NUMBIFY TROPICAL GOLD- lidocaine hcl 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.

Directions:

Test skin for sensitivity prior to use. Apply thick layers every 3 minutes until desired level of pain relief is reached. Discontinue use if sensitivity occurs. Not for use near the eyes, or on the face.

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.

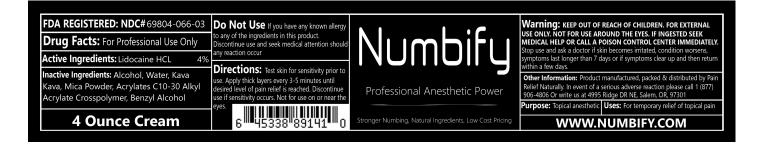
Warning: Keep out of reach of children. For external use only. Not for use around the eyes

Purpose: Topical Anesthetic

Uses: For Temporary Pain Relief

Active Ingredient: Lidocaine HCL 4%

Inactive Ingredients: Alcohol, Water, Kava Kava, Mica Powder, Acrylates C10-30 Alkyl Acrylate Crosspolymer, Benzyl Alcohol



NUMBIFY TROPICAL GOLD

lidocaine hcl cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69804-066
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
LIDO CAINE HYDRO CHLO RIDE (UNII: V13007Z41A) (LIDO CAINE - UNII:98 PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1000 mg	

Inactive Ingredients			
Ingredient Name	Strength		
MICA (UNII: V8A1AW0880)	90 mg in 1000 mg		
ALCOHOL (UNII: 3K9958V90M)	461 mg in 1000 mg		
WATER (UNII: 059QF0KO0R)	361 mg in 1000 mg		
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	25 mg in 1000 mg		
BENZYL ALCOHOL (UNII: LKG8494WBH)	10 mg in 1000 mg		
PIPER METHYSTICUM ROOT (UNII: BOW48C81XP)	13 mg in 1000 mg		

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:69804-066-05	28500 mg in 1 JAR; Type 0: Not a Combination Product	05/29/2017		
2	NDC:69804-066-02	56700 mg in 1 JAR; Type 0: Not a Combination Product	05/29/2017		
3	NDC:69804-066-03	113400 mg in 1 JAR; Type 0: Not a Combination Product	05/29/2017		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	05/29/2017	

Labeler - ridge properties (029478762)

Establishment				
Name	Address	ID/FEI	Business Operations	
ridge properties		029478762	manufacture(69804-066)	

Revised: 4/2020 ridge properties