OKAY FACIAL SERUM- salicylic acid gel
Xtreme Tools International, 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.

OKAY Facial Serum- SALICYLIC ACID ACNE MEDICATION

Drug Facts

Active Ingredient

Salicylic Acid (2%)

Purpose

Acne Treatment

Use

For the treatment of acne.

Warning

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 medication at a time. Avoid contact with eyes. If contact occurs, flush thoroughly with water.

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), Propanediol, Xylitylglucoside, Anhydroxylitol, Xylitol, C15-19 Alkane (Natural), Caprylyl/Capryl Glucoside, Polyacrylate Crosspolymer-6, Sodium Hydroxide, Carica Papaya (Papaya) Fruit Extract, Psidium Guajava Fruit Extract, Phenoxyethanol, Ethylhexylglycerin.

Questions or comments?
Call 1-305-622-7474 Mon-Fri 9 am-5 pm

PACKAGE LABEL

NDC 74553-014-01

OKAY FACIAL SERUM

salicylic acid gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74553-014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Salicylic Acid (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	Salicylic Acid	2 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPANEDIOL (UNII: 5965N8W85T)	
XYLITYLGLUCOSIDE (UNII: O0IEZ166FB)	
ANHYDROXYLITOL (UNII: 8XWR7NN42F)	
XYLITOL (UNII: VCQ006KQ1E)	
C15-19 ALKANE (UNII: CI87N1IM01)	
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE (UNII: KBC00G95HI)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PAPAYA (UNII: KU94FIY6JB)	
GUAVA (UNII: 74O70D6VG0)	
Phenoxyethanol (UNII: HIE492ZZ3T)	
Ethylhexylglycerin (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74553-014-01	1 in 1 BOX	09/11/2023	
1		30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M006	09/11/2023	

Labeler - Xtreme Tools International, Inc (125398904)

Revised: 9/2023 Xtreme Tools International, Inc