APPROVED
The Order of the Ministry
of Health of Ukraine
14.11.2019 No.2283
Registration Certificate
No.UA/3682/01/01
INSTRUCTION
for medical use of medicinal product
LEKVOLEK®

УТВЕРЖДЕНО
Приказ Министерства
здравоохранения Украины
14.11.2019 №2283
Регистрационное удостоверение
№UA/3682/01/01
ИНСТРУКЦИЯ
по медицинскому применению
лекарственного средства
ЭВКОЛЕК

Composition:
active ingredient: chlorophyllipt extract dense;
1 suppository contains: chlorophyllipt extract dense (Eucalypti) (1:15.3-10.76:1) (extractant ethanol 93-96%) (in recalculation on dry substance) – 0.05 g (50 mg)
excipients: polysorbate, hard fat.

Pharmaceutical form. Suppositories.
Basic physical and chemical properties: dark green, bullet-shaped suppositories. Presence of white bloom on the surface of a suppository is allowed.

Pharmacotherapeutic group. Gynecological antiinfectives and antiseptics.
ATC Code G01A X.

Pharmacological properties.
The active substance of the drug is chlorophyllipt extract dense. Chlorophyllipt extract has a stimulating effect on the receptors of the mucous membranes and shows local anti-inflammatory and antiseptic activity.
The drug has antibacterial (bacteriostatic and bactericidal) activity against staphylococci.

Pharmacokinetic properties.
Not described.

Clinical particulars.
Indications.
Treatment of infectious and inflammatory processes of female genital organs (colpitis, vaginitis, vulvovaginitis) and intestines caused by antibiotic-resistant strains of staphylococci.

Contraindications.
Hypersensitivity to chlorophyllipt or other components of the drug.

Interaction with other medicinal products and other forms of interaction.
Data on drug interactions are not available.

Special warnings and precautions for use.
Consult a doctor before treatment.
After insertion of the first suppository wait 6-8 hours before beginning treatment.
A course of treatment is possible after confirming the absence of allergic reactions to chlorophyllipt extract spissum (probable swelling of the lips, nasal mucous membranes or throat and other allergic reactions). If any of the following symptoms occur, discontinue the drug and consult a doctor.

Use during pregnancy or lactation.
The efficacy and safety of the drug during pregnancy and lactation have not been studied, therefore the drug should not be administered to this group of patients.

Effects on ability to drive and use machines.
No data.

Posology and method of administration.
Before suppository administration it is necessary to:
– tear one suppository in the primary packaging along the perforation line of the blister pack;
– pull the edges of the film tearing it apart and releasing the suppository from the primary packaging.
The drug is intended for vaginal or rectal use.
Dose for adults: 1 suppository twice daily. The course of treatment is 10-14 days.

Paediatric patients.
There are no data on the safety and efficacy of the drug in children.

Overdose.
Major symptoms: swelling of the lips, nasal mucous membranes and throat.
Treatment: drug discontinuation. Treatment is symptomatic.

Undesirable effects.
Immune system disorders: allergic reactions, including itching, rash, flushing, angioedema (probable swelling of the lips, nasal mucous membranes, throat, etc.).

Shelf life. 2 years.

Storage. Keep away from children. Store in original package at temperature not exceeding 25 °C.

Package. 5 suppositories in a blister, 1or 2 blister in a pack.

Prescription status. Without prescription.

Manufacturer. Joint Stock Company «Lekhim-Kharkiv».

Location. Ukraine, 61115, Kharkiv region, Kharkiv, Severyna Pototskoho street, building 36.

Date of last revision. 14.11.2019