

HUSH ANESTHETIC- lidocaine gel

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.

Derma Numb Tattoo Numbing Gel

Active Ingredients

Lidocaine

□ Purpose

Topical Anesthetic

Warnings

For external use only · Keep out of reach of children

Uses

Before skin is broken apply a thick layer of HUSH Gel in the area to be Tattooed. Spread evenly throughout. Reapply along the edges. Wrap the area with a plastic wrap and allow 1 hour for absorption. Remove the plastic wrap and prep skin for procedure.

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Do Not Use

In Blistered Areas · In the eyes · If allergic to ingredients

□ Stop Use and ask a doctor if

· skin becomes irritated · condition worsens or symptoms last more than 7 days · symptoms clear up and reoccur within a few days

□ Inactive ingredients

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

Other Information

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

Drug Facts

Active Ingredients	Purpose
Lidocaine 4%.....	Topical Anesthetic

Uses For the temporary relief of discomfort and pain associated with
• tattooing • tattoo removal • piercings

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an·es·thet·ic

HUSH
TATTOO NUMBING GEL

4 OZ.

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an·es·thet·ic

HUSH

TATTOO NUMBING GEL

2 OZ.



crueltyfree
andvegan



HUSH ANESTHETIC

lidocaine gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49947-002
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:49947-002-04	113.4 g in 1 BOTTLE		
2	NDC:49947-002-02	56.7 g in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	12/01/2012	

Labeler - HUSH Anesthetic (012011309)

Revised: 1/2010

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