DAILY FACE WASH CVS- salicylic acid cream CVS

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.

Drug Facts

Active ingredientPurposeSalicylic Acid 2%......Acne Medication

Use For the treatment of acne.

Warnings

For external use only.

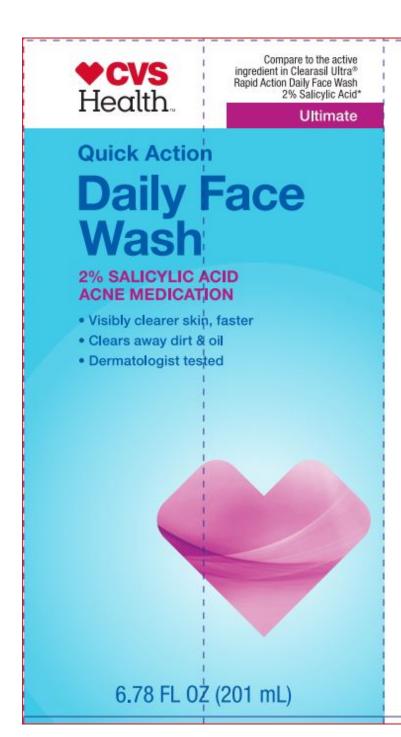
When using this product • skin irritation and dryness are more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time. • to prevent discoloration, avoid contact with clothing

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions • Cleanse twice a day. • Wet face. Apply to hands, add water and work into a lather. • Massage face gently. • Rinse thoroughly.

Inactive ingredients

Water, Sodium Cocoyl Isethionate, Cetearyl Alcohol, Glycerin, Sodium Laureth Sulfate, Sodium Cocoamphoacetate, Cocoamidopropyl Betaine, Sodium Hydroxide, Acrylates/C10-30 Alkyl Acrylate Corsspolymer, Fragrance, Disodium EDTA, Lavendula Stoechas Extract, Helichrysum Italicum Extract, Cistus Monspeliensis Extract.



CVS Health™ Ultimate Quick Action Daily Face Wash is oil free and formulated with 2% salicylic acid acne medication to fight breakouts all day long for

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Distributed by: CVS Pharmacy, Inc.

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Made in the U.S.A. with U.S. and foreign parts V-12412





DAILY FACE WASH CVS

salicylic acid cream

Route of Administration

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:59779-834

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)	
Sodium Cocoyl Isethionate (UNII: 518 XTE 8493)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
Glycerin (UNII: PDC6 A3C0 OX)	
Sodium Laureth Sulfate (UNII: BPV390UAP0)	
Sodium Hydroxide (UNII: 55X04QC32I)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)	
LAVANDULA STOECHAS FLOWERING TOP (UNII: 70759G2U6A)	
HELICHRYSUM ITALICUM FLO WER (UNII: P62Y550 X24)	

	Packagi	ng			
l	# Item	Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:597	79-834-05	201 mL in 1 TUBE; Type 0: Not a Combination Product	09/15/2016	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	09/15/2016	

Labeler - CVS (062312574)

Registrant - Product Quest Mfg (927768135)

Establishment				
Name	Address	ID/FEI	Business Operations	
Product Quest Mfg		927768135	manufacture(59779-834), label(59779-834)	

Revised: 12/2017 CVS