SHOPKO HYDROCORTISONE - hydrocortisone cream UNITED EXCHANGE CORP.

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 Fact

Uses for the temporary relief of itching associated with minor skin irritations, inflammation and rashes due to:

- eczema
- seborrheic dematitis
- psoriasis
- insect bites
- poison ivy, oak, sumac
- soaps
- detergents
- cosmetics
- jewelry
- external genital, feminine and anal itching

Other uses or the use of this product for more than 7 days should be undertaken only under the advice and supervision of a doctor

Warnings

- for external use only
- avoid contact with eyes

Stop using this product 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
- before you begin using any other hydrocortisone product

Do not use this product and ask a doctor

- if you have a vaginal discharge
- before treating diaper rash
- before using on children under 2 years of age

Keep this and all drugs out of the reach of children.

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

Before using any medication, read all label directions. Keep this carton. It contains important information.

Directions

- for 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

Other information

- store at room temperature
- Lot No. and Exp. Date: see box or see crimp of tube

Inactive ingredients

glycerin, propylene glycol, sodium chloride, white petrolatum, cetyl alcohol, dimethicone, isopropyl palmitate, distearyldimonium chloride, methylparaben, propylparaben, purified water

Distributed by:

United Exchange Corp.

17211 Valley View Ave.

Cerritos, CA 90703

www.ueccorp.com

Made in Korea



SHOPKO HYDROCORTISONE hydrocortisone cream **Product Information** HUMAN OTC DRUG Product Type Item Code (Source) NDC:55264-001 **Route of Administration TOPICAL**

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
HYDROCORTISONE (UNII: WI4X0 X7BPJ) (HYDROCORTISONE - UNII:WI4X0 X7BPJ)	HYDROCORTISONE	1 g in 100 g	

Inactive Ingredients	
Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PETROLATUM (UNII: 4T6H12BN9U)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIMETHICO NE (UNII: 92RU3N3Y1O)	
ISOPROPYL PALMITATE (UNII: 8 CRQ2TH63M)	
DISTEARYLDIMONIUM CHLORIDE (UNII: OM9573ZX3X)	
METHYLPARABEN (UNII: A218 C7HI9 T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:55264-001-86	1 in 1 CARTON		
1	28 g in 1 TUBE		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part348	02/08/2011		

Labeler - UNITED EXCHANGE CORP. (840130579)

Registrant - UNITED EXCHANGE CORP. (840130579)

Establishment				
Name	Address	ID/FEI	Business Operations	
TAI GUK PHARM. CO., LTD.		631101656	manufacture	

Revised: 2/2011 UNITED EXCHANGE CORP.