

POST TREATMENT BALM- hydrocortisone cream
Allure Labs Inc.

Drug Facts

Active Ingredient: Hydrocortisone 1%

Purpose: Anti-itch

Uses:

- Promotes healing of itchy skin.
- Keeps skin hydrated.

Warnings:

- For external use only.
- Ask a doctor if bleeding occurs or conditions worsens.

Do not use - If rashes appear on skin.

When using this Product:

- Avoid contact with the eyes, lips and mouth.
- Do not use more than directed unless told to do so by a doctor.

Stop use and ask a doctor - If conditions worsens.

Keep out of reach of children:

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

Directions:

- Occlude eyebrows, outer corners of eyes, nose and lips prior to chemical peel.

Inactive Ingredients:

Petrolatum, Carthamus Tinctorius (Safflower) Seed Oil, Tocopheryl Acetate, Zinc Oxide , Titanium Dioxide, Caprylic/Capric Triglyceride, Glyceryl Isostearate, Polyhydroxystearic Acid , Glycosaminoglycans, Dimethicone, Triethoxycaprylylsilane, PEG-8 , Tocopherol , Ascorbyl Palmitate , Ascorbic Acid, Citric Acid

Manufactured for DermaQuest®, Inc.

Hayward, CA 94544

1272 GK, NL Made in USA

dermaquestinc.com

Drug Facts

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Hydrocortisone 1%	Anti-itch

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DERMAQUEST

Post
Treatment
Balm

Professional

1 FL OZ / 29.6 mL

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DERMAQUEST

Post
Treatment
Balm

Professional



2 OZ/56.7 g

POST TREATMENT BALM

hydrocortisone cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62742-4077
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ)	HYDROCORTISONE	1 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GAG (UNII: U5T7CTT5KM)	
ASCORBYL PALMITATE (UNII: QN83US2B0N)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
SAFFLOWER (UNII: 4VBL71TY4Y)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ACETATE ION (UNII: 569DQM74SC)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
ISOSTEARIC ACID (UNII: X33R8U0062)	
TOCOPHEROL (UNII: R0ZB2556P8)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
POLYGLYCERYL-3 PENTARICINOLEATE (UNII: 7Q0OK5DOT4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
PETROLATUM (UNII: 4T6H12BN9U)	
SAFFLOWER OIL (UNII: 65UEH262IS)	
ZINC OXIDE (UNII: SOI2LOH54Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4077-2	29.6 g in 1 TUBE; Type 0: Not a Combination Product	05/12/2015	
2	NDC:62742-4077-1	56.7 g in 1 TUBE; Type 0: Not a Combination Product	05/12/2015	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
OTC Monograph Drug	M017	05/12/2015	

Labeler - Allure Labs Inc. (926831603)

Registrant - Allure Labs (926831603)

Establishment

Name	Address	ID/FEI	Business Operations
Allure Labs		926831603	manufacture(62742-4077)

Revised: 12/2024

Allure Labs Inc.