

Patient Information Leaflet

Sodium Chloride Injection BP 0.9% w/v

Important information about your medicine

- ▶ Your doctor or nurse will give you the injection
- ▶ If this injection causes you any problems talk to your doctor, nurse or pharmacist
- ▶ Please tell your doctor or pharmacist, if you have any other medical conditions or have an allergy to any of the ingredients of this medicine
- ▶ Please tell your doctor or pharmacist, if you are taking any other medicines

- **Read all of this leaflet carefully** before you start using this medicine. In some circumstances **this may not be possible** and this leaflet will be kept in a safe place should you wish to read it.
- **Keep this leaflet.** You may need to read it again
- If you have any further questions, please **ask your doctor or your pharmacist.**
- This medicine has been prescribed for you personally and you should not pass it on to others. It may harm them, even if their symptoms are the same as yours.

Where to find information in this leaflet

1. What Sodium Chloride Injection BP 0.9% w/v is and what it is used for
2. Before you are given Sodium Chloride Injection BP 0.9% w/v
3. How to use Sodium Chloride Injection BP 0.9% w/v
4. Possible side effects
5. Storing Sodium Chloride Injection BP 0.9% w/v
6. Further information

1. What Sodium Chloride Injection BP 0.9% w/v is and what it is used for

Sodium (common salt) occurs naturally in your body. A solution of 0.9% sodium chloride in water for injections is the same strength as your blood. It is used to:

- prime giving sets before blood or other medicines are given to you.
- replace the loss of sodium from your body.
- make up medicines that may be injected into you (because it occurs naturally in the body).
- irrigate (wash) surfaces of your body.

2. Before you are given Sodium Chloride Injection BP 0.9% w/v

You should NOT be given Sodium Chloride Injection BP 0.9% w/v if you:

- Are **sensitive** or **allergic** to Sodium Chloride Injection BP 0.9% w/v or any of the other ingredients in this injection.

Please tell your doctor or nurse before being given the injection if you:

- suffer from **Heart** disease or heart failure
- have impaired **kidney** function
- have **Diabetes**
- have **Pre-eclampsia** (high blood pressure during pregnancy)
- have **Fluid retention** resulting in **swelling** of parts of the body, particularly your feet and ankles.
- have **Pseudohyponatraemia** (a low level of salt in your blood caused by levels of fat or protein in your blood that are too high)
- have **Hyperlipaemia** (a raised level of fat in your blood)
- have **Hyperproteinaemia** (a raised level of protein in your blood).

Using other medicines:

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

Pregnancy or breast feeding:

Please tell your **doctor or nurse** before being given this injection if you are **pregnant** or **breast feeding**. The doctor will then decide if the injection is suitable for you.

Driving and using machines:

You should **not drive or use machinery** if you are **affected** by the administration of Sodium Chloride Injection BP 0.9% w/v.

3. How to use Sodium Chloride Injection BP 0.9% w/v

Your nurse or doctor will give you the injection.

Your doctor will decide the **correct dosage** for you and **how and when** the injection will be given.

Since the injection will be given to you by a doctor or nurse, it is **unlikely** that you will be given too much. **If you think you have been given too much**, you must tell the person giving you the injection.

4. Possible side effects

Like all medicines, Sodium Chloride Injection BP 0.9% w/v can cause side effects, although not everybody gets them.

- **Pain** at the **site** of **injection** if administered underneath the skin.
- **Dehydration** of internal organs, particularly the brain, which may result in the development of blood clots and internal bleeding.

Someone who has too much sodium chloride in their body may also feel or be sick, have diarrhoea, stomach pain, low blood pressure, an elevated heart rate, excess sweating, headache, dizziness or fever. Other symptoms of excess sodium chloride include thirst, dry mouth or eyes,

swollen legs or chest pain, difficulty breathing, irritability, restlessness, feeling weak, twitching or other unusual muscle movements.

If you think this injection is causing you **any problems**, or you are at all worried, **talk to your doctor, nurse or pharmacist**.

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** at: www.mhra.gov.uk/yellowcard. By reporting side effects you can help provide more information on the safety of this medicine.

5. Storing Sodium Chloride Injection BP 0.9% w/v

Your injection will be stored at less than 25°C and protected from light. The nurse or doctor will check that the injection is not past its expiry date before giving you the injection.

6. Further information

What Sodium Chloride Injection BP 0.9% w/v contains:

This injection contains the **active ingredient** sodium chloride (common salt) in a sterile solution. Each 1 ml of solution contains 9 mg of sodium chloride. This injection contains the following **inactive ingredients**: sodium hydroxide, hydrochloric acid and water for injections.

What Sodium Chloride Injection BP 0.9% w/v looks like and contents of the pack:

Sodium Chloride Injection BP 0.9% w/v is supplied in:

Clear glass ampoules, 2 ml, 5 ml, 10 ml and 20 ml. Packed in cardboard cartons to contain 10 ampoules.

Clear glass vials (33 ml, 100 ml and 200 ml) with bromobutyl rubber stopper, plastic outer cap and inner aluminium ring.

Clear glass vials 50 ml with chlorbutyl rubber stopper, plastic outer cap and inner aluminium ring.

Not all pack sizes may be marketed.

The marketing authorisation number of this medicine is: PL 01502/0006R

Marketing Authorisation Holder:

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For any information about this medicine, please contact the Marketing Authorisation Holder

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