ULTRADERM®

2003

Cream - Ointment

COMPOSITION: Each gram contains:

Triamcinolone acetonide B.P	1.00 mg.
Neomycin (as neomycin sulphate B.P.)	2.50 mg.
Gramicidin U.S.P	0.25 mg.
Nystatin B.P.	100.000 I.U.

PROPERTIES: ULTRADERM possesses strong topical anti-inflammatory and antipruritic activities by virtue of its component triamcinolone acetonide, the potent fluorinated corticosteroid. It has also a broad spectrum antibacterial activities due to the presence of two antibiotics neomycin and gramicidin which make Ultraderm useful in the management of secondarily infected dermatoses.
Ultraderm contains also nystatin, the fungistatic and fungicidal antibiotic which is very effective in the treatment of candidiasis or as a prophylactic agent against the overgrowth of candida.

INDICATIONS: ULTRADERM is successfully used for the relief of inflammatory and pruritic manifestations of corticosteroids-responsive dermatoses which are accompanied by bacterial infection and / or cutaneous candidiasis such as: Eczema, infantile eczema, atopic dermatitis, dermatitis herpetiformis; contact dermatitis; stasis dermatitis; seborrheic dermatitis and dermatitis venenata; neurodermatoses including lichen simplex; lichen planus; chronic discoid lupus erythematosus, anal and valval pruritus.

PRECAUTIONS: The use of Ultraderm should be avoided in patients with extensive skin damage, because the absorption of neomycin can possibly lead to nephrotoxicity or ototoxicity. Furthermore, the prolonged application, or the use of exaggerated quantities of the product, even on the intact skin, may also lead to systemic absorption of triamcinolone acetonide and thus to the usual side effects of corticosteroids.
In case superinfection with other organisms or fungi occurs, a suitable concomitant antimicrobial therapy should be applied and if the response is not promptly remarkable, the product should be discontinued until the infection is completely cleared-up.

SIDE EFFECTS: As with other topical corticosteroids, the use of Ultraderm may cause the following side effects: Skin atrophy; hypertrichosis; skin hypopigmentation; folliculitis; maceration of the skin; secondary infections and hypersensitivity to the product's components. Ototoxicity and/or nephrotoxicity may occur due to the presence of neomycine.

CONTRAINDICATIONS: As with all other topical fluorinated corticosteroids, Ultraderm should not be used in case of: Rosacea, acne vulgaris, skin ulcer, viral infections such as herpes simplex or zoster, varicella, vaccinia and warts.
Ultraderm can not be given also to patients who have a history of hypersensitivity to any of its components nor should it be applied on certain sites such as mucous membranes, flexures, eyes and eyelids.

DOSAGE AND APPLICATION: Ultraderm is available in a water miscible base (cream form) for soft and oozing lesions and an oily base (ointment form) for dry and scaly lesions.
A small quantity should be applied 2-3 times daily.
In some resistant skin diseases, Ultraderm can be applied under an occlusive dressing in order to have a better penetration of the active ingredients.

PRESENTATION: Ultraderm is available as:
Ultraderm cream: tubes of 15 gm.
Ultraderm ointment: tubes of 15 gm.

This is a medicament - keep medicaments out of reach of children

- A medicament is a product which affects your health, and its consumption contrary to the instructions is dangerous for you.
- Follow strictly the doctor's prescription, the method of use, and the instructions of the pharmacist who dispensed the medicament.
- The doctor and the pharmacist are experts in medicine, its benefits and risks.
- Do not by yourself interrupt the period of treatment prescribed for you.
- Do not repeat the same prescription without consulting your doctor.

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