MEDPRIDE- hydrocortisone cream Shield Line LLC

MedPride Hydrocortisone

Active Ingredient

Hydrocortisone Acetate USP (1% w/w)

Purpose

Anti-itch

Uses

■ for temporary relief of minor skin irritations, itching and rashes due to eczema, insect bites, poison ivy, poison oak, poison sumac, soaps, detergents, cosmetics, jewelry, and for external genital, feminine and anal itching. Other uses of this product should be only under the advice and supervision of a doctor

Warnings

For external use only

Do not use

 \blacksquare in the eyes \blacksquare for diaper rash \blacksquare if you have vaginal discharge \blacksquare more than the recommended dosage

Ask a doctor before use

■ if you are pregnant or breast feeding

Stop use and ask a doctor if

■ the condition worsens, or if symptoms persist for more than 7 days or clear up and occur again wihin a few days

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Directions

- adults and children over 2 years of age
- apply evenly to affected area no more than 3 to 4 times daily

- children under 2 years of age
- do not use, consult a doctor
- adults and children over 2 years of age
- apply evenly to affected area no more than 3 to 4 times daily
- children under 2 years of age
- do not use, consult a doctor

■ Adults

- when practical, cleanse the affected area with mild soap and warm water and rinse thoroughly
- gently dry, patting or blotting with bathroom tissue or soft cloth before applying
- apply externally to the area up to 6 times a daily or after a bowel movement
- after application discard pad
- do not flush in toilet

Other information

- store at 20-25C (68-77F)
- avoid excessive heat and humidity

Inactive Ingredients

cetostearyl alcohol, chlorocresol, ceteth-20, edetate disodium, liquid paraffin, propylene glycol, purified water, sodium metabisulphite, White white soft paraffin

Package Label



MEDPRIDE

hydrocortisone cream

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Prod	luct	Inform	ation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:52410-3050

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII: W4X0X7BPJ)

Basis of Strength

HYDROCORTISONE ACETATE

HYDROCORTISONE ACETATE

10 mg
in 1 g

Inactive Ingredients				
Ingredient Name	Strength			
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)				
CHLOROCRESOL (UNII: 36W53O7109)				
CETETH-20 (UNII: 1835H2IHHX)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
MINERAL OIL (UNII: T5L8T28FGP)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
SODIUM METABISULFITE (UNII: 4VON5FNS3C)				
PETROLATUM (UNII: 4T6H12BN9U)				

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:52410- 3050-2	0.9 g in 1 PACKET; Type 0: Not a Combination Product	05/23/2013				
2	NDC:52410- 3050-3	28.3 g in 1 TUBE; Type 0: Not a Combination Product	05/23/2013				
3	NDC:52410- 3050-6	454 g in 1 JAR; Type 0: Not a Combination Product	05/23/2013				

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	M017	05/23/2013				

Labeler - Shield Line LLC (078518916)

Revised: 11/2023 Shield Line LLC