

PACKAGE LEAFLET: INFORMATION FOR THE PATIENT

Erlotinib Zentiva 25 mg film-coated tablets Erlotinib Zentiva 100 mg film-coated tablets Erlotinib Zentiva 150 mg film-coated tablets

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
- This medicine has been prescribed for you. 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 serious side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet (see section 4).

What is in this leaflet:

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

1. WHAT ERLOTINIB IS AND WHAT IT IS USED FOR

Erlotinib Zentiva contains the active substance erlotinib and is a medicine used to treat cancer by preventing the activity of a protein called epidermal growth factor receptor (EGFR). This protein is known to be involved in the growth and spread of cancer cells.

Erlotinib is indicated for adults. This medicine can be prescribed to you if you have non-small cell lung cancer at an advanced stage. It can be prescribed as initial therapy or as therapy if your disease remains largely unchanged after initial chemotherapy, provided your cancer cells have specific EGFR mutations. It can also be prescribed if previous chemotherapy has not helped to stop your disease.

This medicine can also be prescribed to you in combination with another treatment called gemcitabine if you have cancer of the pancreas at a metastatic stage.

2. WHAT YOU NEED TO KNOW BEFORE YOU TAKE ERLOTINIB

Do not take Erlotinib if you:

- are allergic to erlotinib or any of the ingredients of this medicine (listed in section 6).

Warnings and precautions

Talk to your doctor before using Erlotinib if you:

- are taking other medicines that may increase or decrease the amount of erlotinib in your blood or influence its effect (for example antifungals like ketoconazole, protease inhibitors, erythromycin, clarithromycin, phenytoin, carbamazepine, barbiturates, rifampicin, ciprofloxacin, omeprazole, ranitidine, St. John's Wort or proteasome inhibitors). Talk to your doctor as in some cases these medicines may reduce the efficacy or increase the side effects of Erlotinib and your doctor may need to adjust your treatment. Your doctor might avoid treating you with these medicines while you are receiving Erlotinib.
- are taking anticoagulants (a medicine which helps to prevent thrombosis or blood clotting e.g. warfarin) as Erlotinib may increase your tendency to bleed. Talk to your doctor, they will need to regularly monitor you with some blood tests.
- are taking statins (medicines to lower your blood cholesterol) as Erlotinib may increase the risk of statin related muscle problems, which on rare occasions can lead to serious muscle breakdown (rhabdomyolysis) resulting in kidney damage.
- use contact lenses and/or have a history of eye problems such as severe dry eyes, inflammation of the front part of the eye (cornea) or ulcers involving the front part of the eye. See also the section "Other medicines and Erlotinib" below.

Talk to your doctor if you:

- have sudden difficulty in breathing associated with coughing or fever because your doctor may need to treat you with other medicines and interrupt your Erlotinib treatment.
- have diarrhoea because your doctor may need to treat you with anti-diarrhoeal medicines (for example loperamide).
- have severe or persistent diarrhoea, nausea, loss of appetite, or vomiting – talk to your doctor immediately because your doctor may need to interrupt your Erlotinib treatment and may need to treat you in the hospital.
- have ever had problems with your liver. Erlotinib may cause serious liver problems and some cases have been fatal. Your doctor may perform blood tests while you are taking this medicine to monitor whether your liver functions properly.
- have severe pain in the abdomen, severe blistering or peeling of the skin. Your doctor may need to interrupt or stop your treatment.
- develop acute or worsening redness and pain in the eye, increased eye watering, blurred vision and/or sensitivity to light, tell your doctor or nurse immediately as you may need urgent treatment (see section 4 - Possible side effects below).
- are also taking a statin and experience unexplained muscle pain, tenderness, weakness or cramps. Your doctor may need to interrupt or stop your treatment. See also section 4 "Possible side effects".

Liver or kidney disease

It is not known whether Erlotinib has a different effect if your liver or kidneys are not functioning normally. The treatment with this medicine is not recommended if you have severe liver disease or severe kidney disease.

Glucuronidation disorder like Gilbert's syndrome

Your doctor must treat you with caution if you have a glucuronidation disorder like Gilbert's syndrome.

Smoking

You are advised to stop smoking if you are treated with Erlotinib as smoking could decrease the amount of your medicine in the blood.

Children and adolescents

Erlotinib has not been studied in patients under the age of 18 years. The treatment with this medicine is not recommended for children and adolescents.

Other medicines and Erlotinib

Tell your doctor or pharmacist if you are taking, have recently taken or might take any other medicines.

Erlotinib with food and drink

Do not take this medicine with food (see also section 3 "How to take Erlotinib").

Pregnancy and breast-feeding

Avoid pregnancy while being treated with Erlotinib. If you could become pregnant, use adequate contraception during treatment, and for at least 2 weeks after taking the last tablet. If you become pregnant while you are being treated with Erlotinib, immediately inform your doctor who will decide if the treatment should be continued.

Do not breast-feed if you are being treated with Erlotinib, and for at least 2 weeks after taking the last tablet.

If you are pregnant or breast-feeding, think you may be pregnant or are planning to have a baby, ask your doctor or pharmacist for advice before taking this medicine.

Driving and using machines

Erlotinib has not been studied for its possible effects on the ability to drive and use machines but it is very unlikely that your treatment will affect this ability.

Erlotinib 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 tablet, that is to say essentially 'sodium-free'.

3. HOW TO TAKE ERLOTINIB

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

The tablet should be taken at least one hour before or two hours after the ingestion of food.

The usual dose is one tablet of Erlotinib 150 mg each day if you have non-small cell lung cancer.

The usual dose is one tablet of Erlotinib 100 mg each day if you have metastatic pancreatic cancer. Erlotinib is given in combination with gemcitabine treatment.

Your doctor may adjust your dose in 50 mg steps. For the different dose regimens Erlotinib is available in strengths of 25 mg, 100 mg or 150 mg.

Erlotinib 100 mg, the tablet can be divided into equal doses.

If you take more Erlotinib than you should

Contact your doctor or pharmacist immediately. You may have increased side effects and your doctor may interrupt your treatment.

If you forget to take Erlotinib

If you miss one or more doses of Erlotinib, contact your doctor or pharmacist as soon as possible. Do not take a double dose to make up for a forgotten dose.

If you stop taking Erlotinib

It is important to keep taking erlotinib every day, as long as your doctor prescribes it for you.

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

4. POSSIBLE SIDE EFFECTS

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

Contact your doctor as soon as possible if you suffer from any of the side effects listed below. In some cases your doctor may need to reduce your dose of Erlotinib or interrupt treatment:

- Diarrhoea and vomiting (very common: may affect more than 1 in 10 people). Persistent and severe diarrhoea may lead to low blood potassium and impairment of your kidney function, particularly if you receive other chemotherapy treatments at the same time. If you experience more severe or persistent diarrhoea **contact your doctor immediately** as your doctor may need to treat you in the hospital.
- Eye irritation due to conjunctivitis/keratoconjunctivitis (very common: may affect more than 1 in 10 people) and keratitis (common: may affect up to 1 in 10 people).
- A form of lung irritation called interstitial lung disease (uncommon in European patients; common in Japanese patients: may affect up to 1 in 100 people in Europe and up to 1 in 10 in Japan). This disease can also be linked to the natural progression of your medical condition and can have a fatal outcome in some cases. If you develop symptoms such as sudden difficulty in breathing associated with cough or fever contact your doctor immediately as you could be suffering from this disease. Your doctor may decide to permanently stop your treatment with Erlotinib.
- Gastrointestinal perforations have been observed (uncommon: may affect up to 1 in 100 people). Tell your doctor if you have severe pain in your abdomen. Also, tell your doctor if you had peptic ulcers or diverticular disease in the past, as this may increase this risk.
- In rare cases inflammation of the liver (hepatitis) was observed (may affect up to 1 in 1,000 people). Symptoms may include a general feeling of being unwell, with or without possible jaundice (yellowing of the skin and eyes), dark urine, nausea, vomiting and abdominal pain. In rare cases liver failure was observed. This can potentially be fatal. If your blood tests indicate severe changes in your liver function, your doctor may need to interrupt your treatment.

Very common side effects (affects more than 1 in 10 people)

- Rash which may occur or worsen in sun exposed areas. If you are exposed to the sun, protective clothing and/or the use of sunscreen (e.g. mineral-containing sunscreen) may be advisable
- Infection
- Loss of appetite, decreased weight
- Depression
- Headache, altered skin sensation or numbness in the extremities
- Difficulty in breathing, coughing
- Nausea
- Mouth irritation
- Stomach pain, indigestion and flatulence
- Abnormal blood tests for liver function
- Itching, dry skin and loss of hair
- Tiredness, fever, rigors.

Common side effects (affects less than 1 in 10 people)

- Bleeding from the nose
- Bleeding from the stomach or the intestines
- Inflammatory reactions around the fingernails
- Infection of hair follicles
- Acne
- Cracked skin (skin fissures)
- Reduced kidney function (when given outside the approved indications in combination with chemotherapy).

Uncommon side effects (may affect up to 1 in 100 people)

- Eyelash changes
- Excess body and facial hair (of a male distribution pattern)
- Eyebrow changes
- Brittle and loose nails

Rare side effects (may affect up to 1 in 1,000 people)

- Flushed or painful palms or soles of the feet (Palmar plantar erythrodysesthesia syndrome)

Very rare side effects (may affect up to 1 in 10,000 people)

- Cases of perforation or ulceration of the cornea
- Severe blistering or peeling of skin (suggestive of Stevens-Johnson syndrome)
- Inflammation of the coloured part of the eye

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 Yellow Card Scheme at: www.mhra.gov.uk/yellowcard or search for MHRA Yellow Card in the Google Play or Apple App Store. By reporting side effects you can help provide more information on the safety of this medicine.

5. HOW TO STORE ERLOTINIB

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 blister and 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 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 Erlotinib Zentiva contains

- The active substance is erlotinib hydrochloride. Each film-coated tablet contains 25 mg, 100 mg or 150 mg erlotinib (as erlotinib hydrochloride)
- The other ingredients are:

Tablet core: lactose monohydrate; cellulose, microcrystalline (E460) and calcium hydrogen phosphate; sodium starch glycolate; silica, colloidal anhydrous; sodium laurilsulfate; magnesium stearate (E470 b).

Tablet coating: hypromellose (E464); hydroxypropylcellulose (E463); titanium dioxide (E171); macrogol.

What Erlotinib Zentiva looks like and contents of the pack

The 25 mg tablet is a white, round, biconvex tablet with "E9OB" debossed on one side and "25" on the other, with a diameter of approximately 6 mm.

The 100 mg tablet is a white, round, biconvex tablet with a score line on both sides, on one side the tablet is debossed with "E9OB" above the score line and "100" below the score line, with a diameter of approximately 10 mm.

The 150 mg tablet is a white, round, biconvex tablet with "E9OB" debossed on one side and "150" below the other, with a diameter of approximately 10.4 mm.

Each box contains 30 film-coated tablets in a single blister.

Marketing Authorisation Holder and Manufacturer

Marketing Authorisation Holder:

Zentiva Pharma UK Limited, 12 New Fetter Lane, London EC4A 1JP, United Kingdom

Manufacturer:

Synthon B.V., Microweg 22, Nijmegen, Gelderland 6545 CM, The Netherlands

Or

Synthon Hispania, .L., Calle Castello 1, Poligono, Las Salinas, Sant Boi De Llobregat, Barcelona 08830, Spain

This leaflet was last updated in September 2022

ZENTIVA

1065034711

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GENERAL INFO:

PM CODE:	1065034711
PRODUCT NAME:	COMON ERLOTINIB TABCO BL5X6 M24 GB
SAP ID / GMID:	904483; 904484; 904473
AW VERSION:	V2
CREATION DATE:	22.09.2022
AW BY:	MaU
SUPPLIER:	SYNTHON

TECHNICAL INFO:

FORMAT (size):	300 x 380 mm
LAETUS (pharma code):	N/A
FONT + MIN. SIZE:	Helvetica Neue LT Std 9 pt
MATERIAL TYPE (TS):	PILL

COLOURS: [1]

■ Black

TECH. COLOURS: [1]

■ DieCut

ZENTIVA

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