RITE AID PAIN RELIEF CREAM- lidocaine hydrochloride cream Rite Aid Hdgrts Corp

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.

Riteaid Lidocaine pain relieving cream

Lidocaine Pain Relieving Creme

Drug Facts

Active ingredient

Lidocaine HCI 4%

Purpose

Topical anesthetic

Uses

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 carton.
- do not allow contact with the eyes
- do not bandage or apply local heat (such as heating pads) to the area of use

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children and pets.

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

Directions

adults and children over 12 years:

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

children 12 years and younger: ask a doctor

Inactive ingredients

Butylated hydroxyl toluene, cetostearyl alcohol, cetomacrogol 1000, cetyl alcohol, disodium EDTA, disodium hydrogen phosphate, light liquid paraffin, propylene glycol, sorbic acid, transquitol P, white petroleum jelly

Keep Carton As It Contains Important Information

Close cap tightly between uses.

PRINCIPAL DISPLAY PANEL



1 2 25 10 75 100



RITE AID PAIN RELIEF CREAM

lidocaine hydrochloride cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-3789
Route of Administration	TOPICAL		

Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 g
	LIDOCAINE HYDROCHLORIDE

Inactive Ingredie	nts			
	Ingredient	Name		Strength
BUTYLATED HYDRO	XYTOLUENE (UNII: 1P9D0Z171K)			
CETETH-2 (UNII: 7H8)	/AM7778)			
CETYL ALCOHOL (U	NII: 936JST6JCN)			
CETOSTEARYL ALC	OHOL (UNII: 2DMT128M1S)			
EDETATE DISODIUM	(UNII: 7FLD91C86K)			
SO DIUM PHO SPHAT	E, DIBASIC, ANHYDROUS (UNII: 22A	LDO 53M6 F)		
LIGHT MINERAL OII	L (UNII: N6K5787QVP)			
PROPYLENE GLYCO	L (UNII: 6DC9Q167V3)			
SORBIC ACID (UNII: X	(045WJ989B)			
PETROLATUM (UNII:	4T6H12BN9U)			
PETROLATUM (UNII:	4T6H12BN9U)			
PETROLATUM (UNII: Packaging	4T6H12BN9U) Package Descripti	ion Marke	ting Start Date	Marketing End Date
PETROLATUM (UNII: Packaging # Item Code		ion Marke 07/02/20	0	Marketing End Date
PETROLATUM (UNII: Packaging # Item Code 1 NDC:11822-3789-6	Package Descripti	07/02/20	0	Marketing End Date
PETROLATUM (UNII: Packaging # Item Code 1 NDC:11822-3789-6	Package Descripti 1 in 1 CARTON	07/02/20	0	Marketing End Date
PETROLATUM (UNII: Packaging	Package Descripti 1 in 1 CARTON	07/02/20	0	Marketing End Date
PETROLATUM (UNII: Packaging # Item Code 1 NDC:11822-3789-6	Package Descripti 1 in 1 CARTON 133 g in 1 TUBE; Type 0: Not a Comb	07/02/20	0	Marketing End Date
PETROLATUM (UNII: Packaging # Item Code 1 NDC:11822-3789-6 1	Package Descripti 1 in 1 CARTON 133 g in 1 TUBE; Type 0: Not a Comb Ormation	07/02/20	0	

Labeler - Rite Aid Hdgrts Corp (014578892)

Revised: 9/2020

Rite Aid Hdgrts Corp