CARING MILL ACNE DRYING- sulfur lotion Clixit, Llc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

caring mill™ Acne Drying Lotion

Drug Facts

Active Ingredients

Sulfur 10%

Purpose

Acne Treatment

Use:

For the treatment of acne. Healing and drying of acne. Penetrates pores to control acne. Helps prevent breakouts.

Warnings:

For external use only.

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. Rinse right away with water if product gets in eyes.

Keep out of reach of children.

Directions:

DO NOT SHAKE BOTTLE.

Clean the skin thoroughly before using at night. Apply only to areas with acne. Allow solution to dry and rinse off in the morning.

Do not use on broken skin or large areas of skin. If bothersome dryness or peeling occurs, reduce application to every other day.

Inactive Ingredients:

Allantoin, Aqua (Deionized Water), Camphor, Ethylhexylglycerin, Iron Oxide, Isopropyl Alcohol, Magnesium Aluminium Silicate, Phenoxyethanol, Salicylic Acid, Zinc Oxide.

Store at room temperature.

OVERNIGHT TREATMENT FOR STUBBORN ACNE

Benefits

- Powerful sulfur and salicylic acid formula penetrates pores and dries out stubborn acne overnight, revealing smaller less noticeable blemishes.
- Clears acne and promotes healing
- Emergency spot treatment
- Reduces size of stubborn acne
- Fast results
- Ideal for all skin types

Powerful overnight acne treatment

NOT TESTED ON ANIMALS

Made in the USA

Manufactured for **FSA Store Inc.** 240 W 37th St. 6th Floor New York, NY 10018 Customer Help Line: **(844)** 425-4948

Packaging

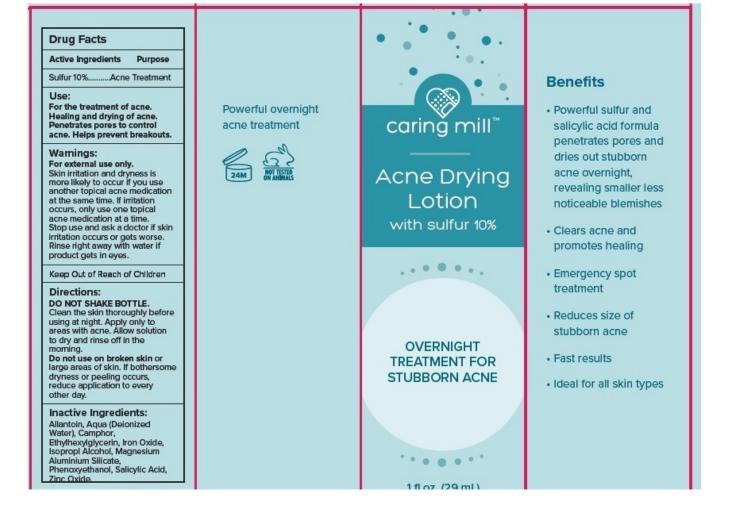
INNER PACKAGE LABEL

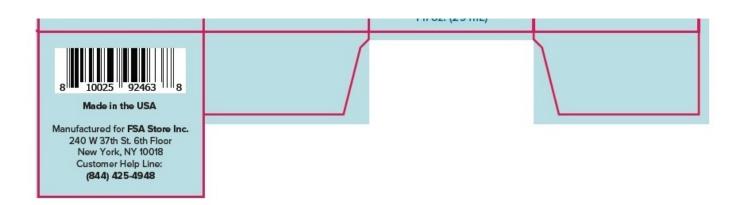






OUTER PACKAGE LABEL





CARING MILL ACNE DRYING

sulfur lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71328-060

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)

SULFUR

100 mg in 1 mL

Inactive Ingredients

Ingredient Name

ALLANTOIN (UNII: 344S277G0Z)

WATER (UNII: 059QF0K00R)

CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)

ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)

FERRIC OXIDE RED (UNII: 1K09F3G675)

ISOPROPYL ALCOHOL (UNII: ND2M416302)

MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)

PHENOXYETHANOL (UNII: HIE492ZZ3T)

SALICYLIC ACID (UNII: HIE492ZZ3T SALICYLIC ACID (UNII: O414PZ4LPZ) ZINC OXIDE (UNII: SOI2LOH54Z)

Product Characteristics

Color	pink	Score
Shape		Size
Flavor		Imprint Code
Contains		

Packaging

7	tem Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:71328-060- 02	1 in 1 BOX	06/21/2021	
:	NDC:71328-060- 01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part333D	06/21/2021				

Labeler - Clixit, Llc (014971694)

Revised: 6/2021 Clixit, Llc