

**CLEAR AND COVER ACNE TREATMENT CONCEALER CLEAR AND COVER ACNE
TREATMENT CONCEALER 13 FUDGE- salicylic acid cream
S+**

**CLEAR AND COVER ACNE TREATMENT CREAM CONCEALER/CLEAR AND COVER
ACNE TREATMENT CREAM CONCEALER 13 fudge**

Drug Facts

Active ingredient

Salicylic Acid 2%

Purpose

Acne Treatment

Use

- for the treatment of acne
- clears acne blemishes
- helps prevent new acne blemishes

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

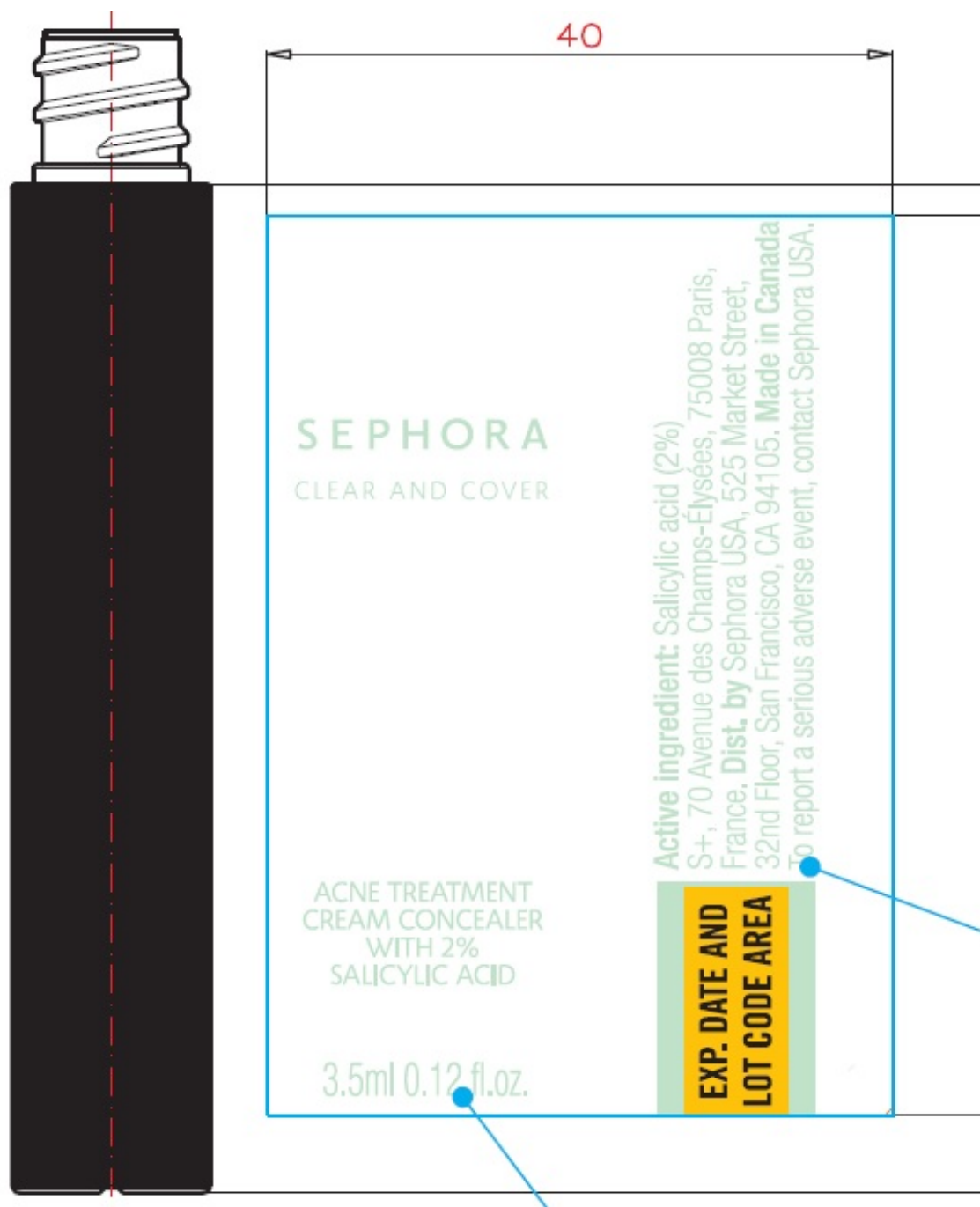
Inactive ingredients

Water (Aqua), Cyclopentasiloxane, Butylene Glycol, Dimethicone, Trisiloxane, Kaolin, Polyisobutene, PEG/PPG-18/18 Dimethicone, Boron Nitride, Cyclodextrin, Talc, Cetyl PEG/PPG-10/1 Dimethicone, Distearidimonium Hectorite, Magnesium Sulfate, Methyl Methacrylate Crosspolymer, Trimethylsiloxysilicate, Phenoxyethanol, Bisabolol, Caprylyl Glycol, Glycerin, Triethoxycaprylylsilane, Chamomilla Recutita (Matricaria) Flower Extract, Tocopherol, Tocopheryl Acetate, Pentaerythrityl Tetra-Di-T-Butyl Hydroxyhydrocinnamate.

May Contain (+/-): Zinc Oxide (CI 77947), Iron Oxides (CI 77491/CI 77492/CI 77499).

Package Labeling:





**CLEAR AND COVER ACNE TREATMENT CONCEALER CLEAR AND
COVER ACNE TREATMENT CONCEALER 13 FUDGE**

salicylic acid cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:31720-965
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
TRISILOXANE (UNII: 9G1ZW13R0G)	
KAOLIN (UNII: 24H4NWX5CO)	
PEG/PPG-18/18 DIMETHICONE (UNII: 9H0AO7T794)	
BORON NITRIDE (UNII: 2U4T60A6YD)	
TALC (UNII: 7SEV7J4R1U)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)	
METHYL METHACRYLATE/GLYCOL DIMETHACRYLATE CROSSPOLYMER (UNII: EG97988M5Q)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
LEVOMENOL (UNII: 24WE03BX2T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
GLYCERIN (UNII: PDC6A3C0OX)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
CHAMOMILE (UNII: FGL3685T2X)	
TOCOPHEROL (UNII: R0ZB2556P8)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:31720-965-18	1 in 1 BOX	06/01/2020	
1		3.5 mL in 1 BOTTLE, WTH APPLICATOR; 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	06/01/2020	

Labeler - S+ (572406531)

Revised: 12/2023

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