

HUSH ANESTHETIC- lidocaine spray
HUSH Anesthetic

Hush Spray

Active Ingredients

Lidocaine HCL 4% w/w

Purpose

Pain Relieving Liquid

Uses

For temporary relief of pain and itching associated with minor cuts or minor skin irritations.

Warnings

For external use only • Avoid contact with eyes

Keep out of reach of children

Stop use and ask a doctor if condition worsens or if symptoms persist for more than 7 days or clear up and occur again with a few days.

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

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.

Other Information

Protect this product from excessive heat and direct sun.

Inactive ingredients

Acrylates/c10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Extract, Aqua (Deionized Water), Calendula Officinale Extract, Caprylyl Glycol, Chamomile (Chamomile Recutita) Extract, Comfrey (Symphytum Officinale) Extract, Disodium EDTA, Glycerin, Green Tea (Camellia Sinensis) Extract, Menthol, Methylisothiazolinone, Propylene Glycol, SD Alcohol 40B, Triethanolamine

Questions or Comments?

Call 305-231-7229 or visit www.hushanesthetic.com

Spray



4oz | 113 ml

Spray



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Drug Facts	
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Inactive ingredients: Water (Aqua), SD Alcohol 40-B, Aloe Barbadosis Leaf Extract, Propylene Glycol, Glycerin, Comfrey (Symphytum Officinale) Extract, Chamomile (Anthemis Nobilis) Flower Extract, Calendula Officinalis Flower Extract, Green Tea (Camellia Sinensis) Leaf Extract, Menthol, Disodium EDTA, Triethanolamine, Methylisothiazolinone, Caprylyl Glycol, Acrylate/c10-30 Alkyl Acrylate Copolymer.	

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49947-001
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

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL (UNII: L7T10EIP3A)	
CHAMAEMELUM NOBILE FLOWER (UNII: O2T154T6OG)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
COMFREY ROOT (UNII: M9VVZ08EKQ)	
CARBOMER INTERPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 132584PQMO)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49947-001-04	113.4 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/01/2012	
2	NDC:49947-001-02	56.7 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/01/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	12/01/2012	

Labeler - HUSH Anesthetic (012011309)

Revised: 1/2023

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