PLEO PEF- candida parapsilosis ointment Sanum Kehlbeck GmbH & Co. KG

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Pleo® Pef OINTMENT 3X

for topical use only Homeopathic Antipruritic / External Analgesic Medicine 1.05 oz (30g)

Indications

For temporary relief of external anal itching and irritation.

DOSAGE

Apply thinly on the affected area 1–3 times daily.

WARNING

If symptoms persist more than a few days, contact a licensed practitioner. As with any other drugs, if you are pregnant or nursing a baby, seek the advice of a health care professional before using this product.

Keep this and all medications out of the reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Protect from light and heat.

Tamper Evident

Do not use this product if metal seal under the tube cap is missing or broken.

INGREDIENTS

Active Ingredients

1 g ointment contains; 0.1 g Candida parapsilosis 3X.

Inactive Ingredients

lactic acid, lanolin alcohol, glycerylmonostearate, propylene glycol, polyglyceryl-3, cocoate magnesium sulfate, water.

Made in Germany

Manufactured by:

SANUM-Kehlbeck

GmbH & Co. KG, Hoya, Germany

Distributed by:

SANUM USA Corp., 1465 Slater Road, Ferndale, WA 98248

Rev. 06/2009

PRINCIPAL DISPLAY PANEL - 30 g Ointment Carton

Pleo® Pef OINTMENT 3X

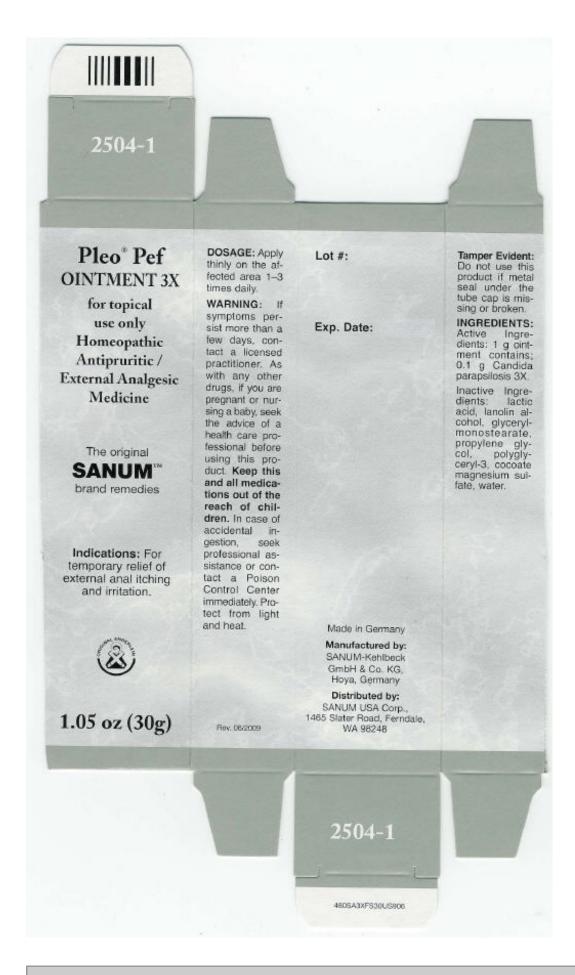
for topical
use only
Homeopathic
Antipruritic /
External Analgesic
Medicine

The original **SANUM**TM brand remedies

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ORIGINAL ENDERLEIN

1.05 oz (30g)



PLEO PEF

candida parapsilosis ointment

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60681-2504		
Route of Administration	TOPICAL				

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
candida parapsilosis (UNII: 0KZ676D44N) (candida parapsilosis - UNII:0KZ676D44N)	candida parapsilosis	3 [HP_X] in 30 g	

Inactive Ingredients			
Ingredient Name	Strength		
glyceryl monostearate (UNII: 230 OU9 XXE4)			
lactic acid (UNII: 33X04XA5AT)			
propylene glycol (UNII: 6DC9Q167V3)			
water (UNII: 059QF0KO0R)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:60681-2504-1	1 in 1 CARTON				
1		30 g in 1 TUBE				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
Unapproved homeopathic		03/19/1996		

Labeler - Sanum Kehlbeck GmbH & Co. KG (318386133)

Revised: 10/2009 Sanum Kehlbeck GmbH & Co. KG