# HYDROCORTISONE ACETATE- hydrocortisone acetate suppository Proficient Rx LP

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

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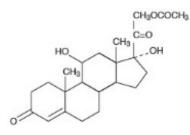
Hydrocortisone Acetate Suppositories 25 mg

For Rectal Administration

**Rx Only** 

#### DESCRIPTION

Hydrocortisone acetate is a corticosteroid designed chemically as pregn-4-ene-3, 20-dione, 21-(acetyloxy)-11, 17-dihydroxy-( $11\beta$ ) with the following structural formula:



Each suppository for rectal administration contains hydrocortisone acetate, USP 25 mg in a specially blended hydrogenated vegetable base.

#### CLINICAL PHARMACOLOGY

In normal subjects, about 26% of hydrocortisone acetate is absorbed when the suppository is applied to the rectum. Absorption of hydrocortisone acetate may vary across abraded or inflamed surfaces.

Topical steroids are primarily effective because of their anti-inflammatory, anti-pruritic and vasoconstrictive action.

#### INDICATIONS AND USAGE

Hydrocortisone acetate suppositories are indicated for use in inflamed hemorrhoids, post-irradiation (factitial) proctitis, as an adjunct in the treatment of chronic ulcerative colitis, cryptitis, other inflammatory conditions of anorectum, and pruritus ani.

#### CONTRAINDICATIONS

Hydrocortisone acetate suppositories are contraindicated in those patients having a history of hypersensitivity to hydrocortisone acetate or any of the components.

#### PRECAUTIONS

Do not use hydrocortisone acetate suppositories unless adequate proctologic examination is made.

If irritation develops, the product should be discontinued and appropriate therapy instituted.

In the presence of an infection, the use of an appropriate antifungal or antibacterial agent should be

instituted. If a favorable response does not occur promptly, hydrocortisone acetate should be discontinued until the infection has been adequately controlled.

#### Carcinogenesis

No long-term studies in animals have been performed to evaluate the carcinogenic potential of corticosteroid suppositories.

#### **Pregnancy Category C**

In laboratory animals, topical steroids have been associated with an increase in the incidence of fetal abnormalities when gestating females have been exposed to rather low dosage levels. There are no adequate and well-controlled studies in pregnant women.

Hydrocortisone acetate suppositories should only be used during pregnancy if the potential benefit justifies the risk of the fetus. Drugs of this class should not be used extensively on pregnant patients, in large amounts, or for prolonged periods of time.

It is not known whether this drug is excreted in human milk, and because many drugs are excreted in human milk and because of the potential for serious adverse reactions in nursing infants from hydrocortisone acetate suppositories, a decision should be made whether to discontinue nursing or to discontinue the drug, taking into account the importance of the drug to the mother.

#### **ADVERSE REACTIONS**

The following local adverse reactions have been reported with hydrocortisone acetate suppositories; burning, itching, irritation, dryness, folliculitis, hypopigmentation, allergic contact dermatitis, secondary infection.

To report an adverse event, please contact Westminster Pharmaceuticals, LLC at 1-844-221-7294.

#### DRUG ABUSE AND DEPENDENCE

Drug abuse and dependence have not been reported in patients treated with hydrocortisone acetate suppositories.

#### OVERDOSAGE

If signs and symptoms of systemic overdosage occur, discontinue use.

#### DOSAGE AND ADMINISTRATION

#### FOR RECTAL ADMINISTRATION

Detach one suppository from strip of suppositories. Hold suppository upright and carefully separate tabs at top opening and pull downward from the pointed end to expose the suppository. Remove the suppository from the pocket. Avoid excessive handling of suppository which is designed to melt at body temperature. Insert one suppository rectally, pointed end first. Insert one suppository in the rectum twice daily, morning and night for two weeks, in nonspecific proctitis. In more severe cases, one suppository three times a day or two suppositories twice daily. In factitial proctitis, the recommended duration of therapy is six to eight weeks or less, according to the response of the individual case.

#### HOW SUPPLIED

Hydrocortisone acetate suppositories 25mg are off-white, smooth surfaced and bullet shaped with one pointed end.

Box of 12 suppositories, NDC 71205-976-12 Box of 24 suppositories, NDC 71205-976-24

#### STORAGE

Store at 20°-25°C (68°-77°F) [See USP Controlled Room Temperature]. Excursions permitted to 15°-30°C (59°-86°F). Store away from heat. Protect from freezing. Avoid contact with eyes.

**KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN.** In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

**PHARMACIST:** This product is not an Orange Book rated product, therefore all prescriptions using this product shall be subject to state and federal statutes as applicable. This product has not been subjected to FDA therapeutic or other equivalency testing. There are no claims of bioequivalence or therapeutic equivalence. Each person recommending a prescription substitution using this product shall make such recommendation based on his/her professional knowledge and opinion, upon evaluating the active ingredients, inactive ingredients, excipients and chemical information contained within the enclosed prescribing information.

Rx Only

#### Manufactured for:

Westminster Pharmaceuticals, LLC Nashville, TN 37217

#### **Relabeled by:**

Proficient Rx LP

Thousand Oaks, CA 91320

Rev. 12/19

#### PRINCIPAL DISPLAY PANEL - 12 Suppository Carton

NDC 71205-976-12 Rx Only

Hydrocortisone Acetate Suppositories 25 mg

FOR RECTAL USE ONLY

12 Suppositories Unit Dose





NDC 71205-976-12

Relabeled By: Proficient Rx LP Thousand Oaks, CA 91320

**RX Only** 

## Hydrocortisone Acetate 25mg

#### #12 Suppositories

Each suppository contains: hydrocortisone acetate 25 mg in a specially blended hydrogenated vegetable oil base.

See package insert

Product ID: SH097612

#### FOR RECTAL USE ONLY

Mfr. For: Westminster Pharmaceuticals, LLC. Nashville, TN 37217 Store at 20°-25°C (68°-77°F) Keep me

HYDROCORTISONE ACETATE

Keep medication out of the reach of children

Hydrocortisone Acetate 25mg #12 Suppositories Lot #:00000 SN# MASTER NDC 71205-976-12 Exp:00/00/00

Hydrocortisone Acetate 25mg #12 Suppositories Lot #.00000 SN# MASTER NDC 71205-976-12 Exp.00/00/00

Hydrocortisone Acetate 25mg #12 Suppositories Lot #.00000 SN#MASTER NDC 71205-976-12 Exp:00/00/00



GTIN: 00371205976125 SN# MASTER Exp. 00/00/00 Lot #:00000

hydrocortisone acetate suppository				
Product Information				
Product T ype	HUMAN PRESCRIPTION DRUG	Item Code (S		

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:71205-976(NDC:69367-243)
Route of Administration	RECTAL		

Active Ingredient/Active Moiety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
HYDROCORTISONE ACETATE (UNII: 3X7931PO74) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE ACETATE	25 mg		
Inactive Ingredients				
Ingredient Name	St	Strength		
HYDRO GENATED PALM KERNEL OIL (UNII: FM8 D1RE2VP)				

Product Characteristics					
Color	WHITE	Score			
Shape	BULLET	Size			
Flavor		Imprint	Code		
Contains					
Packaging					
# Item Code	Package Description		Marketing Start Date	Marketing l	End Date

Unapproved drug other		02/05/2020		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
Marketing Information				
<b>2</b> NDC:71205-976-24	24 in 1 CARTON; Type 0: Not a Combination Product	02/07/2020		
1 NDC:71205-976-12	12 in 1 CARTON; Type 0: Not a Combination Product	05/07/2020		

### Labeler - Proficient Rx LP (079196022)

Establishment				
Name	Address	ID/FEI	Business Operations	
Proficient Rx LP		079196022	REPACK(71205-976), RELABEL(71205-976)	

Revised: 5/2020

Proficient Rx LP