MYDERM MEDICAL 5 LIDOCAINE NUMBING ROLL-ON- lidocaine stick
Inspec Solutions

MyDerm 5% Numbing Roll-On (IS0340)

Active Ingredient

Lidocaine 5%

When using this product avoid conatct with the eyes or muccous membranes.

Lidocaine 5% ----------Local Anesthetic

For external use only.

Stop use and ask doctor if condition worsens, or symtoms persist for more than 7 days,or clear up and reoccur again within a few days.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Acrylates/C10-30 Alkyl Acrylate Crosspolymer

Aloe Vera Barbadensis Leaft Juice

Inactive ingredients

2-animo-2methyl-1propanol

2-Methylamino-2-methyl-1-propanol

Cetearyl Alcohol

Citric Acid

Dimethicone

Disodium EDTA

Ethanol

Ethyhexylglycerin

Glyceril Stearate

Phenoxyethanol

Potassium Sorbate

Silicone Gel

Sodium Benzoate

Steareth-21

Water

Apply to affected area not more than 3 to 4 times daily

For external use only

150340

myDerm™
MEDICAL

5% LIDOCAINE

NUMBING ROLL-ON

- Fast-Acting Topical Anesthetic
- Child Resistant Packaging
- No-Touch Application

TRUE Maximum Strength

3 FL OZ (89 mL)

Drug Facts

Active ingredient	Purpose
Lidocaine HCl 5%	Local Anesthetic

Uses Helps relieve anorectal symptoms (pain, soreness, burning, itching). Temporarily reduces the swelling associated with irritated hemorrhoidal tissue and other anorectal disorders.

Warnings

For external use only.

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

When using this product • avoid contact with the eyes or mucous membranes.

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

Do not use if pregnant or breast-feeding.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Directions • Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily. • Children under 2 years of age: Consult physician.

Inactive ingredients Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Extract, Aminomethyl Propanol, C30-45 Alkyl Cetearyl Dimethicone Crosspolymer, Caprylyl Methicone, Cetearyl Alcohol, Ceteth-20 Phosphate, Dicetyl Phosphate, Dimethicone, Disodium EDTA, Ethylhexylglycerin, Glyceryl Stearate, Phenoxyethanol, SD Alcohol 40, Steareth-21, Water.

Questions? 1-855-MYDERM1

Distributed by: myDerm
Daytona Beach, FL 32117
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MYDERM MEDICAL 5 LIDOCAINE NUMBING ROLL-ON

lidocaine stick

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72667-072
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**
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
DIMETHICONE 100 (UNII: RO266O364U)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
ALCOHOL (UNII: 3K9958V90M)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
CITRIC ACID ACETATE (UNII: DSO12WL7AU)	
MAGNESIUM DISODIUM EDTA (UNII: NDT563S5VZ)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
STEARETH-21 (UNII: 53J3F32P58)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72667-072-01	89 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/06/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	03/06/2024	

Labeler - Inspec Solutions (081030372)

Establishment

Name	Address	ID/FEI	Business Operations
Inspec Solutions		081030372	manufacture(72667-072)