Gyno-Daktarin

NAME OF THE MEDICINAL PRODUCT

GYNO-DAKTARIN CREAM

Miconazole nitrate 20 mg/g cream
GYNO-DAKTARIN VAGINAL CAPSULES

Miconazole nitrate 200 mg, 400 mg, and 1200 mg vaginal capsules GYNO-DAKTARIN VAGINAL OVULES

Miconazole nitrate 100 mg vaginal ovules

QUALITATIVE AND QUANTITATIVE COMPOSITION

GYNO-DAKTARIN Cream: Each gram contains 20 mg of the active

substance miconazole nitrate. GYNO-DAKTARIN Vaginal Capsules: Each vaginal capsule contains

200 mg, 400 mg or 1200 mg of the active substance miconazole nitrate.

GYNO-DAKTARIN Vaginal Ovules: Each vaginal ovule contains 100 mg of the active substance miconazole nitrate.

For excipients, see List of Excipients.

PHARMACEUTICAL FORM

GYNO-DAKTARIN Cream: White, homogenous cream for vulvar and

GYNO-DAKTARIN Vaginal Capsules: White to off-white, egg-shaped capsules for vaginal use

GYNO-DAKTARIN Vaginal Ovules: White to beige colored, egg-shaped ovules for vaginal use.

CLINICAL PARTICULARS Therapeutic Indications

Local treatment of vulvovaginal candidosis and superinfections due to

GYNO-DAKTARIN Cream may also be used for the treatment of mycotic

Posology And Method Of Administration GYNO-DAKTARIN Cream

Once daily before bedtime, administer the contents of 1 applicator (about 5 g of cream) deeply into the vagina (see Instructions for Use and Handling). Repeat this procedure for 7 days, even if symptoms (e.g. pruritus and leukorrhea) have disappeared or menstruation begins. Treatment of concurrent symptoms of mycotic balanitis of the male partner: apply the cream twice daily on the glans penis. The treatment duration is the same as for the female partner.

GYNO-DAKTARIN Vaginal Capsules

200 mg capsules:

Once daily before bedtime, insert one vaginal capsule deeply into the vagina. This is best done in the reclining position. Repeat this procedure for 7 days. The treatment can be shortened by beginning with one vaginal capsule on the first day and continuing with two vaginal capsules (one in the morning, one before bedtime) for the next three days. Complete the entire treatment, even if symptoms (e.g., pruritus and leukorrhea) have disappeared or menstruation begins

400 mg capsules: Once daily before bedtime, insert one vaginal capsule deeply into the

vagina. This is best done in the reclining position. Repeat this procedure for 3 days. The treatment can be repeated if necessary.

Complete the entire treatment even if symptoms (e.g., pruritus and

leukorrhea) have disappeared or menstruation begins In case of severe infections it may be advisable to prescribe a 6-day

treatment course right from the start. 1200 mg capsules:

Insert the vaginal capsule deeply into the vagina, preferably at bedtime.

This is best done in the reclining position. The treatment may be repeated if necessary. In case of severe infections it may be advisable to prescribe a longer

treatment course right from the start. GYNO-DAKTARIN Vaginal Ovules

Once daily before bedtime, insert one ovule deeply into the vagina. This is best done in the reclining position. Repeat this procedure for 14 days, even if symptoms (e.g., pruritus and leukorrhea) have disappeared or menstruation begins

GYNO-DAKTARIN Cream, Vaginal Capsules, and Vaginal Ovules are contraindicated in individuals with a known hypersensitivity to miconazole nitrate or another ingredient of the formulations

Special Warnings and Special Precautions for Use

Should local sensitization or an allergic reaction occur, the treatment should be discontinued. Appropriate therapy is indicated when the sexual partner is also

infected GYNO-DAKTARIN products do not stain skin or clothes

The concurrent use of latex condoms or diaphragms with vaginal anti-infective preparations may decrease the effectiveness of latex contra-

ceptive agents. Therefore, GYNO-DAKTARIN products should not be used concurrently with a latex condom or latex diaphragm. Interactions with Other Medicinal Products and Other Forms of Interaction

Miconazole administered systemically is known to inhibit CYP3A4/2C9. Due to the limited systemic availability after vaginal application, clinically

relevant interactions occur very rarely. In patients on oral anticoagulants, such as warfarin, caution should be exercised and the anticoagulant effect should be monitored. The effects and side effects of some other drugs (e.g., oral hypoglycemics and phenytoin), when co-administered with miconazole, can be increased and caution should be exercised. Contact should be avoided between latex products such as contraceptive diaphragms or condoms and GYNO-DAKTARIN since the constituents of GYNO-DAKTARIN may damage the latex (see Special Warnings and Special Precautions for Use).

Pregnancy and Lactation Use during pregnancy Although intravaginal absorption is limited, GYNO-DAKTARIN Cream, Vaginal Capsules, and Vaginal Ovules should be used in the first trimes

ter of pregnancy only if, in the judgement of the physician, the potential benefits outweigh the possible risks. Use during lactation It is not known whether miconazole nitrate is excreted in human milk. Caution should be exercised when using GYNO-DAKTARIN Cream,

Vaginal Capsules, and Vaginal Ovules during lactation (see Interactions with Other Medicinal Products and Other Forms of Interaction). Effects on Ability to Drive and Use Machines Not applicable.

Undesirable Effects Clinical trial data

Reproductive System and Breast Disorders

The safety of GYNO-DAKTARIN was evaluated in a total of 537 women with microbiologically confirmed candidiasis and symptoms (e.g., vulvovaginal itching, burning/irritation), or signs of vulvar erythema, edema,

Preferred Term

Infections and Infestations Urinary tract infection

Gastrointestinal Disorders

Abdominal pain upper

Abdominal pain lower

Abdominal pain

Nausea

blind clinical trials. Subjects were treated with miconazole intravaginally, randomly assigned to either a single 1200 mg capsule, or a 7-day application of 2% vaginal cream. Adverse Drug Reactions (ADRs) reported by ≥1% of GYNO-DAKTARIN-treated subjects in these trials are shown in Table 1 Table 1 Adverse Drug Reactions Reported by ≥1% of GYNO-**DAKTARIN-treated Subjects in 2 Single Blind Clinical Trials**

excoriation, or vaginal erythema or edema who participated in 2 single-

Miconazole Miconazole 2% System/Organ Vaginal Cream 1200 mg Capsule (n=272) %

Genital pruritus 16.5 23 female Vaginal burning 22.8 22.6 Vulvovaginal 16.2 14.3 discomfort Dysmenorrhoea 3.3 3.4 Vaginal discharge 0.4 3.7 Vaginal haemorrhage 0.4 1.1 Vaginal pain 0.4 1.5 **Nervous System Disorders** Headache 9.6 13.6

1.1

1.8

1.5

1.5

1.5

DAKTARIN-treated Subjects in 2 Single Blind Clinical Trials

Table 1 Adverse Drug Reactions Reported by ≥1% of GYNO-

System/Organ Class Preferred Term	1200 mg Capsule (n=272) %	Vaginal Cream 7 Days (n=265)%
Skin and subcutane	ous Tissue Disorders	
Rash	1.1	0.4
Renal and Urinary D	isorders	
Dysuria	1.1	0.4

Additional ADRs that occurred in <1% of GYNO-DAKTARIN-treated subjects (n = 537 women) in the single-blind clinical studies are listed

Table 2. Adverse Drug Reactions Reported by <1% of GYNO-**DAKTARIN-treated Subjects in 2 Single Blind Clinical Trials**

System/Organ Class Preferred Term	Miconazole 1200 mg Capsule (n=272) %	Miconazole 2% Vaginal Cream 7 Days (n=265) %
Skin and subcutaneou	s tissue disorders	
Rash pruritic	0	0.4
Rosacea	0.4	0
Swelling face	0.7	0
Urticaria	0.4	0

severity.

Postmarketing Data

Adverse drug reactions first identified during postmarketing experience with GYNO-DAKTARIN are included in Table 3. In each table, the frequencies are provided according to the following convention: Very common ≥1/10

Common $\geq 1/100$ and < 1/10≥1/1000 and <1/100 Uncommon $\geq 1/10000, <1/1000$ Rare Very rare <1/10000, including isolated reports

In Table 3, ADRs are presented by MedDRA System organ class and frequency category based on spontaneous reporting rates

Table 3. Adverse Drug Reactions Identified During Postmarketing Experience with GYNO-DAKTARIN by Frequency Category **Estimated from Spontaneous Reporting Rates Immune System Disorders**

Very Rare Hypersensitivity including Anaphylactic and Anaphylactoid reactions, Angioedema

Skin and Subcutaneous Tissue Disorders

Reproductive System and Breast Disorders

Very Rare

Very rare Vaginal irritation

General Disorders and Administrative Site Conditions

Pruritis

Application site reaction

GYNO-DAKTARIN products, an appropriate method of gastric emptying may be used, if considered necessary. See also Interactions with Other Medicinal Products and Other Forms of Interaction. PHARMACOLOGICAL PROPERTIES

GYNO-DAKTARIN products are intended for local application and not

for oral use. In the event of accidental ingestion of large quantities of

Pharmacodynamic Properties

Pharmacotherapeutic classification: (Antiinfectives and antiseptics, excl. combinations with corticosteroids, imidazole derivative) ATC code: G01A F04 Miconazole combines a potent antifungal activity against common

dermatophytes and yeasts with an antibacterial activity against certain gram-positive bacilli and cocci. Miconazole inhibits the biosynthesis of ergosterol in fungi and changes

the composition of other lipid components in the membrane, resulting in fungal cell necrosis. In general, miconazole exerts a very rapid effect on pruritus, a symptom

that frequently accompanies dermatophyte and yeast infections **Pharmacokinetic Properties**

GYNO-DAKTARIN Vaginal Capsules After the capsule has been inserted into the vagina, the outer covering

rapidly disintegrates and the active suspension is almost instantane

ously released GYNO-DAKTARIN Cream, Vaginal Capsules, and Vaginal Ovules **Absorption**: Miconazole persists in the vagina for up to 72 hours after

a single dose. Systemic absorption of miconazole after intravaginal administration is limited, with a bioavailability of 1 to 2% following intravaginal administration of a 1200 mg dose. Plasma concentrations of miconazole are measurable within 2 hours of administration in some subjects, with maximal levels seen 12 to 24 hours after administration. Plasma concentrations decline slowly thereafter and were still measurable in most subjects 96 hours post-dose. A second dose administered 48 hours later resulted in a plasma profile similar to that of the first dose. **Distribution**: Absorbed miconazole is bound to plasma proteins (88.2%) and red blood cells (10.6%). Metabolism and Excretion: The small amount of miconazole that is absorbed is eliminated predominantly in feces as both unchanged drug and metabolites over a four-day post-administration period. Smaller

amounts of unchanged drug and metabolites also appear in urine. The mean apparent elimination half-life is 57 hours.

Preclinical Safety Data Preclinical data reveal no special hazard for humans based on conventional studies of local irritation, single and repeated dose toxicity, geno-

toxicity, and toxicity to reproduction. PHARMACEUTICAL PARTICULARS

List of Excipients GYNO-DAKTARIN Cream: The cream formulation consists of PEG-6 (and) PEG-32 (and) glycol stearate, oleoyl macrogolglycerides, liquid

paraffin, benzoic acid, butylated hydroxyanisole and purified water. GYNO-DAKTARIN Vaginal Capsules: The inactive ingredients of the vaginal capsules are liquid paraffin, white petrolatum. The 1200 mg vaginal capsules also contain lecithin. The capsule itself contains gelatin, glycerol, titanium dioxide, sodium ethylparahydroxybenzoate and sodium propylparahydroxybenzoate. GYNO-DAKTARIN Vaginal Ovules: The inactive ingredient of the ovules

Incompatibilities None Known

Observe expiry date on the outer pack

Special Precautions for Storage GYNO-DAKTARIN Cream: Store at 25°C or below.

7 Days

(n=265)%

0.4

2.3 1.1

1.1

0

is hard fat.

GYNO-DAKTARIN Vaginal Capsules: Store between 15 and 30° C and GYNO-DAKTARIN Vaginal Ovules: Store between 15 and 30° C.

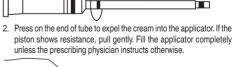
Keep GYNO-DAKTARIN Cream, Vaginal Capsules, and Vaginal Ovules out of reach of children.

Nature and Contents of Container GYNO-DAKTARIN Cream is supplied in tubes containing 40 g and 78 g and with 8 and 16 disposable applicators.

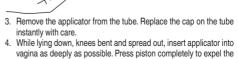
GYNO-DAKTARIN Vaginal Capsules 200 mg are supplied in boxes of 7, 400 mg in boxes of 3, 1200 mg in boxes of 1. GYNO-DAKTARIN Vaginal Ovules are supplied in boxes of 8 or 15

Instructions for Use and Handling <and Disposal> GYNO-DAKTARIN Cream

To open the tube unscrew the cap. Then pierce the seal of the tube using the pin on the top of the cap. Replace the cap with the applicator.



piston shows resistance, pull gently. Fill the applicator completely



cream. Remove the applicator and throw it away. GYNO-DAKTARIN Vaginal Capsules

GYNO-DAKTARIN Vaginal Ovules Not applicable.

Not applicable

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DATE OF REVISION OF THE TEXT