PRAX- pramoxine hydrochloride lotion AvPAK

Pramoxine HCI 1% Lotion

Active Ingredient

pramoxine HCl 1% w/w

Purpose

local anesthetic

Use

for the temporary relief of discomfort and itch in the perianal area

Warnings

For external use only.

Do not

- exceed the recommended daily dosage unless directed by a doctor
- put this product into the rectum by using fingers or any mechanical device or applicator

Stop use and ask a doctor if

- condition worsens
- symptoms do not improve within 7 days
- allergic reactions develop to ingredients in this product
- symptom being treated does not subside or if redness, irritation, swelling, pain, bleeding, or other symptoms develop or increase

Keep out of reach of children.

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

Directions

- Shake well before use.
- When practical, cleanse the affected area with mild soap and warm water and rinse thoroughly.
- Adults and children 12 years of age and older: apply to affected area up to 5 times daily.
- Children under 12 years of age: consult a doctor.

Inactive Ingredients

Aloe Barbadensis Leaf Juice, Cetyl Alcohol, Glycerin, Glyceryl Stearate SE, Mineral Oil, PEG-100 Stearate, Phenoxyethanol, Purified Water, Stearyl Alcohol, White Petrolatum, Xanthan Gum

Package Label

Questions or comments call 1-855-361-3993.

Manufactured for:

AVKARE

Pulaski, TN 38478

www.avkare.com

AV 01/2022



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Other Information

- Store at room temperature 15°-30°C (59°-86°F)
- Do not use if safety seal is torn or missing.

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Manufactured for: AvKARE Pulaski, TN 38478 www.avkare.com AV 01/2022



PRAX

pramoxine hydrochloride lotion

Product	Inform	ation
Product	morm	auon

Product Type HUMAN OTC DRUG Item Code (Source) NDC:50268-682

Route of Administration TOPICAL

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

Inactive Ingredients				
Ingredient Name	Strength			
CETYL ALCOHOL (UNII: 936JST6JCN)				
GLYCERIN (UNII: PDC6A3C0OX)				
PETROLATUM (UNII: 4T6H12BN9U)				
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)				
MINERAL OIL (UNII: T5L8T28FGP)				
WATER (UNII: 059QF0KO0R)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)				
XANTHAN GUM (UNII: TTV12P4NEE)				
PEG-100 STEARATE (UNII: YD01N1999R)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:50268-682- 15	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2022		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	04/11/2022		

Labeler - AVPAK (832926666)

Revised: 3/2024 AvPAK