RESINOL- petrolatum and resorcinol ointment
ResiCal, Inc.

Resinol®

Active ingredients

Petrolatum 55%, Resorcinol 2%

Uses

- temporarily protects minor
  - cuts
  - scrapes
  - burns
- temporarily relieves pain and itching associated with
  - minor skin irritations
  - minor burns
  - sunburn
  - minor cuts
  - scrapes
  - insect bites
  - rashes due to poison ivy, oak, and sumac

Warnings

For external use only

Do not use on
- deep or puncture wounds
- animal bites
- serious burns

When using this product
- do not get into eyes
- do not apply over large areas of the body

Stop use and ask a doctor if
- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- adults and children 2 years and over: apply to affected area not more than 3 to 4 times daily
- children under 2 years: ask a doctor

Inactive ingredients

calamine, corn starch, lanolin, zinc oxide

Questions or comments?

1-800-204-6434 www.resinol.com

Distributed by: ResiCal, Inc.
Orchard Park, NY 14127

PRINCIPAL DISPLAY PANEL - 85.1 g Jar Label

Resinol®

Topical Analgesic/Skin Protectant
MEDICATED OINTMENT

For fast soothing relief of
MINOR SKIN IRRITATIONS

NET WT 3 OZ (85.1 g)

RESINOL

petrolatum and resorcinol ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67492-105
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	55 g in 100 g
RESORCINOL (UNII: YUL4LO94HK) (RESORCINOL - UNII:YUL4LO94HK)	RESORCINOL	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
LANOLIN (UNII: 7EV65EAW6H)	
ZINC OXIDE (UNII: SOI2LOH54Z)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67492-105-13	25.5 g in 1 JAR; Type 0: Not a Combination Product	06/29/2002	12/31/2024
2	NDC:67492-105-14	85.1 g in 1 JAR; Type 0: Not a Combination Product	06/29/2002	
3	NDC:67492-105-15	50 g in 1 TUBE; Type 0: Not a Combination Product	09/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M017	06/29/2002	

Labeler - ResiCal, Inc. (927081369)

Establishment

Name	Address	ID/FEI	Business Operations
Aurora Labs LLC.		176110547	MANUFACTURE(67492-105)

Establishment

Name	Address	ID/FEI	Business Operations
DSC Laboratories Inc.		097807374	MANUFACTURE(67492-105)

Establishment

Name	Address	ID/FEI	Business Operations
Unipack Inc.		009248480	MANUFACTURE(67492-105)

Establishment

Name	Address	ID/FEI	Business Operations
Pharmacal		800873986	MANUFACTURE(67492-105)

Revised: 12/2025 ResiCal, Inc.