

NUMBIFY SANITIZE- benzalkonium chloride cream

Ridge Properties

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.

Warning: KEEP OUT OF REACH OF CHILDREN. FOR EXTERNAL USE ONLY. NOT FOR USE AROUND THE EYES. IF INGESTED SEEK MEDICAL HELP OR CALL A POISON CONTROL CENTER IMMEDIATELY.

Stop use and ask a doctor if skin becomes irritated, condition worsens, symptoms last longer than 7 days or if symptoms clear up and then return within a few days.

Active Ingredients

Purpose

Benzalkonium Chloride - 0.13%.....First Aid Antiseptic
Lidocaine HCL - 4%.....Temporary Relief of Pain & Itching

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Active Ingredients


Purpose

Benzalkonium Chloride - 0.13%.....First Aid Antiseptic
Lidocaine HCL - 4%.....Temporary Relief of Pain & Itching

Uses: First aid to help prevent bacterial contamination or skin infection, and for temporary relief of pain and itching

Inactive Ingredients: Propylene Glycol, Water, Kava Kava, Avocado Oil, Cetyl Alcohol, Sodium Polyacrylate, Polysorbate 80, Glyceryl Stearate, Dimethicone / Divinyldimethicone / Silsesquioxane crosspolymer

Directions: Adults and children 2 years old and over: clean the affected area; spray a small amount on the area 1-3 times daily; may be covered with a sterile bandage (let dry first) * children under 2 years, ask a doctor.

FDA REGISTERED: NDC# 69804-086-03	Do not Use if you have any known allergy to any of the ingredients in this product. Use on, in, or around the eyes. Apply over large areas of the body or in large quantities over raw surfaces or blistered areas. Discontinue use and seek medical attention should any reaction occur.	Numbify Sanitize First Aid Antiseptic & Pain Reliever	Warning: KEEP OUT OF REACH OF CHILDREN. FOR EXTERNAL USE ONLY. NOT FOR USE AROUND THE EYES. IF INGESTED SEEK MEDICAL HELP OR CALL A POISON CONTROL CENTER IMMEDIATELY. Stop use and ask a doctor if skin becomes irritated, condition worsens, symptoms last longer than 7 days or if symptoms clear up and then return within a few days.
Drug Facts: For Professional Use Only Uses: First aid to help prevent bacterial contamination or skin infection, and for temporary relief of pain and itching.	Directions: Adults and children 2 years old and over: clean the affected area; spray a small amount on the area 1-3 times daily; may be covered with a sterile bandage (let dry first) * children under 2 years, ask a doctor.		Other Information: Product manufactured, packed & distributed by Pain Relief Naturally. In event of a serious adverse reaction please call 1 (877) 906-4806 Or write us at 4995 Ridge DR NE, Salem, OR, 97301.
Inactive Ingredients: Propylene Glycol, Water, Kava Kava, Avocado Oil, Cetyl Alcohol, Sodium Polyacrylate, Polysorbate 80, Glyceryl Stearate, Dimethicone / Divinyldimethicone / Silsesquioxane crosspolymer		Active Ingredients Benzalkonium Chloride - 0.13% Lidocaine HCL - 4%	Purpose First Aid Antiseptic Temporary Relief of Pain & Itching
4 Ounce Cream			*Avoid Excessive Heat* ASK A DR BEFORE USE IF YOU HAVE: Deep or puncture wounds, animal bites, or serious burns. WWW.NUMBIFY.COM

NUMBIFY SANITIZE

benzalkonium chloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69804-086
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 10000 mg
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	13 mg in 10000 mg

Inactive Ingredients

Ingredient Name	Strength
CETYL DIMETHICONE/BIS-VINYLDIMETHICONE CROSSPOLYMER (UNII: AE7QA6TW0Q)	900 mg in 10000 mg
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	4030 mg in 10000 mg
WATER (UNII: 059QF0K00R)	3687 mg in 10000 mg
AVOCADO OIL (UNII: 6VNO72PFC1)	900 mg in 10000 mg

PIPER METHYSTICUM ROOT (UNII: BOW48C81XP)

70 mg in 10000 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69804-086-05	28500 mg in 1 JAR; Type 0: Not a Combination Product	06/19/2020	
2	NDC:69804-086-02	56700 mg in 1 JAR; Type 0: Not a Combination Product	06/19/2020	
3	NDC:69804-086-03	113400 mg in 1 JAR; Type 0: Not a Combination Product	06/19/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	06/19/2020	

Labeler - Ridge Properties (029478762)

Establishment

Name	Address	ID/FEI	Business Operations
Ridge Properties		029478762	manufacture(69804-086)

Revised: 4/2020

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