

SODIUM CHLORIDE (concentrated solutions)

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| Indication | <ul style="list-style-type: none"> Treatment of hyponatraemia¹ | |
| ORAL | Sodium chloride 6% | |
| | Presentation | <ul style="list-style-type: none"> Oral solution (sachet): 1 mmol in 1 mL |
| | Dosage | <ul style="list-style-type: none"> Daily requirement 2–6 mmol/kg/day <ul style="list-style-type: none"> Divide into 4 or more doses to reduce gastric upset² |
| | Preparation | <ul style="list-style-type: none"> Nil required |
| | Administration | <ul style="list-style-type: none"> Draw up prescribed dose into oral/enteral syringe Oral/OGT/NGT with feed |
| ORAL | Sodium chloride 23.4% | |
| | Presentation | <ul style="list-style-type: none"> Compounded oral solution: 4 mmol in 1 mL <ul style="list-style-type: none"> Available: Mater Pharmacy Production Services Vial (solution): 23.4% (40 mmol in 10 mL) <ul style="list-style-type: none"> If glass ampoule, use a filtered needle (due to risk of glass shard contamination) |
| | Dosage | <ul style="list-style-type: none"> Daily requirement 2–6 mmol/kg/day <ul style="list-style-type: none"> Divide into 4 or more doses to reduce gastric upset² |
| | Preparation | <ul style="list-style-type: none"> Nil required |
| | Administration | <ul style="list-style-type: none"> Draw up prescribed dose into oral/enteral syringe Oral/OGT/NGT with feed² |
| INTRAVENOUS | Sodium chloride 23.4% (must be diluted before administration³) | |
| | Presentation | <ul style="list-style-type: none"> Vial (solution): 23.4% (equivalent to 40 mmol in 10 mL) |
| | Dosage | <ul style="list-style-type: none"> Daily requirement 2–6 mmol/kg/day |
| | Burette (100 mL) | |
| | Dilution | <ul style="list-style-type: none"> Add prescribed dose of sodium chloride 23.4% to compatible 24-hour maintenance fluid |
| | Administration | <ul style="list-style-type: none"> Calculate infusion rate/hour to deliver prescribed daily dose volume over 24 hours Refer to worked example for 100 mL burette calculation below |
| | Concentrated infusion (via CVL recommended) requires sufficient volume of maintenance fluid | |
| Dilution | <ul style="list-style-type: none"> Preparation is according to weight Refer to Quick Guide for CVL below | |
| Administration | <ul style="list-style-type: none"> Prime the infusion line and reduce total syringe volume to prescribed dose IV infusion via syringe driver pump at prescribed rate On completion, disconnect syringe and infusion line Flush access port at same rate as infusion Refer to worked example for concentrated infusion | |
| Contraindication Caution | <ul style="list-style-type: none"> Contraindication <ul style="list-style-type: none"> Nil known Caution <ul style="list-style-type: none"> Nil known | |
| Special considerations | <ul style="list-style-type: none"> Dosage varies according to degree of hyponatraemia² Manage doses above 6–8 mmol/kg/day cautiously (refer to side effects) in consultation with SMO or NNP CVL recommended for administration of any hypertonic solution greater than 0.9 %⁴ with consideration of catheter placement | |

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| Monitoring | <ul style="list-style-type: none"> • Electrolytes, 6–24 hourly to evaluate rate of sodium level rise (or as requested by SMO)² • Extravasation risk, if administered via PIVC⁵ |
| Compatibility | <ul style="list-style-type: none"> • 5% glucose⁵, 10% glucose⁶, 0.9% sodium chloride⁶ |
| Incompatibility | <ul style="list-style-type: none"> • No information: check individual drug or solution³ |
| Interactions | <ul style="list-style-type: none"> • Nil known |
| Stability | <ul style="list-style-type: none"> • Oral solution (sachet) <ul style="list-style-type: none"> ◦ Store below 25 °C⁷ • Oral solution (compounded) <ul style="list-style-type: none"> ◦ Store in refrigerator 2–8 °C ◦ Discard 4 weeks after opening or as per local infection control policy (limited evidence) • Vial <ul style="list-style-type: none"> ◦ Store at room temperature below 30 °C³ |
| Side effects | <ul style="list-style-type: none"> • Appropriate parenteral use for electrolyte replacement is unlikely to result in adverse effects³ • Mild hypernatraemia symptoms include³: <ul style="list-style-type: none"> ◦ Reduced salivation and lacrimation, fever, tachycardia, hypertension, restlessness, irritability and weakness • Excessive administration may lead to hypernatraemia, causing³: <ul style="list-style-type: none"> ◦ Organ dehydration and dysfunction, hypokalaemia and acidosis ◦ Congestive heart failure and pulmonary oedema particularly if cardiovascular disease or receiving corticosteroids or drugs that give rise to sodium retention ◦ Loss of bicarbonate with an acidifying effect due to excessive chloride ions • Inadvertent administration of concentrated sodium chloride solutions³: <ul style="list-style-type: none"> ◦ May result in sudden hypernatraemia and potential complications (e.g. somnolence, confusion progressing to convulsions, cardiovascular shock, CNS disorders, extensive haemolysis, respiratory failure and cortical necrosis of the kidney) • Rapid administration (a change in serum sodium of more than 12 mmol over 24 hours) may cause osmotic stress and subsequent demyelination injury to the brain¹ |
| Actions | <ul style="list-style-type: none"> • Sodium is the principle cation of extracellular fluid and chloride is the principle anion of extracellular fluid • Sodium content normally determines the volume of extracellular fluid, and is important in the regulation of osmolarity, acid-base balance, and the membrane potential of cells |
| Abbreviations | CNS: central nervous system, CVL: central venous line, IV: intravenous, NGT: nasogastric tube, NNP: neonatal nurse practitioner, OGT: orogastric tube, PIVC: peripheral intravenous catheter, SMO: senior medical officer |
| Keywords | electrolyte, hyponatraemia, hypernatraemia, hypersal 6%, neonatal medicine, neonatal monograph, sodium chloride, strong salt, sodium chloride 23.4% |

The Queensland Clinical Guideline [Neonatal medicines](#)⁸ is integral to and should be read in conjunction with this monograph. Refer to the disclaimer. Destroy all printed copies of this monograph after use.

100 mL burette: worked example calculation

To calculate volume of intravenous sodium chloride 23.4% (mL) to add to each burette for 24 hours

| | Worked example scenario | 100 mL burette example |
|--------|---|---|
| Step 1 | Identify <ul style="list-style-type: none"> Baby weight=2.5 kg Sodium dose required in 24 hours =4 mmol/kg in 24 hours | $2.5 \text{ kg} \times 4 \text{ mmol} = 10 \text{ mmol/day}$ |
| Step 2 | Calculate required daily dose of sodium chloride 23.4% in mL $\frac{\text{calculated dose (mmol)}}{\text{stock strength (mmol)}}$ | $\frac{10 \text{ mmol/day}}{4 \text{ mmol/mL}} = 2.5 \text{ mL per day}$ |
| Step 3 | Identify total maintenance fluid volume (e.g. glucose 10% and usually excludes drug/lipid/arterial and venous catheter patency infusion) <i>Example: 12.5 mL/hour of 10% glucose</i> | $12.5 \text{ mL/hour} \times 24 \text{ hours} = 300 \text{ mL in 24 hours}$ |
| Step 4 | Calculate number of 100 mL burettes required in 24 hours | $\frac{300 \text{ mL}}{100 \text{ mL}} = 3 \text{ burettes in 24 hours}$ |
| Step 5 | Calculate the volume (mL) of sodium chloride 23.4% to add to each burette | $\frac{2.5 \text{ mL}}{3 \text{ burettes}} = 0.83 \text{ mL}$ |
| Step 6 | Add 0.83 mL of sodium chloride 23.4 % to each 99.17 mL (total 100 mL) of 10% glucose until requirement change | |

Quick Guide for CVL

Sodium chloride 23.4% via CVL according to weight to achieve a concentration of less than 0.5 mmol/mL.

| Weight of baby | Draw up sodium chloride 23.4% | Make up to total volume | Rate of infusion | Delivers (mmol/kg/day) |
|------------------|-------------------------------|-------------------------|------------------|------------------------|
| Less than 1.5 kg | 10 mmol/kg (2.5 mL/kg) | 30 mL | 0.5 mL/hour | 4 mmol/kg/day |
| 1.5–3 kg | 5 mmol/kg (1.25 mL/kg) | 30 mL | 1 mL/hour | 4 mmol/kg/day |
| More than 3 kg | 2.5 mmol/kg (0.63 mL/kg) | 30 mL | 2 mL/hour | 4 mmol/kg/day |

References

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Document history

| ID number | Effective | Review | Summary of updates |
|--------------------|------------|------------|--|
| NMedQ21.070-V1-R26 | 20/07/2021 | 20/07/2026 | Endorsed by Queensland Neonatal Services Advisory Group (QNSAG) |
| NMedQ21.070-V2-R26 | 13/08/2021 | 20/07/2026 | <ul style="list-style-type: none"> Amended to clarify use of glass ampoule for oral route |
| NMedQ26.070-V3-R31 | 24/02/2026 | 24/02/2031 | Full review. Endorsed by QNSAG <ul style="list-style-type: none"> Added: Sodium chloride 6% oral No change to dose or frequency Amended: if glass ampoule use filter Added: worked example for burette preparation Updated compatibility, incompatibility, references, presentation |

QR code

