

Package leaflet: Information for the patient

Atorvas 10 mg film-coated tablets

Atorvas 20 mg film-coated tablets

Atorvas 40 mg film-coated tablets

Atorvas 80 mg film-coated tablets

atorvastatin

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

1. What Atorvas is and what it is used for

Atorvas belongs to a group of medicines known as statins, which are lipid (fat) regulating medicines.

Atorvas is used to lower lipids known as cholesterol and triglycerides in the blood when a low fat diet and lifestyle changes on their own have failed. If you are at an increased risk of heart disease, Atorvas can also be used to reduce such risk even if your cholesterol levels are normal. You should maintain a standard cholesterol lowering diet during treatment.

2. What you need to know before you take Atorvas

DO NOT take Atorvas:

- if you are allergic to atorvastatin or any of the other ingredients of this medicine (listed in section 6)
- if you have or have ever had a disease which affects the liver
- if you have had any unexplained abnormal blood tests for liver function
- if you are a woman able to have children and not using reliable contraception
- if you are pregnant or trying to become pregnant
- if you are breast-feeding
- if you use the combination of glecaprevir/pibrentasvir in the treatment of hepatitis C.

Warnings and precautions

Talk to your doctor, pharmacist or nurse before taking Atorvas:

- if you have severe respiratory failure
- if you are taking or have taken in the last 7 days a medicine called fusidic acid, (a medicine for bacterial infection) orally or by injection. The combination of fusidic acid and Atorvas can lead to serious muscle problems (rhabdomyolysis)
- if you have had a previous stroke with bleeding into the brain, or have small pockets of fluid in the brain from previous strokes
- if you have kidney problems
- if you have an under-active thyroid gland (hypothyroidism)
- if you have had repeated or unexplained muscle aches or pains, a personal history or family history of muscle problems
- if you have had previous muscular problems during treatment with other lipid-lowering medicines (e.g. other ‘-statin’ or ‘-fibrate’ medicines)
- if you have or have had myasthenia (a disease with general muscle weakness including in some cases muscles used when breathing), or ocular myasthenia (a disease causing eye muscle weakness) as statins may sometimes aggravate the condition or lead to the occurrence of myasthenia (see section 4)
- if you regularly drink a large amount of alcohol
- if you have a history of liver disease
- if you are older than 70 years.

If any of these apply to you, your doctor will need to carry out a blood test before and possibly during your Atorvas treatment to predict your risk of muscle related side effects. The risk of muscle related side effects e.g. rhabdomyolysis is known to increase when certain medicines are taken at the same time (see section 2 “Other medicines and Atorvas”).

Also tell your doctor or pharmacist if you have a muscle weakness that is constant. Additional tests and medicines may be needed to diagnose and treat this.

While you are on this medicine your doctor will monitor you closely if you have diabetes or are at risk of developing diabetes. You are likely to be at risk of developing diabetes if you have high levels of sugars and fats in your blood, are overweight and have high blood pressure.

Other medicines and Atorvas

Tell your doctor or pharmacist if you are taking, have recently taken or might take any other medicines. There are some medicines that may change the effect of Atorvas or their effect may be changed by Atorvas. This type of interaction could make one or both of the medicines less effective. Alternatively, it could increase the risk or severity of side-effects, including the important muscle wasting condition known as rhabdomyolysis described in section 4:

- Medicines used to alter the way your immune system works, e.g. ciclosporin
- Certain antibiotics or antifungal medicines, e.g. erythromycin, clarithromycin, telithromycin, ketoconazole, itraconazole, voriconazole, fluconazole, posaconazole, rifampin, fusidic acid
- Other medicines to regulate lipid levels, e.g. gemfibrozil, other fibrates, colestipol
- Some calcium channel blockers used for angina or high blood pressure, e.g. amlodipine, diltiazem, medicines to regulate your heart rhythm e.g. digoxin, verapamil, amiodarone

- Letemovir, a medicine that helps stop you from getting ill from cytomegalovirus
- Medicines used in the treatment of HIV e.g. ritonavir, lopinavir, atazanavir, indinavir, darunavir, the combination of tipranavir/ritonavir etc.
- Some medicines used in the treatment of hepatitis C e.g. telaprevir, boceprevir and the combination of elbasvir/grazoprevir, ledipasvir/sofosbuvir
- Other medicines known to interact with Atorvas include ezetimibe (which lowers cholesterol), warfarin (which reduces blood clotting), oral contraceptives, stiripentol (an anti-convulsant for epilepsy), cimetidine (used for heartburn and peptic ulcers), phenazone (a painkiller), colchicine (used to treat gout), and antacids (indigestion products containing aluminium or magnesium)
- Medicines obtained without a prescription: St John's Wort
- Daptomycin (a medicine used to treat complicated skin and skin structure infections and bacteria present in the blood)
- **If you need to take oral fusidic acid to treat a bacterial infection you will need to temporarily stop using this medicine. Your doctor will tell you when it is safe to restart Atorvas. Taking Atorvas with fusidic acid may rarely lead to muscle weakness, tenderness or pain (rhabdomyolysis). See more information regarding rhabdomyolysis in section 4.**

Atorvas with food, drink and alcohol

See section 3 for instructions on how to take Atorvas. Please note the following:

Grapefruit juice

Do not take more than one or two small glasses of grapefruit juice per day because large quantities of grapefruit juice can change the effects of Atorvas.

Alcohol

Avoid drinking too much alcohol while taking this medicine. See section 2 "Warnings and precautions" for details.

Pregnancy and breast-feeding

Do not take Atorvas if you are pregnant, or if you are trying to become pregnant.

Do not take Atorvas if you are able to become pregnant unless you use reliable contraceptive measures.

Do not take Atorvas if you are breast-feeding.

The safety of Atorvas during pregnancy and breast-feeding has not yet been proven. Ask your doctor or pharmacist for advice before taking any medicine.

Driving and using machines

Normally this medicine does not affect your ability to drive or operate machines. However, do not drive if this medicine affects your ability to drive. Do not use any tools or machines if your ability to use them is affected by this medicine.

Atorvas contains sodium

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

Atorvas 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 Atorvas

Before starting treatment, your doctor will place you on a low-cholesterol diet, which you should maintain also during therapy with Atorvas.

The usual starting dose of Atorvas is 10 mg once a day in adults and children aged 10 years or older. This may be increased if necessary, by your doctor until you are taking the amount you need. Your doctor will adapt the dose at intervals of 4 weeks or more. The maximum dose of Atorvas is 80 mg once a day.

Atorvas tablets or tablet parts should be swallowed with one glass of water, and can be taken at any time of day, with or without food. However, try to take your tablet at the same time every day.

20 mg, 40 mg and 80 mg film-coated tablets:

The tablet can be divided into equal doses.

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 duration of treatment with Atorvas is determined by your doctor.

Please ask your doctor if you think that the effect of Atorvas is too strong or too weak.

If you take more Atorvas than you should

If you accidentally take too many Atorvas tablets (more than your usual daily dose), contact your doctor or nearest hospital for advice.

If you forget to take Atorvas

If you forget to take a dose, just take your next scheduled dose at the correct time. Do not take a double dose to make up for a forgotten dose.

If you stop taking Atorvas

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

4. Possible side effects

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

If you experience any of the following serious side effects or symptoms, stop taking your tablets and tell your doctor immediately or go to the nearest hospital accident and emergency department.

Rare: may affect up to 1 in 1,000 people:

- Serious allergic reaction which causes swelling of the face, tongue and throat that can cause great difficulty in breathing.
- Serious illness with severe peeling and swelling of the skin, blistering of the skin, mouth, eyes genitals and fever. Skin rash with pink-red blotches especially on palms of hands or soles of feet which may blister.

- Muscle weakness, tenderness, pain or rupture or red-brown discolouration of urine and particularly, if at the same time, you feel unwell or have a high temperature it may be caused by an abnormal muscle breakdown (rhabdomyolysis). The abnormal muscle breakdown does not always go away, even after you have stopped taking atorvastatin, and it can be life-threatening and lead to kidney problems.

Very rare: may affect up to 1 in 10,000 people:

- if you experience problems with unexpected or unusual bleeding or bruising, tiredness, loss of appetite, stomach pain, feeling sick, yellowing of your skin or eyes (jaundice), this may be suggestive of a liver complaint. You should consult your doctor as soon as possible.
- lupus-like disease syndrome (including rash, joint disorders and effects on blood cells).

Other possible side effects with Atorvas:

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

- Inflammation of the nasal passages, pain in the throat, nosebleed
- Allergic reactions
- Increases in blood sugar levels (if you have diabetes continue careful monitoring of your blood sugar levels), increase in blood creatine kinase
- Headache
- Nausea, constipation, wind, indigestion, diarrhoea
- Joint pain, joint swelling, muscle pain, muscle spasms and back pain
- Blood test results that show your liver function can become abnormal.

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

- Anorexia (loss of appetite), weight gain, decreases in blood sugar levels (if you have diabetes you should continue careful monitoring of your blood sugar levels)
- Having nightmares, insomnia
- Dizziness, numbness or tingling in the fingers and toes, reductions of sensation to pain or touch, change in sense of taste, loss of memory
- Blurred vision
- Ringing in the ears and/or head
- Vomiting, belching, abdominal pain upper and lower, pancreatitis (inflammation of the pancreas leading to stomach pain)
- Hepatitis (liver inflammation)
- Rash, skin rash and itching, hives, hair loss
- Neck pain, muscle fatigue
- Fatigue, feeling unwell, weakness, chest pain, swelling especially in the ankles (oedema), raised temperature
- Urine tests that are positive for white blood cells.

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

- Visual disturbance
- Unexpected bleeding or bruising
- Cholestasis (yellowing of the skin and whites of the eyes)
- Tendon injury
- Rash that may occur on the skin or sores in the mouth (lichenoid drug reaction)

- Purple skin lesions (signs of blood vessel inflammation, vasculitis).

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

- An allergic reaction - symptoms may include sudden wheezing and chest pain or tightness, swelling of the eyelids, face, lips, mouth, tongue or throat, difficulty breathing, collapse
- Hearing loss
- Gynaecomastia (breast enlargement in men).

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

- Muscle weakness that is constant
- Myasthenia gravis (a disease causing general muscle weakness including in some cases muscles used when breathing)
- Ocular myasthenia (a disease-causing eye muscle weakness).

Talk to your doctor if you experience weakness in your arms or legs that worsens after periods of activity, double vision or drooping of your eyelids, difficulty swallowing, or shortness of breath.

Possible side effects reported with some statins (medicines of the same type):

- Sexual difficulties
- Depression
- Breathing problems including persistent cough and/or shortness of breath or fever
- Diabetes. This is more likely if you have high levels of sugars and fats in your blood, are overweight and have high blood pressure. Your doctor will monitor you while you are taking this medicine.

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 Atorvas

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 the blister after “EXP”. The expiry date refers to the last day of that month.

This medicinal product does not require any special storage conditions.

HDPE bottles:

Shelf life after first opening of the bottle: 6 months.

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 Atorvas contains

The active substance is atorvastatin (as atorvastatin calcium trihydrate).

Each film-coated tablet contains 10 mg of atorvastatin (as atorvastatin calcium trihydrate).

Each film-coated tablet contains 20 mg of atorvastatin (as atorvastatin calcium trihydrate).

Each film-coated tablet contains 40 mg of atorvastatin (as atorvastatin calcium trihydrate).

Each film-coated tablet contains 80 mg of atorvastatin (as atorvastatin calcium trihydrate).

The other ingredients are microcrystalline cellulose, calcium carbonate, lactose (as monohydrate), croscarmellose sodium, hydroxypropylcellulose, polysorbate 80, magnesium stearate, hypromellose, macrogol 6000, titanium dioxide (E171).

What Atorvas looks like and contents of the pack

10 mg film-coated tablets:

White to off-white, round, biconvex film-coated tablet with debossment "A10" on one side, plain on the other side.

20 mg film-coated tablets:

White to off-white, round, biconvex film-coated tablet with debossment "A20" on one side and breaking notch on the other side.

40 mg film-coated tablets:

White to off-white, round, biconvex film-coated tablet with debossment "A40" on one side and cross scored breaking notch on the other side.

80 mg film-coated tablets:

White to off-white, round, biconvex film-coated tablet with debossment "A80" on one side and cross scored breaking notch on the other side.

The film-coated tablets are packed in aluminium/aluminium blisters or PVC/PE/PVDC/aluminium blisters and inserted in a carton or packed in HDPE bottles closed with a cap with desiccant.

Pack sizes:

aluminium/aluminium blister:

10 mg film-coated tablets: 7, 14, 28, 30, 56, 60, 63, 90, 91, 100

20 mg film-coated tablets: 7, 14, 28, 30, 56, 60, 63, 90, 91, 100

40 mg film-coated tablets: 7, 14, 28, 30, 56, 60, 63, 90, 91, 100

PVC/PE/PVDC/aluminium blister:

10 mg film-coated tablets: 7, 14, 28, 30, 56, 60, 63, 90, 100

20 mg film-coated tablets; 7, 14, 28, 30, 56, 60, 63, 90, 100

40 mg film-coated tablets; 7, 14, 28, 30, 56, 60, 63, 90, 100

80 mg film-coated tablets; 30, 50, 60, 90, 100

Bottles:

100, 250 film-coated tablets

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 57, 1526 Ljubljana, Slovenia.

Lek S.A., 16 Podlipie Street, 95-010 Stryków, Poland.
Lek Pharmaceuticals d.d., Trimlini 2D, 9220 Lendava, Slovenia.

This medicine is authorised in the Member States of the European Economic Area and in the United Kingdom (Northern Ireland) under the following names:

Austria	Atorvastatin Sandoz 10 mg – Filmtabletten Atorvastatin Sandoz 20 mg – Filmtabletten Atorvastatin Sandoz 40 mg – Filmtabletten Atorvastatin Sandoz 80 mg – Filmtabletten
Czechia	TULIP NEO
Denmark	Atorvastatin Sandoz
Estonia	Tulip 10 mg Tulip 20 mg Tulip 40 mg Tulip 80 mg
France	Atorvastatine Sandoz 80 mg, comprimé pelliculé sécable
Germany	Atorvastatin HEXAL 80 mg Filmtabletten
Greece	Rotacor 10 mg επικαλυμμένα με λεπτό υμένιο δισκία Rotacor 20 mg επικαλυμμένα με λεπτό υμένιο δισκία Rotacor 40 mg επικαλυμμένα με λεπτό υμένιο δισκία
Ireland	Atorvas 10 mg film-coated tablets Atorvas 20 mg film-coated tablets Atorvas 40 mg film-coated tablets Atorvas 80 mg film-coated tablets
Italy	ATORVASTATINA SANDOZ GMBH
Norway	Atorvastatin Sandoz 80 mg tablett, filmdrasjerte
Poland	Tulip 40 mg Tulip 80 mg
Portugal	Atorvastatina Sandoz 10 mg Atorvastatina Sandoz 20 mg Atorvastatina Sandoz 40 mg Atorvastatina Sandoz 80 mg
Slovakia	Tulip 10 mg filmom obalené tablety Tulip 20 mg filmom obalené tablety Tulip 40 mg filmom obalené tablety Tulip 80 mg filmom obalené tablety
Slovenia	Tulip 10 mg filmsko obložene tablete Tulip 20 mg filmsko obložene tablete Tulip 40 mg filmsko obložene tablete Tulip 80 mg filmsko obložene tablete
Spain	Atorvastatina Placasod 10 mg comprimidos recubiertos con película EFG Atorvastatina Placasod 20 mg comprimidos recubiertos con película EFG Atorvastatina Placasod 40 mg comprimidos recubiertos con película EFG Atorvastatina Placasod 80 mg comprimidos recubiertos con película EFG
Sweden	Atorbir 80 mg Filmdragerad tablett
United Kingdom (Northern Ireland)	Atorvastatin 80 mg film-coated Tablets

This leaflet was last approved in 09/2024.