DCH PAIN RELIEF- lidocaine hcl 4% cream Derma Care Research Labs, LLC

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.

DCH Pain Relief Cream, Lidocaine HCI 4%

Lidocaine HCI 4%

Topical Analgesic

Temporarily relieves minor 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 directions on this carton, avoid contact with eyes and mucous membranes, do not apply to wounds or damaged, broken, or irritated skin, do not bandage tightly or apply local heat (such as heating pads) or a medicated patch to the area of use, do not use at the same time as other topical analgesics, a transient burning sensation may occur upon application but generally disappears in several days, avoid applying into skin folds.

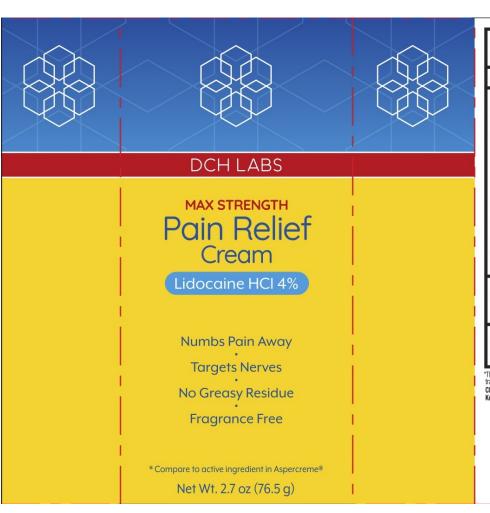
Stop use and ask a doctor if the condition worsens, redness is present, irritation develops, symptoms persist for more than 7 days or clear up and occur again within a few days, you experience signs of skin injury, such as pain, swelling, or blistering where the product was applied.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Adults and children over 12 years: apply a thin layer to the affected area every 6 to 8 hours, not to exceed 3 applications in a 24-hour period. AFTER APPLYING, WASH HANDS WITH SOAP AND WATER.

Children under 12 years or younger: ask a doctor.

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Alcohol Denat., Aloe Barbadensis Leaf Extract, Aminomethyl Propanol, C30-45 Alkyl Cetearyl Dimethicone Crosspolymer, Caprylyl Methicone, Cetearyl Alcohol, Ceteth-20 Phosphate, Dicetyl Phosphate, Dimethicone, Disodium EDTA, Ethylhexylglycerin, Glyceryl Stearate, Methylparaben, Steareth-21, Water.



Drug Facts

Active ingredient

Purnose

rarily 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 punicture wounds
 for more than one week without consulting a doctor

- Iro more than one week without consulting a doctor

 When using this product
 use only as directed. Read and follow directions and warnings on this carton
 avaid contact with eyes and mucous membranes.
 do not apply to wounds or damaged, broken or irritated skin
 do not banked bightly or apply local heat (such as heating pads) or a medicated patch to the airsa of use
 do not use and the same time as other topical analysesics
 a transient burning sensation may occur upon application but generally disappears in several days.
 avoid applying into skin folds

- condition was as a contract with the contract was a contract within a few days.
- 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 care professional before use.
Keep out of reach of children. If swallowed, get medical help or contact a Poison
Control Center immediately.

- 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 pendio, WASH HANDS WITH SOAP AND WATER
 Children 12 years or younger: ask a doctor

Inactive ingredients acrylates/C10-30 alkyl acrylate crosspolymer, alcohol denat, also barbadensis leaf juice, aminomethyl propanol, C30-45 alkyl ceteraryl dimethioner crosspolymer, caprylyl methicone, cetearyl checklon, cleeth-20 phosphate, dicetyl phosphate, dimethicone, disadium EUTA, ethylhexylglycerin, gycycryl stearart, emethylograden, staeenth-21, water

This product is not manufactured or distributed by Chattem, Inc., owner of the registered

Child-resistant packaging. Close cap tightly between uses. Keep carton as it contains important information.



Manufactured by: DermaCare Research Labs 11 C 440 Fentress Blvd., Daytona Beach, FL 32114

DCH PAIN RELIEF

lidocaine hcl 4% cream

Product Information

Product Type HUMAN OTC DRUG **Item Code (Source)** NDC:72839-361

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
STEARETH-21 (UNII: 53J3F32P58)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	

CETETH-20 PHOSPHATE (UNII: 921FTA1500)

CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 809Y72KV36)

ALOE VERA LEAF (UNII: ZY81Z83H0X)

AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
DIHEXADECYL PHOSPHATE (UNII: 2V6E5WN99N)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
WATER (UNII: 059QF0KO0R)	
C30-45 ALKYL CETEARYL DIMETHICONE CROSSPOLYMER (UNII: 4ZK9VP326R)	
DIMETHICONE 200 (UNII: RGS4T2AS00)	
ALCOHOL (UNII: 3K9958V90M)	

ı	P	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:72839- 361-02	76.5 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/10/2021		

Marketing Information					
ting End ate					

Labeler - Derma Care Research Labs, LLC (116817470)

Registrant - Derma Care Research Labs, LLC (116817470)

Establishment							
Name	Address	ID/FEI	Business Operations				
Derma Care Research Labs, LLC		116817470	manufacture(72839-361)				

Revised: 7/2023 Derma Care Research Labs, LLC