ACNE TINTED SPOT TREATMENT- resorcinol, sulfur liquid
GLYTONE LLC

Glytone Acne Tinted Spot Treatment

Drug Facts

Active Ingredients

Resorcinol 2%

Sulfur 8%

Purpose

Acne Treatment

Use

For the treatment of acne.

- Dries and clears acne blemishes and allows skin to heal.
- Penetrates pores to eliminate most acne blemishes, blackheads, and whiteheads.
- Helps prevent the development of new acne blemishes, blackheads, and whiteheads.

Warnings

For external use only.

When using this product

- apply only to areas with acne
- rinse right away with water if it gets in 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 skin irritation occurs or gets worse

Keep out of reach of children

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

Do not use

on

- broken skin
- large areas of the skin

Directions

- Cleanse the skin thoroughly before applying this product.
- Cover the 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 to 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, PROPYLENE GLYCOL. ALCOHOL DENAT., POLYSORBATE 20, CAPRYLIC/CAPRIC TRIGLYCERIDE, MAGNESIUM ALUMINUM SILICATE, SODIUM STEAROYL, GLUTAMATE, CETEARYL ALCOHOL, GLYCERYL SEARATE, POLYSORBATE 60, CHLOROXYLENOL, ETHYLHEXYLGLYCERIN, FRAGRANCE, PHENOXYETHANOL, POLYGLYCERYL-6 PALMITATE.SUCCINATE, XANTHAN GUM, IRON OXIDES, TITANIUM DIOXIDE

Principal Display Panel

GLYTONE

ACNE TINTED SPOT TREATMENT

ACNE THERAPY

2% RESORCINOL

8% SULFURE

30ML / 1 FL. OZ.

ACNE TINTED SPOT TREATMENT

resorcinol, sulfur liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84262-023
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	0.08 mg
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	0.02 mg

Inactive Ingredients	
Ingredient Name	**Strength**
POLYSORBATE 60 (UNII: CAL22UVI4M)	
CHLOROXYLENOL (UNII: 0F32U78V2Q)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)	
IRON OXIDES (UNII: 1K09F3G675)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

FARNESOL (UNII: EB41QIU6JL)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
XANTHAN GUM (UNII: TTV12P4NEE)	
POLYGLYCERYL-4 PALMITATE (UNII: J6LAJ6FC55)	
WATER (UNII: 059QF0KO0R)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
ALCOHOL (UNII: 3K9958V90M)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84262-023-01	1 in 1 BOTTLE	03/01/2024	
1		30 in 1 CARTON; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	03/01/2024	

Labeler - GLYTONE LLC (119226548)

Registrant - GLYTONE LLC (119226548)

Revised: 12/2024 GLYTONE LLC