NANO PURE- benzalkonium chloride liquid Nano Global

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.

Nano Pure

Drug Facts

Active ingredient

Benzalkonium Chloride 0.10%

Purpose

Antiseptic

Uses

• Helps reduce bacteria on skin that potentially cause disease.

Warnings

For external use only.

Do not use

- in the eyes; if contact occurs, rinse thoroughly with water.
- over large areas of the body.
- if you are allergic to any ingredients.

Ask a doctor before use if you have

- deep puncture wounds.
- animal bites.
- serious burns.

Stop use and ask a doctor if

- condition persists or gets worse.
- skin irritation or redness develop.

Keep out of reach of children.

• In case of accidental ingestion, get medical help or contact Poison Control Center immediately.

Directions

•Spray onto hands or skin and apply evenly until dry.

Other information

• Store at 15° to 30°C (59° to 86°F). Avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients

Ethyl Alcohol, Fragrance (Parfum), Purified Water.

Questions?

1-833-777-NANO





Learn more at MYNANO.COM

GERM BARRIER

kin Protection

Active Protection

Most hand sanitizers only protect for seconds

Safe³

Nano Pure™ forms a safe germ barrier

Effective

Kills 99.99% of germs

Be Well, Do Good

Your purchase supports the fight against infectious diseases worldwide

Dermatologist Tested

*when used as directed

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Purpose Antiseptic

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Warnings

Made in the U.S.A.

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Global Corp.

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PATENTS PENDING



NANO PURE

benzalkonium chloride liquid

Kills 99.99%

of Germs

.

2 FL OZ (59.15ML)

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71530-025

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

	BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM -	BENZALKONIUM	0.1 g
ı	UNII:7N6JUD5X6Y)	CHLORIDE	in 100 mL

active Ingredients			
Ingredient Name	Strength		
LCOHOL (UNII: 3K9958 V90M)			
WATER (UNII: 059QF0KO0R)			

Ш	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:71530-025- 01	47.32 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/13/2020		
	NDC:71530-025- 02	59.15 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/13/2020		

Marketing Inform	Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	05/13/2020				

Labeler - Nano Global (079594740)

Revised: 5/2020 Nano Global