HUSH ANESTHETIC- lidocaine, benzalkonium chloride soap
HUSH Anesthetic

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.

HUSH Foam Soap

Active Ingredients

Lidocaine 4% w/w

Benzalkonium Chloride (0.13%) w/w

Purpose

Pain Relieving Liquid

First Aid Antiseptic

Uses

First aid to help prevent bacterial contamination or skin infection and for temporary relief of pain and itching associated with minor cuts or minor skin irritations.

Directions

Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily. Children under 2 years of age, consult a physician.

Warnings

For external use only • Avoid contact with the eyes

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

Do not use in large quantities, particularly over raw surfaces or blistered areas

In case of deep or puncture wounds, animal bites, or serious bites, consult a doctor • If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a doctor

Inactive ingredients

Aloe Barbadensis Leaf Extract, Aqua (Deionized Water), Calendula Officials Flower Extract, Caprylyl Glycol, Chamomile (Chamomilla Recutita) Flower Extract, Citric Acid, Cocamidopropyl Betaine, Comfrey (Symphytum Officinale) Root Extract, Disodium EDTA,

Disodium Laureth Sulfosuccinate, Glycerin, Green Tea (Camellia Sinensis) Leaf Extract, Methylisothiazollnone, PEG-80 Sorbitan Laurate, Propylene Glycol

Other Information

Questions or Comments? Call 305-231-7229 or visit www.hushanesthetic.com

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49947-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	40 mg in 1 g

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PEG-80 SORBITAN LAURATE (UNII: 239B50Y732)	
DISODIUM LAURETH SULFOSUCCINATE (UNII: D6DH1DTN7E)	
MATRICARIA CHAMOMILLA FLOWERING TOP OIL (UNII: SA8AR2W4ER)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
CAMELLIA SINENSIS WHOLE (UNII: C5M4585ZBZ)	
SYMPHYTUM OFFICINALE WHOLE (UNII: H8FJJ6KX5Y)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CITRIC ACID ACETATE (UNII: DSO12WL7AU)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49947-003-01	48.2 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	04/01/2014	

Labeler - HUSH Anesthetic (012011309)

Revised: 1/2022 HUSH Anesthetic