

PACKAGE LEAFLET
Package leaflet: Information for the user

Pirfenidone Rowex 267 mg film-coated tablets
Pirfenidone Rowex 801 mg film-coated tablets
Pirfenidone

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 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. This includes any possible side effects not listed in this leaflet. See section 4.

What is in this leaflet

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

1. What Pirfenidone Rowex is and what it is used for

Pirfenidone Rowex contains the active substance pirfenidone and it is used for the treatment of Idiopathic Pulmonary Fibrosis (IPF) in adults.

IPF is a condition in which the tissues in your lungs become swollen and scarred over time, and as a result makes it difficult to breathe deeply. This makes it hard for your lungs to work properly. Pirfenidone Rowex helps to reduce scarring and swelling in the lungs, and helps you breathe better.

2. What you need to know before you take Pirfenidone Rowex

Do not take Pirfenidone Rowex

- if you are allergic to pirfenidone or any of the other ingredients of this medicine (listed in section 6)
- if you have previously experienced angioedema with pirfenidone, including symptoms such as swelling of the face, lips and/or tongue which may be associated with difficulty breathing or wheezing
- if you are taking a medicine called fluvoxamine (used to treat depression and obsessive compulsive disorder [OCD])
- if you have severe or end stage liver disease
- if you have severe or end stage kidney disease requiring dialysis.

If any of the above affects you, do not take Pirfenidone Rowex. If you are unsure ask your doctor or pharmacist.

Warnings and precautions

Talk to your doctor or pharmacist before taking Pirfenidone Rowex

- You may become more sensitive to sunlight (photosensitivity reaction) when taking Pirfenidone Rowex. Avoid the sun (including sunlamps) whilst taking Pirfenidone Rowex. Wear sunblock daily and cover your arms, legs and head to reduce exposure to sunlight (see section 4: Possible side effects).
- You should not take other medicines, such as tetracycline antibiotics (such as doxycycline), which may make you more sensitive to sunlight.
- You should tell your doctor if you suffer from kidney problems
- You should tell your doctor if you suffer from mild to moderate liver problems.
- You should stop smoking before and during treatment with Pirfenidone Rowex. Cigarette smoking can reduce the effect of Pirfenidone Rowex.
- Pirfenidone Rowex may cause dizziness and tiredness. Be careful if you have to take part in activities where you have to be alert and co-ordinated.
- Pirfenidone Rowex can cause weight loss. Your doctor will monitor your weight whilst you are taking this medicine.
- Stevens-Johnson syndrome, toxic epidermal necrolysis, and drug reaction with eosinophilia and systemic symptoms (DRESS), have been reported in association with pirfenidone treatment. Stop using Pirfenidone Rowex and seek medical attention immediately if you notice any of the symptoms related to these serious skin reactions described in section 4.

Pirfenidone Rowex may cause serious liver problems and some cases have been fatal. You will need a blood test before you start taking Pirfenidone Rowex and at monthly intervals for the first 6 months and then every 3 months thereafter whilst you are taking this medicine to check whether your liver is working properly. It is important that you have these regular blood tests for as long as you are taking Pirfenidone Rowex.

Children and adolescents

Do not give Pirfenidone Rowex to children and adolescents under the age of 18.

Other medicines and Pirfenidone Rowex

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

This is especially important if you are taking the following medicines, as they may change the effect of Pirfenidone Rowex.

Medicines that may increase side effects of Pirfenidone Rowex:

- enoxacin (a type of antibiotic)
- ciprofloxacin (a type of antibiotic)
- amiodarone (used to treat some types of heart disease)
- propafenone (used to treat some types of heart disease)
- fluvoxamine (used to treat depression and obsessive compulsive disorder (OCD)).

Medicines that may reduce how well Pirfenidone Rowex works:

- omeprazole (used in the treatment of conditions such as indigestion, gastroesophageal reflux disease)
- rifampicin (a type of antibiotic).

Pirfenidone Rowex with food and drink

Do not drink grapefruit juice whilst taking this medicine. Grapefruit may prevent Pirfenidone Rowex from working properly.

Pregnancy and breast-feeding

As a precautionary measure, it is preferable to avoid the use of Pirfenidone Rowex if you are pregnant, planning to become pregnant, or think you might be pregnant as the potential risks to the unborn child are unknown.

If you are breast-feeding or plan to breast-feed speak to your doctor or pharmacist before taking Pirfenidone Rowex. As it is unknown whether Pirfenidone Rowex passes into breast milk, your doctor will discuss the risks and benefits of taking this medicine while breast-feeding if you decide to do so.

Driving and using machines

Do not drive or use machines if you feel dizzy or tired after taking Pirfenidone Rowex.

Pirfenidone Rowex contains sodium

Pirfenidone Rowex contains less than 1 mmol sodium (23 mg) per film-coated tablet, that is to say essentially 'sodium-free'.

3. How to take Pirfenidone Rowex

Treatment with Pirfenidone Rowex should be started and overseen by a specialist doctor experienced in the diagnosis and treatment of IPF.

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

Your medicine will usually be given to you in increasing doses as follows:

- for the first 7 days take a dose of 267 mg (1 yellow tablet), 3 times a day with food (a total of 801 mg/day)
- from day 8 to 14 take a dose of 534 mg (2 yellow tablets), 3 times a day with food (a total of 1,602 mg/day)
- from day 15 onwards (maintenance), take a dose of 801 mg (3 yellow tablets or 1 dark pink tablet), 3 times a day with food (a total of 2,403 mg/day).

The recommended maintenance daily dose of Pirfenidone Rowex is 801 mg (3 yellow tablets or 1 dark pink tablet) three times a day with food, for a total of 2403 mg/day.

Swallow the tablets whole with a drink of water, during or after a meal to reduce the risk of side effects such as nausea (feeling sick) and dizziness. If symptoms continue, see your doctor.

Dose reduction due to side effects

Your doctor may reduce your dose if you suffer from side effects such as, stomach problems, any skin reactions to sunlight or sun lamps, or significant changes to your liver enzymes.

If you take more Pirfenidone Rowex than you should

Contact your doctor, pharmacist or nearest hospital casualty department immediately if you have taken more tablets than you should, and take your medicine with you.

If you forget to take Pirfenidone Rowex

If you forget a dose, take it as soon as you remember. Do not take a double dose to make up for a forgotten dose. Each dose should be separated by at least 3 hours. Do not take more tablets each day than your prescribed daily dose.

If you stop taking Pirfenidone Rowex

In some situations, your doctor may advise you to stop taking Pirfenidone Rowex. If for any reason you have to stop taking Pirfenidone Rowex for more than 14 consecutive days, your doctor will restart your treatment with a dose of 267 mg 3 times a day, gradually increasing this to a dose of 801 mg 3 times a day.

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.

Stop taking Pirfenidone Rowex and seek medical attention tell your doctor immediately if you notice any of the following symptoms or signs:

- Swelling of the face, lips and/or tongue, itching, hives, difficulty breathing or wheezing, or feeling faint, which are signs of angioedema, a serious allergic reaction or anaphylaxis.
- Yellowing of the eyes or skin, or dark urine, potentially accompanied by itching of the skin, pain on the upper right side of your stomach area (abdomen), loss of appetite, bleeding or bruising more easily than normal, or feeling tired. These may be signs of abnormal liver function and could indicate liver injury, which is an uncommon side effect of Pirfenidone Rowex.
- Reddish non-elevated, or circular patches on the trunk, often with central blisters, skin peeling, ulcers of mouth, throat, nose, genitals and eyes. These serious skin rashes can be preceded by fever and flu-like symptoms (Stevens-Johnson syndrome or toxic epidermal necrolysis).
- Widespread rash, high body temperature and enlarged lymph nodes (DRESS syndrome or drug hypersensitivity syndrome).

Other side effects may include

Talk to your doctor if you get any side effects.

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

- infections of the throat or the airways going into the lungs and/or sinusitis
- feeling sick (nausea)

- stomach problems such as acid reflux, vomiting, and feeling constipated
- diarrhoea
- indigestion or stomach upset
- weight loss
- decreased appetite
- difficulty sleeping
- tiredness
- dizziness
- headache
- shortness of breath
- cough
- aching joints/joint pains.

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

- bladder infections
- feeling sleepy
- changes in taste
- hot flushes
- stomach problems such as feeling bloated, abdominal pain and discomfort, heart burn, and passing wind
- blood tests may show increased levels of liver enzymes
- skin reactions after going out in the sun or using sunlamps
- skin problems such as itchy skin, skin redness or red skin, dry skin, skin rash
- muscle pain
- feeling weak or feeling low in energy
- chest pain
- sunburn.

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

- low levels of sodium in the blood. This may cause headache, dizziness, confusion, weakness, muscle cramps or nausea and vomiting.
- blood tests may show decrease in white blood cells.

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: HPRA Pharmacovigilance; website: www.hpra.ie. By reporting side effects, you can help provide more information on the safety of this medicine.

5. How to store Pirfenidone Rowex

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, bottle and blister after EXP. The expiry date refers to the last day of that month. Do not store this medicine above 30°C.

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 Pirfenidone Rowex contains

Pirfenidone Rowex 267 mg film-coated tablets

- The active substance is pirfenidone. Each film-coated tablet contains 267 mg pirfenidone.
- The other ingredients are:
Tablet core: pregelatinised starch, croscarmellose sodium (E468), hydroxypropyl cellulose (E463), silicon dioxide (E551), magnesium stearate (E572)
Tablet coating (Opadry yellow 85F220100): polyvinyl alcohol – part. hydrolysed (E1203), titanium dioxide (E171), macrogol 3350, talc (E553B) and iron oxide yellow (E172).

Pirfenidone Rowex 801 mg film-coated tablets

- The active substance is pirfenidone. Each film-coated tablet contains 801 mg pirfenidone.
- The other ingredients are:
Tablet core: pregelatinised starch, croscarmellose sodium (E468), hydroxypropyl cellulose (E463), silicon dioxide (E551), magnesium stearate (E572)
Tablet coating (Opadry pink 85F240048): polyvinyl alcohol – part. hydrolysed (E1203), titanium dioxide (E171), macrogol 3350, talc (E553B), iron oxide yellow (E172), iron oxide red (E172) and iron oxide black (E172).

What Pirfenidone Rowex looks like and contents of the pack




Pirfenidone Rowex 267 mg film-coated tablets are yellow, oval, approximately 1.2 x 0.7 cm biconvex film-coated tablets, debossed with 'SD267' on one side.

Pirfenidone Rowex 801 mg film-coated tablets are dark pink, oval, approximately 1.8 x 0.9 cm biconvex film-coated tablets, debossed with 'SD801' on one side.

Pirfenidone Rowex 267 mg film-coated tablets are available in PVC/PE/PVDC-Alu blisters in

- Blister packs containing 63 or 252 film-coated tablets
- Unit-dose blister packs of 63x1 or 252x1 film-coated tablets
- 2-week treatment initiation packs:
 - Blister multipack containing 63 (1 pack of 21 and 1 pack of 42) film-coated tablets or
 - Unit-dose blister multipack containing 63 (1 pack of 21x1 and 1 pack of 42x1) film-coated tablets
- Continuation packs:
 - Blister multipack containing 252 (3 packs of 84) film-coated tablets or
 - Unit-dose blister multipack containing 252 (3 packs of 84x1) film-coated tablets

The 267 mg blister packs containing 63 film-coated tablets, the unit-dose blister packs containing 63x1 film-coated tablets, and the 2-week treatment initiation packs are marked with the following symbols and abbreviated names of the day as a reminder to take a dose three times a day:




 (sunrise; morning dose)  (sun; daytime dose) and  moon; evening dose).
Mon. Tue. Wed. Thu. Fri. Sat. Sun.

Pirfenidone Rowex 801 mg film-coated tablets are available in PVC/PE/PVDC-Alu blisters in

- Blister packs containing 84 or 252 film-coated tablets
- Unit-dose blister packs of 84x1 or 252x1 film-coated tablets

- Continuation packs:
 - Blister multipack containing 252 (3 packs of 84) film-coated tablets or
 - Unit-dose blister multipack containing 252 (3 packs of 84x1) film-coated tablets

The 801 mg blisters are marked with the following symbols and abbreviated names of the day as a reminder to take a dose three times a day:

 (sunrise; morning dose)  (sun; daytime dose) and  moon; evening dose).
Mon. Tue. Wed. Thu. Fri. Sat. Sun.

Not all pack sizes may be marketed.

Marketing Authorisation Holder and Manufacturers

Marketing Authorisation Holder

Rowex Ltd., Bantry, Co. Cork, Ireland.

Manufacturers

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

Lek Pharmaceuticals d.d., Trimlini 2d, 9220 Lendava, Slovenia.

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

DK	Pirfenidone Sandoz
EL	Pirfenidone/Sandoz 267 mg επικαλυμμένα με λεπτό υμένιο δισκία Pirfenidone/Sandoz 801 mg επικαλυμμένα με λεπτό υμένιο δισκία
ES	Pirfenidona Sandoz 267 mg comprimidos recubiertos con película EFG Pirfenidona Sandoz 801 mg comprimidos recubiertos con película EFG
FI	Pirfenidone Sandoz 267 mg tabletti, kalvopäällysteinen Pirfenidone Sandoz 801 mg tabletti, kalvopäällysteinen
FR	PIRFENIDONE SANDOZ 267 mg, comprimé pelliculé PIRFENIDONE SANDOZ 801 mg, comprimé pelliculé
HR	Pirfenidon Sandoz 267 mg filmom obložene tablete Pirfenidon Sandoz 801 mg filmom obložene tablete
IE	Pirfenidone Rowex 267 mg film-coated tablets Pirfenidone Rowex 801 mg film-coated tablets
IT	Pirfenidone Sandoz
NL	Pirfenidon Sandoz 267 mg, filmomhulde tabletten Pirfenidon Sandoz 801 mg, filmomhulde tabletten

NO Pirfenidone Sandoz
PL Pirfenidone Sandoz
PT Pirfenidona Sandoz
SE Pirfenidone Sandoz

This leaflet was last revised in 11/2025.