POST PEEL PROTECTANT VITALITY INSTITUTE- hydrocortisone cream Vi Medical Products 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.

Post Peel Protectant Anti-Itch Cream

Active Ingredients Purpose

Hydrocortisone 1% Anti-itch

Uses

- For the temporary relief of of itching associated with minor skin irritations and rashes due to
- Eczema
- insect bites
- poison ivy, poison oak, poison sumac
- soaps and detergents cosmetics
- jewelry
- Keep out of reach of children. If swallowed, get medial help or contact a Poison Control Center rigth away

Stop use and ask a doctor

if condition worsens, or if symptoms persist for more than 7 day or clear up and occur again within a few days, do not use this or any other hydrocortisone products unless you hav consulted a doctor.

Warnings

- ©For external use only
- Avoid Contact with eyes

Directions

- Adults and children under 2 years of age and older: Appyl to affected area not more than 3 to 4 times daily
- Children under 2 years of age: Do not use, consult a doctor

Inactive Ingredients

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Water (Aqua), Cyclopentasiloxane, Demthicone/Vinyl Dimethicone Crosspolymer, Aloe Barbadensis Leaf Juice, Glycerin, Cetearyl Alcohol, Ceteareth-20, Caprylic/Capric Triglyceride, Cetyl Alcohol, Ammonium Acryloyldimethyltaurate/VP Copolymer, Avena Sativa (Oat) Kernel Extract, Chrysanthemum Pathenium (Feverfew) Extract, Tocophoneryl Acetate, Allantoint, Ethylhexylglycerin, Phenoxyethanol

Vitality Institute

Skin Care System

Post Peel Protectant

Anti-itch Cream

30 mL/1 FL OZ



POST PEEL PROTECTANT

ANTI-ITCH **CREAM**

Active Ingredient: Hydrocortisone 1%

Drug Facts

ACTIVE INGREDIENTS

Hydrocortisone 1% Anti-itch

PURPOSE

USES • For the temporary relief of itching associated with minor skin imitations and rashes due to *eczema *insect bites *poison ivy, poison oak, or poison sumac *soaps and detergents cosmetics • jewelry

WARNINGS

•For External use only •Avoid contact with eyes •Stop use and ask a doctor if •condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, do not use this or any other hydrocortisone product unless you have consulted a doctor. • Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS

•Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily • Children under 2 years of age: Do not use, consult a doctor.

INACTIVE INGREDIENTS

Water (Aqua), Caprylic/Capric Triglyceride, Cetearyl Alcohol, Glycerin, Ethylhexyl Palmitate, Colloidal Silver, Butyrospermum, Parkii (Shea Butter), Dimethicone, Glyceryl Stearate, PEG-100 Stearate, Myristyl Myristate, Aloe Barbadensis Leaf Juice, Avena Sativa (Oat) Kernel Extract, Chrysanthemum Parthenium (Feverfew) Extract, Bisabolol, Glycine Soja (Soybean) Oil, Tocopherol, Allantoin, Panthenol, Sodium Polyacrylate, Disodium EDTA, Ethylhexylglycerin, Phenoxyethanol

OTHER INFORMATION •store at 20°C (66° - 77°F)

Sample Size Not For Resale Distributed by: VI Aesthetics™ Made in the USA Los Angeles, CA 90230 | 855.VI.PEELS | www.vipeel.com



30 mL/1 FL OZ

POST PEEL PROTECTANT VITALITY INSTITUTE

hydrocortisone cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70484-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength HYDROCORTISONE (UNII: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ) HYDROCORTISONE 7.5 g in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
CYCLOMETHICONE 5 (UNII: 0 THT5PCI0R)		
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)		

ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528 SWUY)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
AMMONIUM ACRYLO YLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
OAT (UNII: Z6J799EAJK)	
TANACETUM PARTHENIUM (UNII: 6 GE7Z076 1K)	
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALLANTO IN (UNII: 344S277G0Z)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:70484-001-02	1 in 1 BOX	03/25/2015	
1 NDC:70484-001-01	30 mL 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 not final	part348	03/25/2015	

Labeler - Vi Medical Products Inc. (063910521)

Registrant - VEGE-KURL, INC (021072509)

Establishment				
Name	Address	ID/FEI	Business Operations	
VEGE-KURL, INC		021072509	manufacture(70484-001)	

Revised: 4/2016 Vi Medical Products Inc.