

DERMACLEAR PADS- salicylic acid liquid

Allure Labs

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 2%

Purpose: Acne Treatment

Uses: For treatment of Acne

Warnings:

- For external use only.
- Overuse of this product will cause dryness.

Do not use if you have sensitive skin.

When using this product:

- Avoid contact with the eyes, lips and mouth.
- 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.
- If sensitive to the sun or when increased sun exposure is expected, use a sunscreen.

Stop use and ask a doctor:

- If severe skin irritation occurs.
- When using other topical medication at the same time.

Keep out of reach of children:

- If swallowed, get medical help or contact a Poison Control Center right away.

Gently rub pad over any area in need. Use every other night or as recommended by your skin care professional. Keep the lid securely closed to prevent evaporation. To avoid dryness, do not overuse. Not recommended for those with aspirin / NASID allergies.

Inactive Ingredients:

Alcohol Denat., Water (Aqua), Glycerin, PEG-8/SMDI Copolymer, Glycolic Acid, Aloe Barbadensis Leaf Juice, Ammonium Hydroxide, Melaleuca Alternifolia Leaf Oil, Allantoin, Limonene.

Manufactured for DermaQuest®, Inc.

Hayward, CA 94544

1272 GK, NL Made in USA

dermaquestinc.com



DERMACLEAR PADS			
salicylic acid liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62742-4076
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 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
PEG-8/SMDI COPOLYMER (UNII: CCX72L6NY6)	
GLYCOLIC ACID (UNII: 0WT12SX38S)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
AMMONIA (UNII: 5138Q19F1X)	
TEA TREE OIL (UNII: VIF565UC2G)	
ALLANTOIN (UNII: 344S277G0Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4076-2	1 in 1 CARTON	05/08/2015	
1	NDC:62742-4076-1	85 g in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	05/08/2015	

Labeler - Allure Labs (926831603)

Registrant - Allure Labs (926831603)

Establishment

Name	Address	ID/FEI	Business Operations
Allure Labs		926831603	manufacture(62742-4076)

Revised: 11/2020

Allure Labs