# ACNE TREATMENT GEL- salicylic acid gel S+

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#### **Drug Facts**

## Active ingredient

Salicylic acid 2%

#### **Purpose**

Acne treatment

#### Use

For the treatment of acne

#### Warnings

#### For external use only

## When using this product

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.

# Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

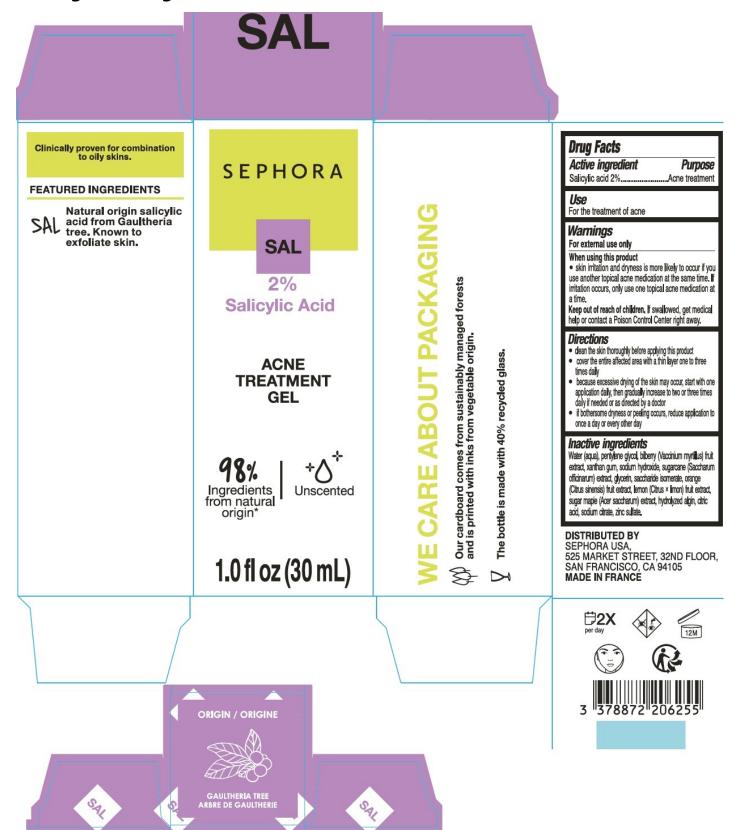
#### **Directions**

- clean the skin thoroughly before applying this product
- cover the entire 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 or 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 (aqua), pentylene glycol, biberry (Vaccinium myrtilus) fruit extract, xanthan gum, sodium hydroxide, sugarcane (Saccharum officinarum) extract, glycerin, saccharide isomerate, orange (Citrus sinensis) fruit extract, lemon (Citrus x limon) fruit extract, sugar maple (Acer saccharum) extract, hydrolyzed algin, citric acid, sodium citrate, zinc sulfate.

## **Package Labeling:**



# SEPHORA

SAL

2%
Salicylic Acid
ACNE TREATMENT GEL

1.0 fl oz (30 mL)

#### THIS UNIT NOT LABELED FOR RETAIL SALE

ACTIVE INGREDIENT: SALICYLIC ACID 2%
WARNINGS: FOR EXTERNAL USE ONLY.
WHEN USING THIS PRODUCT SKIN IRRITATION AND
DRYNESS IS MORE LIKELYTO OCCUR IFYOU USE ANOTHER
TOPICAL ACNE MEDICATION ATTHE SAMETIME. IF IRRITATION
OCCURS, ONLY USE ONE TOPICAL ACNE MEDICATION AT A TIME.
KEEPOUT OF REACH OF CHILDREN. IF SWALLOWED, GET
MEDICAL HELPOR CONTACT A POISON CONTROL CENTER RIGHT
AWAY.

DISTRIBUTED BY SEPHORA USA, 525 MARKET STREET, 32ND FLOOR, SAN FRANCISCO, CA 94105. MADE IN FRANCE.







#### ACNE TREATMENT GEL

salicylic acid gel

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:31720-301

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient Name

Basis of Strength

SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)

SALICYLIC ACID

20 mg in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
PENTYLENE GLYCOL (UNII: 50C1307PZG)		
MYRICA CERIFERA FRUIT (UNII: 2VZ27D14UH)		
XANTHAN GUM (UNII: TTV12P4NEE)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		
SUGARCANE (UNII: 81H2R5AOH3)		
GLYCERIN (UNII: PDC6A3C0OX)		
SACCHARIDE ISOMERATE (UNII: W8K377W98I)		
ORANGE (UNII: 5EVU04N5QU)		
<b>LEMON</b> (UNII: 24RS0A988O)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
SODIUM CITRATE (UNII: 1Q73Q2JULR)		
ZINC SULFATE (UNII: 89DS0H96TB)		

Packaging				
#	tem Code	Package Description	Marketing Start Date	Marketing End Date
	NDC-21720 201			

1	08 NDC:31/20-301-	1 in 1 BOX	12/05/2022	
1		30 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 Drug	M006	12/05/2022		

# **Labeler -** S+ (572406531)

Revised: 12/2023 S+