



5/12613455/0916

Directions for use
Read carefully!



0.9% Sodium Chloride Injection B.P.

Composition

100 ml of solution contain

Active ingredients:

Sodium Chloride 0.9 g

Excipients:

Water for injections

Theoretical osmolarity: 308 mOsm/l

Titration acidity: < 0.3 mmol/l

pH: 4.5 - 7.0

Electrolyte concentrations:

Sodium 154 mmol/l

Chloride 154 mmol/l

Pharmaceutical form

Solution for injection

Pharmaco-therapeutic group

Vehicle solution

Indications

Solvent or diluent for compatible electrolyte concentrates or drugs.

Contraindications

None

Precautions for use and special warnings

0.9% Sodium Chloride Injection B.P. should only be administered with particular caution to patients with

- hypernatraemia
- hyperchloraemia

Interactions

When mixing with other medicaments, possible incompatibilities should be considered.

Dosage

The quantity to be chosen depends on the desired concentration of the medicament to be dissolved.

For the use of this solution as solvent/diluent for compatible electrolyte concentrates or medicaments, the instructions for use relating to the medicament to be added should be observed.

Method of administration

Intravenous or subcutaneous injection.

Overdose

Symptoms

Overdose of 0.9% Sodium Chloride solution may result in hypernatraemia, hyperchloraemia, hyperhydration, hyperosmolarity of the serum, and metabolic acidosis.

Emergency treatment, antidotes

Immediate stop of administration, administration of diuretics with continuous monitoring of serum electrolytes, correction of electrolyte and acid-base imbalances.

Undesirable effects

Administration of larger amounts of the solution may lead to hypernatraemia and hyperchloraemia.

Note: Patients are advised to inform their doctor or pharmacist of any adverse reaction they experience in connection with the administration of this drug.

Usage during pregnancy

No adverse reaction has been reported. However, risk versus benefits should be assessed before using.

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Storage

The product should not be stored above the temperature stated on the label.

Shelf life

The product must not be used beyond the expiry date stated on the label.

Presentation

In plastic ampoules "Mini-Plasco®" and "Mini-Plasco® Connect" of 5 ml, 10 ml, 20 ml in box of 20's.

Not all presentation is available locally.

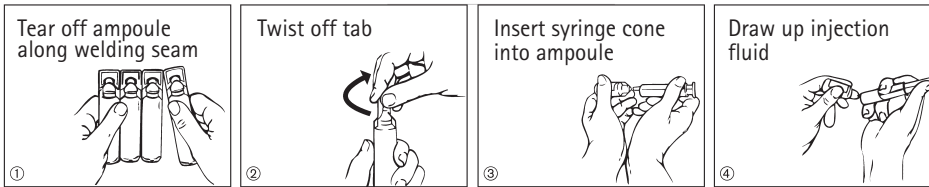
Instruction for use/handling

The solution is supplied in single-dose containers. Discard unused contents.

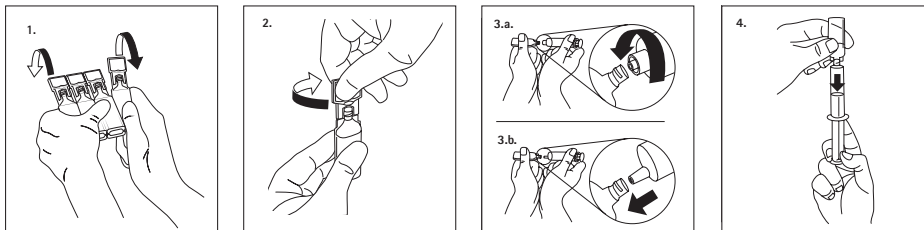
Use contents immediately after opening the container.

Only to be used if solution is clear and the container undamaged.

Mini-Plasco® Handling



Mini-Plasco® Connect Handling



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Manufactured by:
**B. Braun Medical
Industries Sdn. Bhd.**
(Company No. 19051-M)
11900 Bayan Lepas,
Penang, Malaysia.

