been reported in patients with large burned body surface in which silver sulfadiazine can be absorbed. In hypersensitive patients reactions similar to those produced by sulfoderivatives may occur.

DOSAGE AND METHOD OF ADMINISTRATION

After cleaning the lesions, a thin layer is applied directly to the lesions. The dose is determined by the extent of the lesions. Cure of the lesions and application of **Hebermin®** on alternate days is recommended. In infected lesions, local cleaning and complementary antibiotic treatment is required.

INTERACTIONS WITH OTHER MEDICATIONS AND OTHER FORMS OF INTERACTION

No symptoms have been described for incompatibilities or interactions with other medications.

USE IN PREGNANCY AND LACTATION

Hebermin®should not be used in women with full-term pregnancies, or newborns during the first month of life, since silver sulfadiazine can produce kemicterus.

Center for Genetic Engineering and Biotechnology

Distinctive company of Cuban biotechnology which develops, produces, markets and exports innovative products, for key areas of the biomedical, veterinary, agricultural, aquaculture and industrial sectors, for one health. It has a portfolio of research and development (R&D) projects and products, protected by patents. Its more than 30 products marketed in more than 35 countries, include first and only drugs of its kind, to treat diseases that do not have other effective therapeutic solutions. Several of its medicines are inserted into national programs to offer comprehensive health care. We work with social responsibility and in harmony with the environment.

CIGB CENTRO DE INGENIERÍA GENÉTICA Y BIOTECNOLOGÍA

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Hebermin®

HEALING CREAM FOR THE TREATMENT OF BURNS AND OTHER SKIN LESIONS

RESTORE YOUR DAMAGED SKIN

Hebermin®

Healing cream for the treatment of burns and other skin lesions

RESTORE YOUR DAMAGED SKIN

- REDUCES HEALING AND EPITHELIALIZATION TIME IN BURNS, BEDSORES AND SURGICAL PROCEDURES.
- REGULATES CELL GROWTH.
- PROTECTS THE DAMAGE SKIN OF INFECTIONS POSSIBLE, BECAUSE IT IS AN ANTIMICROBIAL OF BACTERIOSTATIC ACTION.

PHARMACEUTICAL FORM

Cream

STRENGTH

10 µg of FCEhr/g cream

PRESENTATION

White-opaque polyethylene bottles with capacity for 30 and 200 g, respectively.

COMPOSITION

Each 100 g contains recombinant human epidermal growth factor (FCEhr) 0.001 g; silver sulfadiazine 1.00 g; 4-methyl hydroxybenzoate 0.180 g; 4-propyl hydroxybenzoate 0.020 g.

SHELF LIFE AND STORAGE CONDITIONS

Hebermin® is sable during 24 months. Store below 25 °C. Protect from light.

INDICATIONS

Hebermin® is used in the treatment of superficial and deep dermal and hypodermal burns and bedsores. It can be used in other surgical processes that require healing or tissue regeneration, such as lesions caused by radiation, cytostatic extravasation ulcers and ulcers due to circulatory insufficiency, as well as in the prophylaxis of superficial radiotherapy lesions.

CONTRAINDICATIONS

Hebermin® is contraindicated in patients with hypersensitivity or allergy to sulfonamides. It should not be used in newborns, or during pregnancy and lactation.

PRECAUTIONS

Hebermin® should be used with care in patients with hepatic and renal impairment and in patients with glucose 6 phosphate dehydrogenase deficiency.

SPECIAL WARNINGS AND PRECAUTIONS FOR USE

The surface where the cream is applied should not be exposed directly to the sun, since it alters the medication. In such cases, the treated surface should be covered or bandaged.

UNDESIRABLE EFFECTS

Reactions characterized by intense burning that tend to disappear, rash, pruritus and eventually crystalluria have