REPLENIX BP ACNE- benzoyl peroxide gel
Topiderm, Inc.

Replenix® BP Acne

Drug Facts

Active ingredient

Benzoyl Peroxide, USP 10%

Purpose

Anti-acne

Uses

- Topical acne medication.

Warnings

- **When using this product avoid unnecessary sun exposure and use a sunscreen.**
- For external use only.
- Keep away from eyes, lips, and mouth.
- If irritation develops, discontinue use and consult a doctor.
- Using other topical acne medications at the same time or immediately following use of this product may increase dryness or irritation of the skin. If this occurs, consult a doctor.
- May bleach fabrics.
- **Keep out of reach of children.** If swallowed, seek professional assistance or contact a Poison Control Center immediately.

Directions

- Cleanse the skin thoroughly before applying.
- Apply a thin layer daily, then gradually increase to two or three times daily.
- If bothersome drying or peeling occurs, reduce applications.
- **If going outside, use a sunscreen.** If irritation or sensitivity develops, discontinue use of both products and consult a doctor.

Inactive ingredients

Carbomer, Purified Water, Sodium Hydroxymethylglycinate, Sodium Lauroyl Sarcosinate, Stearic Acid.

PRINCIPAL DISPLAY PANEL - 57 g Bottle Label

REPLENiX®
Acne

BP ACNE GEL
10% SPOT
TREATMENT

MEDICAL GRADE INGREDIENTS
Benzoyl Peroxide USP 10%

DERMATOLOGIST
• RECOMMENDED •

40 YRS

Net 2 oz (57 g)

REPLENIX BP ACNE

benzoyl peroxide gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51326-841
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	100 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
SODIUM LAUROYL SARCOSINATE (UNII: 632GS99618)	
SODIUM HYDROXYMETHYLGLYCINATE (UNII: DIG6BWZ9XT)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51326-841-20	57 g in 1 BOTTLE; Type 0: Not a Combination Product	05/13/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH DRUG	M006	05/13/2021	

Labeler - Topiderm, Inc. (049121643)

Registrant - Topiderm, Inc. (049121643)

Establishment

Name	Address	ID/FEI	Business Operations
Topiderm, Inc.		049121643	MANUFACTURE(51326-841)

Revised: 6/2024 Topiderm, Inc.