CVS TRIPLE ACTION ASTRINGENT - salicylic acid liquid CVS Pharmacy

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 ingredient Purpose

Salicylic Acid......Acne treatment

Uses

- treats acne clears blackheads
- helps prevent new acne blemishes and blackheads

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

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- treats acne clears blackheads
- helps prevent new acne blemishes and blackheads

Warnings

For external use only

Flammable: Keep away from fire or flame

When using this product

- avoid contact with eyes. If contact occurs, flush thoroughly with water.
- using other topical acne medication at the same time or right after 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.

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Directions

- cleanse skin thoroughly before applying medication
- moisten cotton ball or pad and pat over face and neck
- because too much drying of teh skin may occur, start with 1 application daily, then gradually increase to 2 to 3 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

Inactive Ingredients

purified water, SD alcohol 40-B, glycerin, sodium citrate, hamamelis virginiana (witch hazel) leaf extract, menthol, fragrance, sodium hydroxide, disodium EDTA, benzophenone-4, citric acid, isopropyl alcohol, polysorbate 60, yellow 5 blue 1



CVS TRIPLE ACTION ASTRINGENT

salicylic acid liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-807
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20.5 mg in 1 mL		

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
ALCOHOL (UNII: 3K9958V90M)			
ISOPROPYL ALCOHOL (UNII: ND2M416302)			
POLYSORBATE 60 (UNII: CAL22UVI4M)			
MENTHOL (UNII: L7T10EIP3A)			
SODIUM CITRATE (UNII: 1Q73Q2JULR)			
BENZOPHENONE (UNII: 701M4TTV9O)			
EDETATE DISO DIUM (UNII: 7FLD91C86K)			
HAMAMELIS VIRGINIANA LEAF (UNII: T07U1161SV)			
GLYCERIN (UNII: PDC6 A3C0 OX)			
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
SO DIUM HYDRO XIDE (UNII: 55X04QC32I)			
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)			

Packaging				
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-807-31	250 mL in 1 BOTTLE, PLASTIC		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part333D	07/08/2010		

Labeler - CVS Pharmacy (062312574)

Registrant - Pharma Pac, LLC (140807475)

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

Revised: 7/2010 CVS Pharmacy