

Detrunorm[®] 15mg coated tablets

(propiverine hydrochloride)

Patient Information Leaflet

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, pharmacist or nurse.
- * 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, pharmacist or nurse. This includes any possible side effects not listed in this leaflet. See section 4.

The name of your medicine is Detrunorm 15 mg Coated Tablets (referred to as Detrunorm throughout this leaflet).

What is in this leaflet:

- 1 What Detrunorm are and what they are used for
- 2 What you need to know before you take Detrunorm
 - How to take Detrunorm
- 3 Possible side effects
- 4 How to store Detrunorm
- 5 Contents of the pack and other information
- 6

1 What Detrunorm are and what they are used for

Detrunorm is used for the treatment of people who have difficulty in controlling their bladders due to bladder overactivity or, in some cases, problems with the spinal cord. Detrunorm contains the active substance propiverine hydrochloride.

This substance prevents the bladder from contracting and increases the amount that the bladder can hold. Detrunorm is used to treat the symptoms of overactive bladder.

2 What you need to know before you take Detrunorm

Do not take Detrunorm:

- * if you are allergic to propiverine hydrochloride or any of the other ingredients of this medicine listed in section 6 (allergic reactions include mild symptoms such as itching and/or rash. More severe symptoms include swelling of the face, lips, tongue and/or throat with difficulty in swallowing or breathing);
- * if you have obstruction of the bowel;
- * if you have obstruction to the bladder outlet (difficulty in passing urine);
- * if you have myasthenia gravis (a disease causing muscle weakness);
- * if you have a loss of function of the muscles controlling your bowel movements (intestinal atony);
- * if you have severe inflammation of the bowel (ulcerative colitis) that may lead to diarrhoea containing blood and mucus and abdominal pains;
- * if you have toxic megacolon (a condition involving enlargement of the bowel);
- * if you have increased pressure in the eye (uncontrolled angle closure glaucoma);
- * if you have moderate or severe liver disease;
- * if you have fast or irregular heartbeat.

Warnings and precautions

Talk to your doctor, pharmacist or nurse before taking Detrunorm

- * if you have damage to the nerves that control blood pressure, heart rate, bowel and bladder movement and other bodily functions (autonomic neuropathy);
- * if you have liver problems;
- * if you have kidney problems;

- * if you have severe heart failure;
- * if you have enlargement of the prostate gland;
- * if you have recurrent urinary tract infection;
- * if you have tumours of the urinary tract;
- * if you have glaucoma;
- * if you have heartburn and indigestion due to back flow of gastric juice into the throat (hiatus hernia with reflux oesophagitis);
- * if you have irregular heartbeat;
- * if you have fast heartbeat.

Other medicines and Detrunorm

Tell your doctor or pharmacist if you are taking, have recently taken or might

take any of the following medicines as they may interact with Detrunorm:

- * antidepressants (e.g. imipramine, clomipramine, amitriptyline);
- * sleeping tablets (e.g. benzodiazepines);
- * anticholinergics taken by mouth or injection (usually used to treat asthma, stomach cramps, eye problems or urinary incontinence);
- * amantadine (used to treat flu and Parkinson's disease);
- * neuroleptics such as promazine, olanzapine, quetiapine (drugs used to treat psychotic disorders like schizophrenia and anxiety);
- * beta stimulants (drugs used to treat asthma);
- * cholinergics (e.g. carbachol, pilocarpin);
- * isoniazid (a treatment for tuberculosis);
- * metoclopramide (used to treat nausea and vomiting);
- * concomitant treatment with methimazole (used to treat hyperfunction of the thyroid gland) and medicines used to treat fungal diseases (e.g. ketoconazole, itraconazole).

Nevertheless, it may still be all right for you to take Detrunorm. Your doctor will be able to decide what is suitable for you.

Please tell your doctor or pharmacist if you are taking or have recently taken any other medicines, including medicines obtained without a prescription.

Taking Detrunorm with food and drink

The tablets should be swallowed before meals.

Pregnancy, breast-feeding and fertility

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.

Do not take Detrunorm if you are pregnant, likely to become pregnant or are breast-feeding.

Driving and using machines

Detrunorm can sometimes cause sleepiness and blurred vision. You should not drive or operate machinery until you are sure you are not affected.

Detrunorm contains glucose, lactose and sucrose

Detrunorm contains glucose, lactose and sucrose (sugars). If you have been told by your doctor that you have an intolerance to some sugars, contact your doctor before taking this medicine.

Detrunorm also contain the colouring agent cochineal red A (E124)

May cause allergic reactions.

3 How to take Detrunorm

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 recommended dose is

Adults and the elderly: the recommended dose of Detrunorm is two to three tablets daily.

Use in children and adolescents

Detrunorm is not recommended for children.

Detrunorm[®] 15mg coated tablets

(propiverine hydrochloride)

Patient Information Leaflet (continued)

Method of administration

Take your tablets at the same time each day. Swallow your tablets whole before meals.

If you take more Detrunorm than you should

If you accidentally take more than your prescribed dose, contact your nearest casualty department or tell your doctor or pharmacist immediately. Remember to take the pack and any remaining tablets with you. Overdosage can cause symptoms such as restlessness, dizziness, disorders in speech and vision, muscular weakness, dry mouth, faster heartbeat and problems passing urine.

If you forget to take Detrunorm

Do not worry. Simply leave out that dose completely. Then take your next dose at the right time. Do not take a double dose to make up for a forgotten dose.

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

4 Possible side effects

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

All medicines can cause allergic reactions although serious allergic reactions are very rare. The following symptoms are first signs for such reactions:

- * Any sudden wheeziness, difficulty in breathing or dizziness, swelling of the eyelids, face, lips or throat;
- * Peeling and blistering of the skin, mouth, eyes and genitals;
- * Rash affecting your whole body.

If you get any of these symptoms during treatment, you should stop taking the tablets and contact your doctor **immediately**.

You might suffer an acute attack of glaucoma. If you have been seeing coloured rings around lights or if you should develop severe pain in and around either eye **you should seek medical attention urgently**.

The following side effects have been reported:

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

Dry mouth.

Common (may affect up to 1 in 10 people)

Abnormal vision and difficulty in focussing, fatigue, headache, abdominal pain, indigestion, constipation.

Uncommon (may affect up to 1 in 100 people)

Feeling sick and vomiting, dizziness, trembling (tremor), inability to empty the bladder, difficulty in passing urine (urinary retention), flushing, altered sense of taste, decreased blood pressure with drowsiness, itching.

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

Rash;
Faster heart beat.

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

Feeling your heartbeat, restlessness and confusion.

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

Sensing things that are not real (hallucinations);
Speech disorder.

All possible side effects are transient and recede after a dose reduction or termination of the therapy after maximum 1-4 days.

During long-term therapy hepatic enzymes should be monitored, because reversible changes of liver enzymes might occur in rare cases.

Reporting of side effects

If you get any side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed on 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 Detrunorm

- * **Keep out of the sight and reach of children.**
- * For oral use.
- * Do not take Detrunorm after the expiry date which is stated on the blister label or carton. The expiry date refers to the last day of the month.
- * If your doctor tells you to stop taking this medicine, return any unused tablets to your pharmacist (chemist) for safe disposal. Only keep this medicine if your doctor tells you to.
- * If the medicine become discoloured or show any other signs of deterioration, you should seek the advice of your pharmacist who will tell you what to do.
- * Medicines should not be disposed of via wastewater or household waste. Ask your pharmacist how to dispose of medicines that are no longer required. These measures will help to protect the environment.

6 Contents of the pack and other information

What Detrunorm contains:

Each coated tablet contains 15 mg propiverine hydrochloride (equivalent to 13.64 mg propiverine). Also contains lactose monohydrate, cellulose powder, magnesium stearate, sucrose, talc, kaolin, calcium carbonate, titanium dioxide (E171), arabic gum, colloidal anhydrous silica, macrogol 6000, glucose monohydrate, Ponceau R (E124), montan glycol wax.

What Detrunorm looks like and contents of the pack

Detrunorm are rose-coloured, biconvex, round sugar-coated tablets. Each pack contains 60 Tablets.

Manufacturer and Licence Holder

Manufactured by APOGEPHA Arzneimittel GmbH, Kyffhäuserstraße 27, 01309 Dresden, Germany and is procured from within the EU and repackaged by the Product Licence Holder: Lexon (UK) Limited, Unit 18, Oxleasow Road, East Moons Moat, Redditch, Worcestershire, B98 0RE.

If you have any questions or are not sure about anything, ask your doctor or pharmacist. They will have additional information about this medicine and will be able to advise you.

POM

PL 15184/1755

Detrunorm 15mg coated tablets

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Revision date: 02/07/19

**Blind or partially sighted?
Is this leaflet hard to see or read?
Phone Lexon (UK) Limited,
Tel: 01527 505414 to obtain the leaflet
in a format suitable for you**

Propiverine Hydrochloride 5mg coated tablets

Patient Information Leaflet

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- * 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, pharmacist or nurse. This includes any possible side effects not listed in this leaflet. See section 4.

The name of your medicine is Propiverine 15 mg Coated Tablets (referred to as Propiverine throughout this leaflet).

What is in this leaflet:

- 1 What Propiverine are and what they are used for
- 2 What you need to know before you take Propiverine
- 3 How to take Propiverine
- 4 Possible side effects
- 5 How to store Propiverine
- 6 Contents of the pack and other information

1 What Propiverine are and what they are used for

Propiverine is used for the treatment of people who have difficulty in controlling their bladders due to bladder overactivity or, in some cases, problems with the spinal cord. Propiverine contains the active substance propiverine hydrochloride. This substance prevents the bladder from contracting and increases the amount that the bladder can hold. Propiverine is used to treat the symptoms of overactive bladder.

2 What you need to know before you take Propiverine

Do not take Propiverine:

- * if you are allergic to propiverine hydrochloride or any of the other ingredients of this medicine listed in section 6 (allergic reactions include mild symptoms such as itching and/or rash. More severe symptoms include swelling of the face, lips, tongue and/or throat with difficulty in swallowing or breathing);
- * if you have obstruction of the bowel;
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- * if you have myasthenia gravis (a disease causing muscle weakness);
- * if you have a loss of function of the muscles controlling your bowel movements (intestinal atony);
- * if you have severe inflammation of the bowel (ulcerative colitis) that may lead to diarrhoea containing blood and mucus and abdominal pains;
- * if you have toxic megacolon (a condition involving enlargement of the bowel);
- * if you have increased pressure in the eye (uncontrolled angle closure glaucoma);
- * if you have moderate or severe liver disease;
- * if you have fast or irregular heartbeat.

Warnings and precautions

- Talk to your doctor, pharmacist or nurse before taking Propiverine
- * if you have damage to the nerves that control blood pressure, heart rate, bowel and bladder movement and other bodily functions (autonomic neuropathy);
 - * if you have liver problems;
 - * if you have kidney problems;
 - * if you have severe heart failure;
 - * if you have enlargement of the prostate gland;
 - * if you have recurrent urinary tract infection;
 - * if you have tumours of the urinary tract;

- * if you have glaucoma;
- * if you have heartburn and indigestion due to back flow of gastric juice into the throat (hiatus hernia with reflux oesophagitis);
- * if you have irregular heartbeat;
- * if you have fast heartbeat.

Other medicines and Propiverine

Tell your doctor or pharmacist if you are taking, have recently taken or might take any of the following medicines as they may interact with Propiverine:

- * antidepressants (e.g. imipramine, clomipramine, amitriptyline);
- * sleeping tablets (e.g. benzodiazepines);
- * anticholinergics taken by mouth or injection (usually used to treat asthma, stomach cramps, eye problems or urinary incontinence);
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Nevertheless, it may still be all right for you to take Propiverine. Your doctor will be able to decide what is suitable for you.

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Taking Propiverine with food and drink

The tablets should be swallowed before meals.

Pregnancy, breast-feeding and fertility

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.

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Driving and using machines

Propiverine can sometimes cause sleepiness and blurred vision. You should not drive or operate machinery until you are sure you are not affected.

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Propiverine also contain the colouring agent cochineal red A (E124)

May cause allergic reactions.

3 How to take Propiverine

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The recommended dose is

Adults and the elderly: the recommended dose of Propiverine is two to three tablets daily.

Use in children and adolescents

Propiverine is not recommended for children.

Method of administration

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Propiverine Hydrochloride 5mg coated tablets

Patient Information Leaflet (continued)

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