

**DHC ACNE SPOT THERAPY- sulfur cream**  
**DHC USA Incorporated**

*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.*

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**DHC Acne Spot Therapy**

**Drug Facts**

**Active Ingredient**

Sulfur 3%

**Purpose**

Acne treatment

**Uses**

For the management of acne. Helps clear up acne blemishes.

**Warnings**

**For external use only**

- **When using this product** skin irritation and dryness is more likely to occur if used with another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time.
- Avoid contact with eyes. If excessive skin irritation develops or increases, discontinue use and consult a doctor.
- **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

**Do not use on**

- Broken skin
- Large areas of the skin

**When using this product**

- Apply only to areas with acne

**Directions**

Cleanse skin thoroughly and tone before applying. 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, butylene glycol, isononyl isononanoate, cetearyl alcohol, sorbitan stearate, cetareth-20, phenoxyethanol, sucrose cocoate, xanthan gum, sodium citrate, citric acid, allantoin, disodium EDTA, magnesium ascorbyl phosphate, tocopherol, sodium hydroxide, brassica campestris (rapeseed) seed oil, glycyrrhiza glabra (licorice) root extract, camellia sinensis leaf extract, royal jelly extract, scutellaria baicalensis root extract, houttuynia cordata extract, perilla ocymoides leaf extract, aloe barbadensis leaf extract

### **Questions or Comments?**

**800-DHC-CARE (342-2273)**

dhccare.com

Distributed by DHC USA Inc.  
Mechanicsburg, PA 17050

### **PRINCIPAL DISPLAY PANEL - 15 g Tube Box**

DHC

Acne

Spot Therapy

Acne Treatment

.52 oz. (15 g) Net wt.

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Acne Treatment  
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### Drug Facts (continued)

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### Drug Facts (continued)

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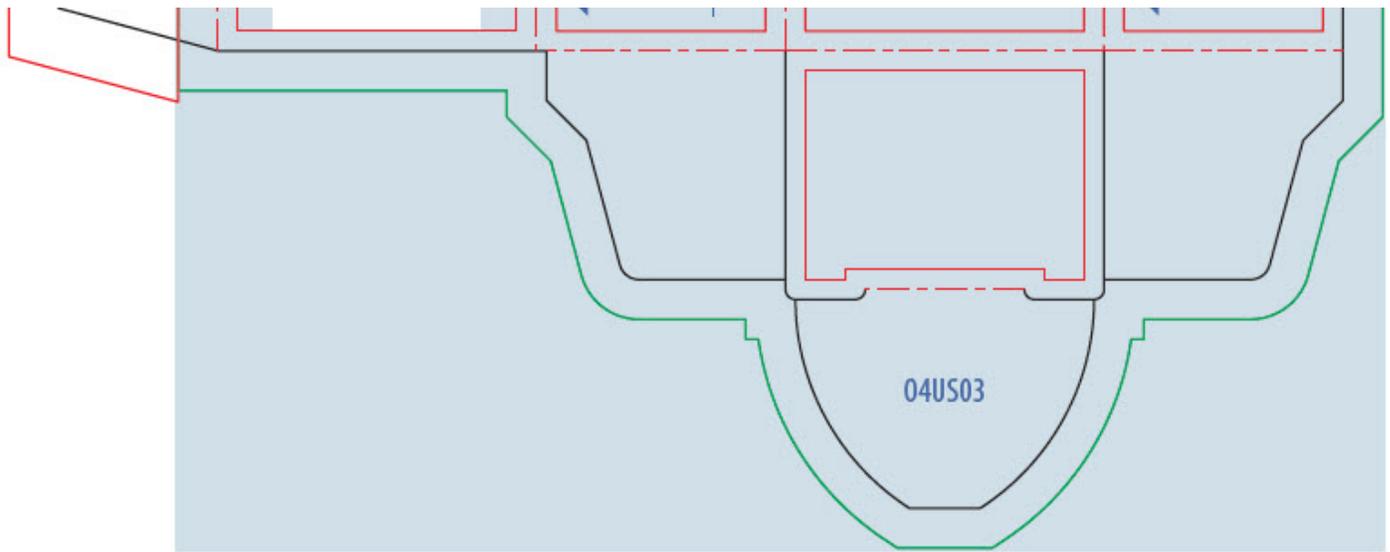
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DHC

Distributed by DHC USA Inc.  
Mechanicsburg, PA 17050  
Made in USA

**Questions or Comments?**  
800-DHC-CARE (342-2273)  
DHCare.com





## DHC ACNE SPOT THERAPY

sulfur cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63433-389
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>Sulfur</b> (UNII: 70FD1KFU70) (Sulfur - UNII:70FD1KFU70)	Sulfur	6 mg in 0.2 g

### Inactive Ingredients

Ingredient Name	Strength
<b>Water</b> (UNII: 059QF0KO0R)	
<b>Butylene Glycol</b> (UNII: 3XUS85K0RA)	
<b>Isononyl Isononanoate</b> (UNII: S4V5BS6GCX)	
<b>Cetostearyl Alcohol</b> (UNII: 2DMT128M1S)	
<b>Sorbitan Monostearate</b> (UNII: NVZ4I0H58X)	
<b>Sodium Citrate, Unspecified Form</b> (UNII: 1Q73Q2JULR)	
<b>Citric Acid Monohydrate</b> (UNII: 2968PHW8QP)	
<b>Phenoxyethanol</b> (UNII: HIE492ZZ3T)	
<b>Xanthan Gum</b> (UNII: TTV12P4NEE)	
<b>Allantoin</b> (UNII: 344S277G0Z)	
<b>.Alpha.-Tocopherol</b> (UNII: H4N855PNZ1)	
<b>Brassica Rapa Subsp. Oleifera Oil</b> (UNII: N4G8379626)	
<b>Magnesium Ascorbyl Phosphate</b> (UNII: 0R822556M5)	
<b>Glycyrrhiza Glabra</b> (UNII: 2788Z9758H)	
<b>Royal Jelly</b> (UNII: L497I37FOC)	
<b>Green Tea Leaf</b> (UNII: W2ZU1RY8B0)	
<b>Houttuynia Cordata Flowering Top</b> (UNII: RH041UUZ22)	

<b>Scutellaria Lateriflora Top</b> (UNII: C6CNB75R61)	
<b>Perilla Frutescens Leaf</b> (UNII: T4L5881Y68)	
<b>Aloe Vera Leaf</b> (UNII: ZY81Z83H0X)	
<b>Edetate Disodium</b> (UNII: 7FLD91C86K)	
<b>Sodium Hydroxide</b> (UNII: 55X04QC32I)	

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63433-389-00	15 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2018	

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part333D	01/01/2018	

**Labeler** - DHC USA Incorporated (004087554)

**Registrant** - ABBE Laboratories, Inc. (781745286)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
ABBE Laboratories, Inc.		781745286	MANUFACTURE(63433-389)

Revised: 2/2022

DHC USA Incorporated