DIASINC- benzalkonium chloride spray
PureTek Corporation

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.

DiaSinc

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Uses

■ a no-rinse topical antiseptic that protects against bacterial contamination

Warnings

For external use only

Do not use on

■ deep or puncture wounds ■ animal bites ■ serious burns

When using this product

■ do not get into eyes ■ If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if

■ If irritation or redness develops ■ condition persists for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

■ spray soiled and/or odorous (body and/or perineal) areas ■ gently wipe clean ■ repeat as necessary until all soils are removed and skin is clean ■ pat dry (no rinsing necessary)

Other information

■ Store at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature]. ■ protect from freezing ■ avoid excessive heat

Inactive ingredients

aloe barbadensis (aloe vera) leaf juice, aqua (purified water), butylene glycol, disodium EDTA, fragrance, GenRx Complex® [consisting of: bisabolol, calcium pantothenate (vitamin B5), Carthamus tinctorius (safflower) oleosomes, maltodextrin, niacinamide (vitamin B3), pyridoxine HCl (vitamin B6), silica, sodium ascorbyl phosphate (vitamin C), sodium starch octenylsuccinate, tocopheryl acetate (vitamin E), Zingiber officinale (ginger) root extract], glycerin, phenoxyethanol, polysorbate 20, sodium hyaluronate, sodium hydroxide.

DiaSinc™

DIASINC

benzalkonium chloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59088-596
Route of Administration	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**
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	

ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
LEVOMENOL (UNII: 24WE03BX2T)	
CALCIUM PANTOTHENATE (UNII: 568ET80C3D)	
CARTHAMUS TINCTORIUS SEED OLEOSOMES (UNII: 9S60Q72309)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GINGER (UNII: C5529G5JPQ)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
WATER (UNII: 059QF0KO0R)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59088-596-16	237 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/24/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/24/2023	

Labeler - PureTek Corporation (785961046)

Revised: 3/2023 PureTek Corporation