LIDOCAINE HCL 4%- lidocaine hcl 4% ointment
CHAIN DRUG MARKETING ASSOCIATION INC

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.

QC Pain Relieving Cream + Lidocaine
Lidocaine HCl 4%

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benzyl alcohol, carbomer, hydrogenated lecithin, polysorbate 80, propylene glycol, purified water, triethanolamine, vitamin E

Adults and children 12 years and older: apply a thin layer to affected area every 6 to 8 hours, not to exceed 3 to 4 applications in a 24-hour period.

BEFORE AND AFTER APPLYING, WASH HANDS WITH SOAP AND WATER.

Children under 12 years: ask a doctor

temporary relieves minor joint and muscle pain

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.

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Topical anesthetic

LIDOCAINE HCL 4%

lidocaine hcl 4% ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-451
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
WATER (UNII: 059QF0KO0R)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63868-451-72	76.5 g in 1 BOTTLE; Type 0: Not a Combination Product	10/28/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	10/28/2020	

Labeler - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

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