

PCA SKIN SKIN PROCEDURE- petrolatum ointment
CP Skin Health Group, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

PCA Skin Skin Procedure Ointment

Drug Facts

Active ingredient

Petrolatum 60%

Purpose

Skin protectant (ointment)

Uses

• temporarily protects minor: ° cuts ° scrapes ° burns • helps prevent and temporarily protects and helps relieve chafed, chapped or cracked skin • helps prevent and protect skin from the drying effects of wind and cold weather

Warnings

For external use only

When using this product

do not get into eyes

Stop use and ask a doctor if

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

Do not use on

- deep or puncture wounds
- animal bites
- serious burns

Do not use on damaged and broken skin

Keep out of reach of children

Directions

- Apply as needed or as directed by physician
- for best results, apply liberally to moist skin.

Other information

- Do not use if package is broken or open
- protect this product from excessive heat and direct sun
- noncomedogenic
- fragrance-free
- hypoallergenic

Inactive ingredients

Butyrospermum Parkii (Shea) Butter, Caprylic/Capric Triglyceride, Carthamus Tinctorius (Safflower) Seed Oil, Helianthus Annuus (Sunflower) Extract, Helianthus Annuus (Sunflower) Seed Wax, Oryza Sativa (Rice) Bran Extract, Panthenol, Phenoxyethanol, Rosmarinus Officinalis (Rosemary) Leaf Extract, Silybum Marianum Fruit Extract, Stearyl Glycyrrhetinate, Tocopherol, Tocopheryl Acetate.

Questions?

Please call **877.722.7546**

Package Labeling:

skin procedure ointment

occlusion, soothing, and healing

pommade pour la peau après intervention occlusion, calmante, et apaisante



2.1 fl oz (62.1 mL)

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 Scottsdale, AZ 85251, USA

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PCA SKIN SKIN PROCEDURE

petrolatum ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68726-680
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	600 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SHEA BUTTER (UNII: K49155WL9Y)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
SAFFLOWER OIL (UNII: 65UEH262IS)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
HELIANTHUS ANNUUS SEED WAX (UNII: 42DG15CHXV)	
RICE BRAN (UNII: R60QEP13IC)	
PANTHENOL (UNII: WW9CM0067Z)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ROSEMARY (UNII: IJ67X351P9)	
MILK THISTLE (UNII: U946SH95EE)	
STEARYL GLYCYRRHETINATE (UNII: 3YYE6VJS0P)	
TOCOPHEROL (UNII: R0ZB2556P8)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68726-680-01	62.1 mL in 1 TUBE; Type 0: Not a Combination Product	09/10/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M016	09/10/2019	

Labeler - CP Skin Health Group, Inc. (611921669)

Revised: 1/2023

CP Skin Health Group, Inc.