

**SULFO LO- sulfur 3% soap bar soap**  
**Crown Laboratories**

-----  
**Sulfo-Lo Cleansing Bar Soap**

***Active ingredients***

Sulfur 3%

***Purpose***

Acne Treatment

***Uses***

Deep cleansing antibacterial soap for management of acne, blackheads and comedones.

***Warnings***

- **For external use only** • Avoid contact with eyes, wash with water if exposed
- **Do not use on**
  - broken skin • large areas of skin
- **When using this product**
  - apply only to areas with acne • 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 • If undue irritation persists consult a physician
- **Keep out of reach of children.**
  - If swallowed, get medical help or contact a Poison Control Center.

***Directions***

- Work into a lather and rinse with warm water • Use daily as skin cleanser or as directed by a physician.

***Other information***

Store at 20° -25° C (68° -77° F) [see USP Controlled Room Temperature].

***Inactive ingredients***

Sodium Tallowate, Sodium Cocoate/Sodium Palm Kernelate, Water (Aqua), Glycerin,

Fragrance (Parfum), Sodium Chloride, Butyrospermum parkii (Shea) Butter, Yellow 5 (CI19140), Titanium Dioxide (CI77891), Pentasodium Pentetate, Tetrasodium Etidronate, Iron Oxides (CI77492), Tocopheryl Acetate

NDC 0316-0118-35

sulfo lo®

Cleansing Bar Soap

Helps Treat and Manage Acne, Blackheads and Comedones

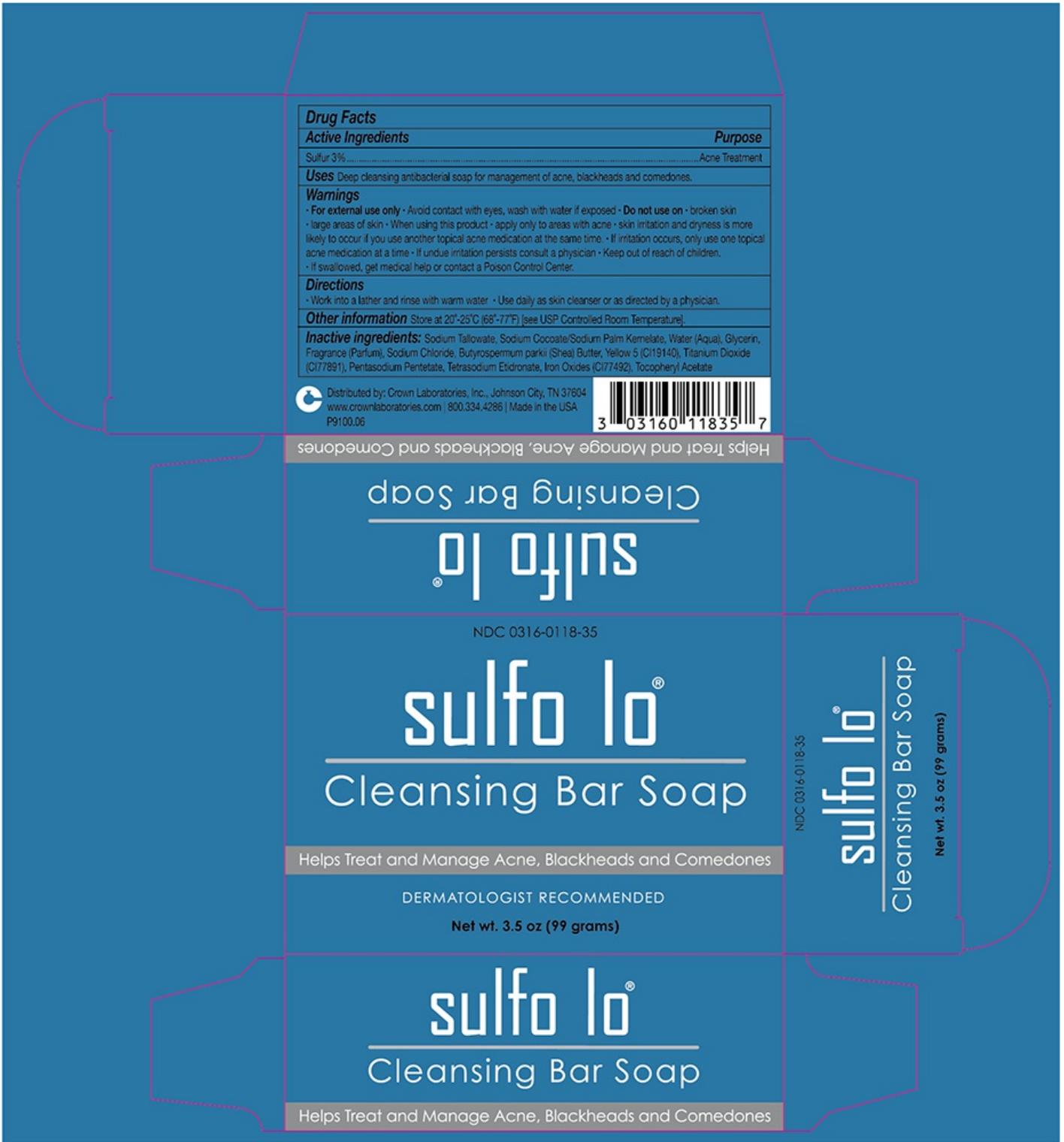
DERMATOLOGIST RECOMMENDED

Net wt. 3.5oz (99 grams)

Distributed by: Crown Laboratories, Inc., Johnson City, TN 37604

[www.crownlaboratories.com](http://www.crownlaboratories.com) | 800.334.4286 | Made in the USA

P9100.06



## SULFO LO

sulfur 3% soap bar soap

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0316-0118
<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	30 mg in 1 g

**Inactive Ingredients**

Ingredient Name	Strength
<b>.ALPHA.-TOCOPHEROL ACETATE, D-</b> (UNII: A7E6112E4N)	
<b>SODIUM TALLOWATE, BEEF</b> (UNII: 07RIK6QMEW)	
<b>SODIUM COCOATE</b> (UNII: R1TQH25F4I)	
<b>SODIUM PALM KERNELATE</b> (UNII: 6H91L1NXTW)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>PENTASODIUM PENTETATE</b> (UNII: 961TOZ5L7T)	
<b>ETIDRONATE TETRASODIUM</b> (UNII: CZZ9T1T1X4)	
<b>BROWN IRON OXIDE</b> (UNII: 1N032N7MFO)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>SHEA BUTTER</b> (UNII: K49155WL9Y)	
<b>BASIC YELLOW 5</b> (UNII: 07BP340B4T)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0316-0118-35	99 g in 1 CARTON; Type 0: Not a Combination Product	11/01/1965	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	11/01/1965	

**Labeler** - Crown Laboratories (119508400)**Registrant** - Crown Laboratories (119508400)