MYDERM- lidocaine gel
Inspec Solutions LLC.

MyDerm Pain Reliever Aloe Gel Lidocaine HCl 0.5%

Active Ingredient

Lidocaine 0.5%

Lidocaine 0.5% ..... Pain Reliever

Temporary relief of pain and itching due to

- sunburn - minor burn - insect bites - minor cuts - scraps

Warnings

For external use only

When using this product keep out of eyes

Rinse with warter to remove

Stop use and ask a doctor

system persist for more than 7 days.

Keep out of reach of children. If product is swallowed, get medical help or contact a poison control center right away.

Direction

Adults and children 2 years and older: aooly to affected area not more that 3-4 times a day

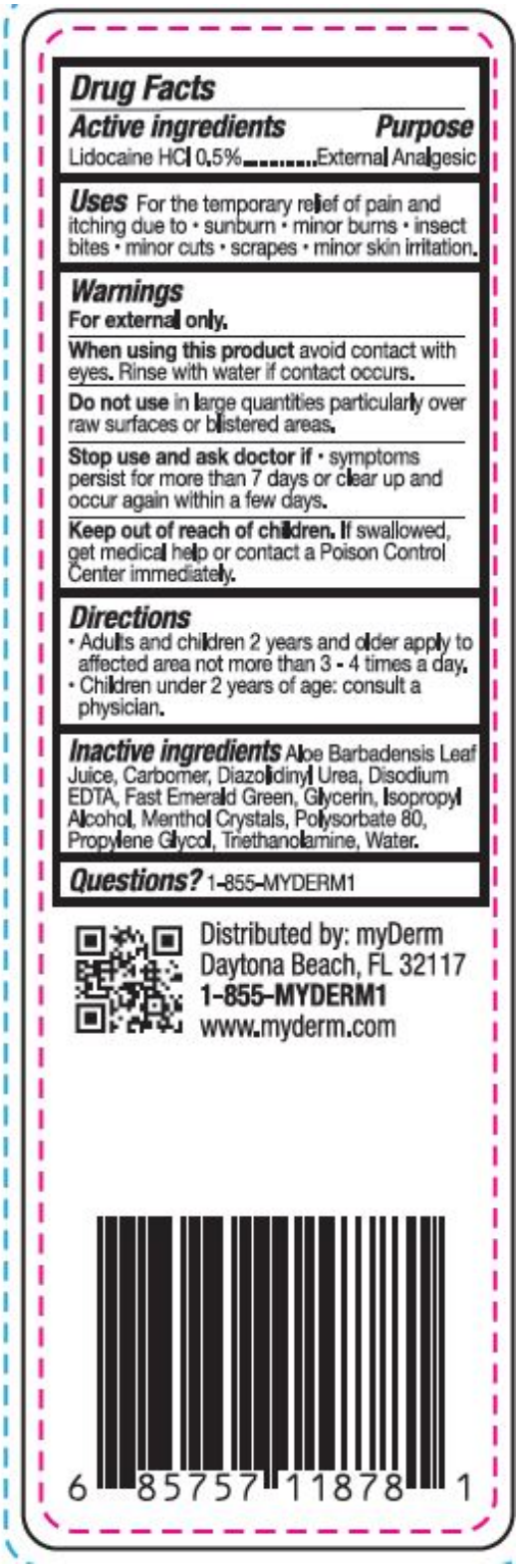
Children under 2 years of age: consult a physician.

Inactive Ingredients: Aloe Barbadensis Leaf Extract, Carbomer, Diazolidinyl Urea, Disodium EDTA, Glycerin, Isopropyl Alcohol, Menthol, Polysorbate 80, Propylene Glycol, Triethanolamine, Water, Yellow 5.

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

Questions? 1-855-MYDERM1

5.5 in



MYDERM
lidocaine gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72667-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	0.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
MENTHOL (UNII: L7T10EIP3A)	
PROPYLENE GLYCOL 2-STEARATE (UNII: 5317AOR1RX)	
TROLAMINE SALICYLATE (UNII: H8O4040BHD)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER 1342 (UNII: 809Y72KV36)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
DISODIUM HEDTA (UNII: KME849MC7A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
FD&C YELLOW NO. 5 FREE ACID (UNII: 6TP696149N)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72667-100-01	592 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/26/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/26/2019	

Labeler - Inspec Solutions LLC. (081030372)

Registrant - Inspec Solutions LLC. (081030372)

Establishment

Name	Address	ID/FEI	Business Operations
Inspec Solutions LLC.		081030372	manufacture(72667-100)

Revised: 4/2025 Inspec Solutions LLC.