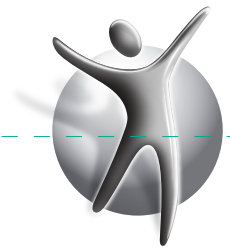
PACKAGE LEAFLET: INFORMATION FOR THE USER

Voltarol

140 mg Medicated Plaster



Diclofenac sodium

Anti-inflammatory • Relieves pain

For use in adolescents from 16 years of age and adults.
Active substance: diclofenac sodium

Read all of this leaflet carefully before you start using this medicinal product because it contains important information for you.
Always use this medicine exactly as described in this leaflet or as your doctor or pharmacist have told you.
- Keep this leaflet. You may need to read it again.
- Ask your pharmacist if you need more information or advice.
- If you get any side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet.
- You must talk to a doctor if you do not feel better or if you feel worse after 7 days.

What is in this leaflet

1. What Voltarol 140 mg Medicated Plaster is and what it is used for
2. What you need to know before you use Voltarol 140 mg Medicated Plaster
3. How to use Voltarol 140 mg Medicated Plaster
4. Possible side effects
5. How to store Voltarol 140 mg Medicated Plaster
6. Contents of the pack and other information

1. WHAT VOLTAROL 140 MG MEDICATED PLASTER IS AND WHAT IT IS USED FOR

Voltarol 140 mg Medicated Plaster is a medicine that relieves pain. It belongs to a group of medicines known as non-steroidal anti-inflammatory drugs (NSAIDs).

Voltarol 140 mg Medicated Plaster is used for the local symptomatic and short term treatment of pain associated with acute strains, sprains or bruises on the arms and legs as a result of injuries, e.g. sports injuries in adolescents from 16 years of age and adults.

2. WHAT YOU NEED TO KNOW BEFORE YOU USE VOLTAROL 140 MG MEDICATED PLASTER

Do not use Voltarol 140 mg Medicated Plaster
- if you are **allergic** to **diclofenac** or any of the **other ingredients** in this medicine (listed in section 6);
- if you are **allergic** to any **other non-steroidal anti-inflammatory drug** (NSAID, e.g. acetylsalicylic acid or ibuprofen);
- if you have **ever developed asthma, hives** or **swelling and irritation inside the nose** after taking acetylsalicylic acid or any other NSAID;
- if you are suffering from an active **stomach or duodenal ulcer;**
- if you are in the last three months of pregnancy.
- if you are a child or an adolescent younger than 16 years of age.

Do not use Voltarol 140 mg Medicated Plaster **on injured skin** (e.g. skin abrasions, cuts, burns), **infected skin or skin affected by exudative dermatitis or eczema;**

Warnings and precautions

Talk to your doctor or pharmacist before using Voltarol 140 mg Medicated Plaster
- if you suffer or have previously suffered from bronchial asthma or allergies; you may experience a bronchial muscle cramp (bronchospasm), which makes breathing difficult
- if you notice a skin rash, blistering, burning sensation, tightness of breath or wheezing that develops after applying the medicated plaster. If this happens, immediately remove the medicated plaster and stop treatment.
- if you suffer from disorders of the **kidneys, heart** or **liver**
- if you have previously suffered from a **stomach** or **intestinal ulcer**, **intestinal inflammation** or a **tendency to bleed**.

Side effects can be reduced by using the lowest effective dose for the shortest possible period of time.

IMPORTANT precautions
- The medicated plaster must not come into contact with or be applied to the eyes, lips, mouth or throat.
- Elderly patients should use Voltarol 140 mg Medicated Plaster with caution because they are more likely to experience side effects.

After taking off the medicated plaster, avoid exposing the treated area to direct sunlight or tanning lamps in order to reduce the risk of sensitivity to light.

Children and adolescents
Voltarol 140 mg Medicated Plaster should not be used in children and adolescents under 16 years of age because no adequate experience is available for this age group.

Other medicines and Voltarol 140 mg Medicated Plaster
Tell your doctor or pharmacist if you are taking or have recently taken or might take any other medicines.

Voltarol 140 mg Medicated Plaster may interact with blood pressure lowering drugs and may possibly enhance the effects of blood thinning drugs, although the chance of either of these occurring with a topically administered preparation is extremely low.

Provided that Voltarol 140 mg Medicated Plaster is used correctly, only a small amount of diclofenac is absorbed into the body so that the interactions described for diclofenac-containing medicines taken orally are unlikely to happen.

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 using this medicine.

Pregnancy
In the last 3 months of pregnancy, Voltarol 140 mg Medicated Plaster must not be used because an increased risk of complications for the mother and the child cannot be ruled out (see "Do not use Voltarol 140 mg Medicated Plaster").

In the first 6 months of pregnancy or if you want to become pregnant, Voltarol 140 mg Medicated Plaster should be used only after talking to your doctor.

Breast-feeding
Small quantities of diclofenac pass into the breast milk. Talk to your doctor before using Voltarol 140 mg Medicated Plaster during breast-feeding. In any case, if you are breast-feeding Voltarol 140 mg Medicated Plaster should not be applied directly onto the breast area.

Driving and using machines
Voltarol 140 mg Medicated Plaster has no influence on your ability to drive and use machines.

3. HOW TO USE VOLTAROL 140 MG MEDICATED PLASTER

Always use this medicine exactly as described in this leaflet or as your doctor or pharmacist have told you. Check with your doctor or pharmacist if you are not sure.

The recommended dose is one medicated plaster twice daily.

Attach one medicated plaster to the painful area twice daily, in the morning and in the evening. The maximum total daily dose is 2 medicated plasters, even if there is more than one injured area to be treated. Treat only one painful area at a time.

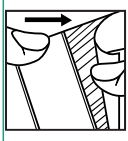
Use in children and adolescents
Voltarol 140 mg Medicated Plaster is contraindicated for use in children and adolescents under 16 years of age.

There are insufficient data of efficacy and safety available for children and adolescents below 16 years (see section 2).

In adolescents aged 16 years and over, if this medicine is required for more than 7 days for pain relief or if the symptoms worsen, please consult a doctor.

Method of administration
Use this medicine only on the skin.

Instructions for use:
1. Tear open the envelope containing the medicated plaster on the notch and remove the medicated plaster.



To apply the plaster:
2. Remove one of the two protective films.

3. Apply to the area to be treated and remove the remaining protective film.

4. Apply slight pressure with the palms of your hand until complete adhesion to the skin is achieved.

To remove the plaster:
5. Moisten the plaster with water and peel away an edge of the plaster and pull smoothly away from the skin.
6. To remove any product residues, wash the affected area with water gently rubbing the area with your fingers using a circular movement.

If necessary, the medicated plaster can be held in place using a net bandage.

Use the medicated plaster only on intact, healthy skin.

The medicated plaster should not be applied to the face, eyes, lips, mouth or throat.

Do not use the medicated plaster together with an air-tight (occlusive) bandage.

Do not wear it when bathing or showering.

Do not divide the medicated plaster, by cutting with scissors, for example.

Duration of use
Do not use Voltarol 140 mg Medicated Plaster for longer than 7 days.
If symptoms worsen or persist for longer than 7 days, you should consult a doctor.

If you have the impression that the effect of Voltarol 140 mg Medicated Plaster is too strong or too weak, please talk to your doctor or pharmacist.

If you apply more Voltarol 140 mg Medicated Plaster than you should
Please tell your doctor if you experience side effects after incorrect use of this medicine, if you apply more patches than you should or if a patch is accidently applied to a child. They will be able to advise you of any action that may need to be taken.

If you forget to use Voltarol 140 mg Medicated Plaster
You should apply a new patch to the affected area when you remember. Do not apply more than one patch to make up for the missed patch.

Do not use in children and adolescents under 16 years of age.

Do not use more than 2 plasters within 24 hours.

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.

Tell your doctor immediately and stop using the plaster if you notice any of the following:
sudden itchy rash (hives); swelling of the hands, feet, ankles, face, lips, mouth or throat; difficulty breathing; drop in blood pressure (feeling lightheaded) or weakness.

You may experience the following side effects:

Common side effects (may affect up to 1 in 10 people):
local skin reactions, such as skin redness, burning sensation, itching, inflamed skin redness, skin rash, sometimes with pustules or wheals.

Very rare side effects (may affect up to 1 in 10,000 people):
Hypersensitivity reactions or local allergic reactions (contact dermatitis). Photosensitivity (sensitivity to sunlight).

In patients externally using drugs from the same drug group as diclofenac, there have been isolated reports of generalised skin rash, hypersensitivity reactions such as swelling of the skin, lips, eyes or throat and anaphylactic-type (severe allergic) reactions. Including problems with blood circulation and light sensitivity reactions.

Absorption of diclofenac into the body by the skin is very low compared to the drug concentration in the blood following diclofenac taken by mouth. Therefore, the likelihood of side effects occurring in the body as a whole (such as stomach or kidney problems or difficulty breathing) is very low.

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 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 VOLTAROL 140 MG MEDICATED PLASTER

Keep out of the sight and reach of children.

Do not use this medicine after the expiry date which is stated on the outer carton and the sachet after "EXP". The expiry date refers to the last day of that month.

Store below 30 °C.
Store in the original package in order to protect from desiccation and light.
Keep the sachet tightly closed in order to protect from desiccation and light.

Do not use Voltarol 140 mg Medicated Plaster if you notice that it is damaged.

Used plasters should be folded in half with the sticky side inwards.

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 Voltarol 140 mg Medicated Plaster contains
The active substance is diclofenac sodium.
Each medicated plaster contains 140 mg diclofenac sodium.

The other ingredients are:
Supporting layer:
Polyester non-woven fabric

Adhesive layer:
Basic butylated methacrylate copolymer
Copolymer acrylate vinyl acetate
PEG 12 stearate
Sorbitan oleate

Liner:
Mono silicone coated paper

What Voltarol 140 mg Medicated Plaster looks like and contents of the pack
Voltarol 140 mg Medicated Plaster are white 10x14 cm sized self-adhesive plasters made of non-woven fabric on one and paper on other side.
Voltarol 140 mg Medicated Plaster is available in packs of 2, 5 and 10 plasters, each in a single sachet.

Not all pack sizes may be marketed.

Marketing Authorisation Holder and Manufacturer

Marketing Authorisation Holder:
GlaxoSmithKline Dungarvan Limited, Knockbrack, Dungarvan, County Waterford, Ireland.

Manufacturer:
GlaxoSmithKline Consumer Healthcare (UK) Trading Limited, Brentford, TW8 9GS, U.K.

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20060 Cassina de' Pecchi (MI), Itália.

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