DR. PIERCING AFTERCARE- benzalkonium chloride swab
Broadway Products 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.

BROADWAY PRODUCTS (as PLD) - Dr. Piercing - Aftercare Swab (81926-001)

ACTIVE INGREDIENTS

Benzalkonium Chloride 0.11%

Purpose

Antiseptic

Use

FDA Registered antiseptic for the care and cleaning of piercings.

WARNINGS:

For external use only. If redness, irritation, infection or swelling occurs, discontinue use and consult a physician. Keep out of eyes. Do not insert into the ear canal. Store at room temprature.

Keep out of reach of children.

Directions:

1. Hold swab with the colored round marker end up.
2. Bend the colored round marker tip until it "snaps".
3. Liquid flows down into the white tip.
4. Apply medicated tip to piercing. Dispose.

OTHER INGREDIENTS

Isotonic Saline Solution, Aloe Vera, Allantoin, Panthenol/Pro-Vitamin B5.

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DISTRIBUTED BY: Broadway Products LLC, Sellersburg, Indiana.
LEARN MORE: www.drpiercing.com

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Dr. Piercing Afterca...roduct, 36 Swab Piece New

DR. PIERCING AFTERCARE

benzalkonium chloride swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81926-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.1 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
PANTHENOL (UNII: WV9CM0O67Z)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
CUPRIC NITRATE (UNII: 9TC879S2ZV)	
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
TROLAMINE (UNII: 9O3K93S3TK)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ALLANTOIN (UNII: 344S277G0Z)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81926-001-36	36 in 1 CASE	01/01/2021	
1		0.15 mL 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	part333A	01/01/2021	

Labeler - Broadway Products LLC (002672936)

Revised: 7/2023 Broadway Products LLC