PERSONAL CARE DEEP CLEANING ASTRINGENT- salicylic acid liquid Delta Brands & Products 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.

Drug Facts

Active Ingredient

Salicylic Acid 0.5%

Purpose

Acne treatment

Use

for the treatment of acne

Warnings

■ **For external use only** Keep out of eyes ■ If contact with eyes occurs, immediately flush with water.

Using other topical acne drugs at the same time or right after use of this product may increase dryness or irritation of the skin. If this occurs, only one drug should be used unless directed by a doctor.

Keep out of reach of children

If accidentally ingested seek medical help immediately or contact your local Poison Control Center.

Directions

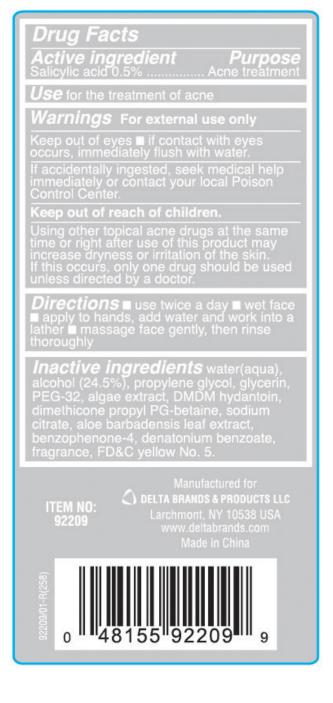
■ use twice a day ■ wet face ■ apply to hands, add water and work into a lather ■ massage face gently, then rinse thoroughly.

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, fragrance, FD&C yellow No. 5.

Package Label





PERSONAL CARE DEEP CLEANING ASTRINGENT

salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72133-200
Route of Administration	TOPICAL		

Active Ingredient/Active	Moiety		
	Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4	4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.5 g in 100 mL
	8	8	

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6 DC9 Q 16 7 V3)	
DMDM HYDANTO IN (UNII: BYR0546 TOW)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
SULISOBENZONE (UNII: 1W6L629B4K)	
DENATO NIUM BENZO ATE (UNII: 4YK5Z54AT2)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
POLYETHYLENE GLYCOL 1500 (UNII: 1212Z7S33A)	

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:72133-200- 08	237 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/09/2018	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	04/09/2018	

Labeler - Delta Brands & Products LLC (080999173)

Revised: 4/2018 Delta Brands & Products LLC