

DERMEND MOISTURIZING ANTI-ITCH- pramoxine hydrochloride lotion
Ferndale Laboratories, Inc.

DerMend Moisturizing Anti-Itch Lotion

Active ingredient

Pramoxine HCl 1% w/w

Purpose

external analgesic

Use

for the temporary relief of itching associated with minor skin irritations

Warnings

For external use only.

Avoid contact with the eyes.

Stop use and ask a doctor if

- conditions worsens
- symptoms last more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

If swallowed, seek medical attention or contact a Poison Control Center right away.

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 a doctor

Inactive ingredients

cetyl alcohol, di-isopropyl adipate, dimethicone, glycerin, FORLAN-L (Contains: petrolatum, lanolin, hydrogenated coconut oil, sorbitan sesquioleate, stearyl alcohol, and cetyl alcohol), mineral oil, polyoxyl 40 stearate, potassium sorbate, povidone, purified water, sorbic acid, stearic acid, and trolamine

Package Label

NDC 0496-0586-08

8oz (237mL)

**Dermatologist
Recommended**



**moisturizing
anti-itch
lotion**

Pramoxine Hydrochloride 1%

Targeted care to relieve
dry, itchy mature skin.

STEROID FREE

NET WT. 8 fl oz (237 mL)

DerMend.com

Drug Facts

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polyoxyl 40 stearate, potassium sorbate,
povidone, purified water, sorbic acid, stearic acid,
and trolamine



Manufactured for

Ferndale Healthcare, Inc.

Ferndale, MI 48220 USA (888) 548-0900

DerMend® is a registered trademark of

Clark Pharmaceuticals, LLC, Fort Worth, TX 76107

Rev: 11/16

DERMEND MOISTURIZING ANTI-ITCH

pramoxine hydrochloride lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0496-0586
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIISOPROPYL ADIPATE (UNII: P7E6YFV72X)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

GLYCERIN (UNII: PDC6A3C0OX)
PETROLATUM (UNII: 4T6H12BN9U)
LANOLIN (UNII: 7EV65EAW6H)
HYDROGENATED COCONUT OIL (UNII: JY81OXM1OM)
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)
MINERAL OIL (UNII: T5L8T28FGP)
POLYOXYL 40 STEARATE (UNII: 13A4J4NH9I)
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)
POVIDONE (UNII: FZ989GH94E)
WATER (UNII: 059QF0KO0R)
SORBIC ACID (UNII: X045WJ989B)
STEARIC ACID (UNII: 4ELV7Z65AP)
TROLAMINE (UNII: 9O3K93S3TK)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0496-0586-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2017	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/01/2017	

Labeler - Ferndale Laboratories, Inc. (005320536)

Establishment			
Name	Address	ID/FEI	Business Operations
Ferndale Laboratories, Inc.		005320536	manufacture(0496-0586)

Revised: 12/2024

Ferndale Laboratories, Inc.