

**REZAMID- sulfur and resorcinol lotion**  
**Summers Laboratories Inc**

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**SUMMERS LABS (as PLD) - REZAMID (11086-022)**

**ACTIVE INGREDIENTS**

SULFUR 5%

RESORCINOL 2%

**PURPOSE**

ACNE TREATMENT LOTION

**USE**

DRIES UP ACNE PIMPLES, HELPS PREVENT NEW PIMPLES

**WARNINGS**

- FOR EXTERNAL USE ONLY

DO NOT USE

- ON BROKEN SKIN
- ON LARGE AREAS OF THE BODY

WHEN USING THIS PRODUCT

- APPLY TO AFFECTED AREAS ONLY
- DO NOT GET INTO EYES
- skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time

STOP USE AND ASK A DOCTOR IF

- IF SKIN IRRITATION OCCURS OR GETS WORSE

KEEP OUT OF REACH OF CHILDREN. IF SWALLOWED, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

**DIRECTIONS**

- shake very well before using
- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer one to three times daily
- because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor
- if bothersome dryness or peeling occurs, reduce application to once a day or every

other day

## INACTIVE INGREDIENTS

water, SD-40 alcohol 28%, zinc oxide, talc, titanium dioxide, propylene glycol, attapulgitte, lauramide DEA, iron oxides, sodium bisulfite, PEG-8 laurate, parachlorometaxylenol, hydroxyethylcellulose, sodium chloride, sodium polynaphthalene sulfonate, EDTA, methyl paraben, xanthan gum, butylparaben, fragrance, simethicone.



## REZAMID

sulfur and resorcinol lotion

### Product Information

**Product Type**

HUMAN OTC DRUG

**Item Code (Source)**

NDC:11086-022

**Route of Administration** TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>SULFUR</b> (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	5 g in 100 mL
<b>RESORCINOL</b> (UNII: YUL4LO94HK) (RESORCINOL - UNII:YUL4LO94HK)	RESORCINOL	2 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>ATTAPULGITE</b> (UNII: U6V729APAM)	
<b>LAURIC DIETHANOLAMIDE</b> (UNII: I29I2VHG38)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>SODIUM BISULFITE</b> (UNII: TZ X5469Z 6I)	
<b>PEG-8 LAURATE</b> (UNII: 762O8IWA10)	
<b>CHLOROXYLENOL</b> (UNII: 0F32U78V2Q)	
<b>HYDROXYETHYL CELLULOSE (140 MPA.S AT 5%)</b> (UNII: 8136Y38GY5)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM NAPHTHALENESULFONATE</b> (UNII: D3F8YRX7TP)	
<b>EDETIC ACID</b> (UNII: 9G34HU7RV0)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>BUTYLPARABEN</b> (UNII: 3QPI1U3FV8)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>SILICON</b> (UNII: Z4152N8IUI)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11086-022-01	56.7 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/30/2013	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	10/30/2013	

**Labeler** - Summers Laboratories Inc (002382612)

