DCH NERVE PAIN RELIEF- lidocaine 4%, menthol 1% spray Derma Care Research Labs, LLC

DCH Nerve Pain Relief Spray, Lidocaine 4%, Menthol 1%

Lidocaine 4%, Menthol 1%

Topical anesthetic, Topical analgesic

Temporarily relieves minor pain

For external use only.

Flammable--Keep away from fire or flame.

Do not use if you have had an allergic reaction to lidocaine or other local anesthetics, on large areas of he body or on cut, irritated, blistered or swollen skin, on puncture wounds, for more than one week without consulting a doctor.

When using this product only use as directed, avoid contact with eyes and mucous membranes, rare cases of serious burns have been reported with products of this type, 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 path to the area of use, a transient burning sensation may occur upon application but generally disappears in several days, avoid applying into skin folds.

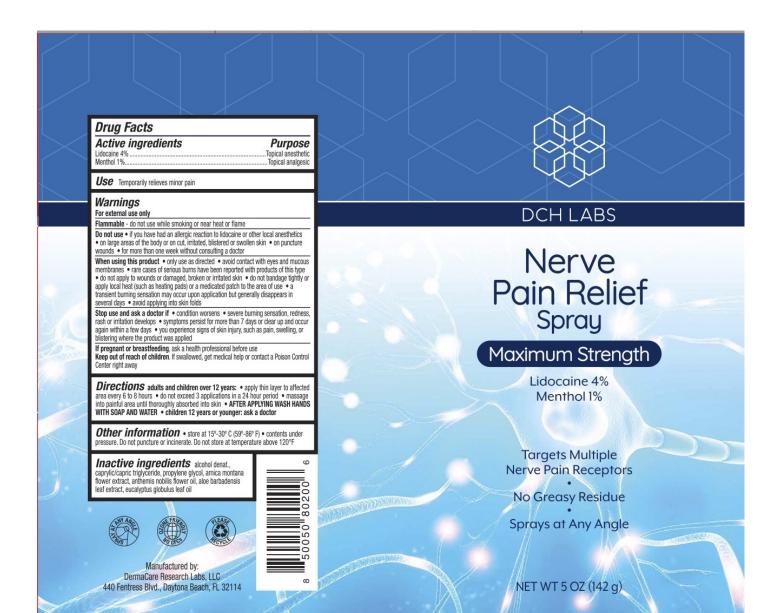
Stop use and ask a doctor if condition worsens, severe burning sensation, redness, rash or irritation develops, symptoms persist for more than 7 days or clear up and occur again within a few days, you experience signs of skin injury, such as pain, swelling, or blistering where the product was applied.

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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.

Alcohol denat., caprylic/capric triglyceride, propylene glycol, arnica montana flower extract, anthemis nobilis flower oil, aloe barbadensis leaf extract, eucalyptus globulus leaf oil.



DCH NERVE PAIN RELIEF

lidocaine 4%, menthol 1% spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72839-223

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
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength

ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	
CHAMAEMELUM NOBILE FLOWER (UNII: O2T154T6OG)	
EUCALYPTUS OIL (UNII: 2R040NI662)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	

	Packaging			
1	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:72839-223- 05	142 g in 1 CAN; Type 0: Not a Combination Product	03/24/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/24/2023	

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(72839-223)

Revised: 12/2024 Derma Care Research Labs, LLC