

NASOCLENZ- benzalkonium chloride 0.13% gel
Silicon Valley Innovations, Inc.

NasoClenz Cleansing Kit Drug Facts

Active Ingredient

Benzalkonium chloride (0.13%)

Uses

- Helps reduce bacteria that can cause skin infections
- Antiseptic preparation to decrease bacteria on skin

Warnings

For external use only, do not ingest.

Do Not Use

- If you are allergic to any of the ingredients
- In or near the eyes
- On children under 2 years of age

Ask a doctor before use if you have

- Deep or puncture wounds
- Serious burns

Stop use and ask a doctor if

- Irritation, swelling, pain or other symptoms develop

Keep out of reach of children.

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

Directions

1. Apply gel to tip of Cleansing Wand
2. Insert tip with gel into nostril, rotating it in each direction several times
3. Wipe any excess gel from nose with a tissue

- Use every 4-8 hours as needed
- Children under age 12 use with adult supervision

Other Information

Protect from excessive heat (temperatures about 104°F/40°C)

Inactive Ingredients - 81900-013

Carbomer, Diethylene Glycol Monoethyl Ether, Glycerin, Propylene Glycol, Triethanolamine, Water

Inactive Ingredients - 81900-014

Carbomer, Diethylene Glycol Monoethyl Ether, Glycerin, Peppermint Oil, Propylene Glycol, Triethanolamine, Water

Questions?

Call 1-844-4MY-NASO

OR (1-844-469-6276)

M-F 9am to 5pm PST

NasoClenz.com

Principal Display Panel - 81900-013

nasoclenz[®] Unscented

NasoClenz[®] Cleansing Kit



Drug Facts

Active Ingredient	Purpose
Benzalkonium Chloride 0.13%.....	Antiseptic

Uses

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- Helps reduce bacteria that can cause skin infections

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SVDOC-024 Rev B

Principal Display Panel - 81900-014

nasoclenz[®] Refreshing Mint

NasoClenz[®] Cleansing Kit



Drug Facts

Active Ingredient	Purpose
Benzalkonium Chloride 0.13%.....	Antiseptic

Uses

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- Helps reduce bacteria that can cause skin infections

Warnings

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Do not use

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Drug Facts (Continued)

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SVDOC-032 Rev A

NASOCLENZ

benzalkonium chloride 0.13% gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81900-014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)		BENZALKONIUM CHLORIDE	1.19 mg in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A1I8X02B)				
TROLAMINE (UNII: 9O3K93S3TK)				
CARBOMER HOMOPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: F68VH75CJC)				
PEPPERMINT OIL (UNII: AV092KU4JH)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81900-014-03	1 in 1 BOX	09/25/2023	
1		12 mL in 1 TUBE; Type 1: Convenience Kit of Co-Package		
2	NDC:81900-014-04	2 in 1 BOX	09/25/2023	
2		1 in 1 POUCH		
2		1 mL in 1 POUCH; Type 1: Convenience Kit of Co-Package		
3	NDC:81900-014-05	4 in 1 BOX	09/25/2023	
3		1 in 1 POUCH		
3		1 mL in 1 POUCH; Type 1: Convenience Kit of Co-Package		
4	NDC:81900-014-06	1 in 1 BOX	02/16/2024	
4		5 mL in 1 TUBE; Type 1: Convenience Kit of Co-Package		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	505G(a)(3)	09/25/2023		

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benzalkonium chloride 0.13% gel

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TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER HOMOPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: F68VH75CJC)	
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A1I8X02B)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81900-013-03	1 in 1 BOX	07/15/2022	
1		12 mL in 1 TUBE; Type 1: Convenience Kit of Co-Package		
2	NDC:81900-013-04	2 in 1 BOX	07/15/2022	
2		1 in 1 POUCH		
2		1 mL in 1 POUCH; Type 1: Convenience Kit of Co-Package		
3	NDC:81900-013-05	4 in 1 BOX	07/15/2022	
3		1 in 1 POUCH		
3		1 mL in 1 POUCH; Type 1: Convenience Kit of Co-Package		
4	NDC:81900-013-60	60 in 1 BOX	11/15/2022	
4		1 in 1 POUCH		
4		1 mL in 1 POUCH; Type 1: Convenience Kit of Co-Package		
5	NDC:81900-013-01	40 in 1 BOX	11/01/2021	
5		1 in 1 POUCH		
5		1 mL in 1 POUCH; Type 1: Convenience Kit of Co-Package		
6	NDC:81900-013-02	8 in 1 BOX	11/01/2021	
6		1 in 1 POUCH		
6		1 mL in 1 POUCH; Type 1: Convenience Kit of Co-Package		
7	NDC:81900-013-06	1 in 1 BOX	02/16/2024	
		5 mL in 1 TUBE; Type 1: Convenience Kit of Co-		

7	5 mL in 1 TUBE; Type 1: Convenience Kit or Co-Package		
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	11/01/2021	

Labeler - Silicon Valley Innovations, Inc. (118048729)

Registrant - Silicon Valley Innovations, Inc. (118048729)

Establishment			
Name	Address	ID/FEI	Business Operations
Silicon Valley Innovations, Inc.		118048729	label(81900-013, 81900-014) , pack(81900-013, 81900-014)

Revised: 2/2024

Silicon Valley Innovations, Inc.