LEVELINA SOLUTION

COMPOSITION
Per 100ml
Bifonazol (D.C.I) 1g
Excipients: Ethanol, isopropyl myristate

PHARMACEUTICAL FORM AND CONTENTS OF PACKET
Levelina solution, 30ml bottle

ACTION
LEVELINA (Bifonazol) is a broad spectrum antimycotic which deals with dermatophytes, yeasts, mold and other fungal infections such Malessezzia furfus; it is also effective against Corynebacterium minutissimum. Action against dermatophytes (e.g. Trichophyton spp.) is fundamentally fungicidal in vitro, and in yeasts it is strongly fungistatic.

Name of person or entity authorized for commercialization and production of this product:
Laboratorios ERN, S.A. Pedro IV, 499-08020 Barcelona, España

INDICATIONS
Skin mycosis caused by dermatophytes, yeasts, molds, and other fungi, such as Malessezia furfur and Corynebacterium minutissimum. These could be, for example; Mycosis of the feet and hands (e.g. athlete´s foot, ringworm of the hands) Mycosis of the body and cataneous complaints (e.g. ringworm), Pitriasis versicolar and Erythrasma.

COUNTERINDICATIONS
Hypersensibility to Bifonazol.

INCOMPATABILITIES
None have been described.

WARNING
Frequent applications may cause dry skin and irritation.

POSOLOGY
Unless advised otherwise by a doctor: Levelina (Bifonazol) is applied once a day, preferably before going to bed, spreading a small amount of solution on the affected area. Generally it is sufficient to apply a few drops for an area the size of your hand. To hygienically support the treatment it is advisable to carefully wash and dry the affected area. To obtain lasting results treatment with LEVELINA (Bifonazol) should not be interrupted after the symptoms have disappeared, but should be continued for a determined period of time:
Interdigital mycosis of the foot and athlete's foot: 3 weeks
Mycosis of the body, hands and cutaneous complaints (ringworm etc.): 2-3 weeks
Pitiriasis versicolor and Erythrasma: 2 weeks
Superficial cutaneous candiadiasis: 2-4 weeks

OVERDOSE AND ITS TREATMENT
Given the concentration of the active principle and the form of administration, intoxication is not possible even in the case of accidental ingestion.
In the event of overdose or accidental ingestion consult El Servicio de Información Toxocológica. Tel 91 562 04 20.

ADVERSE REACTIONS
In rare cases light cutaneous reactions may appear. Any adverse reaction which is not described in the prospectus should be reported to your doctor.

EXPIRY
This medicine should not be used after the expiry date indicated on the packaging.

PRESCRIPTION ONLY
KEEP OUT OF THE REACH AND SIGHT OF CHILDREN.

TITULAR AUTHORIZED FOR COMMERCIALIZATION
Laboratorios ERN, S.A. Pedro IV, 499-08020 Barcelona, España

RESPONSIBLE FOR MANUFACTURING
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