GLY-SAL 5-2- salicylic acid liquid
Topiderm, Inc.

Gly-Sal 5-2

Drug Facts

Active ingredient

Salicylic Acid USP, 2%

Purpose

Acne medication

Uses

Skin cleanser for the treatment of acne.

Warnings

For external use only

Keep away from eyes, lips, and mouth.

Using other topical medications, while using this product or immediately thereafter, may increase dryness or irritation. If this occurs discontinue use and see your doctor.

Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center.

Sunscreen use is recommended with any glycolic acid product and for an additional week thereafter because some individuals may be more sensitive to sunlight.

Directions

Wet affected area, apply, and rinse well.

- Because excessive drying of the skin can occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a physician
- If bothersome dryness occurs, reduce application to once a day or every other day.

Inactive ingredients

Camellia sinensis (Green Tea) Polyphenols, Cocamidopropyl Betaine, Disodium EDTA, Fragrance, Glycolic Acid, Linoleamidopropyl PG-Dimonium Chloride Phosphate, Polysorbate-20, Purified Water, Red #40 [CI-16035], Sodium C12-15 Pareth-15 Sulfonate, Sodium C14-16 Olefin Sulfonate, Triethanolamine, Yellow #5 [CI-19140], Zinc PCA.

PRINCIPAL DISPLAY PANEL - 200 ml Bottle Label

COMPLIMENTS OF

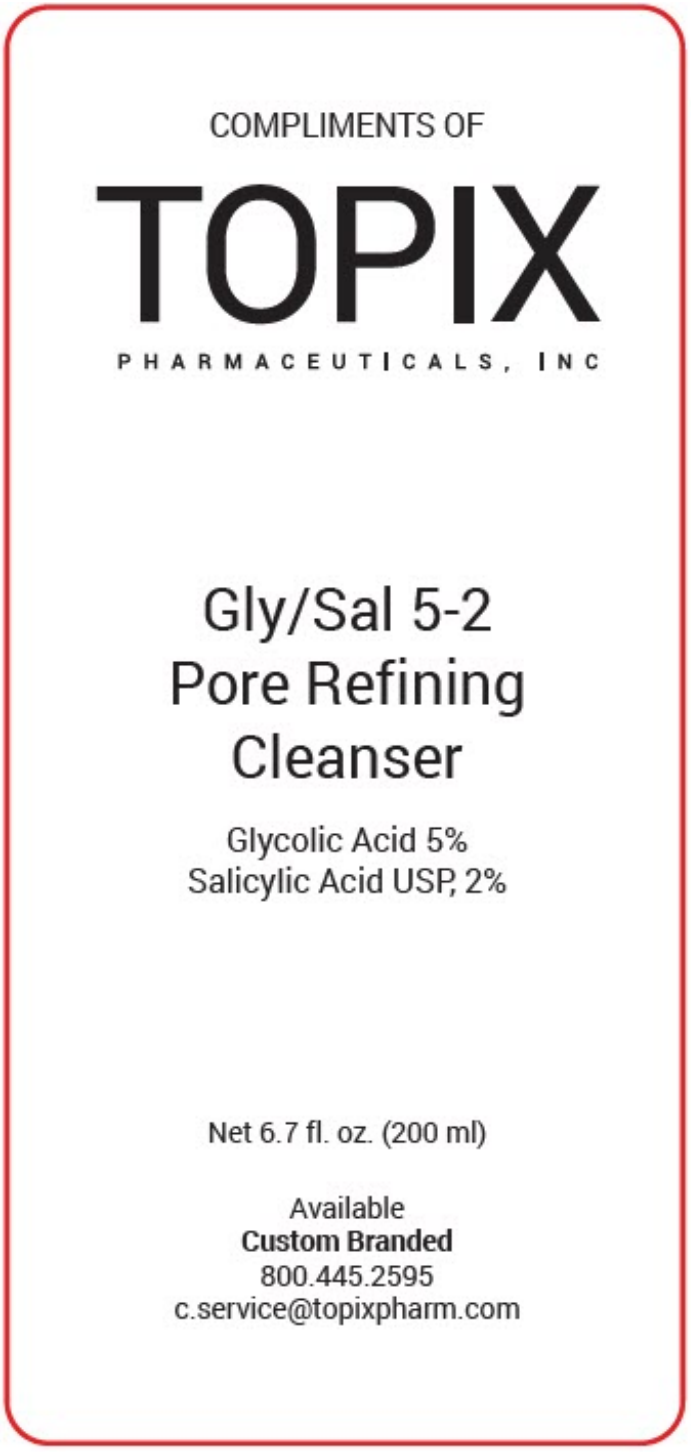
TOPIX
PHARMACEUTICALS, INC

Gly/Sal 5-2
Pore Refining
Cleanser

Glycolic Acid 5%
Salicylic Acid USP, 2%

Net 6.7 fl. oz. (200 ml)

Available
Custom Branded
800.445.2595
c.service@topixpharm.com



GLY-SAL 5-2

salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51326-927
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCOLIC ACID (UNII: 0WT12SX38S)	
LINOLEAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE (UNII: 5Q87K461JO)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
WATER (UNII: 059QF0KO0R)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
SODIUM C12-15 ALKETH-15 SULFONATE (UNII: 353VA59XH8)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
TROLAMINE (UNII: 9O3K93S3TK)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
ZINC PIDOLATE (UNII: C32PQ86DH4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51326-927-01	200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH DRUG	M006	07/30/2020	

Labeler - Topiderm, Inc. (049121643)

Registrant - Topiderm, Inc. (049121643)

Establishment

Name	Address	ID/FEI	Business Operations
Topiderm, Inc.		049121643	MANUFACTURE(51326-927)

Revised: 7/2024 Topiderm, Inc.