POST TREATMENT BALM- hydrocortisone cream
Allure Labs Inc.

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: 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

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DERMAQUEST

Post Treatment Balm

Professional

1 FL OZ / 29.6 mL

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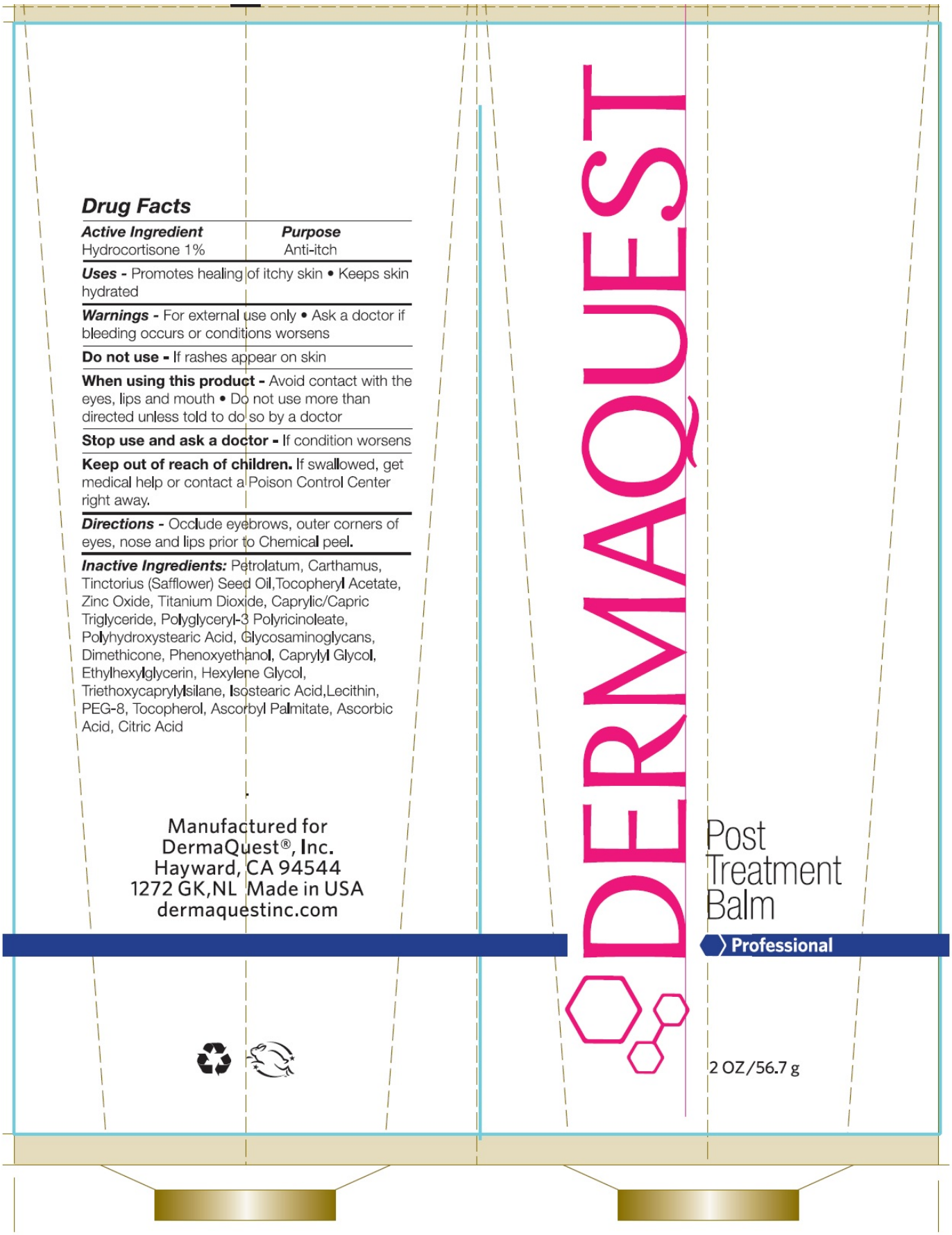

POST TREATMENT BALM

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: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE	1 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
GAG (UNII: U5T7CTT5KM)	
ASCORBYL PALMITATE (UNII: QN83US2B0N)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SAFFLOWER (UNII: 4VBL71TY4Y)	
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)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	

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 Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	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: 12/2020 Allure Labs Inc.