MEDICATED APRICOT SCRUB- salicylic acid gel
United Natural Foods, Inc. dba UNFI

Equaline 041.006/041AK
Medicated Apricot Scrub

Active ingredient

Salicylic acid 2%

Purpose

Acne medication

Use

for the treatment of acne

Warnings

For external use only

When using this product

- skin irritation and dryness is more likely to occure 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

- moisten face with water. Apply product with wet fingertips and masage over face. Rinse well and pat dry
- avoid contact with the eyes. If contact occurs, flush thoroughly with water.

Other information

store at room temperature

Inactive ingredients

water, Juglans regia (walnut) shell powder, glyceryl stearate SE, glycerin, cetearyl alcohol, cetyl alcohol, decyl glucoside, Zea mays (corn) kernel meal, sodium hydroxide,

cocamidopropyl betaine, glyceryl stearate, PEG-100 stearate, Prunus armeniaca (apricot) fruit extract, ceteareth-20, carbomer, polysorbate 60, fragrance, PPG-2 methyl ether, phenethyl alcohol, methylisothiazolinone, benzyl alcohol, limonene, titanium dioxide

Questions?

Call 1-855-423-2630

Disclaimer

*This product is not manufactured or distributed by Unilever, distributor of St. Ives ®Blemish Control Apricot Scrub.

Adverse reaction

Like it or let us make it right.

That's our quality promise.

855-423-2630

DISTRIBUTED BY UNFI

PROVIDENCE, RI 02908 USA

Principal Panel Display

EQUALINE®

COMPARE TO St. Ives Blemish Control Apricot Scrub*

BLEMISH AND BLACKHEAD CONTROL

Medicated Apricot Scrub

salicylic acid acne medication with natural exfoliants

dermatologist tested

exfoliates to prevent blemishes and cleanse pores

NET WT 6 0Z (170 g)

MEDICATED APRICOT SCRUB

salicylic acid gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41163-041
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	19.8 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
JUGLANS REGIA SHELL (UNII: PJ10MT7VKA)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
CORN GRAIN (UNII: C1Z9U7094Z)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
APRICOT (UNII: 269CJD5GZ9)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PPG-2 METHYL ETHER (UNII: RQ1X8FMQ9N)	
PHENYLETHYL ALCOHOL (UNII: ML9LGA7468)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-041-30	170 g in 1 TUBE; Type 0: Not a Combination Product	07/06/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	07/06/2015	

Labeler - United Natural Foods, Inc. dba UNFI (943556183)

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