTRY PROFESSIONALS

One Step Hand Wash & Sanitizer

KILLS 99.99% of most germs

NET CONTENTS: 33.8 FL. OZ. (1000 ML)

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LAA125M

BIOCENTRIS E2 HANDWASH

benzalkonium chloride liquid

Product Information	Produ	ıct lı	nforr	natior
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:81232-402

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - CHLORIDE UNII: 7N6JUD5X6Y) BENZALKONIUM CHLORIDE UNII: F5UM2KM3W7) (BENZALKONIUM - CHLORIDE UNII: 7N6JUD5X6Y)

Inactive Ingredients				
Ingredient Name	Strength			
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)				
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E)				
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)				
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)				
ALCOHOL (UNII: 3K9958V90M)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
POLYQUATERNIUM-10 (1000 MPA.S AT 2%) (UNII: GMR4PEN8PK)				
POLYQUATERNIUM-11 (1000000 MW) (UNII: 0B44BS5IJS)				
WATER (UNII: 059QF0KO0R)				

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:81232- 402-03	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018				
2	NDC:81232- 402-10	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018				
3	NDC:81232- 402-01	1000 mL in 1 POUCH; Type 0: Not a Combination Product	04/20/2021				
4	NDC:81232- 402-17	532 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2021				

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333E	01/01/2018				

Revised: 6/2022 Biocentris Pharmaceuticals, LLC