POST TREATMENT REPAIR CREAM- hydrocortisone cream
Vege Labs LLC.

ACTIVE INGREDIENTS

Hydrocortisone 1%

PURPOSE

Anti-itch

USES

For the temporary relief 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.

WARNIGS

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.

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, Caprylic/Capric Triglyceride, Cetearyl Alcohol, Ethylhexyl Palmitate, Glycerin, Glyceryl Stearate, PEG-100 Stearate, Pentylene Glycol, Butyrospermum Parkii (Shea) Butter, Dimethicone, Lauryl Glucoside, Myristyl Myristate, 4-t-Butylcyclohexanol, Phenoxyethanol, Coconut Alcohol, Sodium Polyacrylate, Colloidal Silver, Sclerocarya Birrea Seed Oil, Allantoin, Panthenol, Tocopheryl Acetate, Bisabolol, Ethylhexylglycerin, Maltodextrin, Glucose, Zingiber Officinale (Ginger) Root Extract, Aloe Barbadensis Leaf Juice

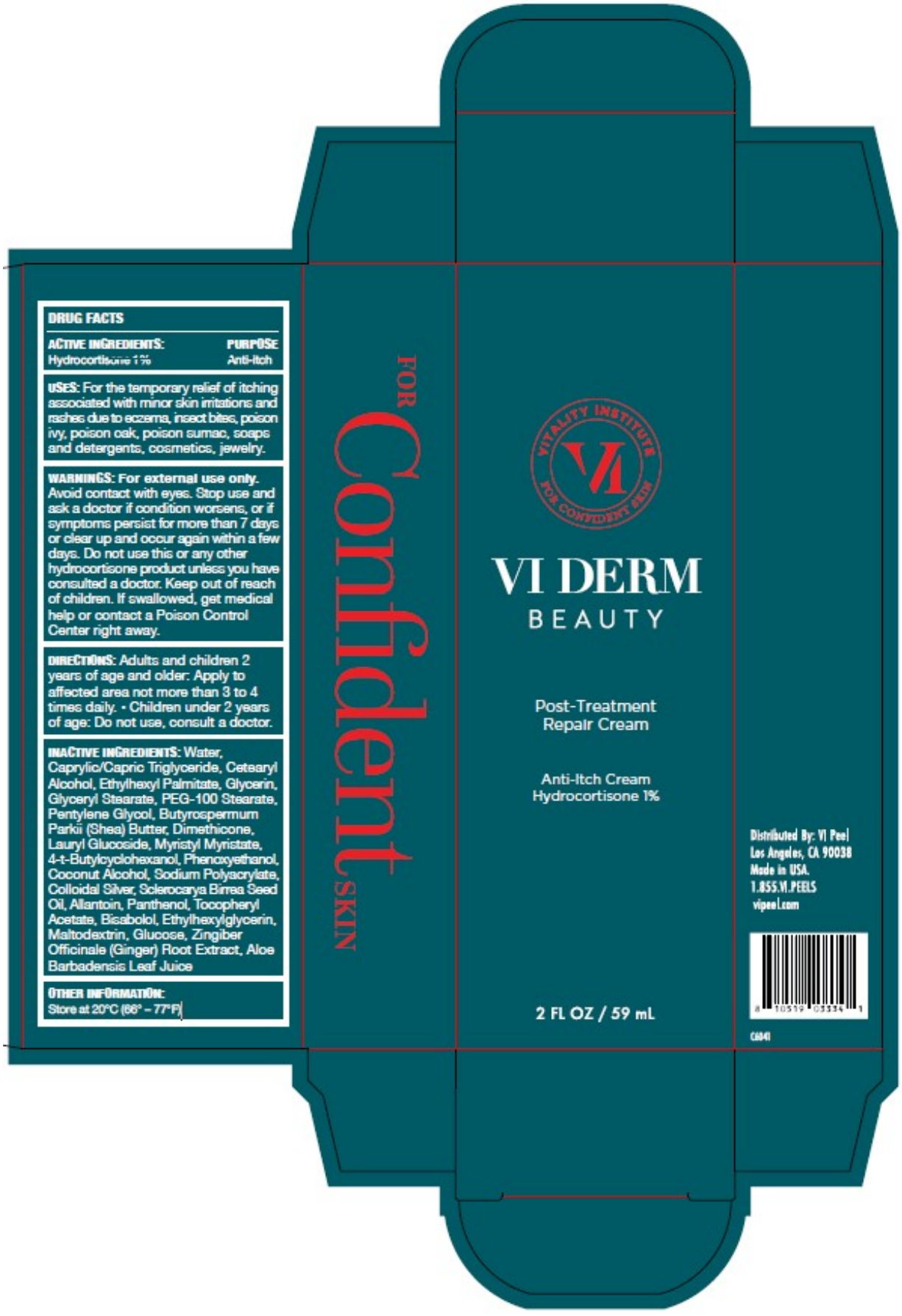
OTHER INFORMATION

Store at 20°C (66° – 77°F)

Keep out of reach of children.

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

Vitality Post-Treatment Repair Cream 2 fl. oz.



POST TREATMENT REPAIR CREAM

hydrocortisone cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67923-105
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE ACETATE	1 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
COCONUT ALCOHOL (UNII: 13F4MW8Y9K)	
GLUCOSE (UNII: 5SL0G7R0OK)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
WATER (UNII: 059QF0KO0R)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
PEG-100 STEARATE (UNII: YD01N1999R)	
BUTYROSPERMUM PARKII (SHEA) BUTTER (UNII: K49155WL9Y)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
MYRISTYL MYRISTATE (UNII: 4042ZC00DY)	
COLLOIDAL SILVER (UNII: 3M4G523W1G)	
SCLEROCARYA BIRREA SEED OIL (UNII: WDO4TLS35F)	
GINGER (UNII: C5529G5JPQ)	
ALLANTOIN (UNII: 344S277G0Z)	
PANTHENOL (UNII: WV9CM0O67Z)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
4-T-BUTYLCYCLOHEXANOL (UNII: K0H1405S9C)	
GLYCERIN (UNII: PDC6A3C0OX)	
BISABOLOL (UNII: 24WE03BX2T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67923-105-04	1 in 1 CARTON	04/19/2022	
1		59 mg in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/19/2022	

Labeler - Vege Labs LLC. (117878620)

Revised: 3/2025 Vege Labs LLC.