SIGNATURE CARE BLEMISH AND BLACKHEAD CONTROL- salicylic acid emulsion SAFEWAY, 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.

SIGNATURE CARE BLEMISH AND BLACKHEAD CONTROL APRICOT SCRUB

Active ingredient

Salicylic Acid 2%

Purpose

Acne Treatment

Uses

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.
- 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

- cover the entire affected area with a thin layer and rinse thoroughly 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.

Other information

- may stain some fabrics
- store at room temperature

Inactive ingredients

water, juglans regia (walnut) shell powder, glyceryl stearate SE, propylene glycol, sodium laureth sulfate, zea mays (corn) kernel meal, cocamidopropyl betaine, cetearyl alcohol, triethanolamine, cetyl alcohol, PEG-100 stearate, glcyeryl stearate, cetyl acetate, ceteareth-20, polysorbate 60, carbomer, acetylated lanolin alcohol, fragrance (parfum), phenethyl alcohol, PPG-2-methyl ether, limonene, linalool, methylisothiazolinone, glycerin, prunus armeniaca (apricot) fruit extract, titanium dioxide (CI177891).

SIGNATURE CARE BLEMISH AND BLACKHEAD CONTROL

salicylic acid emulsion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21130-720
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 g

Inactive Ingredients	
Ingredient Name	**Strength**
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
JUGLANS REGIA SHELL (UNII: PJ10MT7VKA)	
CORN GRAIN (UNII: C1Z9U7094Z)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL STEARATE/PEG-100 STEARATE (UNII: RD25J5V947)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWJY)	
CETYL ACETATE (UNII: 4Q43814HXS)	
ACETYLATED LANOLIN ALCOHOLS (UNII: SNN716810P)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PPG-2 METHYL ETHER (UNII: RQ1X8FMQ9N)	
APRICOT (UNII: 269CJD5GZ9)	
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMOPOLYMER TYPE B (ALLYL SUCROSE CROSSLINKED) (UNII: Z135WT9208)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
TROLAMINE (UNII: 9O3K93S3TK)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
PHENYLETHYL ALCOHOL (UNII: ML9LGA7468)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
LIMONENE, (+/-)- (UNII: 9MC3I34447)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:21130-720-12	227 g in 1 TUBE; Type 0: Not a Combination Product	08/13/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	08/13/2014	

Labeler - SAFEWAY, INC. (009137209)

Revised: 7/2021 SAFEWAY, INC.