

RESURFIX PLUS- petrolatum ointment
Topiderm, Inc.

Resurfix Ointment Plus

Drug Facts

Active ingredients

Petrolatum U.S.P

Purpose

Skin protectant

Uses

- A unique dressing formulated to aid in the Skin Barrier healing process. Also provides protection and helps heal stressed skin conditions such as chapped, dry and windburned skin.

Warnings

- Not to be applied over puncture wounds, infections, or lacerations.
- If condition worsens or does not improve within 7 days, contact a physician.
- **Keep out of reach of children.** In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.
- For external use only, not to be swallowed.
- Avoid contact with eyes.

Directions

- Apply as directed by your physician.

Inactive ingredients

Allantoin, Benzyl Alcohol, Camellia Sinensis (Green Tea) Leaf Extract, Ceramide 2, Ceramide-3, C12-15 Alkyl Benzoate, Ethylhexyl Palmitate, Glycine Soja (Soybean) Seed Extract, Lanolin Alcohol, Mineral Oil, Palmitoyl Oligopeptide, Panthenol, PEG-10 Rapeseed Sterol, Phospholipids, Saccharomyces Lysate Extract, Sorbitan Isostearate, Squalane, Tocopheryl Acetate, Tribehenin, Yucca Glauca Root Extract, Water.

PRINCIPAL DISPLAY PANEL - 100 g Tube Label

REPLENIX®

RESURFIX+SKIN BARRIER
HEALING OINTMENT
SKIN PROTECTANT

Post Aesthetic • Post Procedure • Post Treatment
Ceramide 2 & 3 • LYCD
Green Tea Extract • Squalane NF
Panthenol • Vitamin E • Allantoin
Petrolatum USP

Net wt. 3.5 oz. (100 g.)

Topix Pharmaceuticals, Inc. • N. Amityville, NY 11701

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rev 02/16

Made in U.S.A.

884

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RESURFIX PLUS

petrolatum ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51326-884
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	86 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALLANTOIN (UNII: 344S277G0Z)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CERAMIDE NG (UNII: C04977SRJ5)	
CERAMIDE NP (UNII: 4370DF050B)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
SOYBEAN (UNII: L7HT8F1ZOD)	
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)	
MINERAL OIL (UNII: T5L8T28FGP)	
PALMITOYL TRIPEPTIDE-1 (UNII: RV743D216M)	
PANTHENOL (UNII: WW9CM0067Z)	
PEG-10 RAPESEED STEROL (UNII: 258O76T85M)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
SACCHAROMYCES LYSATE (UNII: R85W246Z1C)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
SQUALANE (UNII: GW89575KF9)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TRIBEHENIN (UNII: 8OC9U7TQZ0)	
YUCCA GLAUCA ROOT (UNII: 1A15YBH7N1)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:51326-884-03	100 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2000	
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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M016	01/01/2000	

Labeler - Topiderm, Inc. (049121643)

Registrant - Topiderm, Inc. (049121643)

Establishment

Name	Address	ID/FEI	Business Operations
Topiderm, Inc.		049121643	MANUFACTURE(51326-884)

Establishment

Name	Address	ID/FEI	Business Operations
Topix Pharmaceuticals, Inc.		117745066	PACK(51326-884)

Revised: 11/2019

Topiderm, Inc.