

NURINSE HAND SANITIZER-Q- benzalkonium chloride liquid NuGenTec

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.

NuRinse Hand Sanitizer Q

Active Ingredient(s)

Benzalkonium Chloride 0.13%

Purpose

Antiseptic, Hand Sanitizer, Antibacterial

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only.

When using this product: Keep out of eyes. In case of contact with eyes, flush thoroughly with water. Avoid contact with broken skin. Do not inhale or ingest.

Stop use and ask a Doctor: If skin irritation develops.

Keep out of reach of children.: If ingested, get medical help or call the Poison Control Center immediately.

Do not use

- in children less than 2 months of age
- on open skin wounds

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

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Directions

- When using NuRinse Hand Sanitizer-Q, apply a palmful of the product to one hand and rub the product all over the surfaces of both hands until your hands are dry. Do not wipe away excess sanitizer. For children under 6, use only with adult supervision. Not recommended for infants.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)
- Do not store above 120°F.
- May discolor some fabrics.
- Harmful to wood finishes and plastics.

Inactive ingredients

- Purified water USP
- Humectant
- Moisturizer
- Foaming agent

Package Label - Principal Display Panel

750mL is one bottle NDC: 77368-901-75

NuRinse[®]
Hand Sanitizer-Q

Alcohol Free Hand Sanitizer
Designed in The USA. Made in The USA.™

Product Description:
Foaming moisturizing, water-based, non-flammable hand sanitizer that soothes while sanitizing

Active Ingredients:
Benzalkonium Chloride 0.13%....Antimicrobial

Inactive Ingredients:
Water, humectant, moisturizer, and foaming agent

California Prop 65 Compliant

NuRinse[®]
Hand Sanitizer-Q

Foaming, Moisturizing Water-based Hand Sanitizer

**NO ALCOHOL
NON-FLAMMABLE
NON-HAZARDOUS
NO BLEACH
NO AMMONIA
FDA APPROVED**

Directions:
When using NuRinse Hand Sanitizer-Q, apply a palmful of the product to one hand and rub the product all over the surfaces of both hands until your hands are dry. Do not wipe away excess sanitizer. For children under 6, use only with adult supervision. Not recommended for infants.

WARNINGS:
Use: For external use only on hands.
When using this product: Keep out of eyes. In case of contact with eyes, flush thoroughly with water. Avoid contact with broken skin. Do not inhale or ingest.
Stop use and ask a Doctor: If skin irritation develops.
Keep out of reach of children.: If ingested, get medical help or call the Poison Control Center immediately.

Other Information:
Do not store above 120°F. May discolor some fabrics. Harmful to wood finishes and plastics.

Manufactured and Distributed By:
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Redefining Chemistry™
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0 51497 17914 4

1-Gallon (~3.8L)

NURINSE HAND SANITIZER-Q

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77368-903
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	0.08 mg in 100 mL
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	0.25 mL in 100 mL
WATER (UNII: 059QF0KO0R)	
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	0.25 mL in 100 mL
ACETAMIDOETHOXYETHANOL (UNII: LVX2APC4XR)	0.25 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77368-903-75	750 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/14/2021	
2	NDC:77368-903-02	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/14/2021	
3	NDC:77368-903-10	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2021	
4	NDC:77368-903-05	18927 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2021	
5	NDC:77368-903-01	3786 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - NuGenTec (090331927)

Registrant - NuGenTec (090331927)

Establishment

Name	Address	ID/FEI	Business Operations
NuGenTec		090331927	manufacture(77368-903)

Revised: 12/2022

NuGenTec