

REPLENIX ACNE SOLUTION BENZOYL PEROXIDE WASH- benzoyl peroxide liquid
Topiderm, Inc.

REPLENIX® ACNE SOLUTIONS 5% BENZOYL PEROXIDE WASH

Drug Facts

Active ingredients

Benzoyl Peroxide USP, 5%

Purpose

Acne treatment

Uses

BP Wash is a therapeutic combination of sudsing cleanser and benzoyl peroxide for the treatment of acne.

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 or sensitivity 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, only one medication should be used unless directed by a doctor.
- May bleach fabrics.
- **Keep out of reach of children.** If swallowed, seek professional assistance or contact a Poison Control Center immediately.

Directions

- Shake well. Wet affected area, wash, and rinse well.
- Use once or twice daily or as directed by a physician.
- 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, Phenoxyethanol, Purified Water, Sodium Benzoate, Sodium C14-16 Olefin

Sulfonate, Stearic Acid USP.

PRINCIPAL DISPLAY PANEL - 200 ml Tube Label

REPLENIX®

ACNE SOLUTIONS

Acne Wash

Benzoyl Peroxide USP, 5%

Net 6.7 fl. oz. (200 ml)

Topix Pharmaceuticals, Inc.

N. Amityville, NY 11701

REPLENiX[®]
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N. Amityville, NY 11701

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R0519

Made in U.S.A.

1110

REPLENIX ACNE SOLUTION BENZOYL PEROXIDE WASH

benzoyl peroxide liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51326-111
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)		BENZOYL PEROXIDE	50 mg in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
WATER (UNII: 059QF0KO0R)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51326-111-01	200 mL in 1 TUBE; Type 0: Not a Combination Product	11/01/2017	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug		M006	11/01/2017	

Labeler - Topiderm, Inc. (049121643)

Registrant - Topiderm, Inc. (049121643)

Establishment

Name	Address	ID/FEI	Business Operations
Topiderm, Inc.		049121643	MANUFACTURE(51326-111)

Establishment

Name	Address	ID/FEI	Business Operations
Topix Pharmaceuticals, Inc.		117745066	PACK(51326-111)

Revised: 12/2019

Topiderm, Inc.