

## Package leaflet: Information for the user

### Imatinib Actavis Group 400 mg film-coated tablets

#### Imatinib

**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, 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, pharmacist or nurse. This includes any possible side effects not listed in this leaflet. See section 4.

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

1. What Imatinib Actavis Group is and what it is used for
2. What you need to know before you take Imatinib Actavis Group
3. How to take Imatinib Actavis Group
4. Possible side effects
5. How to store Imatinib Actavis Group
6. Contents of the pack and other information

#### **1. What Imatinib Actavis Group is and what it is used for**

Imatinib Actavis Group is a medicine containing an active substance called imatinib. This medicine works by inhibiting the growth of abnormal cells in the diseases listed below. These include some types of cancer.

#### **Imatinib Actavis Group is a treatment for:**

- **Chronic myeloid leukaemia (CML).** Leukaemia is a cancer of white blood cells. These white cells usually help the body to fight infection. Chronic myeloid leukaemia is a form of leukaemia in which certain abnormal white cells (named myeloid cells) start growing out of control.

In adult patients, Imatinib Actavis Group is intended for use in the most advanced phase of the disease (blast crisis). In children and adolescents, Imatinib Actavis Group can be used in different phases of the disease (chronic, accelerated phase and blast crisis).

#### **Imatinib Actavis Group is also a treatment for adults for:**

- **Philadelphia chromosome positive acute lymphoblastic leukaemia (Ph-positive ALL).** Leukaemia is a cancer of white blood cells. These white cells usually help the body to fight infection. Acute lymphoblastic leukaemia is a form of leukaemia in which certain abnormal white cells (named lymphoblasts) start growing out of control. Imatinib Actavis Group inhibits the growth of these cells.
- **Myelodysplastic/myeloproliferative diseases (MDS/MPD).** These are a group of blood diseases in which some blood cells start growing out of control. Imatinib Actavis Group inhibits the growth of these cells in a certain subtype of these diseases.
- **Hypereosinophilic syndrome (HES) and/or chronic eosinophilic leukaemia (CEL).** These are blood diseases in which some blood cells (named eosinophils) start growing out of control. Imatinib Actavis Group inhibits the growth of these cells in a certain subtype of these diseases.
- **Dermatofibrosarcoma protuberans (DFSP).** DFSP is a cancer of the tissue beneath the skin in which some cells start growing out of control. Imatinib Actavis Group inhibits the growth of these cells.

In the rest of this leaflet, we will use the abbreviations when talking about these diseases.

If you have any questions about how Imatinib Actavis Group works or why this medicine has been prescribed for you, ask your doctor.

## **2. What you need to know before you take Imatinib Actavis Group**

Imatinib Actavis Group will only be prescribed to you by a doctor with experience in medicines to treat blood cancers or solid tumours.

Follow all your doctor's instructions carefully, even if they differ from the general information contained in this leaflet.

### **Do not take Imatinib Actavis Group:**

- if you are allergic to imatinib or any of the other ingredients of this medicine (listed in section 6).

If this applies to you, **tell your doctor without taking Imatinib Actavis Group.**

If you think you may be allergic but are not sure, ask your doctor for advice.

### **Warnings and precautions**

Talk to your doctor before taking Imatinib Actavis Group:

- if you have or have ever had a liver, kidney or heart problem.
- if you are taking the medicine levothyroxine because your thyroid has been removed.
- if you have ever had or might now have a hepatitis B infection. This is because Imatinib Actavis Group could cause hepatitis B to become active again, which can be fatal in some cases. Patients will be carefully checked by their doctor for signs of this infection before treatment is started.

If any of these apply to you, **tell your doctor before taking Imatinib Actavis Group.**

**During treatment with Imatinib Actavis Group, tell your doctor straight away** if you put on weight very quickly. Imatinib Actavis Group may cause your body to retain water (severe fluid retention).

While you are taking Imatinib Actavis Group, your doctor will regularly check whether the medicine is working. You will also have blood tests and be weighed regularly.

### **Children and adolescents**

Imatinib Actavis Group is also a treatment for children and adolescents with CML. There is no experience in children with CML below 2 years of age. There is limited experience in children and adolescents with Ph-positive ALL and very limited experience in children and adolescents with MDS/MPD, DFSP and HES/CEL.

Some children and adolescents taking Imatinib Actavis Group may have slower than normal growth. The doctor will monitor the growth at regular visits.

### **Other medicines and Imatinib Actavis Group**

Tell your doctor or pharmacist if you are taking, have recently taken or might take any other medicines, including medicines obtained without a prescription (such as paracetamol) and including herbal medicines (such as St. John's Wort). Some medicines can interfere with the effect of Imatinib Actavis Group when taken together. They may increase or decrease the effect of Imatinib Actavis Group, either leading to increased side effects or making Imatinib Actavis Group less effective. Imatinib Actavis Group may do the same to some other medicines.

Tell your doctor if you are using medicines that prevent the formation of blood clots.

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

- If you are pregnant or breast-feeding, think you may be pregnant or are planning to have a baby, ask your doctor for advice before taking this medicine.
- Imatinib Actavis Group is not recommended during pregnancy unless clearly necessary as it may harm your baby. Your doctor will discuss with you the possible risks of taking Imatinib Actavis Group during pregnancy.
- Women who might become pregnant are advised to use effective contraception during treatment.
- Do not breast-feed during the treatment with Imatinib Actavis Group.
- Patients who are concerned about their fertility while taking Imatinib Actavis Group are advised to consult with their doctor.

### **Driving and using machines**

You may feel dizzy or drowsy or get blurred vision while taking this medicine. If this happens, do not drive or use any tools or machines until you are feeling well again.

### **Imatinib Actavis Group contains lactose**

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

## **3. How to take Imatinib Actavis Group**

Your doctor has prescribed Imatinib Actavis Group because you suffer from a serious condition. Imatinib Actavis Group can help you to fight this condition.

However, always take this medicine exactly as your doctor or pharmacist has told you. It is important that you do this as long as your doctor or pharmacist tells you to. Check with your doctor or pharmacist if you are not sure.

Do not stop taking Imatinib Actavis Group unless your doctor tells you to. If you are not able to take the medicine as your doctor prescribed or you feel you do not need it anymore, contact your doctor straight away.

### **How much Imatinib Actavis Group to take**

#### **Use in adults**

Your doctor will tell you exactly how many tablets of Imatinib Actavis Group to take.

- **If you are being treated for CML in blast crisis:**  
The usual starting dose is 600 mg to be taken as one tablet of 400 mg plus 2 tablets of 100 mg **once** a day.

Your doctor may prescribe a higher or lower dose depending on how you respond to the treatment. If your daily dose is 800 mg (2 tablets), you should take one tablet in the morning and a second tablet in the evening.

- **If you are being treated for Ph-positive ALL:**  
The starting dose is 600 mg to be taken as one tablet of 400 mg plus 2 tablets of 100 mg **once** a day.
- **If you are being treated for MDS/MPD:**  
The starting dose is 400 mg to be taken as one tablet **once** a day.
- **If you are being treated for HES/CEL:**

The starting dose is 100 mg, to be taken as one tablet of 100 mg **once** a day. Your doctor may decide to increase the dose to 400 mg, to be taken as one tablet of 400 mg **once** a day, depending on how you respond to treatment.

- **If you are being treated for DFSP:**

The dose is 800 mg per day (2 tablets), to be taken as one tablet in the morning and a second tablet in the evening.

**Use in children and adolescents**

The doctor will tell you how many tablets of Imatinib Actavis Group to give to your child. The amount of Imatinib Actavis Group given will depend on your child's condition, body weight and height. The total daily dose in children and adolescents must not exceed 800 mg with CML. The treatment can either be given to your child as a once-daily dose or alternatively the daily dose can be split into two administrations (half in the morning and half in the evening).

**When and how to take Imatinib Actavis Group**

- **Take Imatinib Actavis Group with a meal.** This will help protect you from stomach problems when taking Imatinib Actavis Group.
- **Swallow the tablets whole with a large glass of water.**

If you are unable to swallow the tablets, you can dissolve them in a glass of still or mineral water or apple juice:

- Use about 200 ml for each 400 mg tablet.
- Stir with a spoon until the tablets have completely dissolved.
- Once the tablet has dissolved, drink everything in the glass straight away. Traces of the dissolved tablets may be left behind in the glass.

**How long to take Imatinib Actavis Group**

Keep taking Imatinib Actavis Group every day for as long as your doctor tells you.

**If you take more Imatinib Actavis Group than you should**

If you have accidentally taken too many tablets, talk to your doctor **straight away**. You may require medical attention. Take the medicine pack with you.

**If you forget to take Imatinib Actavis Group**

- If you forget a dose, take it as soon as you remember. However if it is nearly time for the next dose, skip the missed dose.
- Then continue with your normal schedule.
- Do not take a double dose to make up a forgotten dose.

If you have any further questions on the use of this medicine, ask your doctor, pharmacist or nurse.

**4. Possible side effects**

Like all medicines, this medicine can cause side effects, although not everybody gets them. They are usually mild to moderate.

**Some side effects may be serious. Tell your doctor straight away if you get any of the following:**

**Very common** (may affect more than 1 in 10 people) **or common** (may affect up to 1 in 10 people) **side effects:**

- Rapid weight gain. Imatinib Actavis Group may cause your body to retain water (severe fluid retention).
- Signs of infection such as fever, severe chills, sore throat or mouth ulcers. Imatinib Actavis Group can reduce the number of white blood cells, so you might get infections more easily.
- Unexpected bleeding or bruising (when you have not hurt yourself).

**Uncommon** (may affect up to 1 in 100 people) **or rare** (may affect up to 1 in 1,000 people) **side effects:**

- Chest pain, irregular heart rhythm (signs of heart problems).
- Cough, having difficulty breathing or painful breathing (signs of lung problems).
- Feeling light-headed, dizzy or fainting (signs of low blood pressure).
- Feeling sick (nausea), with loss of appetite, dark-coloured urine, yellow skin or eyes (signs of liver problems).
- Rash, red skin with blisters on the lips, eyes, skin or mouth, peeling skin, fever, raised red or purple skin patches, itching, burning sensation, pustular eruption (signs of skin problems).
- Severe abdominal pain, blood in your vomit, stools or urine, black stools (signs of gastrointestinal disorders).
- Severely decreased urine output, feeling thirsty (signs of kidney problems).
- Feeling sick (nausea) with diarrhoea and vomiting, abdominal pain or fever (signs of bowel problems).
- Severe headache, weakness or paralysis of limbs or face, difficulty speaking, sudden loss of consciousness (signs of nervous system problems such as bleeding or swelling in skull/brain).
- Pale skin, feeling tired and breathlessness and having dark urine (signs of low levels of red blood cells).
- Eye pain or deterioration in vision, bleeding in the eyes.
- Pain in your hips or difficulty walking.
- Numb or cold toes and fingers (signs of Raynaud's syndrome).
- Sudden swelling and redness of the skin (signs of a skin infection called cellulitis).
- Difficulty hearing.
- Muscle weakness and spasms with an abnormal heart rhythm (signs of changes in the amount of potassium in your blood).
- Bruising.
- Stomach pain with feeling sick (nausea).
- Muscle spasms with a fever, red-brown urine, pain or weakness in your muscles (signs of muscle problems).
- Pelvic pain sometimes with nausea and vomiting, with unexpected vaginal bleeding, feeling dizzy or fainting due to low blood pressure (signs of problems with your ovaries or womb).
- Nausea, shortness of breath, irregular heartbeat, clouding of urine, tiredness and/or joint discomfort associated with abnormal laboratory test results (e.g. high potassium, uric acid and calcium levels and low phosphorous levels in the blood).

**Not known** (frequency cannot be estimated from the available data):

- Combination of a widespread severe rash, feeling sick, fever, high level of certain white blood cells or yellow skin or eyes (signs of jaundice) with breathlessness, chest pain/discomfort, severely decreased urine output and feeling thirsty etc. (signs of a treatment-related allergic reaction).
- Chronic renal failure.

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

**Other side effects may include:**

**Very common side effects** (may affect more than 1 in 10 people):

- Headache or feeling tired.
- Feeling sick (nausea), being sick (vomiting), diarrhoea or indigestion.
- Rash.
- Muscle cramps or joint, muscle or bone pain.
- Swelling such as round your ankles or puffy eyes.
- Weight gain.

If any of these affect you severely, **tell your doctor.**

**Common side effects** (may affect up to 1 in 10 people):

- Anorexia, weight loss or a disturbed sense of taste.
- Feeling dizzy or weak.
- Difficulty in sleeping (insomnia).
- Discharge from the eye with itching, redness and swelling (conjunctivitis), watery eyes or having blurred vision.
- Nose bleeds.
- Pain or swelling in your abdomen, flatulence, heartburn or constipation.
- Itching.
- Unusual hair loss or thinning.
- Numbness of the hands or feet.
- Mouth ulcers.
- Joint pain with swelling.
- Dry mouth, dry skin or dry eye.
- Decreased or increased skin sensitivity.
- Hot flushes, chills or night sweats.

If any of these affect you severely, **tell your doctor**.

**Not known** (frequency cannot be estimated from the available data):

- Reddening and/or swelling on the palms of the hands and soles of the feet which may be accompanied by tingling sensation and burning pain.
- Slowing of growth in children and adolescents.
- Recurrence (reactivation) of Hepatitis B infection when you have had hepatitis B in the past (a liver infection).

If any of these affect 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 HPRA Pharmacovigilance, Earlsfort Terrace, IRL – Dublin 2; Tel: +353 1 6764971; Fax: + 353 1 6762517. Website: [www.hpra.ie](http://www.hpra.ie); E-mail: [medsafety@hpra.ie](mailto:medsafety@hpra.ie).

By reporting side effects you can help provide more information on the safety of this medicine.

## **5. How to store Imatinib Actavis Group**

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 after EXP. The expiry date refers to the last day of that 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 Imatinib Actavis Group contains**

- The active substance is imatinib (as mesilate). Each film-coated tablet contains 478 mg imatinib mesilate corresponding to 400 mg of imatinib.
- The other ingredients are hypromellose (E464), cellulose microcrystalline, lactose monohydrate, crospovidone, silica (colloidal anhydrous), magnesium stearate, macrogol (E1521), talc (E553b), yellow iron oxide (E172) and red iron oxide (E172).

### **What Imatinib Actavis Group looks like and contents of the pack**

Brownish-orange, capsule-shape, 22 mm in length and 10 mm in width, biconvex, film-coated tablets with no scorings or markings.

*Pack sizes:*

The film-coated tablets are supplied in blister packs of 30, 60, 90 and 120 film-coated tablets.

Not all pack sizes may be marketed.

**Marketing Authorisation Holder**

Actavis Group PTC ehf.  
Reykjavíkurvegi 76-78  
220 Hafnarfjörður  
Iceland

**Manufacturer**

Actavis Group PTC ehf.  
Reykjavíkurvegur 76-78  
IS-220 Hafnarfjörður  
Iceland

**This medicinal product is authorised in the Member States of the EEA under the following names:**

SE	Imatinib Actavis Group
AT	Imatinib Actavis Group 400 mg Filmtabletten
BG	Imatinib Actavis Group
CY	Imatinib Actavis Group
CZ	Imatinib Actavis Group
DK	Imatinib Actavis Group
EE	Imatinib Actavis Group
EL	Imatinib Actavis Group
FI	Imatinib Actavis Group
HR	Imatinib Actavis Group 400 mg filmom obložene tablete
HU	Imatinib Actavis Group 400 mg filmtabletta
IE	Imatinib Actavis Group 400 mg Film-coated Tablets
IS	Imatinib Actavis Group
LT	Imatinib Actavis Group 400 mg plėvele dengtos tabletės
LV	Imatinib Actavis Group 400 mg apvalkotās tabletes
MT	Imatinib Actavis Group
NO	Imatinib Actavis Group
PL	Imatinib Actavis Group
RO	Imatinib Actavis Group 400 mg comprimate filmate
SI	Imatinib Actavis Group Filmsko obložene tablete
SK	Imatinib Actavis Group 400mg
UK	Imatinib Actavis Group 400 mg Film-coated Tablets

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