

BENZALKONIUM CHLORIDE- benzalkonium chloride cloth
ASP Global, 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.

BENZALKONIUM CHLORIDE

Drug Facts

Active Ingredients

Benzalkonium chloride 0.13%

Purpose

Antiseptic

Use

for preparation of the skin prior to an injection.

Warnings

For external use only. Flammable, keep away from fire or flame. Do not use with electrocautery procedures or near eyes. Stop use and ask a doctor if irritation or redness develops.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

Apply topically as needed to clean intended area. Discard after single use.

Inactive Ingredients

Aloe Barbadensis Leaf Juice, Disodium EDTA, DMDM Hydantoin, Ethylparaben, Methylparaben, Phenoxyethanol, Polysorbate 20, Sodium Citrate, Water

PRINCIPAL DISPLAY PANEL - 10 Applicator Package

BaylorScott&White

HEALTH

OPEN



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Wipe Size 15 x 20 cm

Manufactured for:
ASP Global, LLC
 7800 Third Flag Parkway,
 Austell, GA 30168, USA
 Made in China

BENZALKONIUM CHLORIDE

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59448-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (Benzalkonium - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
Water (UNII: 059QF0KO0R)	
ETHYLPARABEN (UNII: 14255EXE39)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59448-010-01	10 in 1 PACKAGE	02/28/2022	
1		6.2 g in 1 APPLICATOR; 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	part333E	02/28/2022	

Labeler - ASP Global, LLc (080361159)**Registrant** - ASP Global, LLc (080361159)

Revised: 3/2022

ASP Global, LLc