

**DERMACLEAR BHA SPOT TREATMENT- salicylic acid gel**  
**Allure Labs**

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**Drug Facts**

Active Ingredient: Salicylic Acid - 2%

Purpose: Acne Treatment

Uses:

- For the treatment of acne.

Warnings

- For external use only

When using this product

- Avoid contact with eyes, lips and mouth.
- Using other topical acne medication 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 Centre right away.

Directions

- Apply a small portion over the affected area with finger tips and gently massage onto the skin.
- Reapply as needed.

Other ingredients:

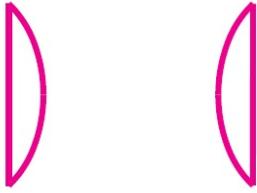
Water (aqua), SD Alcohol 40-B, Butylene Glycol, Hydroxypropylcellulose

Manufactured For Dermaquest ®, Inc.

Hayward, CA 94544

1272 GK, NL Made in USA

dermaquestinc.com



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**DERMAQUEST**

DermaClear BHA Spot Treatment **DermaClear**

0.17 OZ / 4.8 g

**DIRECTIONS:**  
Apply product to affected area using applicator wand. Avoid the eyes, lips and mouth. Use disposable wand for professional use.

**DERMACLEAR BHA SPOT TREATMENT**

salicylic acid gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:62742-4195
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>SALICYLIC ACID</b> (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	2 mg in 1 g

**Inactive Ingredients**

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>HYDROXYPROPYL CELLULOSE, UNSPECIFIED</b> (UNII: 9XZ8H6N6OH)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4195-2	1 in 1 CARTON	12/10/2020	
1	NDC:62742-4195-1	4.8 g in 1 BOTTLE, WTH APPLICATOR; 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	12/10/2020	

**Labeler** - Allure Labs (926831603)**Registrant** - Allure Labs (926831603)**Establishment**

Name	Address	ID/FEI	Business Operations
Allure Labs		926831603	manufacture(62742-4195)

Revised: 12/2024

Allure Labs