

OIL FREE ACNE WASH- salicylic acid 2% lotion
Topco Associates, LLC

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.

Top Care 947.003/947AC, AD

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 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

- use up to twice a daily. Wet face, apply to hands, add water and work into lather. Massage face gently. Rinse well.
- avoid contact with eyes. If contact occurs, flush thoroughly with water.

Inactive ingredients

water, sodium C14-16 olefin sulfonate, cocamidopropyl betaine, glycerin, sodium chloride, decyl glucoside, disodium EDTA, fragrance, linoleamidopropyl PG-dimonium chloride phosphate, propylene glycol, Aloe barbadensis leaf extract, Anthemis nobilis flower extract, Chamomilla recutita (matricaria) flower extract, yellow 5, red 40

disclaimer

This product is not manufactured or distributed by Neutrogena Corporation.

Adverse Reactions Section

DISTRIBUTED BY TOPCO ASSOCIATES LLC

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principal display panel

Top Care

beauty

OIL-FREE

ACNE WASH

Salicylic Acid

Acne treatment

Compare to

NEUTROGENA

Dermatologist tested

6 FL OZ (177 mL)



OIL FREE ACNE WASH

salicylic acid 2% lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:36800-947
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Salicylic acid (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	Salicylic acid	20.6 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
LINOLEAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: 5Q87K461JO)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CHAMAEMELUM NOBILE FLOWER (UNII: O2T154T6OG)	
CHAMOMILE (UNII: FGL3685T2X)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-947-30	177 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/21/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	05/21/2009	

Labeler - Topco Associates, LLC (006935977)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(36800-947)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(36800-947)