PHARMACYS PRESCRIPTION 8 OZ ACNE CONTROL- salicylic acid liquid
American Consumer Products Corp

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.

Pharmacys Prescription 8 OZ Acne Control

Active Ingredient

Active ingredient: Salicylic Acid 0.5%

Purpose

Purpose: Acne treatment

Uses: for the treatment of acne.

Warnings

Warnings: For external use only.

Flammable, keep away from open flame.

Keep out of reach of children

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

Inactive Ingredients

Water (aqua), Alcohol (24.5%), Propylene Glycol, Glycerin, PEG-32, Algae Extract, DMDM Hydantoin, Dimethicone Propyl PG-Betaine, Sodium Citrate, Aloe Barbadensis Leaf Extract, Benzophenone-4, Denatonium Benzoate, Fragrance, FD&C Yellow No..5.

Directions

Directions: Wash face as you normally would and pat dry with a clean towel. Apply a small amount to cotton ball or pad and applyu using a dabbing motion. Take care to avoid rubbing in eyes. Use astringent two to three times daily or as directed by your doctor to keep your skin clear and healthy. Apply a light moisturizer after using astringent.

When using this product

- avoid contact with eyes. If contact occurs, immediately flush with water.

- 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.

Indications & Usage

If bothersome drying or peeling occurs, reduce application.

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salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72197-015
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.05 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**

ALCOHOL (UNII: 3K9958V90M)	
POLYETHYLENE GLYCOL 1600 (UNII: 1212Z7S33A)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SULISOBENZONE (UNII: 1W6L629B4K)	
PHYMATOLITHON CALCAREUM (UNII: 6J1M3WA0ZK)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72197-015-08	236.58 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/16/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	09/16/2019	

Labeler - American Consumer Products Corp (081101181)

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