SALICYLIC ACID - salicylic acid liquid
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.

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drug facts

Active ingredient	**Purpose**
Salicylic Acid (0.5%)	Acne Medication

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Salicylic Acid (0.5%)	Acne Medication

Use for the management of acne

Keep out of reach of children. In case of accidental ingestion, get medical help or contact a Poison Control Center right away.

Use for the management of acne

Directions
- Cleanse skin thoroughly before applying medication
- cover the entire affected area with a thin layer one to two 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

Warnings
For external use only.

Using other topical acne medications at the same time or immediately following use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used

unless directed by a doctor.

When using this product avoid contact with eyes. If contact occurs, immediately flush with water.

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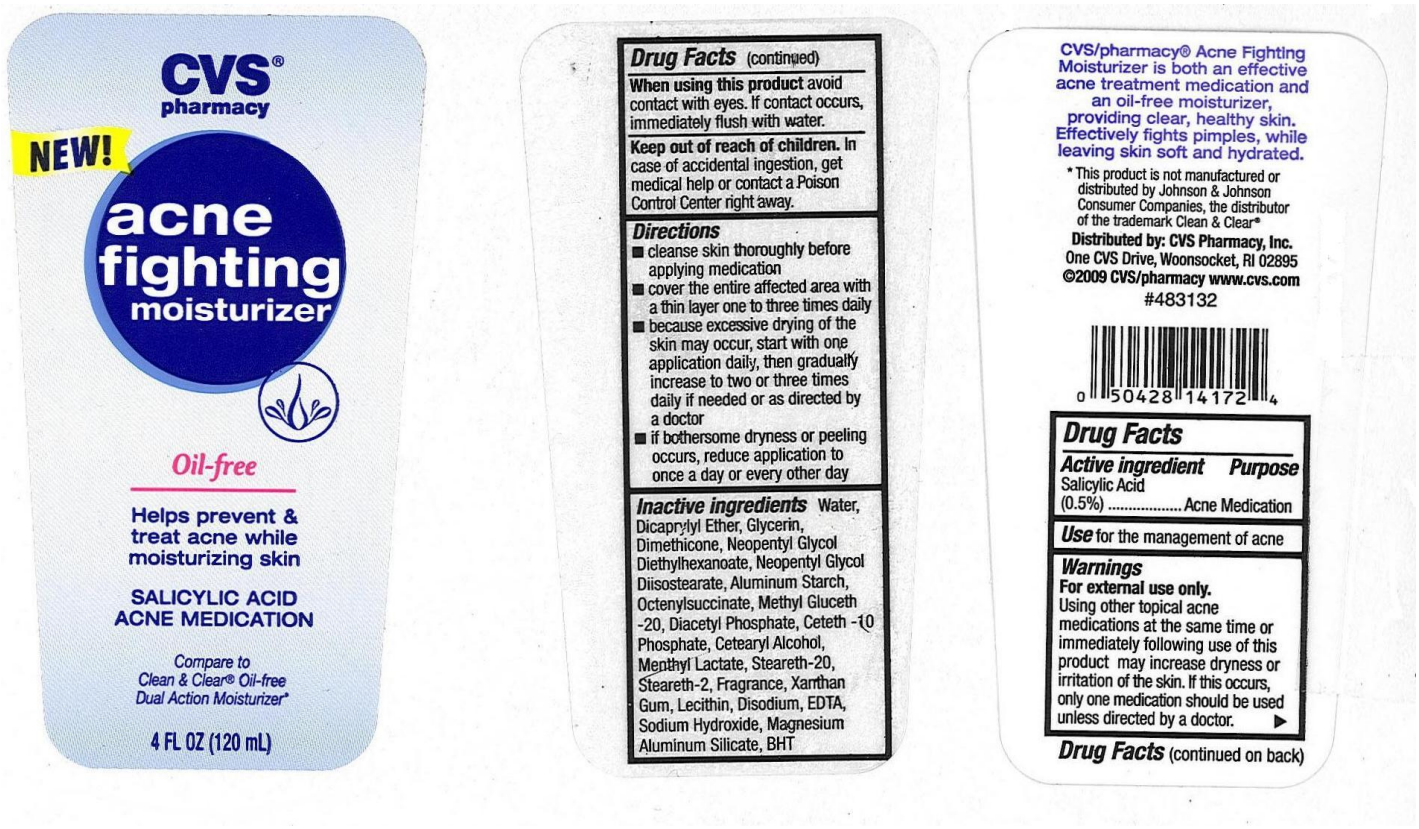
Inactive Ingredients Water, Dicaprylyl Ether, Glycerin, Dimethicone, Neopentyl Glycol Diethylhexanonate, Neopentyl Glycol Diisostearate, Aluminum Starch, Octenylsuccinate, Methyl Gluceth -20, Diacetyl Phosphate, Ceteth-10 Phosphate, Cetearyl Alcohol, Menthyl Lactate, Steareth-20, Steareth-2, Fragrance, Xanthan Gum, Lecithin, Disodium EDTA, Sodium Hydroxide, Magnesium Aluminum Silicate, BHT



SALICYLIC ACID

salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59779-078
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
DICAPRYLYL ETHER (UNII: 77JZM5516Z)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	

MENTHYL LACTATE (UNII: 2BF9E65L7I)	
STEARETH-20 (UNII: L0Q8IK9E08)	
XANTHAN GUM (UNII: TTV12P4NEE)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-078-08	120 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	05/19/2009	

Labeler - CVS (062312574)

Registrant - Pharma Pac, LLC (140807475)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Pac, LLC		140807475	manufacture

Revised: 2/2011 CVS