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.

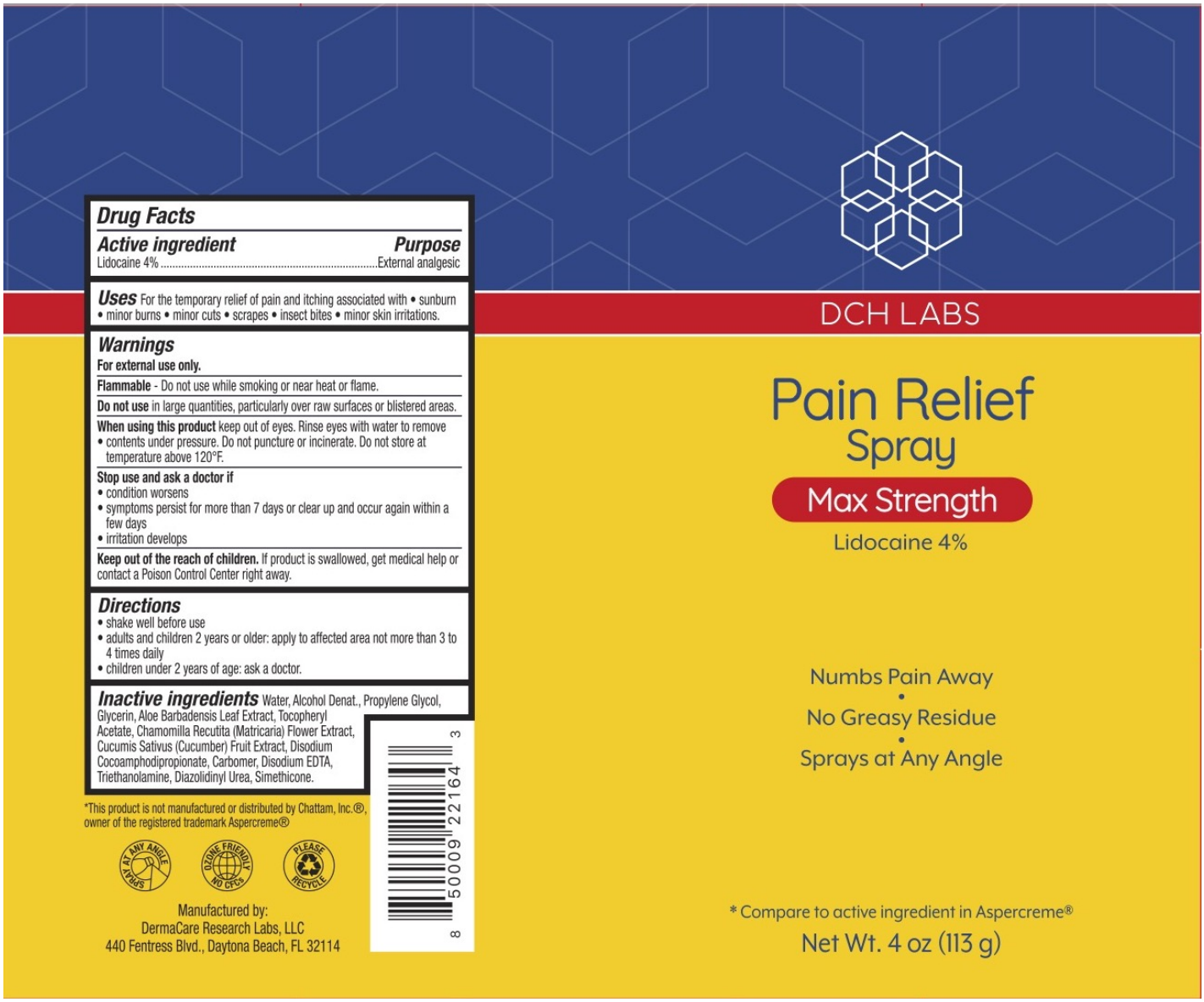
When using this product avoid contact with eyes. Rinse eyes with water to remove. Contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120F.

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.



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