

DCH PAIN RELIEF- lidocaine 4% spray
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 Spray, Lidocaine 4%

Lidocaine 4%

External Analgesic.

For the temporary relief of pain and itching due to sunburn, minor burns, insect bites, minor cuts, scrapes, and minor skin irritations.

For external use only.

Flammable--Do not use while smoking or near heat or flame.

Do not use in large quantities, particularly over raw surfaces or blistered areas.

When using this product avoid contact with eyes. Rinse eyes with water to remove. Contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120F.

Stop use and ask a doctor if the condition worsens or symptoms persist for more than 7 days or clear up and occur again within a few days, irritation develops.

Keep out of reach of children. If the product is swallowed, get medical help or contact a Poison Control Center right away.

Adults and children 2 years and older: apply to the affected area, not more than 3 to 4 times a day. Children under 2 years of age: consult a physician. To apply to face, squeeze into palm of hand and gently apply.

Water, Glycerin, Alcohol Denat., Propylene Glycol, Aloe Barbadensis Leaf Extract, Tocopheryl Acetate, Chamomilla Recutita (Matricaria) Flower Extract, Cucumis Sativus (Cucumber) Fruit Extract, Disodium EDTA, Triethanolamine, Diazolidinyl Urea, Simethicone.



Drug Facts

Active ingredient Lidocaine 4% **Purpose** External analgesic

Uses For the temporary relief of pain and itching associated with • sunburn • minor burns • minor cuts • scrapes • insect bites • minor skin irritations.

Warnings

For external use only.

Flammable - Do not use while smoking or near heat or flame.

Do not use in large quantities, particularly over raw surfaces or blistered areas.

When using this product keep out of eyes. Rinse eyes with water to remove contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120°F.

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
- irritation develops

Keep out of the reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- shake well before use
- adults and children 2 years or older: apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: ask a doctor.

Inactive ingredients Water, Alcohol Denat., Propylene Glycol, Glycerin, Aloe Barbadensis Leaf Extract, Tocopheryl Acetate, Chamomilla Recutita (Matricaria) Flower Extract, Cucumis Sativus (Cucumber) Fruit Extract, Disodium Cocoamphodipropionate, Carbomer, Disodium EDTA, Triethanolamine, Diazolidinyl Urea, Simethicone.

*This product is not manufactured or distributed by Chattam, Inc.®, owner of the registered trademark Aspercreme®



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



DCH LABS

Pain Relief Spray

Max Strength

Lidocaine 4%

Numbs Pain Away
•
No Greasy Residue
•
Sprays at Any Angle

* Compare to active ingredient in Aspercreme®

Net Wt. 4 oz (113 g)

DCH PAIN RELIEF

lidocaine 4% spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72839-226
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
CUCUMBER (UNII: YY7C30VXJT)	

DISODIUM COCOAMPHODIPROPIONATE (UNII: 6K8PRP397M)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
WATER (UNII: 059QF0KO0R)
GLYCERIN (UNII: PDC6A3C0OX)
ALCOHOL (UNII: 3K9958V90M)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
EDETATE DISODIUM (UNII: 7FLD91C86K)
TROLAMINE (UNII: 9O3K93S3TK)
ALOE VERA LEAF (UNII: ZY81Z83H0X)
CARBOMER 940 (UNII: 4Q93RCW27E)
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
CHAMOMILE (UNII: FGL3685T2X)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72839-226-04	113 g in 1 CAN; Type 0: Not a Combination Product	06/10/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	06/10/2021	

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		116817470	manufacture(72839-226)

Revised: 7/2023

Derma Care Research Labs, LLC