UNIVERSAL CLARIFYING PAD- salicylic acid liquid
Allure Labs

Drug Facts

Active Ingredient: Salicylic Acid 2.0%

Purpose:Acne Treatment

Uses:
For the treatment of acne.

Warnings:
• May cause photosensitivity. Always use SPF. For external use only.
• Overuse of this product will cause dryness.
• Do not use if you have a very sensitive skin.

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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.
• 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.

Directions:
• Twice weekly at night, gently swipe treatment pad over clean skin. Allow to fully absorb. If irritation persists, consult your skincare professional. For best results, follow with B5 Moisture Matte.

N/A

Inactive ingredients:
Water (Aqua), SD Alcohol 40-B, Lactic Acid, Aloe Barbadensis Leaf Juice, Glycerin,

Sodium Hydroxide, Hibiscus Sabdariffa Flower Extract, Kojic Acid, Ethylhexylglycerin, Vaccinium Myrtillus (Bilberry) Extract, Saccharum Officinarum (Sugar Cane) Extract, Citrus Aurantium Dulcis (Orange) Fruit Extract, Citrus Limon (Lemon) Fruit Extract, Acer Saccharum (Sugar Maple) Extract, Vaccinium Corymbosum (Blueberry) Fruit Extract, Morus Alba Fruit Extract, Mandelic Acid, Phytic Acid, Phenoxyethanol

UNIVERSAL CLARIFYING PAD

salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62742-4276
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
LACTIC ACID (UNII: 33X04XA5AT)	
KOJIC ACID (UNII: 6K23F1TT52)	
SUGARCANE (UNII: 81H2R5AOH3)	
MANDELIC ACID (UNII: NH496X0UJX)	
ACER SACCHARUM BARK/SAP (UNII: Z120VL0KAC)	
WATER (UNII: 059QF0KO0R)	
VACCINIUM CORYMBOSUM (BLUEBERRY) FRUIT (UNII: DVH063L9QI)	
BILBERRY (UNII: 9P2U39H18W)	
WHITE MULBERRY (UNII: MN25R0HH5A)	
LEMON (UNII: 24RS0A988O)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HIBISCUS SABDARIFFA FLOWER (UNII: 45TGG6IU6M)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	
ORANGE (UNII: 5EVU04N5QU)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PHYTIC ACID (UNII: 7IGF0S7R8I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4276-2	1 in 1 CARTON	01/07/2026	
1	NDC:62742-4276-1	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 Drug	M006	01/07/2026	

Labeler - Allure Labs (926831603)

Registrant - Allure Labs (926831603)

Establishment

Name	Address	ID/FEI	Business Operations

Allure Labs		926831603	manufacture(62742-4276)