

## PACKAGE LEAFLET

### Package leaflet: Information for the user

#### Letrozol Sandoz 2,5 mg, filmomhulde tabletten

letrozole

**Read all of this leaflet carefully before you start taking this medicine because it contains important information for you.**

- Keep this leaflet. You may need to read it again.
- If you have any further questions, ask your doctor or pharmacist or nurse.
- This medicine has been prescribed for you only. Do not pass it on to others. It may harm them, even if their signs of illness are the same as yours.
- If you get any side effects, talk to your doctor or pharmacist or nurse. This includes any possible side effects not listed in this leaflet. See section 4.

#### **What is in this leaflet**

1. What [Nationally completed name] is and what it is used for
2. What you need to know before you take [Nationally completed name]
3. How to take [Nationally completed name]
4. Possible side effects
5. How to store [Nationally completed name]
6. Contents of the pack and other information

#### **1. What [Nationally completed name] is and what it is used for**

[Nationally completed name] contains an active substance called letrozole. It belongs to a group of medicines called aromatase inhibitors. It is a hormonal (or “endocrine”) breast cancer treatment. Growth of breast cancer is frequently stimulated by oestrogens which are female sex hormones. [Nationally completed name] reduces the amount of oestrogen by blocking an enzyme (“aromatase”) involved in the production of oestrogens and therefore may block the growth of breast cancer that needs oestrogens to grow. As a consequence tumour cells slow or stop growing and/or spreading to other parts of the body.

#### **What [Nationally completed name] is used for**

[Nationally completed name] is used to treat breast cancer in women who have gone through menopause i.e cessation of periods.

It is used to prevent cancer from happening again. It can be used as first treatment before breast cancer surgery in case immediate surgery is not suitable or it can be used as first treatment after breast cancer surgery or following five years treatment with tamoxifen. [Nationally completed name] is also used to prevent breast tumour spreading to other parts of the body in patients with advanced breast cancer.

If you have any questions about how [Nationally completed name] works or why this medicine has been prescribed for you, ask your doctor.

#### **2. What you need to know before you take [Nationally completed name]**

Follow all the doctor’s instructions carefully. They may differ from the general information in this leaflet.

### **Do not take [Nationally completed name]**

- if you are allergic to letrozole or to any of the other ingredients of this medicine (listed in section 6),
- if you still have periods, i.e. if you have not yet gone through the menopause,
- if you are pregnant,
- if you are breast-feeding.

### **If any of these conditions apply to you, do not take this medicine and talk to your doctor. Warnings and precautions**

Talk to your doctor or pharmacist before taking [Nationally completed name]

- if you have a severe kidney disease,
- if you have a severe liver disease,
- if you have a history of osteoporosis or bone fractures (see also “Follow-up during [Nationally completed name] treatment” in section 3).

If any of these conditions apply to you, **tell your doctor**. Your doctor will take this into account during your treatment with [Nationally completed name].

Letrozole may cause inflammation in tendons or tendon injury (see section 4). At any sign of tendon pain or swelling – rest the painful area and contact your doctor.

### **Children and adolescents**

Children and adolescents should not use this medicine.

### **Elderly**

People aged 65 years and over can use this medicine at the same dose as for other adults.

### **Other medicines and [Nationally completed name]**

Tell your doctor or pharmacist if you are taking, have recently taken or might take any other medicines, including medicines obtained without a prescription.

### **Pregnancy, breast-feeding and fertility**

- You should only take [Nationally completed name] when you have gone through the menopause. However, your doctor should discuss with you the use of effective contraception, as you may still have the potential to become pregnant during treatment with [Nationally completed name].
- You must not take [Nationally completed name] if you are pregnant or breast feeding as it may harm your baby.

### **Driving and using machines**

If you feel dizzy, tired, drowsy or generally unwell, do not drive or operate any tools or machines until you feel normal again.

### **[Nationally completed name] contains lactose and sodium**

If you have been told by your doctor that you have an intolerance to some sugars, contact your doctor before taking this medicine.

This medicine contains less than 1 mmol sodium (23 mg) per dose, that is to say essentially “sodium free”.

## **3. How to take [Nationally completed name]**

Always take this medicine exactly as your doctor or pharmacist has told you. Check with your doctor or pharmacist if you are not sure.

The usual dose is one tablet of [Nationally completed name] to be taken once a day. Taking [Nationally completed name] at the same time each day will help you remember when to take your tablet.

The tablet can be taken with or without food and should be swallowed whole with a glass of water or another liquid.

#### **How long to take [Nationally completed name]**

Continue taking [Nationally completed name] every day for as long as your doctor tells you. You may need to take it for months or even years. If you have any questions about how long to keep taking [Nationally completed name], talk to your doctor.

#### **Follow-up during [Nationally completed name] treatment**

You should only take this medicine under strict medical supervision. Your doctor will regularly monitor your condition to check whether the treatment is having the right effect. [Nationally completed name] may cause thinning or wasting of your bones (osteoporosis) due to the reduction of oestrogens in your body. Your doctor may decide to measure your bone density (a way of monitoring for osteoporosis) before, during and after treatment.

#### **If you take more [Nationally completed name] than you should**

If you have taken too much [Nationally completed name], or if someone else accidentally takes your tablets, contact a doctor or hospital for advice immediately. Show them the pack of tablets. Medical treatment may be necessary.

#### **If you forget to take [Nationally completed name]**

- If it is almost time for your next dose (e.g. within 2 or 3 hours), skip the dose you missed and take your next dose when you are meant to.
- Otherwise, take the dose as soon as you remember, and then take the next tablet as you would normally.
- Do not take a double dose to make up for the one that you missed.

#### **If you stop taking [Nationally completed name]**

Do not stop taking [Nationally completed name] unless your doctor tells you to. See also the section above "How long to take [Nationally completed name]".

## **4. Possible side effects**

Like all medicines, this medicine can cause side effects, although not everybody gets them.

Most of the side effects are mild to moderate and will generally disappear after a few days to a few weeks of treatment.

Some of these side effects, such as hot flushes, hair loss or vaginal bleeding, may be due to the lack of oestrogens in your body.

Do not be alarmed by this list of possible side effects. You may not experience any of them.

#### **Some side effects could be serious:**

##### **Uncommon** (may affect up to 1 in 100 people):

- Weakness, paralysis or loss of feeling in any part of the body (particularly arm or leg), loss of coordination, nausea, or difficulty speaking or breathing (sign of a brain disorder, e.g. stroke).
- Sudden oppressive chest pain (sign of a heart disorder).

- Swelling and redness along a vein which is extremely tender and possibly painful when touched.
- Severe fever, chills or mouth ulcers due to infections (lack of white blood cells).
- Severe persistent blurred vision.
- Inflammation of a tendon or tendonitis (connective tissues that connect muscles to bones)

**Rare** (may affect up to 1 in 1,000 people):

- Difficulty breathing, chest pain, fainting, rapid heart rate, bluish skin discoloration, or sudden arm, leg or foot pain (signs that a blood clot may have formed).
- Rupture of a tendon (connective tissues that connect muscles to bones)

**If any of the above occurs, tell your doctor straight away.**

You should also inform the doctor straight away if you experience any of the following symptoms during treatment with [Nationally completed name]:

- Swelling mainly of the face and throat (signs of allergic reaction).
- Yellow skin and eyes, nausea, loss of appetite, dark-coloured urine (signs of hepatitis).
- Rash, red skin, blistering of the lips, eyes or mouth, skin peeling, fever (signs of skin disorder).

**Some side effects are very common** (may affect more than 1 in 10 people):

- Hot flushes
- Increased level of cholesterol (hypercholesterolaemia)
- Fatigue
- Increased sweating
- Pain in bones and joints (arthralgia)

If any of these affects you severely, tell your doctor.

**Some side effects are common** (may affect up to 1 in 10 people):

- Skin rash
- Headache
- Dizziness
- Malaise (generally feeling unwell)
- Gastrointestinal disorders such as nausea, vomiting, indigestion, constipation, diarrhoea
- Increase in or loss of appetite
- Pain in muscles
- Thinning or wasting of your bones (osteoporosis), leading to bone fractures in some cases (see also "Follow-up during [Nationally completed name] treatment" in section 3)
- Swelling of arms, hands, feet, ankles (oedema)
- Depression
- Weight increase
- Hair loss
- Raised blood pressure (hypertension)
- Abdominal pain
- Dry skin
- Vaginal bleeding
- Palpitations, rapid heart rate
- Joint stiffness (arthritis)
- Chest pain

If any of these affects you severely, tell your doctor.

**Other side effects are uncommon** (may affect up to 1 in 100 people):

- Nervous disorders such as anxiety, nervousness, irritability, drowsiness, memory problems, somnolence, insomnia
- Pain or burning sensation in the hands or wrist (carpal tunnel syndrome)
- Impairment of sensation, especially that of touch
- Eye disorders such as blurred vision, eye irritation
- Skin disorders such as itching (urticaria)
- Vaginal discharge or dryness
- Breast pain
- Fever
- Thirst, taste disorder, dry mouth
- Dryness of mucous membranes
- Weight decrease
- Urinary tract infection, increased frequency of urination
- Cough
- Increased level of enzymes
- Yellowing of the skin and eyes
- High blood levels of bilirubin (a breakdown product of red blood cells)

**Side effects with frequency not known** (frequency cannot be estimated from the available data)

Trigger finger, a condition in which your finger or thumb catches in bent position.

If any of these affects you severely, tell your doctor.

### Reporting of side effects

If you get any side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. You can also report side effects directly via [the national reporting system](#) listed in [Appendix V](#). By reporting side effects, you can help provide more information on the safety of this medicine.

## 5. How to store [Nationally completed name]

Keep this medicine out of the sight and reach of children.

Do not use this medicine after the expiry date which is stated on the carton and on each blister after EXP. The expiry date refers to the last day of the month.

This medicine does not require any special storage conditions.

Do not use any pack that is damaged or shows signs of tampering.

Do not throw away any medicines via wastewater or household waste. Ask your pharmacist how to throw away medicines you no longer use. These measures will help protect the environment.

## 6. Contents of the pack and other information

### What [Nationally completed name] contains

- The active substance is letrozole. Each film-coated tablet contains 2.5 mg letrozole.
- The other ingredients are lactose monohydrate, croscarmellose sodium, povidone K-29/32 (E1201), silica, colloidal anhydrous (E551) and magnesium stearate. The coating is composed of hypromellose (E464), titanium dioxide (E171), iron oxide yellow (E172), macrogol (E1521) and talc (E553b).

### What [Nationally completed name] looks like and contents of the pack

- [Nationally completed name] is dark yellow, round (6.1 mm in diameter), slightly biconvex film-coated tablet (tablet), debossed with “5” on one side and “H” on the other side.
- [Nationally completed name] is packaged in clear PVC/PE/PVdC//Alu blister.

Pack sizes: 28, 30, 60, 98, 100 or 120 film-coated tablets in each carton.

Not all pack sizes may be marketed.

### Marketing Authorisation Holder and Manufacturer

Vergunninghouder:

Sandoz B.V.  
Hospitaaldreef 29  
1315 RC Almere  
Nederland

Fabrikanten:

Pharmadox Healthcare Limited  
KW20A Kordin Industrial Park  
PLA 3000 Paola  
Malta

Salutas Pharma GmbH  
Otto-Von-Guericke-Allee 1  
39179 Barleben  
Saxony-Anhalt  
Duitsland

Lek Pharmaceuticals d.d.  
Verovškova ulica 57  
Ljubljana 1526,  
Slovenië

**In het register ingeschreven onder:** RVG 133221

**This medicine is authorised in the Member States of the European Economic Area under the following names:**

Nederland	Letrozol Sandoz 2,5 mg, filmomhulde tabletten
Oostenrijk	Letrozol Sandoz GmbH 2,5 mg - Filmtabletten
België	Letrobloxus 2,5 mg filmomhulde tabletten
Tsjechië	Letrozol Sandoz 2,5 mg potahované tablety
Duitsland	Letrozol - 1 A Pharma 2,5 mg Filmtabletten
Spanje	Letrozol Sandoz Farmacéutica 2,5 mg comprimidos recubiertos con película EFG
Finland	Letrozole Hexal 2,5 mg kalvopäällysteiset tabletit
Frankrijk	LETROZOLE GNR 2,5 mg, comprimé pelliculé
Kroatië	Letrozol Sandoz 2,5 mg filmom obložene tablete
Ierland	Letrozole Rowex 2.5 mg film-coated tablets
Italië	Letrozolo Sandoz GmbH
Portugal	Letrozol Sandoz Farmacêutica 2,5 mg Comprimido revestido por película
Zweden	Letrozole Hexal

Slowakije                      Letrozol Sandoz d.d. 2,5 mg

**Deze bijsluiter is voor het laatst goedgekeurd in november 2025.**