

**GLY-SAL 10-2- salicylic acid liquid**  
**Topiderm, Inc.**

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**Gly-Sal 10-2**  
**Spray**

***Drug Facts***

**Active ingredient**

Salicylic Acid 2%

**Purpose**

Acne medication

**Uses**

For the treatment of acne of the body, with the skin enhancement properties of glycolic acid

**Warnings**

- For external use only.
- Direct spray away from face, especially eyes, lips and mouth.
- If irritation develops, discontinue use and consult a doctor.
- Using other topical acne medication 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.
  
- **Keep out of reach of children.**
- If swallowed, seek professional assistance or contact a Poison Control Center immediately.
- Flammable; keep tightly closed, away from flame and heat.
- 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**

- Spray the entire affected area of the body one to three times daily.
- 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 or peeling occurs, reduce application to once a day or every other day.

**Inactive ingredients**

Acetone, Ammonium Hydroxide, Disodium EDTA, Glycolic Acid, Hamamelis Virginiana (Witch Hazel) Water, Imidazolidinyl Urea, Polysorbate-20, Purified Water, SD Alcohol 40B (13% v/v) Sodium Benzoate

**PRINCIPAL DISPLAY PANEL - 89 ml Bottle Label**

COMPLIMENTS OF

TOPIX  
PHARMACEUTICALS, INC

Gly/Sal 10-2  
Spray

Glycolic Acid 10%  
Salicylic Acid USP, 2%

Net 3 fl. oz. (89 ml)

Available  
Custom Branded  
800.445.2595  
[c.service@topixpharm.com](mailto:c.service@topixpharm.com)

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Made in U.S.A.

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**GLY-SAL 10-2**

salicylic acid liquid

**Product Information**

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:51326-938 |
| <b>Route of Administration</b> | TOPICAL        |                           |               |

**Active Ingredient/Active Moiety**

| Ingredient Name   | Basis of Strength | Strength      |
|---|-------------------|---------------|
| <b>SALICYLIC ACID</b> (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ) | SALICYLIC ACID    | 20 mg in 1 mL |

**Inactive Ingredients**

| Ingredient Name  | Strength       |
|--|----------------|
| <b>GLYCOLIC ACID</b> (UNII: 0WT12SX38S)  | 100 mg in 1 mL |
| <b>GREEN TEA LEAF</b> (UNII: W2ZU1RY8B0)   |                |
| <b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)   |                |
| <b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)   |                |
| <b>LINOLEAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE</b> (UNII: 5Q87K461JO) |                |
| <b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)   |                |
| <b>WATER</b> (UNII: 059QF0KO0R)  |                |
| <b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)  |                |
| <b>SODIUM C12-15 ALKETH-15 SULFONATE</b> (UNII: 353VA59XH8)                              |                |
| <b>SODIUM C14-16 OLEFIN SULFONATE</b> (UNII: O9W3D3YF5U)                                 |                |
| <b>TROLAMINE</b> (UNII: 9O3K93S3TK)  |                |
| <b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)  |                |
| <b>ZINC PIDOLATE</b> (UNII: C32PQ86DH4)  |                |

**Packaging**

| # | Item Code        | Package Description   | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:51326-938-01 | 89 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 06/10/2019           |                    |

**Marketing Information**

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC MONOGRAPH DRUG | M006                                     | 06/10/2019           |                    |

**Labeler** - Topiderm, Inc. (049121643)**Registrant** - Topiderm, Inc. (049121643)**Establishment**

| Name           | Address | ID/FEI    | Business Operations    |
|----------------|---------|-----------|------------------------|
| Topiderm, Inc. |         | 049121643 | MANUFACTURE(51326-938) |