

PAIN RELIEF CREAM- lidocream hydrochloride cream

AEC Consumer Products

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.

Drug Facts

Active Ingredient

Lidocaine Hydrochloride 4%.....External Analgesic

Purpose

Purpose

External Analgesic

Warning

For External use only

When using this product

When using this product use only as directed

- avoid 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

Stop use and ask a doctor if condition worsens symptoms persist more than 7 days
symptoms clear up and occur again with a few days

Keep out of use of children

Keep out of use of children. If swallowed, get medical help or contact a Poison Control Center right away

Storage and Handling

Apply as needed to affected areas 3-4 times a day.

Inactive Ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Allantoin, Aqua, Arnica Mountain Flower Extract, Astaxanthin, Benzyl Alcohol, BHT, Bisabolol, Boswellia Serrata Resin Extract, Cannabis Sativa Oil/Extract, Cetyl Alcohol, Citrus Aurantium Bergamia Peel Oil, Curcurria Longa Root Extract, Cyclopentasiloxane, Dimethicone, Dimethicol, Emu Oil, Ethoxydiglycol, Glucosamine, Glycerin, Glyceryl Stearate SE, Isopropyl Palmitate, Melaleuca Alternifolia Leaf Oil, Methylsulfonylmethane, PEG-100 Stearate, Phenoxyethanol, Polyquaterium-10, Potassium Sorbate, Propylene Glycol, Retinyl Palmitate, Salicylic Acid, Sodium Hydroxide, Stearic Acid, Stearyl Alcohol, Tetrasodium EDTA, Tocopherol, Triethanolamine, Vaccinium Myrtillus Fruit/Leaf Extract

Indications

Use

For the temporary relief of pain

Do not use

Do not use in large quantities, particularly over raw or blistered surfaces for more than one week without consulting a doctor

Ask a doctor

Ask a doctor if condition worsens symptoms persist more than 7 days symptoms clear up and occur again with a few days

Directions

Adults and children over 12 years: apply as needed to affected areas 3-4 times a day. Consult a physician if pain persists.

dosage

Apply as needed to affected areas 3-4 times a day.

image of label

PATENT-PENDING FORMULA



Advanced Topical Analgesic
+ Soothing Botanicals

INFUSED WITH 300mg Full Spectrum CBD Oil[†]
Lidocream^{Plus}
4% Lidocaine Maximum Strength

† EXTRACTED FROM INDUSTRIAL HEMP

pain relief cream

NET WT 3.4 OZ (96.4 g)

Printing Area

Drug Facts

Active Ingredient	Purpose
Lidocaine Hydrochloride 4%	External Analgesic
Use	
For the temporary relief of pain	
Warnings	
For external use only	
Do not use: in large quantities, particularly over raw or blistered surfaces for more than one week without consulting a doctor	
When using this product: use only as directed avoid 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 more than 7 days symptoms 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. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions	
Adults and children over 12 years: Apply as needed to affected areas 3-4 times a day. Consult a physician if pain persists. Children 12 years of younger: consult a doctor	
Inactive Ingredients	
Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Allantoin, Aqua (Deionized Water), Arnica Montana Flower Extract, Astaxanthin, Benzyl Alcohol, BHT, Bisabolol, Boswellia Serrata Resin Extract, Cannabis Sativa (Hemp) Oil Extract, Cetyl Alcohol, Citrus Aurantium Bergamia (Bergamot) Peel Oil, Curcuma Longa (Turmeric) Root Extract, Cyclopentasiloxane, Dimethicone, Dimethiconol, Emu Oil, Ethoxydiglycol, Glucosamine, Glycerin, Glyceryl Stearate SE, Isopropyl Palmitate, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Methylsiloxymethane, PEG-100 Stearate, Phenoxyethanol, Polyquaternium-10, Potassium Sorbate, Propylene Glycol, Retinyl Palmitate, Salicylic Acid, Sodium Hydroxide, Stearic Acid, Stearyl Alcohol, Tetrasodium EDTA, Tocopherol, Triethanolamine, Vaccinium Myrtillus Fruit/Leaf Extract	
Questions? 910-987-7964	



Made in the U.S.A.

Manufactured Exclusively for AEC Consumer Products
135 Airport Road, Fayetteville, NC 28306
www.aecconsumerproducts.com

Non Printing Area

PAIN RELIEF CREAM

lidocream hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:86089-222
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	3.86 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALLANTOIN (UNII: 344S277G0Z)	

water (UNII: 059QF0KO0R)
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)
ASTAXANTHIN (UNII: 8XPW32PR7I)
BENZYL ALCOHOL (UNII: LKG8494WBH)
BISABOLOL OXIDE A (UNII: 16AE65F94Y)
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)
CETYL ALCOHOL (UNII: 936JST6JCN)
CITRUS AURANTIIFOLIA FRUIT OIL (UNII: 7937R189CB)
CURCUMA LONGA LEAF (UNII: H2HC4RY52C)
DIMETHICONE (UNII: 92RU3N3Y1O)
EMU OIL (UNII: 344821WD61)
ETHOXYDIGLYCOL BEHENATE (UNII: N76ISC4ZZO)
GLUCOSAMINE (UNII: N08U5BOQ1K)
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)
MELALEUCA ALTERNIFOLIA LEAF (UNII: G43C57162K)
PEG-100 STEARATE (UNII: YD01N1999R)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
POLYQUATERNIUM-10 (1000 MPAS AT 2 %) (UNII: GMR4PEN8PK)
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
SODIUM HYDROXIDE (UNII: 55X04QC32I)
STEARIC ACID (UNII: 4ELV7Z65AP)
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)
TETRASODIUM DIETHYLENETRIAMINEPENTAMETHYLENEPHOSPHONATE (UNII: E6DVO371LC)
TOCOPHEROL (UNII: R0ZB2556P8)
VACCINIUM MYRTILLUS LEAF (UNII: Y4U591OU70)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:86089-222-01	94.6 g in 1 TUBE; Type 0: Not a Combination Product	09/23/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	09/23/2019	

Labeler - AEC Consumer Products (080994102)

Registrant - Cospro Development Corp (785638821)

Establishment

Name	Address	ID/FEI	Business Operations
Cospro Development Corp		785638821	manufacture(86089-222) , pack(86089-222) , label(86089-222)

