

HYDROCORTISONE- hydrocortisone ointment
HF Acquisition Co LLC, DBA HealthFirst

Hydrocortisone acetate (equivalent to Hydrocortisone 1%)

Anti-itch cream

for the temporary relief of itching associated with minor skin irritations and rashes
other uses of this product should be only under the advice and supervision of a doctor

For external use only

Do not use

for the treatment of diaper rash. Consult a doctor

When using this product

avoid contact with eyes

do not begin use of any other hydrocortisone product unless you've consulted a doctor

Stop use and ask a doctor if

condition worsens

symptoms persist more than 7 days

condition clears up and occurs again within a few days

Keep out of the 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; do not use, consult a doctor

clean the affected area

store at room temperature

do not use any opened or torn packets

you may report a serious adverse reaction to this product to 800-275-3433

cetyl alcohol, citric acid, diazolidinyl urea, edetate disodium, glycerin, glyceryl
monostearate, methylparaben, mineral oil, polyethylene glycol, propylene glycol,
propylparaben, stearic acid, trolamine, water

2packs in 1pouch NDC (51662-1284-1)

Anti-itch ointment



(01) 00351662128411
(17) 190630
(21) 271747424508
(10) 8863



See manufacturer's package insert
Distributed by HF Acquisition Co., LLC
Mukilteo, WA 98275

DO NOT USE IF PACKET IS OPENED OR TORN.

NORTH[®]

by Honeywell

CORTISONE 1% CREAM
Anti-Itch Cream

Hydrocortisone Acetate

equivalent to Hydrocortisone 1%) NET WT. 0.9g



HYDROCORTISONE

hydrocortisone ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51662-1284(NDC:59898-800)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name		Strength
HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:W4X0X7BPJ)		1 g in 100 g

Inactive Ingredients	
Ingredient Name	Strength
STEARIC ACID (UNII: 4ELV7Z65AP)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51662-1284-1	2 in 1 POUCH	04/30/2010	
1		0.9 g in 1 PACKET; 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	04/30/2010	

Labeler - HF Acquisition Co LLC, DBA HealthFirst (045657305)

Registrant - HF Acquisition Co LLC, DBA HealthFirst (045657305)

Establishment

Name	Address	ID/FEI	Business Operations
HF Acquisition Co LLC, DBA HealthFirst		045657305	repack(51662-1284)

Revised: 8/2025

HF Acquisition Co LLC, DBA HealthFirst