MAMISAN PAIN RELIEVING CREAM LIDOCAINE PLUS MENTHOL- lidocaine hydrochloride, menthol cream PLANTIMEX DISTRIBUTORS INC

Mamisan Pain Relieving Cream Lidocaine Plus Menthol

Drug Facts

Active Ingredients

Lidocaine HCI 4%

Menthol 1%

Purpose

Topical anesthetic

Topical analgesic

Use

temporarily relieves minor pain

Warnings

For external use only

Do not use

- on large areas of the body or on cut, irritated or swollen skin
- on puncture wounds
- for more than one week without consulting a doctor

When using this product

- use only as directed. Read and follow all directions and warnings on this label
- rare cases of serious burns have been reported with products of this type
- do not bandage tightly or apply local heat (such as heating pads) to the area of use with a medicated patch
- avoid contact with the eyes and mucous membranes.
- a transient burning sensation may occur upon application but generally disappears in several days.

Stop use and ask doctor if

- condition worsens
- redness is present
- irritation develops
- symptoms persist for more than 7 days or clear up and occur again within a few

days

• you experience signs of skin injury, such as pain, swelling, or blistering where the product was applied

If pregnant or breast-feeding,

ask a health-professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

• a dults and children over 12 years of age and older:

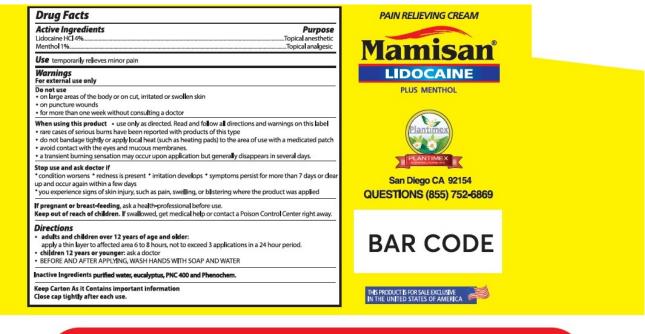
apply a thin layer to affected area 6 to 8 hours, not to exceed 3 applications in a 24 hour period.

- children 12 years or younger: ask a doctor
- BEFORE AND AFTER APPLYING, WASH HANDS WITH SOAP AND WATER

Inactive Ingredients

purified water, eucalyptus, PNC 400 and phenochem.

Package Labeling:





PLUS MENTHOL

DOCAINE

PLUS MENTHOL

NDC-

PAIN RELIEVING CREAM



NET WT. 2.7 oz (76.6g)

SEE CARTON FOR FULL DETAILS	
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		EVING CREAM I	IDO		PLUSI	MENTH	OL	
lidocaine hydroc	hloride, ment	hol cream						
Product Infor	mation							
	mation					NDC-72	016 001	
Product Type		HUMAN OTC DRUG	Item	Code (Sou	rce)	NDC:72816-001		
Route of Admin	istration	TOPICAL						
Active Ingred	ient/Active	Moiety						
	Ingr	edient Name			Basis of Strength		Strength	
LIDOCAINE HYDRO UNII:98PI200987)	OCHLORIDE (U	NII: V13007Z41A) (LIDOCAIN	IE -				40 mg in 1 g	
MENTHOL (UNII: L	7T10EIP3A) (MEN	ITHOL - UNII:L7T10EIP3A)					10 mg in 1 g	
Inactive Ingre	edients							
Ingredient Name					Strength			
WATER (UNII: 0590								
EUCALYPTUS GUN	4 (UNII: 72T9EZ	C2VX)						
Packaging								
# Item Code	Ра	ckage Description			ing Starl ate		eting End Date	
1 NDC:72816-001-01	76.6 g in 1 BO Product	TTLE; Type 0: Not a Combin	nation	06/19/2023				

Marketing End Date

Labeler - PLANTIMEX DISTRIBUTORS INC (185464463)

Revised: 12/2023

PLANTIMEX DISTRIBUTORS INC