UNSCENTED NASALGUARD ANTISEPTIC TOPICAL- benzalkonium chloride gel
TRUTEK CORP.

Unscented NasalGuard Antiseptic Topical Gel

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Use

To decrease bacteria on the skin.

Directions

Apply to the outside of the nose as indicated below.

Begin by squeezing 1 to 2 pin-sized drops on your pointer finger.

Spread the gel between your pointer finger and thumb to ensure an even application.

Apply directly around the nostrils and above the upper lip.

Apply every 4 to 6 hours or as often as needed.

Reapply if nose or face becomes wet.

SAVE THIS (BOX) FOR REFERENCE

Inactive Ingredients

Dimethicone, Glycerin, Hydroxyethyl Cellulose, Purified Water. Alson contains non-toxic, cosmetic grade preservatives and emulsifiers.

Questions or Comments ?

Visit www.NasalGuard.com or call 1-855-NASALGUARD (1-855-627-2548).

NASALGUARD ANTISEPTIC TOPICAL Gel Unscented

For external use only

When using this product keep out of eyes. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if

irritation and redness develop
condition persists for more than 72 hours

If pregnant or breast-feeding, ask a health professional before use.

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

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NasalGuard Unscented Tube and Carton

Proof Date : 20 February 2024

UNSCENTED NASALGUARD ANTISEPTIC TOPICAL

benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81907-102
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 1000 mg

Inactive Ingredients	
Ingredient Name	**Strength**
DIMETHICONE 350 (UNII: 2Y53S6ATLU)	
STEARETH-2 (UNII: V56DFE46J5)	
WATER (UNII: 059QF0KO0R)	
BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
STEARETH-21 (UNII: 53J3F32P58)	
GLYCERIN (UNII: PDC6A3C0OX)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81907-102-01	1 in 1 BOX	02/01/2024	
1	NDC:81907-102-02	3000 mg in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	02/14/2022	

Labeler - TRUTEK CORP. (170729235)

Registrant - TRUTEK CORP. (170729235)

Establishment

Name	Address	ID/FEI	Business Operations
TRUTEK CORP.		170729235	manufacture(81907-102)

Revised: 12/2025 TRUTEK CORP.