DE LA CRUZ SULFUR ACNE MEDICATION- 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.

DE LA CRUZ ® SULFUR ACNE MEDICATION

Drug Facts

Active ingredient

Sulfur, 10%

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 if excessive 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 - 73.7 g Jar Label

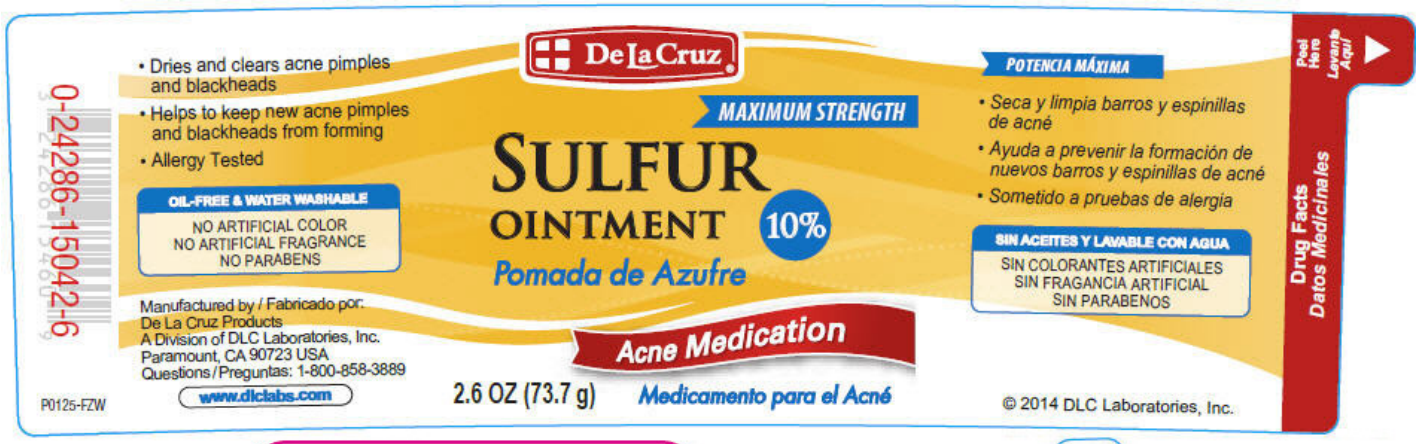
De La Cruz ®

MAXIMUM STRENGTH

SULFUR
OINTMENT
10%

Acne Medication

2.6 OZ (73.7 g)



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Datos Medicinales

Ingrediente activo — **Propósito**
Azufre, 10% ..... Medicamento para el acné

Usos para el tratamiento del acné
- penetra los poros, seca y limpia barros y espinillas de acné
- ayuda a prevenir la formación de nuevos barros y espinillas de acné

Advertencias
Para uso externo únicamente

No lo use
- sobre heridas abiertas
- en áreas grandes de la piel

Cuando use este producto
- evite el contacto con los ojos
- es más probable que ocurra irritación y resequedad de la piel si se usa algún otro medicamento tópico para el acné al mismo tiempo o inmediatamente después del uso de este producto. Si se produce irritación, use solo un medicamento tópico para el acné a la vez.
- aplique únicamente en áreas con acné

Deje de usarlo y consulte con su médico si se desarrolla una excesiva irritación en la piel o si esta aumenta.

Manténgase fuera del alcance de los niños. En caso de ingestión, pida ayuda médica o comuníquese con un Centro de Control de Envenenamientos inmediatamente.

Instrucciones
- limpie completamente la piel antes de aplicar
- cubra el área afectada con una capa fina
- deje actuar por 10 minutos
- retire completamente con agua
- debido a que la piel se puede secar excesivamente, comience con una aplicación al día e incremente gradualmente a dos o tres veces al día si es necesario o si lo recomienda su médico
- si la piel se seca o pela y provoca molestias, reduzca las aplicaciones a una al día, o a días alternos

Ingrediente inactivo
polietilenglicol

¿Preguntas?
1-800-858-3889

DE LA CRUZ SULFUR ACNE MEDICATION

sulfur ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24286-1525
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	10 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:24286-1525-2	73.7 g in 1 JAR; Type 0: Not a Combination Product	08/11/2014	03/26/2021
2	NDC:24286-1525-3	2.8 g in 1 POUCH; Type 0: Not a Combination Product	03/17/2015	12/06/2021
3	NDC:24286-1525-4	6 g in 1 JAR; Type 0: Not a Combination Product	03/17/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	08/11/2014	

Labeler - DLC Laboratories, Inc. (093351930)

Establishment

Name	Address	ID/FEI	Business Operations
DLC Laboratories, Inc.		093351930	manufacture(24286-1525) , label(24286-1525)

Revised: 12/2021 DLC Laboratories, Inc.