

DESOXIMETASONE - desoximetasone cream
Physicians Total Care, Inc.

Desoximetasone
Cream USP, 0.05%, Cream USP, 0.25%
Gel USP, 0.05%

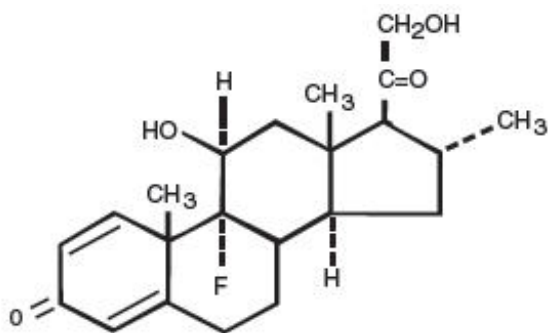
For external use only. Not for ophthalmic use.

Rx only

DESCRIPTION

Desoximetasone Cream USP, 0.05%; Desoximetasone Cream USP, 0.25% and Desoximetasone Gel USP, 0.05% contain desoximetasone a synthetic fluorinated corticosteroid for topical dermatologic use. The corticosteroids constitute a class of primarily synthetic steroids used topically as anti-inflammatory and anti-pruritic agents.

Chemically, desoximetasone is Pregna-1, 4-diene-3, 20-dione, 9-fluoro-11,21-dihydroxy-16-methyl-, (11 β , 16 α)-. It has the molecular formula C₂₂H₂₉FO₄ and a molecular weight of 376.47. The CAS Registry Number is 382-67-2. The chemical structure is:



Desoximetasone is a white to off-white, odorless, crystalline powder and is soluble in alcohol, acetone, chloroform, and hot ethyl acetate; slightly soluble in ether and benzene; and insoluble in water, dilute aqueous acids and alkalis.

Each gram of Desoximetasone Cream USP, 0.05% contains 0.5 mg of desoximetasone in an emollient cream consisting of white petrolatum, purified water, isopropyl myristate, lanolin alcohols, mineral oil, cetostearyl alcohol and edetate disodium.

Each gram of Desoximetasone Cream USP, 0.25% contains 2.5 mg of desoximetasone in an emollient cream consisting of white petrolatum, purified water, isopropyl myristate, lanolin alcohols, mineral oil and cetostearyl alcohol.

Each gram of Desoximetasone Gel USP, 0.05% contains 0.5 mg of desoximetasone in a gel base consisting of purified water, docusate sodium, edetate disodium, isopropyl myristate, carbomer 940, triethylamine, and SDAG-3 95% alcohol.

CLINICAL PHARMACOLOGY

The corticosteroids are a class of compounds comprising steroid hormones secreted by the adrenal cortex and their synthetic analogs. In pharmacologic doses, corticosteroids are used primarily for their

anti-inflammatory and/or immunosuppressive effects.

Topical corticosteroids, such as desoximetasone, are effective in the treatment of corticosteroid-responsive dermatoses primarily because of their anti-inflammatory, antipruritic, and vasoconstrictive actions. However, while the physiologic, pharmacologic, and clinical effects of the corticosteroids are well known, the exact mechanisms of their actions in each disease are uncertain. Desoximetasone, a corticosteroid, has been shown to have topical (dermatologic) and systemic pharmacologic and metabolic effects characteristic of this class of drugs.

Pharmacokinetics

The extent of percutaneous absorption of topical corticosteroids is determined by many factors including the vehicle and the integrity of the epidermal barrier, and the use of occlusive dressings.

Topical corticosteroids can be absorbed through normal intact skin. Inflammation and/or other disease processes in the skin may increase percutaneous absorption. Occlusive dressings substantially increase the percutaneous absorption of topical corticosteroids.

Once absorbed through the skin, topical corticosteroids enter pharmacokinetic pathways similar to systemically administered corticosteroids. Corticosteroids are bound to plasma proteins in varying degrees, are metabolized primarily in the liver and excreted by the kidneys. Some of the topical corticosteroids and their metabolites are also excreted into the bile.

Pharmacokinetic studies in men with Desoximetasone Cream USP, 0.25% with tagged desoximetasone showed a total of $5.2\% \pm 2.9\%$ excretion in urine ($4.1\% \pm 2.3\%$) and feces ($1.1\% \pm 0.6\%$) and no detectable level (limit of sensitivity: $0.005 \mu\text{g/mL}$) in the blood when it was applied topically on the back followed by occlusion for 24 hours. Seven days after application, no further radioactivity was detected in urine or feces. The half-life of the material was 15 ± 2 hours (for urine) and 17 ± 2 hours (for feces) between the third and fifth trial day. Studies with other similarly structured steroids have shown that predominant metabolite reaction occurs through conjugation to form the glucuronide and sulfate ester. Studies performed with Desoximetasone Cream USP, 0.05%; Desoximetasone Cream USP, 0.25% and Desoximetasone Gel USP, 0.05% indicate that they are in the high range of potency as compared with other topical corticosteroids.

INDICATIONS AND USAGE

Topical corticosteroids are high potency corticosteroids indicated for the relief of the inflammatory and pruritic manifestations of corticosteroid responsive dermatoses.

CONTRAINDICATIONS

Topical corticosteroids are contraindicated in those patients with a history of hypersensitivity to any of the components of the preparation.

WARNINGS

Desoximetasone Cream USP, 0.05%; Desoximetasone Cream USP, 0.25% and Desoximetasone Gel USP, 0.05% are not for ophthalmic use.

Keep out of reach of children.

PRECAUTIONS

General

Systemic absorption of topical corticosteroids can produce reversible hypothalamic-pituitary-adrenal

(HPA) axis suppression with the potential for glucocorticosteroid insufficiency after withdrawal of treatment. Manifestations of Cushing's syndrome, hyperglycemia, and glucosuria can also be produced in some patients by systemic absorption of topical corticosteroids while on treatment. Use of more than one corticosteroid-containing product at the same time may increase total systemic glucocorticoid exposure. (See **DOSAGE AND ADMINISTRATION** section.)

Patients applying a topical steroid to a large surface area or to areas under occlusion should be evaluated periodically for evidence of HPA axis suppression. This may be done by using ACTH stimulation and urinary free cortisol tests.

If HPA axis suppression is noted, an attempt should be made to withdraw the drug, to reduce the frequency of application, or to substitute a less potent corticosteroid. Recovery of HPA axis function is generally prompt upon discontinuation of topical corticosteroids. Infrequently, signs and symptoms of glucocorticosteroid insufficiency may occur requiring supplemental systemic corticosteroids. For information on systemic supplementation, see prescribing information for those products.

Pediatric patients may be more susceptible to systemic toxicity from equivalent doses due to their larger skin surface to body mass ratios. (See **PRECAUTIONS – Pediatric Use.**)

If irritation develops, Desoximetasone Cream USP, 0.05%; Desoximetasone Cream USP, 0.25% or Desoximetasone Gel USP, 0.05% should be discontinued and appropriate therapy instituted. Allergic contact dermatitis with corticosteroids is usually diagnosed by observing *failure to heal* rather than noting a clinical exacerbation as with most topical products not containing corticosteroids. Such an observation should be corroborated with appropriate diagnostic patch testing.

If concomitant skin infections are present or develop, an appropriate antifungal or antibacterial agent should be used. If a favorable response does not occur promptly, use of Desoximetasone Cream USP, 0.05%; Desoximetasone Cream USP, 0.25% or Desoximetasone Gel USP, 0.05% should be discontinued until the infection has been adequately controlled.

This medication should not be used on the face, underarm or groin area.

As with other corticosteroids, therapy should be discontinued when control is achieved. If no improvement is seen within two weeks, the diagnosis and therapy should be re-evaluated.

Information for the Patient

Patients using topical corticosteroids should receive the following information and instructions. This information is intended to aid in the safe and effective use of this medication. It is not a disclosure of all possible adverse or unintended effects:

1. This medication is to be used as directed by the physician. It is for external use only. Avoid contact with the eyes.
2. This medication should not be used for any disorder other than that for which it was prescribed.
3. The treated skin area should not be bandaged, otherwise covered or wrapped, so as to be occlusive unless directed by the physician.
4. Patients should report to their physician any signs of local adverse reactions.
5. Other topical corticosteroid-containing products should not be used with Desoximetasone Cream USP, 0.05% or Desoximetasone Cream USP, 0.25% or Desoximetasone Gel USP, 0.05% without first talking to your physician.
6. If no improvement is seen in 2 weeks, contact your physician.

Laboratory Tests

The following tests may be helpful in evaluating patients for HPA axis suppression:

ACTH stimulation test

Urinary free cortisol test

Carcinogenesis, Mutagenesis, and Impairment of Fertility

Long-term animal studies have not been performed to evaluate the carcinogenic potential or the effect on fertility of desoximetasone.

Desoximetasone was nonmutagenic in the Ames test.

Pregnancy

Teratogenic Effects

Pregnancy Category C

Corticosteroids have been shown to be teratogenic in laboratory animals when administered systemically at relatively low dosage levels. Some corticosteroids have been shown to be teratogenic after dermal application in laboratory animals.

Desoximetasone has been shown to be teratogenic and embryotoxic in mice, rats, and rabbits when given by subcutaneous or dermal routes of administration in doses 3 to 30 times the human dose of Desoximetasone Cream USP, 0.25% or in doses 15 to 150 times the human dose of Desoximetasone Cream USP, 0.05% or Desoximetasone Gel USP, 0.05%.

There are no adequate and well-controlled studies in pregnant women. Desoximetasone Cream USP, 0.05%; Desoximetasone Cream USP, 0.25% and Desoximetasone Gel USP, 0.05% should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

Nursing Mothers

It is not known whether topical administration of corticosteroids can result in sufficient systemic absorption to produce detectable quantities in breast milk. Systemically administered corticosteroids are secreted into breast milk in quantities not likely to have a deleterious effect on the infant. Nevertheless, 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.

Pediatric Use

Pediatric patients may demonstrate greater susceptibility to topical corticosteroid-induced HPA axis suppression and Cushing's syndrome than adult patients because of a larger skin surface area to body weight ratio.

Hypothalamic-pituitary-adrenal (HPA) axis suppression, Cushing's syndrome and intracranial hypertension have been reported in pediatric patients receiving topical corticosteroids. Manifestations of adrenal suppression in pediatric patients include linear growth retardation, delayed weight gain, low plasma cortisol levels, and absence of response to ACTH stimulation. Manifestations of intracranial hypertension include bulging fontanelles, headaches, and bilateral papilledema. Chronic corticosteroid therapy may interfere with linear bone growth in pediatric patients.

Geriatric Use

Clinical studies of Desoximetasone Cream USP, 0.05%; Desoximetasone Cream USP, 0.25% and Desoximetasone Gel USP, 0.05% did not include sufficient numbers of subjects aged 65 and over to determine whether they respond differently from younger subjects. Other reported clinical experience has not identified differences in responses between the elderly and younger patients. In generally, dose selection for an elderly patient should be cautious.

ADVERSE REACTIONS

In controlled clinical studies the incidence of adverse reactions was low (0.8%) for Desoximetasone Cream USP, 0.05% and included pruritus, erythema, vesiculation and burning sensation. The incidence

of adverse reactions was also 0.8% for Desoximetasone Cream USP, 0.25% and included burning, folliculitis, and folliculopustular lesions. The incidence of adverse reactions for Desoximetasone Gel USP, 0.05% was 0.3% with one subject reporting stinging and burning at the site of application.

The following local adverse reactions are reported infrequently when topical corticosteroids are used as recommended but may occur more frequently with the use of occlusive dressings and higher potency corticosteroids. These are listed in an approximate decreasing order of occurrence: burning, itching, irritation, dryness, folliculitis, hypertrichosis, acneiform eruptions, hypopigmentation, perioral dermatitis, allergic contact dermatitis, maceration of the skin, secondary infection, skin atrophy, striae, and miliaria.

Systemic absorption of topical corticosteroids has produced hypothalamic-pituitary-adrenal (HPA) axis suppression, manifestations of Cushing's syndrome, hyperglycemia, and glucosuria in some patients.

OVERDOSAGE

Topically applied Desoximetasone Cream USP, 0.05%; Desoximetasone Cream USP, 0.25% and Desoximetasone Gel USP, 0.05% can be absorbed in sufficient amounts to produce systemic effects (see **PRECAUTIONS**).

DOSAGE AND ADMINISTRATION

Apply a thin film of Desoximetasone Cream USP, 0.05% or Desoximetasone Cream USP, 0.25% or Desoximetasone Gel USP, 0.05% to the affected skin areas twice daily. Rub in gently.

HOW SUPPLIED

Desoximetasone Cream USP, 0.25% is supplied in

15 gram tubes	NDC 54868-3041-0
60 gram tubes	NDC 54868-3041-1

Store at 20° - 25°C (68° - 77°F) [see USP Controlled Room Temperature].

Mfd. by: Taro Pharmaceuticals Inc., Brampton, Ontario, Canada L6T 1C1
Revised June, 2009

PK-6352-0

Relabeling of "Additional" barcode label by:

Physicians Total Care, Inc.
Tulsa, OK 74146

PRINCIPAL DISPLAY PANEL - 0.25%, 60 g Tube Carton

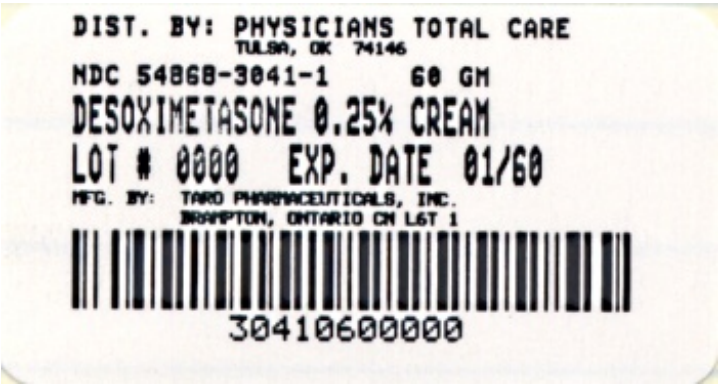
60 g

**Desoximetasone
Cream USP, 0.25%**

FOR EXTERNAL USE ONLY.
NOT FOR OPHTHALMIC USE.

Rx only

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



DESOXIMETASONE				
desoximetasone cream				
Product Information				
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:54868-3041(NDC:51672-1270)	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
DESOXIMETASONE (UNII: 4E07GXB7AU) (DESOXIMETASONE - UNII:4E07GXB7AU)			DESOXIMETASONE	2.5 mg in 1 g
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0KO0R)				
PETROLATUM (UNII: 4T6H12BN9U)				
MINERAL OIL (UNII: T5L8T28FGP)				
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)				
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54868-3041-0	1 in 1 CARTON		
1		15 g in 1 TUBE		
2	NDC:54868-3041-1	1 in 1 CARTON		
2		60 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA073193	06/23/1994	

Labeler - Physicians Total Care, Inc. (194123980)

Establishment

Name	Address	ID/FEI	Business Operations
Physicians Total Care, Inc.		194123980	relabel

Revised: 10/2009

Physicians Total Care, Inc.