

Package leaflet: Information for the user

Olmesartanmedoxomil/Amlodipine STADA 20 mg/5 mg filmomhulde tabletten
Olmesartanmedoxomil/Amlodipine STADA 40 mg/5 mg filmomhulde tabletten
Olmesartanmedoxomil/Amlodipine STADA 40 mg/10 mg filmomhulde tabletten

Olmesartan medoxomil / Amlodipine besilate

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

1. What Olmesartanmedoxomil/Amlodipine STADA is and what it is used for

Olmesartanmedoxomil/Amlodipine STADA contains two substances called olmesartan medoxomil and amlodipine (as amlodipine besilate). Both of these substances help to control high blood pressure.

- *olmesartan medoxomil* belongs to a group of medicines called “angiotensin-II receptor antagonists” which lower blood pressure by relaxing the blood vessels
- *amlodipine* belongs to a group of substances called “calcium channel blockers”. Amlodipine stops calcium from moving into the blood vessel wall which stops them from tightening thereby also reducing blood pressure

The actions of both these substances contribute to stopping the tightening of blood vessels, so that they relax and blood pressure decreases.

Olmesartanmedoxomil/Amlodipine STADA is used for the treatment of high blood pressure in patients whose blood pressure is not controlled enough with either olmesartan medoxomil or amlodipine alone.

2. What you need to know before you take Olmesartanmedoxomil/Amlodipine STADA

DO NOT take Olmesartanmedoxomil/Amlodipine STADA:

- if you are allergic to olmesartan medoxomil or to amlodipine or a special group of calcium channel blockers, the dihydropyridines, or to any of the other ingredients of this medicine (listed in section 6)
- if you think you may be allergic, talk to your doctor before taking Olmesartanmedoxomil/Amlodipine STADA
- if you are more than 3 months pregnant (it is also better to avoid Olmesartanmedoxomil/Amlodipine STADA in early pregnancy - see section “Pregnancy and breast-feeding”)

- if you have diabetes or impaired kidney function and you are treated with a blood pressure lowering medicine containing aliskiren
- if you have severe liver problems, if bile secretion is impaired or drainage of bile from the gallbladder is blocked (e.g. by gallstones), or if you are experiencing any jaundice (yellowing of the skin and eyes)
- if you have very low blood pressure
- if you are suffering from insufficient blood supply to your tissues with symptoms like low blood pressure, low pulse, fast heartbeat. These are signs of shock including shock due to severe heart troubles (cardiogenic shock)
- if the blood flow from your heart is obstructed (e.g. because of the narrowing of the aorta (aortic stenosis))
- if you suffer from low heart output (resulting in shortness of breath or peripheral swellings) after a heart attack (acute myocardial infarction)

Warnings and precautions

Talk to your doctor or pharmacist before taking Olmesartanmedoxomil/Amlodipine STADA.

Tell your doctor if you are taking any of the following medicines used to treat high blood pressure:

- an ACE-inhibitor (for example enalapril, lisinopril, ramipril), in particular if you have diabetes-related kidney problems
- aliskiren

Your doctor may check your kidney function, blood pressure, and the amount of electrolytes (e.g. potassium) in your blood at regular intervals.

See also information under the heading “DO NOT take Olmesartanmedoxomil/Amlodipine STADA”.

Tell your doctor if you have any of the following health problems:

- kidney problems or a kidney transplant
- liver disease
- heart failure or problems with your heart valves or heart muscle
- severe vomiting, diarrhoea, treatment with high doses of “water tablets” (diuretics) or if you are on a low salt diet
- increased levels of potassium in your blood
- problems with your adrenal glands (hormone-producing glands on top of the kidneys)

Contact your doctor if you experience diarrhoea that is severe, persistent and causes substantial weight loss. Your doctor may evaluate your symptoms and decide on how to continue your blood pressure medication.

As with any medicine which reduces blood pressure, an excessive drop in blood pressure in patients with blood flow disturbances of the heart or brain could lead to a heart attack or stroke. Your doctor will therefore check your blood pressure carefully.

You must tell your doctor if you think that you are (or might become) pregnant. Olmesartanmedoxomil/Amlodipine STADA is not recommended in early pregnancy, and must not be taken if you are more than 3 months pregnant, as it may cause serious harm to your baby if used at that stage (see section “Pregnancy and breast-feeding”).

Children and adolescents

Olmesartanmedoxomil/Amlodipine STADA is not recommended for children and adolescents under the age of 18.

Other medicines and Olmesartanmedoxomil/Amlodipine STADA

Tell your doctor or pharmacist if you are taking, have recently taken or might take any of the following medicines, especially:

- other blood pressure lowering medicines, as the effect of Olmesartanmedoxomil/Amlodipine STADA can be increased. Your doctor may need to change your dose and/or to take other precautions:
If you are taking an ACE-inhibitor or aliskiren (see also information under the headings “DO NOT take Olmesartanmedoxomil/Amlodipine STADA” and “Warnings and precautions”)
- potassium supplements, salt substitutes containing potassium, “water tablets” (diuretics) or heparin (for thinning the blood and prevention of blood clots). Using these medicines at the same time as Olmesartanmedoxomil/Amlodipine STADA may raise the levels of potassium in your blood
- lithium (a medicine used to treat mood swings and some types of depression) used at the same time as Olmesartanmedoxomil/Amlodipine STADA may increase the toxicity of lithium. If you have to take lithium, your doctor will measure your lithium blood levels
- Non-Steroidal Anti-Inflammatory Drugs (NSAIDs, medicines used to relieve pain, swelling and other symptoms of inflammation, including arthritis) used at the same time as Olmesartanmedoxomil/Amlodipine STADA may increase the risk of kidney failure. The effect of Olmesartanmedoxomil/Amlodipine STADA can be decreased by NSAIDs
- colessevelam hydrochloride, a drug that lowers the level of cholesterol in your blood, as the effect of Olmesartanmedoxomil/Amlodipine STADA may be decreased. Your doctor may advise you to take Olmesartanmedoxomil/Amlodipine STADA at least 4 hours before colessevelam hydrochloride
- certain antacids (indigestion or heartburn remedies), as the effect of Olmesartanmedoxomil/Amlodipine STADA can be slightly decreased
- medicines used for HIV/AIDS (e.g. ritonavir, indinavir, nelfinavir) or for the treatment of fungal infections (e.g. ketoconazole, itraconazole)
- diltiazem, verapamil, (agents used for heart rhythm problems and high blood pressure)
- rifampicin, erythromycin, clarithromycin (antibiotics – used for the treatment of infections caused by bacteria)
- St. John’s wort (*Hypericum perforatum*), a herbal remedy
- dantrolene (infusion for severe body temperature abnormalities)
- simvastatin, an agent used to lower levels of cholesterol and fats (triglycerides) in the blood
- tacrolimus, cyclosporine (used to control your body’s immune response, enabling your body to accept the transplanted organ)

Olmesartanmedoxomil/Amlodipine STADA with food and drink

Olmesartanmedoxomil/Amlodipine STADA can be taken with or without food. Swallow the tablet with some fluid (such as one glass of water). If possible, take your daily dose at the same time each day, for example at breakfast time.

Grapefruit juice and grapefruit should not be consumed by people who are taking Olmesartanmedoxomil/Amlodipine STADA. This is because grapefruit and grapefruit juice can lead to an increase in the blood levels of the active ingredient amlodipine, which can cause an unpredictable increase in the blood pressure lowering effect of Olmesartanmedoxomil/Amlodipine STADA.

Elderly

If you are over 65 years of age, your doctor will regularly check your blood pressure at any dose increase, to make sure that your blood pressure does not become too low.

Black patients

As with other similar drugs the blood pressure lowering effect of Olmesartanmedoxomil/Amlodipine STADA can be somewhat less in black patients.

Pregnancy and breastfeeding

Pregnancy

You must tell your doctor if you think that you are (or might become) pregnant. Your doctor will normally advise you to stop taking Olmesartanmedoxomil/Amlodipine STADA before you become pregnant or as soon as you know you are pregnant and will advise you to take another medicine instead of Olmesartanmedoxomil/Amlodipine STADA. Olmesartanmedoxomil/Amlodipine STADA is not recommended in early pregnancy, and must not be taken when more than 3 months pregnant, as it may cause serious harm to your baby if used after the third month of pregnancy.

If you become pregnant during therapy with Olmesartanmedoxomil/Amlodipine STADA, please inform and see your doctor without delay.

Breast-feeding

Amlodipine, one of the two active ingredients in <Product name> has been shown to pass into breast milk in small amounts. Tell your doctor if you are breast-feeding or about to start breast-feeding. Olmesartanmedoxomil/Amlodipine STADA is not recommended for mothers who are breast-feeding, and your doctor may choose another treatment for you if you wish to breast-feed, especially if your baby is newborn, or was born prematurely.

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

You may feel sleepy, sick or dizzy or get a headache while being treated for your high blood pressure. If this happens, do not drive or use machines until the symptoms wear off. Ask your doctor for advice.

Olmesartanmedoxomil/Amlodipine STADA contains lactose and sorbitol (E420)

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

This medicine contains 0.45 mg sorbitol per tablet.

This medicine contains 0.90 mg sorbitol per tablet.

This medicine contains 0.90 mg sorbitol per tablet.

3. How to take Olmesartanmedoxomil/Amlodipine STADA

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

Dosage

The recommended dose of Olmesartanmedoxomil/Amlodipine STADA is one tablet per day.

Method of administration

- the tablets can be taken with or without food. Swallow the tablet with some fluid (such as a glass of water). The tablet should not be chewed. Do not take them with grapefruit juice
- if possible, take your daily dose at the same time each day, for example at breakfast time

If you take more Olmesartanmedoxomil/Amlodipine STADA than you should

If you take more tablets than you should you may experience low blood pressure with symptoms such as dizziness, fast or slow heartbeat.

If you take more tablets than you should or if a child accidentally swallows some, go to your doctor or nearest emergency department immediately and take your medicine pack or this leaflet with you.

If you forget to take Olmesartanmedoxomil/Amlodipine STADA

If you forget to take a dose, take your normal dose on the following day as usual. Do not take a double dose to make up for a forgotten dose.

If you stop taking Olmesartanmedoxomil/Amlodipine STADA

It is important to continue to take Olmesartanmedoxomil/Amlodipine STADA unless your doctor tells you to stop.

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. If they do occur, they are often mild and do not require treatment to be stopped.

Although not many people may get them, the following side effects can be serious:

Allergic reactions (itching, rash, swelling of the face, mouth and/or larynx (voice box) together with itching and rash, severe skin reactions including intense skin rash, hives, reddening of the skin over your whole body, severe itching, blistering, peeling and swelling of the skin, inflammation of mucous membranes (Stevens-Johnson syndrome, toxic epidermal necrolysis), some life-threatening) may occur during treatment with Olmesartanmedoxomil/Amlodipine STADA. If this happens stop taking Olmesartanmedoxomil/Amlodipine STADA and talk to your doctor immediately.

Olmesartanmedoxomil/Amlodipine STADA can cause the blood pressure to fall too low in susceptible individuals or as the result of an allergic reaction. This could cause severe light-headedness or fainting. If this happens stop taking Olmesartanmedoxomil/Amlodipine STADA, talk to your doctor immediately and lie down flat.

Frequency not known: If you experience yellowing of the whites of the eyes, dark urine, itching of the skin, even if you started therapy with <Product name> longer time ago, **contact your doctor immediately** who will evaluate your symptoms and decide on how to continue your blood pressure medication.

Other possible side effects with Olmesartanmedoxomil/Amlodipine STADA:

Common (may affect up to 1 in 10 people)

- dizziness
- headache
- swelling of ankles, feet, legs, hands, or arms
- tiredness

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

- dizziness on standing up
- lack of energy
- tingling or numbness of hands or feet
- vertigo
- awareness heart beat
- fast heart beat
- low blood pressure with symptoms such as dizziness, light-headedness
- difficult breathing
- cough
- nausea
- vomiting

- indigestion
- diarrhoea
- constipation
- dry mouth
- upper abdominal pain
- skin rash
- cramps
- pain in arms and legs
- back pain
- feeling more of an urge to pass urine
- sexual inactivity
- inability to get or maintain an erection
- weakness

Some changes in blood test results have also been seen and include the following: Increased as well as decreased blood potassium levels, increased blood creatinine levels, increased uric acid levels, increases in a test of liver function (gamma glutamyl transferase levels).

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

- drug hypersensitivity
- fainting
- redness and warm feeling of the face
- red itchy bumps (hives)
- swelling of face

Side effects reported with use of olmesartan medoxomil or amlodipine alone, but not with Olmesartanmedoxomil/Amlodipine STADA or in a higher frequency:

Olmesartan medoxomil

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

- bronchitis
- sore throat
- runny or stuffy nose
- cough
- abdominal pain
- stomach flu
- diarrhoea
- indigestion
- nausea
- pain in the joints or bones
- back pain
- blood in the urine
- infection of the urinary tract
- chest pain
- flu-like symptoms
- pain
- changes in blood test results as increased fat levels (hypertriglyceridaemia), blood urea or uric acid increased and increase in tests of liver and muscle function

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

- reduced number of a type of blood cells known as platelets, which can result in easily bruising or prolonged bleeding time
- quick allergic reactions that may affect the whole body and may cause breathing problems as well as a rapid fall of blood pressure and even lead to fainting (anaphylactic reactions)

- angina (pain or uncomfortable feeling in the chest, known as angina pectoris)
- itching
- eruption of the skin
- allergic skin rash
- rash with hives
- swelling of the face
- muscular pain
- feeling unwell

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

- swelling of the face, mouth and/or larynx (voice box)
- acute kidney failure and kidney insufficiency
- lethargy

Amlodipine

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

- oedema (fluid retention)

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

- abdominal pain
- nausea
- ankle swelling
- feeling sleepy
- redness and warm feeling of the face
- visual disturbance (including double vision and blurred vision)
- awareness of heartbeat
- diarrhoea
- constipation
- indigestion
- cramps
- weakness
- difficult breathing

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

- trouble sleeping
- sleep disturbances
- mood changes including feeling anxious
- depression
- irritability
- shiver
- taste changes
- fainting
- ringing in the ears (tinnitus)
- worsening of angina pectoris (pain or uncomfortable feeling in the chest)
- irregular heartbeat
- runny or stuffy nose
- loss of hair
- purplish spots or patches on the skin due to small haemorrhages (purpura)
- discolouration of the skin
- excessive sweating
- eruption of the skin
- itching
- red itchy bumps (hives)

- pain of joints or muscles
- problems in passing urine
- urge to pass urine at night
- increased need to urinate (pass urine)
- breast enlargement in men
- chest pain
- pain
- feeling unwell
- increase or decrease in weight

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

- confusion

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

- reduction in the number of white cells in the blood, which could increase the risk of infections
- a reduction in the number of a type of blood cells known as platelets, which can result in bruising easily or prolonged bleeding time
- increase in blood glucose
- increased tightness of muscles or increased resistance to passive movement (hypertonia)
- tingling or numbness of hands or feet
- heart attack
- inflammation of blood vessels
- inflammation of the liver or the pancreas
- inflammation of stomach lining
- thickening of gums
- elevated liver enzymes
- yellowing of the skin and eyes
- increased sensitivity of the skin to light

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

- trembling, rigid posture, mask-like face, slow movements and a shuffling, unbalanced walk

Het melden van bijwerkingen

Krijgt u last van bijwerkingen, neem dan contact op met uw arts of apotheker. Dit geldt ook voor mogelijke bijwerkingen die niet in deze bijsluiter staan. U kunt bijwerkingen ook rechtstreeks melden via het Nederlands Bijwerkingen Centrum Lareb, website: www.lareb.nl. Door bijwerkingen te melden, kunt u ons helpen meer informatie te verkrijgen over de veiligheid van dit geneesmiddel.

5. How to store Olmesartanmedoxomil/Amlodipine STADA

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. The expiry date refers to the last day of that month.

Store in original package in order to protect from moisture.

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 Olmesartanmedoxomil/Amlodipine STADA contains

The active substances are olmesartan medoxomil and amlodipine (as besilate).

Each tablet contains 20 mg of olmesartan medoxomil and 5 mg amlodipine (as besilate).

Each tablet contains 40 mg of olmesartan medoxomil and 5 mg amlodipine (as besilate).

Each tablet contains 40 mg of olmesartan medoxomil and 10 mg amlodipine (as besilate).

Other ingredients are

Tablet core:

Microcrystalline cellulose; crospovidone type A; silica, colloidal anhydrous; magnesium stearate; lactose monohydrate

Tablet coat:

Olmesartanmedoxomil/Amlodipine STADA 20 mg/5 mg filmomhulde tabletten:
AquaPolish P white 919.03 STA (hydroxypropyl pregelatinized starch; stearic acid; sorbitol (E420); titanium dioxide)

Olmesartanmedoxomil/Amlodipine STADA 40 mg/5 mg filmomhulde tabletten:
AquaPolish P yellow 929.05 STA (hydroxypropyl pregelatinized starch; stearic acid; sorbitol (E420); titanium dioxide; iron oxide yellow; iron oxide red)

Olmesartanmedoxomil/Amlodipine STADA 40 mg/10 mg filmomhulde tabletten:
AquaPolish P red 949.06 STA (hydroxypropyl pregelatinized starch; stearic acid; sorbitol (E420); titanium dioxide; iron oxide red)

What Olmesartanmedoxomil/Amlodipine STADA looks like and contents of the pack

Olmesartanmedoxomil/Amlodipine STADA 20 mg/5 mg are white, round biconvex film-coated tablets packed in oPA-Alu-PVC/ Alu blister

Olmesartanmedoxomil/Amlodipine STADA 40 mg/5 mg are yellowish-white, round, biconvex film-coated tablets packed in oPA-Alu-PVC/ Alu blister

Olmesartanmedoxomil/Amlodipine STADA 40 mg/10 mg are brownish-red, round, biconvex film-coated tablets packed in oPA-Alu-PVC/ Alu blister

Olmesartanmedoxomil/Amlodipine STADA filmomhulde tabletten are available in packs of

10, 14, 28, 30, 56, 60, 90, 98, 100, 112 film-coated tablets

Not all pack sizes may be marketed.

Marketing Authorisation Holder and Manufacturer(s):

Marketing Authorisation Holder:

STADA Arzneimittel AG
Stadastrasse 2 – 18
61118 Bad Vilbel
Duitsland

Manufacturers:

STADA Arzneimittel AG
Stadastrasse 2 – 18
61118 Bad Vilbel
Duitsland

STADA Arzneimittel GmbH
Muthgasse 36/2
1190 Wenen

Oostenrijk

Clonmel Healthcare Ltd.
Waterford Road
Clonmel, Co. Tipperary
Ierland

In het register ingeschreven onder:

Olmesartanmedoxomil/Amlodipine STADA 20 mg/5 mg filmomhulde tabletten:
RVG 119876

Olmesartanmedoxomil/Amlodipine STADA 40 mg /5 mg filmomhulde tabletten:
RVG 119877

Olmesartanmedoxomil/Amlodipine STADA 40 mg/10 mg filmomhulde tabletten:
RVG 119878

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

België:	Olmesartan / Amlodipine EG 20mg/5mg; 40mg/5mg; 40mg/10mg filmomhulde tabletten
Duitsland:	Olmesartan/Amlodipin AL 20 mg/5 mg; 40 mg/5 mg; 40 mg/10 mg Filmtabletten
Ierland:	Olmesartan / Amlodipine Clonmel 20 mg/5 mg; 40 mg/5 mg; 40 mg/10 mg film-coated tablets
Italië:	Olmesartan e Amlodipina EG
Luxemburg:	Olmesartan / Amlodipine EG 20mg/5mg; 40mg/5mg; 40mg/10mg comprimé pelliculé
Portugal:	Amlodipina + Olmesartan medoxomilo Ciclum
Slowakije:	Olместad 20 mg/5 mg; 40 mg/5 mg; 40 mg/10 mg
Spanje:	Olmesartan/Amlodipino STADA 20 mg/5 mg; 40 mg/5 mg; 40 mg/10 mg comprimidos recubiertos con película EFG
Nederland:	Olmesartanmedoxomil/Amlodipine STADA 20 mg/5 mg; 40 mg/5 mg; 40 mg/10 mg filmomhulde tabletten

Deze bijsluiter is voor het laatst goedgekeurd in april 2022