

DCH NUMBING RELIEF- lidocaine 5% spray
Derma Care Research Labs, LLC

DCH Numbing Relief Spray, Lidocaine 5%

Lidocaine 5%

Topical anesthetic

For the local itching and pain associated with anorectal disorders or inflammation.

For external use only. Flammable--Keep away from fire or flame.

When using this product avoid contact with eyes, do not exceed recommended dosage unless directed by a doctor, do not put this product into the rectum by using fingers or any mechanical device or applicator, contents under pressure. Do not puncture or incinerate. Do not store at temperatures above 120F.

Stop use and ask a doctor if rectal bleeding occurs, condition worsens or does not improve within 7 days, allergic reaction occurs to ingredients in this product, symptom being treated does not subside or if redness, irritation, swelling, pain or other symptoms develop or increase. i

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

When practical, cleanse the affected area with mild soap and warm water, Rinse thoroughly and gently pat dry. Spray affected area. Wash hands after applying. Adults and children 2 years of age and older, apply externally to the affected area up to 6 times a day. Children under 12 years of age: consult a doctor.

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.



DCH LABS

Maximum Strength Numbing Relief Spray

ANORECTAL

Lidocaine 5%

Numbs Away Pain

No Greasy Residue

Sprays at Any Angle

Net Wt. 5 oz (142 g)

Drug Facts

Active ingredient

Lidocaine 5%

Purpose

Topical anesthetic

Uses • helps relieve the local itching and pain associated with anorectal disorders or inflammation

Warnings

For external use only

Flammable - do not use while smoking or near heat or flame

When using this product • avoid contact with eyes • do not exceed recommended dosage unless directed by a doctor • do not put this product into the rectum by using fingers or any mechanical device or applicator • contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120°F

Stop use and ask a doctor if • rectal bleeding occurs • condition worsens or does not improve within 7 days • allergic reaction occurs to ingredients in this product • symptom being treated does not subside or if redness, irritation, swelling, pain or other symptoms develop or increase

If pregnant or breastfeeding, 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 • when practical, cleanse the affected area with mild soap and warm water • rinse thoroughly and gently pat dry • spray affected area • wash hands after applying • adults and children 12 years and older: apply externally to the affected area up to 6 times a day • children under 12 years of age: consult a doctor

Other information Store at 15°-30° C (59°-86° F)

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



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

DCH NUMBING RELIEF

lidocaine 5% spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72839-173
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CARBOMER 940 (UNII: 4Q93RCW27E)	

TROLAMINE (UNII: 9O3K93S3TK)
GLYCERIN (UNII: PDC6A3C0OX)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
ALCOHOL (UNII: 3K9958V90M)
CHAMOMILE (UNII: FGL3685T2X)
CUCUMBER (UNII: YY7C30VXJT)
DISODIUM COCOAMPHODIPROPIONATE (UNII: 6K8PRP397M)
WATER (UNII: 059QF0KO0R)
EDETATE DISODIUM (UNII: 7FLD91C86K)
ALOE VERA LEAF (UNII: ZY81Z83H0X)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72839-173-05	142 g in 1 CAN; Type 0: Not a Combination Product	05/17/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	01/30/2023	

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(72839-173)

Revised: 12/2024

Derma Care Research Labs, LLC