Proxor 100/6 micrograms per actuation pressurised inhalation, solution

beclometasone dipropionate/ formoterol fumarate dihydrate

For use in adults

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 with your doctor, pharmacist or nurse. This includes any possible side effects not listed in this leaflet. See section 4.
- What is in this leaflet:
- 1. What Proxor is and what it is used for
- 2. What you need to know before you use Proxor 3. How to use Proxor
- 4. Possible side effects
- 5. How to store Proxor
- 6. Contents of the pack and other information 1. What Proxor is and what it is used for

Proxor is a pressurised inhalation solution containing two

active substances which are inhaled through your mouth and delivered directly into your lungs. The two active substances are beclometasone dipropionate

and formoterol fumarate dihydrate. Beclometasone dipropionate belongs to a group of medicines called corticosteroids which have an anti-inflammatory action

reducing the swelling and irritation in your lungs. Formoterol fumarate dihydrate belongs to a group of medicines called long-acting bronchodilators which relax the muscles in

your airways and helps you to breathe more easily. Together these two active substances make breathing easier, by providing relief from symptoms such as shortness of

and also help to prevent the symptoms of asthma. Proxor is indicated in the regular treatment of asthma in adult patients in whom: asthma is not adequately controlled by using inhaled

breath, wheezing and cough in patients with asthma or COPD

needed" corticosteroids "as short-acting bronchodilators or

asthma is responding well to treatment with both

corticosteroids and long-acting bronchodilators. <u>COPD</u> Proxor can also be used to treat the symptoms of severe

chronic obstructive pulmonary disease (COPD) in adult patients. COPD is a long term disease of the airways in the

lungs which is primarily caused by cigarette smoking. 2. What you need to know before you use Proxor Do not use Proxor: if you are allergic to beclometasone dipropionate or

formoterol fumarate dihydrate or any of the other ingredients of this medicine (listed in section 6).

Warnings and precautions Talk to your doctor, pharmacist or nurse before using Proxor: If you have any heart problems, such as angina (heart

- pain, pain in the chest), a recent heart attack (myocardial infarction), heart failure, narrowing of the arteries around your heart (coronary heart disease), valvular heart disease or any other known abnormalities of your heart or if you have a condition known as hypertrophic obstructive cardiomyopathy (also known as HOCM, a condition where the heart muscle is abnormal). • If you have narrowing of the arteries (also known as arteriosclerosis), if you have high blood pressure or if you
- know that you have an aneurysm (an abnormal bulging of the blood vessel wall). If you have disorders of your heart rhythm such as increased or irregular heart rate, a fast pulse rate or palpitations or if
- you have been told that your heart trace is abnormal. If you have an overactive thyroid gland. · If you have low blood levels of potassium.
- If you have any disease of your liver or kidneys.
- If you have diabetes (if you inhale high doses of formoterol your blood glucose may increase and therefore you may
- need to have some additional blood tests to check your blood sugar when you start using this inhaler and from time to time during treatment) • If you have a tumour of the adrenal gland (known as a phaeochromocytoma). • If you are due to have an anaesthetic. Depending on the
- type of anaesthetic, it may be necessary to stop taking Proxor at least 12 hours before the anaesthesia.
- If you are being, or have ever been, treated for tuberculosis (TB) or if you have a known viral or fungal infection of your
- If you must avoid alcohol for any reason. If any of the above applies to you, always inform your

If you have or had any medical problems or any allergies or if you are not sure as to whether you can use Proxor talk to

doctor before you use Proxor.

(hypokalaemia).

your doctor, asthma nurse or pharmacist before using this Treatment with a beta-2-agonist like the formoterol contained in Proxor can cause a sharp fall in your serum potassium level

If you have severe asthma, you should take special care. This is because a lack of oxygen in the blood and some other treatments you may be taking together with Proxor, such as medicines for treating heart disease or high blood pressure,

known as diuretics or "water tablets" or other medicines used

to treat asthma can make the fall in potassium level worse. For this reason, your doctor may wish to measure the potassium levels in your blood from time to time. If you take higher doses of inhaled corticosteroids over long periods, you may have more of a need for corticosteroids in situations of stress. Stressful situations might include being taken to hospital after an accident, having a serious injury or before an operation. In this case, the doctor treating you

corticosteroids and may prescribe some steroid tablets or a steroid injection. Should you need to go to the hospital, remember to take all of your medicines and inhalers with you, including Proxor and any medicines or tablets bought without a prescription, in their original packaging, if possible. Contact your doctor if you experience blurred vision or other visual disturbances.

will decide whether you may need to increase your dose of

Proxor should not be used in children and adolescent less than 18 years old, until further data becomes available. Other medicines and Proxor: Tell your doctor if you are taking, have recently taken or

might take any other medicines including medicines obtained

without a prescription. Some medicines may increase the effects of Proxor and vour

Children and adolescents

doctor may wish to monitor you carefully if you are taking these medicines (including some medicines for HIV: ritonavir,

Do not use beta blockers with this medicine. Beta blockers such as atenolol, propranolol and sotalol are used to treat a number of conditions including high blood pressure and

failure; timolol is used to treat glaucoma. If you need to use beta blockers, and including beta blockers in eye-drops, the effect of formoterol may be reduced or formoterol may not work at all. On the other hand, using other beta adrenergic medicinal products (medicines which work in the same way as formoterol) may increase the effects of formoterol. **Using Proxor together with:**

heart conditions such as abnormal heart rhythms and heart

- medicines for treating abnormal heart rhythms (quinidine, disopyramide, procainamide), medicines used to treat allergic reactions (antihistamines), medicines for treating symptoms of depression or mental disorders such as monoaminoxidase inhibitors (for example phenelzine and isocarboxazid), tricyclic antidepressants (for example amitryptiline and imipramine), phenothiazines can cause some changes in the electrocardiogram (ECG, heart trace). They may also increase the risk of disturbances of heart rhythm (ventricular arrhythmias).
- medicines for treating Parkinson's Disease (L-dopa), to treat an underactive thyroid gland (L-thyroxine), medicines containing oxytocin (which causes uterine contraction) and alcohol can lower your heart's tolerance to beta-2 agonists, such as formoterol. · monoaminoxidase inhibitors (MAOIs), including medicines
- with similar properties like furazolidone and procarbazine, used to treat mental disorders, can cause a rise in blood • medicines for treating heart disease (digoxin) can cause

a fall in your blood potassium level. This may increase the

• other medicines used to treat asthma (theophylline, aminophylline or steroids) and diuretics (water tablets) may

likelihood of abnormal heart rhythms.

cause a fall in your potassium level.

· Some anaesthetics can increase the risk of abnormal heart rhythms Pregnancy, breast- feeding and fertility

There are no clinical data on the use of Proxor during pregnancy. Proxor must not be used if you are pregnant, think that you

might be pregnant or are planning to become pregnant, or if you are breast-feeding, unless you are advised to do so by your doctor. **Driving and using machines**

Proxor is unlikely to affect your ability to drive and use

machines. However, if you experience side effects such as dizziness and/or trembling, your ability to drive or operate machinery may be affected. Proxor contains alcohol

This medicine contains 7 mg of alcohol (ethanol) in each

actuation which is equivalent to 0.19 mg/kg per dose of two actuations. The amount in of this medicine is equivalent to less than 1 ml beer or 1 ml wine. The small amount of alcohol in this medicine will not have any noticeable effects. 3. How to use Proxor

Proxor is for inhalation use. Proxor should be inhaled via your mouth into your lungs. Always use this medicine

exactly as your doctor or pharmacist has told you. Check with your doctor or pharmacist if you are not sure. The pharmacist's label will tell you how many puffs to take and how often they must be taken. <u>Dose</u> **Asthma** Your doctor will give you a regular check-up to make sure

you are taking the optimal dose of Proxor. Your doctor will adjust your treatment to the lowest dose that best controls vour symptoms Proxor can be prescribed by your doctor in two different ways:

a) Use Proxor every day to treat your asthma together with a separate "reliever" inhaler to treat sudden worsening of asthma symptoms, such as shortness of

breath, wheezing and cough Adults and the elderly: The recommended dose of this medicine is one or two puffs twice daily. The maximum daily dose is 4 puffs. Remember: You should always have your quick-acting

"reliever" inhaler with you at all times to treat worsening

symptoms of asthma or a sudden asthma attack.

b) Use Proxor every day to treat your asthma and also use Proxor to treat sudden worsening of your asthma symptoms, such as shortness of breath, wheezing and cough Adults and the elderly:

The recommended dose is one puff in the morning and one puff in the evening. You should also use Proxor as a "reliever" inhaler to treat

sudden asthma symptoms. If you get asthma symptoms, take one puff and wait a few

minutes. If you do not feel better, take another puff.

Do not take more than 6 Proxor "reliever" puffs per day. The maximum daily dose of Proxor as your only asthma

inhaler is 8 puffs. If you feel you need more puffs each day to control your asthma symptoms, contact your doctor to seek his advice. He

may need to change your treatment. Use in children and adolescents less than 18 years of age: Children and adolescents aged less than 18 years must NOT

take this medicine. Chronic obstructive pulmonary disease (COPD) Adults and the elderly:

The recommended dose is two puffs in the morning and two puffs in the evening.

At-risk patients:

Older people do not need to have their dose adjusted. No information is available regarding the use of Proxor in people with liver or kidney problems. Proxor is effective for the treatment of asthma in a dose

of beclometasone dipropionate which may be lower than

that of some other inhalers containing beclometasone dipropionate. If you have been using a different inhaler containing beclometasone dipropionate previously, your doctor will advise you on the exact dose of Proxor you should take for your asthma. Do not increase the dose If you feel that the medicine is not very effective, always talk

to your doctor before increasing the dose. If you use more Proxor than you should:

• Taking more formoterol than you should can have the following effects: feeling sick, being sick, heart racing,

palpitations, disturbances of heart rhythm, certain changes in the electrocardiogram (heart trace), headache, trembling,

- feeling sleepy, too much acid in the blood, low blood potassium levels, high levels of glucose in the blood. Your doctor may wish to carry out some blood tests to check your blood potassium and blood glucose levels. • Taking too much beclometasone dipropionate can lead to short-term problems with your adrenal glands. This will get better within a few days however your doctor may need to carry out some blood tests to check your serum cortisol levels.
- If you forget to use Proxor: Take it as soon as you remember. If it is almost time for your

Tell your doctor if you have any of these symptoms.

next dose, do not take the dose you have missed, just take the next dose at the correct time. Do not double the dose. If you stop using Proxor: Do not lower the dose or stop using the medication.

Even if you are feeling better, do not stop taking Proxor or

lower the dose. If you want to do this, talk to your doctor. It is very important for you to use Proxor regularly even though you

may have no symptoms. If your breathing gets worse: If you develop worsening shortness of breath or wheezing (breathing with an audible whistling sound), straight after

inhaling your medicine, stop using Proxor inhaler immediately and use your quick-acting "reliever" inhaler straightaway. You

Your doctor will assess your symptoms and if necessary may

If your symptoms get worse or are difficult to control (e.g. if you are using a separate "reliever" inhaler or Proxor as reliever inhaler more frequently) or if your "reliever" inhaler or Proxor does not improve your symptoms, see your doctor

immediately. Your asthma may be getting worse and your

doctor may need to increase your dose of Proxor or prescribe alternative treatment.

should contact your doctor straightaway.

See also section 4. Possible side effects

If your asthma gets worse:

start you on a different course of treatment.

If you have any further questions on the use of this product, ask your doctor or pharmacist. Method of administration: This medicine is contained in a pressurised canister in a plastic casing with a mouthpiece. There is a counter for the 120 doses actuations and a dose indicator for the 180 actuations on the back of the inhaler, which tells you how many doses are left. For the pack size containing 120 actuations, each time vou

For the pack size containing 180 actuations, the dose indicator will show the approximate number of actuations (sprays) remaining in the canister. The dose-indicator window displays the number of sprays left in the inhaler in units of twenty (e.g., 180, 120, 100, 80, etc). When 20 actuations remain, so that the display shows the number 20 it indicates that the canister is closer to its end of life.

When 180 actuations have been discharged, the display

press the canister, a puff of medicine is released and the

counter will count down by one. Take care not to drop the

inhaler as this may cause the counter to count down.

shows the number 0. The indicator will stop moving at "0". **Testing your inhaler**

Before using the inhaler for the first time or if you have not used the inhaler for 14 days or more, you should test your inhaler to make sure that it is working properly.

1. Remove the protective cap from the mouthpiece 2. Hold your inhaler upright with the mouthpiece at the bottom 3. Direct the mouthpiece away from yourself 4. If you are using the inhaler for the first time firmly depress

- the canister 1 time to release a puff to depress the canister. 5. If you have not used the inhaler for 14 days or more firmly
- counter. Check the dose counter. If you are testing your inhaler for the first time, the counter should read 120.
- depress the canister once to release one puff.
- 6. For the pack size containing 120 actuation check the dose
- 7. For the pack size containing 180 actuation check the dose indicator. Check the dose counter. If you are testing your inhaler for the first time, the counter should read 180.





How to use your inhaler

Whenever possible, stand or sit in an upright position when inhaling



- 1. Remove the protective cap from the mouthpiece and check that the mouthpiece is clean and free from dust and dirt or any other foreign objects.
- 2. Breathe out as slowly and deeply as possible.
- 3. Hold the canister vertically with its body upwards and put your lips around the mouthpiece. Do not bite on the
- 4. Breathe in slowly and deeply through your mouth and, just after starting to breathe in press down firmly on the top of the inhaler to release one puff. If you have weak hands, it may be easier to hold the inhaler with both hands: hold the upper part of the inhaler with both index fingers and its lower part with both thumbs.
- 5. Hold your breath for as long as possible and, finally, remove the inhaler from your mouth and breathe out slowly. Do not breathe into the inhaler. If you need to take another puff, keep the inhaler in the vertical

position for about half a minute, then repeat steps 2 to 5. **Important:** Do not perform steps 2 to 5 too quickly.

After use, close with the protective cap and check the dose counter for the pack size containing 120 doses and the dose indicator for the pack size containing 180 actuations.

To lower the risk of a fungal infection in the mouth and throat, rinse your mouth, gargle with water or brush your teeth each time you use your inhaler. When to replace your inhaler

You should get a replacement when the counter or the dose indicator shows the number 20. Stop using the inhaler when the counter or the dose indicator shows 0 as any puffs left in the device may not be enough to give you a full dose. If you see 'mist' coming from the top of the inhaler or the

sides of your mouth, this means that Proxor will not be getting into your lungs as it should. Take another puff, following the instruction starting again from step 2. If you think the effect of Proxor is too much or not enough, tell

your doctor or pharmacist. If you find it difficult to operate the inhaler while starting to

breathe in you may use the AeroChamber Plus spacer device. Ask your doctor, pharmacist or a nurse about this device. It is important that you read the package leaflet which is

supplied with your AeroChamber Plus spacer device and that you follow the instructions on how to use and how to clean it, carefully. Cleaning

You should clean your inhaler once a week. When cleaning, do not remove the canister from the actuator and do not use water or other liquids to clean your inhaler. To clean your inhaler:

1. Remove the protective cap from the mouthpiece by pulling

- it away from your inhaler. 2. Wipe inside and outside of the mouthpiece and the actuator
- with a clean, dry cloth or tissue. 3. Replace the mouthpiece cover
- 4. Possible side effects

Like all medicines, this medicine can cause side effects,

although not everybody gets them. As with other inhaler treatments there is a risk of worsening shortness of breath and wheezing immediately after using

Proxor and this is known as paradoxical bronchospasm. If this occurs, you should STOP using Proxor immediately and use your quick-acting "reliever" inhaler straightaway to treat the symptoms of shortness of breath and wheezing. You should contact your doctor straightaway. Tell your doctor immediately if you experience any hypersensitivity reactions like skin allergies, skin itching,

skin rash, reddening of the skin, swelling of the skin or mucous membranes especially of the eyes, face, lips and throat. Other possible side effects are listed below according to their Common (may affect up to 1 in 10 people): • fungal infections (of the mouth and throat), headache,

hoarseness, sore throat Pneumonia in COPD patients: Tell your doctor if you have

any of the following while taking Proxor as they could be symptoms of a lung infection:

 fever or chills • increased mucus production, change in mucus colour increased cough or increased breathing difficulties

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

Palpitations, unusually fast heart beat and heart rhythm disorders, some changes in the electrocardiogram (ECG), flu like symptoms, fungal infections of the vagina, sinus

inflammation rhinitis, inflammation of the ear, throat irritation, cough and productive cough, asthma attack. Nausea, abnormal or impaired sense of taste, burning of the lips, dry mouth, swallowing difficulties, indigestion, upset stomach, diarrhoea. Pain in muscle and muscle cramps, reddening of the face, increased blood flow to some tissues in the body, excessive sweating, trembling, restlessness, dizziness, nettle rash or

Alterations of some constituents of the blood: fall in the number of white blood cells, increase in the number of blood platelets, fall in the level of potassium in the blood, increase in blood sugar level, increase in the blood level of insulin, free

fatty acid and ketones. The following side effects have also been reported as "uncommon" in patients with chronic obstructive pulmonary disease: reduction of the amount of cortisol in the blood; this is caused by the effect of corticosteroids on your adrenal

· irregular heart beat Rare (may affect up to 1 in 1,000 people): Chest tightness, missed heartbeat (caused by premature contraction of the heart's ventricles), decrease and increase

in blood pressure, kidney inflammation, swelling of skin and

with

Problems

mucous membrane persisting for several days Very rare (may affect up to 1 in 10,000 people):

how

Shortness of breath, worsening of asthma, a fall in the number of blood platelets, swelling of the hands and feet Using high-dose inhaled corticosteroids over a long time can cause in very rare cases systemic effects. These include:

your

(adreno-suppression), decrease in bone mineral density

adrenal

glands

(thinning of the bones), increased pressure in your eyes (glaucoma), cataracts, growth retardation in children and adolescents, decrease in bone mineral density (thinning of the bones) Not known (frequency cannot be estimated from the

available data):

 Sleeping problems, depression or feeling worried, restless, nervous, over-excited or irritable. These events are more likely to occur in children · Blurred vision

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, Website: www.mhra.gov.uk/yellowcard, or search for MHRA Yellow Card in the Google Play or Apple By reporting side effects you can help provide more

information on the safety of this medicine. 5. How to store Proxor Keep this medicine out of the sight and reach of children.

Before use: store the inhaler in a refrigerator (at 2-8°C) for a

to protect the environment.

maximum of 18 months. After first use: Use the inhaler for a maximum of three months and do not store above 25 °C. Do not use the inhaler after this

period and never use it after the expiry date which is stated on the carton and label after "EXP". Expiry date refers to the last day of that month. Do not freeze. If the inhaler has been exposed to severe cold, warm it with your hands for a few minutes before using. Never warm it by

artificial means. Warning: The canister contains a pressurised liquid. Do not expose the canister to temperatures higher than 50 °C. Do not pierce the canister. Do not throw away any medicines via waste water or

household waste. Ask your pharmacist how to throw away any medicines you no longer use. These measures will help

6. Contents of the pack and other information What Proxor contains:

The active substances are: beclometasone dipropionate, formoterol fumarate dihydrate.

Each actuation/metered dose from the inhaler contains micrograms of beclometasone dipropionate 6 micrograms of formoterol fumarate dihydrate. corresponds to a delivered dose from the mouthpiece of 84.6 micrograms of beclometasone dipropionate and 5.0 micrograms of formoterol fumarate dihydrate.

The other ingredients are: ethanol anhydrous, hydrochloric acid and norflurane (HFA -134a).

This medicine contains fluorinated greenhouse gases

Each inhaler of 120 actuation contains 8.15 g of HFC-134a corresponding to 0.012 tonne CO2 equivalent (global warming potential GWP = 1,430).

Each inhaler of 180 actuation contains 11.2 g of HFC-134a corresponding to 0.016 tonne CO2 equivalent (global warming potential GWP = 1.430).

To help protect the environment, the inhaler contains the CFC-free propellant, HFA-134a, which replaces completely the chlorofluorocarbon (CFC) propellants present in some other inhalers and appears to have a less damaging effect on the ozone layer.

What Proxor looks like and contents of the pack: Proxor is a pressurised inhalation solution contained in an

aluminium canister with a metering valve, fitted in a plastic actuator which incorporates a dose counter (120 actuations pack) or a dose indicator (180 actuations pack) with a pink plastic protective cap. Each pack contains:

- 1 pressurised container which provides 120 actuations (puffs)
 - 2 pressurised containers which provide 120 actuations each
 - 3 pressurised containers which provide 120 actuations each or 1 pressurised container which provides 180 actuations

Not all pack sizes may be marketed. Marketing authorisation holder:

STADA, Linthwaite, Huddersfield, HD7 5QH, UK Manufacturer responsible for batch release:

Genetic S.p.A., Contrada Canfora, 84084 Fisciano (SA), Italy. This leaflet was last revised in October 2024.