ACNE TREATMENT WITH SULFUR- sulfur ointment
DLC Laboratories, Inc.

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.

Acne Treatment with 5% Sulfur

Drug Facts

Active ingredient

Sulfur, 5%

Purpose

Acne medication

Uses

for the treatment of acne
- penetrates the pores and dries and clears acne pimples and blackheads
- helps to keep new acne pimples and blackheads from forming

Warnings

For external use only

Do not use on
- broken skin
- large areas of the skin

When using this product
- 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 or right after use of this product. If irritation occurs, only use one topical acne medication at a time.
- apply only to areas with acne

Stop use and consult a doctor ifexcessive skin irritation develops or increases.

Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions
- cleanse the skin thoroughly before applying

- cover the entire affected area with a thin layer
- leave on for 10 minutes
- thoroughly remove with water
- 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 ingredient

polyethylene glycol

Questions

1-800-858-3889

Manufactured by: De La Cruz Products
A Division of DLC Laboratories, Inc. | Paramount, CA 90723 USA

PRINCIPAL DISPLAY PANEL - 74 g Tube Label

De La Cruz ®

FOR SENSITIVE SKIN

ACNE
TREATMENT

With 5% Sulfur
- Antibacterial & Anti-inflammatory
- Clears & Helps Prevent Acne

OIL-FREE
WATER-WASHABLE

Ointment

2.6 OZ (74 g)

ACNE TREATMENT WITH SULFUR

sulfur ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24286-1573
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	5 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYETHYLENE GLYCOL 3350 (UNII: G2M7P15E5P)	

Product Characteristics			
Color	yellow	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24286-1573-1	74 g in 1 TUBE; Type 0: Not a Combination Product	08/16/2019	
2	NDC:24286-1573-2	6 g in 1 JAR; Type 0: Not a Combination Product	09/13/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	08/16/2019	

Labeler - DLC Laboratories, Inc. (093351930)

Revised: 6/2023 DLC Laboratories, Inc.