Package leaflet:
Information for the user

Bosulif® 100 mg
Bosulif® 400 mg
Bosulif® 500 mg
film-coated tablets
bosutinib

Read all of this leaflet carefully before you start taking this medicine because it contains important information for you and your carer.
- 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 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. This includes any possible side effects not listed in this leaflet. See section 4.

What is in this leaflet

1. What Bosulif is and what it is used for

Bosulif contains the active substance bosutinib. Bosulif is used to treat adult and paediatric patients aged 6 years and older who have a type of leukaemia called chronic phase (CP) Philadelphia chromosome-positive (Ph-positive) Chronic Myeloid Leukaemia (CML) and are newly-diagnosed or for whom previous medicines to treat CML have either not worked or are not suitable. It is also used to treat adult patients with accelerated phase (AP) and blast phase (BP) Ph+ CML for whom previous medicines to treat CML have not worked or are not suitable.

In patients with Ph-positive CML a change in DNA (genetic material) triggers a signal that tells the body to produce too many of a specific type of white blood cell called granulocytes. Bosulif blocks this signal and therefore stops the production of these cells.

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

2. What do you need to know before you take Bosulif

Do not take Bosulif
- if you are allergic to bosutinib or any of the other ingredients of this medicine (listed in section 6).
- if your doctor has told you that your liver has been damaged and is not working normally.

Warnings and precautions
Talk to your doctor, pharmacist or nurse before taking Bosulif:
- **if you have, or have had in the past, liver problems.** Tell your doctor if you have a history of liver problems including hepatitis (liver infection or inflammation) of any kind, or a history of any signs and symptoms of liver problems (see section 4 “Possible side effects”) because Bosulif may affect your liver function. Your doctor should do blood tests to check your liver function prior to your starting treatment with Bosulif and for the first 3 months of treatment with Bosulif, and as clinically indicated.
- **if you have diarrhoea and vomiting.** Tell your doctor if you develop any signs and symptoms of stomach or intestinal problems (see section 4 “Possible Side Effects”) Your doctor may provide an antidiarrheal or antiemetic product and/or fluids in order to reduce the symptoms. Your doctor may also withhold temporarily, dose reduce, or discontinue Bosulif (see section 3 “How to take Bosulif”). You should ask your doctor if use of your treatment for nausea or vomiting medicines together with Bosulif may result in a greater risk of heart arrhythmias.
- **if you suffer from bleeding problems.** Tell your doctor if you develop any signs and symptoms of blood problem (see section 4 “Possible Side Effects”), because Bosulif reduces the capability of your body to stop bleeding. During the first month, your doctor will perform weekly and then monthly complete blood counts for you. Your doctor may also withhold temporarily, dose reduce or discontinue Bosulif (see section 3 “How to take Bosulif”).
- **if you have an infection.** Tell your doctor if you develop any of the following signs and symptoms such as fever, problems with urine such as burning on urination, a new cough, or a new sore throat because Bosulif reduces the capability of your body to defend from infections.
- **if you have fluid retention.** Tell your doctor if you develop any of the following signs and symptoms of fluid retention during Bosulif treatment such as swelling of the ankles, feet or legs; difficulty breathing chest pain or a cough (these may be signs of fluid retention in the lungs or chest). Your doctor will monitor you for fluid retentiion and will manage your symptoms.
- **if you have heart problems.** Tell your doctor if you have a heart disorder, such as heart failure and decreased blood flow to the heart which can lead to heart attack. Get medical help right away if you get shortness of breath, weight gain, chest pain, or swelling in your hands, ankles or feet.
- **if you have been told you have abnormal heart rhythm**. Tell your doctor if you have arrhythmias or an abnormal electrical signal called “prolongation of the QT interval”. This is always important, but especially if you are experiencing frequent or prolonged diarrhoea as described above. If you faint (loss of consciousness) or have an irregular heartbeat while taking Bosulif, tell your doctor immediately, as this may be a sign of a serious heart condition (see section 2 “What do you need to know before you take Bosulif”). Your doctor will perform an electrocardiogram (ECG) before you start therapy. Your doctor will do a blood test prior to and during therapy and if you have low potassium or magnesium, your doctor will provide a treatment to correct the low blood levels.
- **if you have been told that you have problems with your kidneys.** Tell your doctor if you are urinating more frequently and producing larger amounts of urine with a pale colour or if you are urinating less frequently and producing smaller amounts of urine with a dark colour. Also tell your doctor if you are losing weight or have experienced swelling of your feet, ankles, legs, hands or face. Your doctor will assess how your kidneys are functioning before treatment and will closely monitor how your kidneys are functioning during the course of treatment with bosutinib.
- **if you have ever had or might now have a hepatitis B infection.** This is because Bosulif could cause hepatitis B to become active again, which can be fatal in some cases. Your doctor will test you for this infection before starting treatment. If you have this infection, your doctor will monitor you closely for signs and symptoms of the infection throughout therapy and several months after you have stopped therapy.
- **if you have or have had pancreas problems.** Tell your doctor if you develop abdominal pain or discomfort. If you have abdominal pain and your blood tests show high levels of lipase, an enzyme that helps your body break down fats in food, then your doctor may interrupt your treatment and perform tests to rule out problems with your pancreas.
- **if you have any of these symptoms: serious skin rashes.** Tell your doctor if you develop any of the following signs and symptoms of painful red or purplish rash that spreads and blisters and/or other lesions begin to appear in the mucous membrane (e.g., mouth and lips). If you develop a severe skin reaction during treatment, your doctor will permanently discontinue treatment.
- **if you notice any of these symptoms: pain in your side, blood in your urine or reduced amount of urine.** When your disease is very severe, your body may not be able to clear all the waste products from the dying cancer cells. This is called tumour lysis syndrome and can cause kidney failure and heart problems within 48 hours of the first dose of Bosulif. Your doctor will be aware of this and may ensure you are adequately hydrated and give you other medicines to help prevent it. Your doctor will perform a blood test to check for high uric acid levels and your doctor will provide a treatment to correct the high levels prior to starting therapy.

Sun/UV protection
You may become more sensitive to the sun or UV rays while taking bosutinib. It is important to cover sunlight-exposed areas of skin and use sunscreen with high sun protection factor (SPF).

Patients of Asian origin
If you are of Asian origin, you may have an increased risk of side effects with Bosulif. Your doctor will closely monitor you for serious side effects especially when increasing the dose.

Children and adolescents
Bosulif is not recommended for people whose age is under 6 years. This medicine has not been studied in children below the age of 1 year.

Other medicines and Bosulif
Tell your doctor or pharmacist if you are taking, have recently taken or might take any other medicines, including medicines obtained without a prescription, vitamins, and herbal medicines. Some medicines can affect the levels of Bosulif in your body. You should inform your doctor if you are taking medicines containing active substances such as those listed below:

The following active substances may increase the risk of side effects with Bosulif:
- ketoconazole, itraconazole, voriconazole, posaconazole and fluconazole, used to treat fungal infections.
- clarithromycin, telithromycin, erythromycin, and ciprofloxacin, used to treat bacterial infections.
- nefazodone, used to treat depression.
- mibefradil, diltiazem and verapamil, used to lower blood pressure in people with high blood pressure.
- ritonavir, lopinavir/ritonavir, indinavir, nelfinavir, saquinavir, atazanavir, amprenavir, fosamprenavir and darunavir, used to treat human immunodeficiency virus (HIV)/AIDS.
- boceprevir and telaprevir, used to treat hepatitis C.
- aprepitant, used to prevent and control nausea (feeling sick) and vomiting.
- imatinib, used to treat a type of leukaemia.
- crizotinib, used to treat a type of lung cancer called non-small cell lung cancer.

The following active substances may reduce the effectiveness of Bosulif:
- rifampicin, used to treat tuberculosis.
- phenytoin and carbamazepine, used to treat epilepsy.
- bosentan, used to lower high blood pressure in the lungs (pulmonary artery hypertension).
- nafcillin, an antibiotic used to treat bacterial infections.
- St. John’s Wort (a herbal preparation obtained without a prescription), used to treat depression.
- efavirenz and etravirine, used to treat HIV infections/AIDS.
- modafinil, used to treat certain types of sleep disorders.

These medicines should be avoided during your treatment with Bosulif. If you are taking any of them, tell your doctor. Your doctor may change the dose of these medicines, change the dose of Bosulif, or switch you to a different medicine.

The following active substances may affect the heart rhythm:
- amiodarone, disopyramide, procainamide, quinidine and sotalol used to treat heart disorder.
- chloroquine, halofantrine used to treat malaria.
- clarithromycin and moxifloxacin antibiotics used to treat bacterial infections.
- haloperidol, used to treat psychotic disease such as schizophrenia.
- domperidone, used to treat nausea and vomiting or to stimulate breast milk production.
- methadone, used to treat pain.

These medicines should be taken with caution during your treatment with Bosulif. If you are taking any of them, tell your doctor.

Acid reducing agents
Proton pump inhibitors (PPIs) should be taken with caution during your treatment with Bosulif as they may reduce the effectiveness of Bosulif. Your doctor may consider short acting antacids as an alternative to PPIs and administration times of Bosulif and antacids should be separated (i.e. take Bosulif in the morning and antacids in the evening) whenever possible.

The medicines listed here may not be the only ones that could interact with Bosulif, if you are not sure if the above applies to you or your child, ask your doctor.

Bosulif with food and drink
Do not take Bosulif with grapefruit or grapefruit juice, as it may increase the risk of side effects.

Pregnancy, breast-feeding and fertility
Bosulif is not to be used during pregnancy, unless clearly necessary, because Bosulif could harm an unborn baby. Ask your doctor for advice before taking Bosulif if you are pregnant or might become pregnant.

Women taking Bosulif will be advised to use effective contraception during treatment and for at least 1 month after the last dose. Vomiting or diarrhoea may reduce the effectiveness of oral contraceptives.

There is a risk that treatment with Bosulif will lead to decreased fertility and you may wish to seek advice about sperm storage before the treatment starts.

If you are breast-feeding, tell your doctor. Do not breast-feed during treatment with Bosulif as it could harm your baby.

Driving and using machines
If you experience dizziness, have blurred vision or feel unusually tired, do not drive or operate machines until these side effects have gone away.

Bosulif contains sodium
This medicine contains less than 1 mmol sodium (23 mg) per 100 mg, 400 mg, or 500 mg film-coated tablet, that is to say essentially ‘sodium-free’.

PAA244896

3. How to take Bosulif

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

Bosulif will only be prescribed to you by a doctor with experience in medicines to treat leukaemia.

Dose and method of administration
Adults
The recommended dose is 400 mg once daily for patients with newly-diagnosed CML. The recommended dose is 500 mg once daily for patients whose previous medicines to treat CML have either not worked or are not suitable. In the event you are not able to tolerate the recommended dose or are not responding to Bosulif correctly, your doctor may adjust your dose further.

Children and adolescents (6 years of age and older)
The recommended dose is 300 mg/m^2 body surface area once daily for newly-diagnosed paediatric patients. The recommended dose is 400 mg/m^2 body surface area once daily for resistant or intolerant paediatric patients.
Dose recommendations are provided in the following table. As appropriate, for the recommended dose you may combine different strengths of bosutinib film-coated tablets and/or hard capsules (see Package Leaflet for hard capsules).

Dosing of bosutinib for newly-diagnosed (ND) and for resistant or intolerant (R/I) paediatric patients

Body surface area	ND recommended dose	R/I recommended dose
0.55–<0.63 m^2	200 mg	250 mg
0.63–<0.75 m^2	200 mg	300 mg
0.75–<0.9 m^2	250 mg	350 mg
0.9–<1.1 m^2	300 mg	400 mg
≥ 1.1 m^2	400 mg*	500 mg*

* maximum starting dose (corresponding to maximum starting dose in adult indication)

In the event you are not able to tolerate the recommended dose or are not responding to Bosulif correctly, your doctor may adjust your dose further.

Take the tablet(s) once a day with food. Swallow the tablet(s) whole with water.
For patients who are unable to swallow a tablet, a hard capsule formulation is available.

If you take more Bosulif than you should
If you accidentally take too many Bosulif tablets or a higher dose than you need, contact a doctor for advice right away. If possible, show the doctor the pack, or this leaflet. You may require medical attention.

If you forget to take Bosulif
If dose is missed by less than 12 hours, take your recommended dose. If a dose is missed by more than 12 hours, take your next dose at your regular time on the following day.
Do not take a double dose to make up for the forgotten tablets.

If you stop taking Bosulif
Do not stop taking Bosulif unless your doctor tells you to do so. 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 right away.

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.

Serious side effects

You must immediately contact your doctor if you experience any of those serious side effects (see also section 2 "What you need to know before you take Bosulif"):

Very Common (may affect more than 1 in 10 people):
- reduction in the number of platelets (thrombocytopenia), red blood cells (anaemia) and/or neutrophils (type of white blood cells) (neutropenia) which may cause you to have abnormal bleeding, fever, or easy bruising without having an injury (you might have blood or lymphatic system disorder) see section 2 "What do you need to know before you take Bosulif").
- fluid around the lungs (pleural effusion).

Common (may affect up to 1 in 10 people):
- low white blood cells count (leukopenia).
- bleeding from the stomach or intestine (gastrointestinal haemorrhage) which may include blood in your vomit, stools (bowel movements) or urine, or have black stools (tarry black bowel movements) (see section 2 "What do you need to know before you take Bosulif").
- chest pain.
- toxic damage to the liver (hepatotoxicity), abnormal hepatic function including liver disorder (hepatic function abnormal) which may be accompanied with itching, yellow eyes or skin, dark urine, and pain or discomfort in the right upper stomach area or fever (see section 2 "What do you need to know before you take Bosulif").
- when the heart does not pump blood as well as it should (heart failure).
- when there is decreased blood flow to the heart (cardiac ischaemia).
- infection of the lung (pneumonia).
- defect in cardiac rhythm (electrocardiogram QT prolonged) that predisposes to fainting, dizziness and palpitation.
- increase in blood pressure (hypertension).
- high level of potassium in the blood (hyperkalaemia).
- acute kidney failure, kidney failure (renal failure), kidney impairment (renal impairment).
- fluid around the heart (pericardial effusion).
- allergic reaction (drug hypersensitivity).
- abnormally high blood pressure in the arteries of the lungs (pulmonary hypertension).
- acute inflammation of the pancreas (pancreatitis acute).

Uncommon (may affect up to 1 in 100 people):
- fever associated with low white blood cell count (febrile neutropenia).
- damage to the liver (liver injury).
- life-threatening allergic reaction (anaphylactic shock).
- abnormal build-up of fluid in the lungs (acute pulmonary oedema).
- skin eruption (drug eruption).
- scaly, peeling rash (exfoliative rash).
- inflammation of the sac-like covering of the heart (pericarditis).
- a marked decrease in the number of granulocytes (a type of white blood cells, granulocytopenia).
- severe skin disorder (erythema multiforme).
- nausea, shortness of breath, irregular heartbeat, muscular cramps, seizure, clouding of urine and tiredness associated with abnormal laboratory test results (high potassium, uric acid and phosphorous levels and low calcium levels in the blood) that can lead to changes in kidney function and acute renal failure (tumour lysis syndrome (TLS)).
- respiratory failure.
- inflammation of blood vessels in the skin which may result in a rash or bruising (cutaneous vasculitis).

Not known (frequency cannot be estimated from the available data):
- severe skin disorder (Stevens-Johnson syndrome, toxic epidermal necrolysis) that may include painful red or purplish rash that spreads and blisters and/or other lesions that begin to appear in the mucous membrane (e.g., mouth and lips) due to an allergic reaction.
- interstitial lung disease (disorders causing scarring in the lungs): signs include cough, difficulty breathing, painful breathing.
- recurrence (reactivation) of hepatitis B infection when you have had hepatitis B in the past (a liver infection).

Other side effects with Bosulif may include:

Very common (may affect more than 1 in 10 people):
- diarrhoea, vomiting, stomach pain (abdominal pain), nausea.
- Fever (pyrexia), swelling of hands, feet or face (oedema), fatigue, weakness.
- respiratory tract infection.
- nasopharyngitis.
- changes in blood test to determine if Bosulif is affecting your liver (alanine aminotransferase (ALT) increased, aspartate aminotransferase (AST) increased) and/or pancreas (lipase increased), kidneys (blood creatinine increased).
- decrease of appetite.
- joint pain (arthralgia), back pain.
- headache.
- skin rash, which may be itchy and/or generalised (rash).
- cough.
- shortness of breath (dyspnoea).
- feeling of instability (dizziness).

Common (may affect up to 1 in 10 people):
- stomach irritation (gastritis)
- pain.
- influenza, bronchitis.
- changes in blood test to determine if Bosulif is affecting your heart (blood creatine phosphokinase increased), liver (blood bilirubin increased, gamma glutamyltransferase (GGT) increased), and/or pancreas (amylase increased).
- low level of phosphorus in the blood (hypophosphataemia), excessive loss of body fluid (dehydration).
- pain in the muscles (myalgia).
- alteration of the sense of taste (dysgeusia).
- ringing in the ears (tinnitus).
- Hives (urticaria), acne.
- sensitivity to UV rays from the sun and other light sources (photosensitivity reaction).
- itching (pruritus).

Reporting of side effects
If you get any side effects, talk to your doctor, pharmacist or nurse. 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 Bosulif

- 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 foil 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 use this medicine if you notice that the pack 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. Content of the pack and other information

What Bosulif contains
- The active substance is bosutinib. Bosulif film-coated tablets come in different strengths.
  Bosulif 100 mg: each film-coated tablet contains 100 mg bosutinib (as monohydrate).
  Bosulif 400 mg: each film-coated tablet contains 400 mg bosutinib (as monohydrate).
  Bosulif 500 mg: each film-coated tablet contains 500 mg bosutinib (as monohydrate).
- The other ingredients are: microcrystalline cellulose (E460), croscarmellose sodium (E468), poloxamers 188, povidone (E1201) and magnesium stearate (E470b). The tablet film-coating contains polyvinyl alcohol, titanium dioxide (E171), macrogol 3350, talc (E553b) and iron oxide yellow (E172, for Bosulif 100 mg and 400 mg) or iron oxide red (E172, for Bosulif 400 mg and 500 mg) (see section 2 "Bosulif contains sodium").

What Bosulif looks like and contents of the pack
Bosulif 100 mg film-coated tablets are yellow, oval biconvex, debossed with "Pfizer" on one side and "100" on the other side.
Bosulif 100 mg is available in blisters containing either 14 or 15 film-coated tablets. Each carton contains 28, 30 or 112 film-coated tablets.

Bosulif 400 mg film-coated tablets are orange, oval biconvex, debossed with "Pfizer" on one side and "400" on the other side.
Bosulif 400 mg is available in blisters containing either 14 or 15 film-coated tablets. Each carton contains 28 or 30 film-coated tablets.

Bosulif 500 mg film-coated tablets are red, oval biconvex, debossed with "Pfizer" on one side and "500" on the other side.
Bosulif 500 mg is available in blisters containing either 14 or 15 film-coated tablets. Each carton contains 28 or 30 film-coated tablets.

Not all pack sizes may be marketed.

Marketing Authorisation Holder
Pfizer Limited
Ramsgate Road
Sandwich
Kent
CT13 9NJ
United Kingdom

Manufacturer
Pfizer Manufacturing Deutschland GmbH
Mooswaldallee 1
79108 Freiburg Im Breisgau
Germany

For any information about this medicine, please contact: Medical Information, Pfizer Ltd, Walton Oaks, Dorking Road, Tadworth, Surrey, KT20 7NS.
Telephone 01304 616161.

This leaflet was last revised in 12/2025.

Ref: GB BO 26_0

PAA244896