Insulin neutral (soluble) – hyperKALAEMIA
100units/mL injection

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Note:
This guideline provides advice of a general nature. This statewide guideline has been prepared to promote and facilitate standardisation and consistency of practice, using a multidisciplinary approach. The guideline is based on a review of published evidence and expert opinion.
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If for good clinical reasons, a decision is made to depart from the guideline, the responsible clinician must document in the patient’s medical record, the decision made, by whom, and detailed reasons for the departure from the guideline.
This statewide guideline does not address all the elements of clinical practice and assumes that the individual clinicians are responsible for discussing care with consumers in an environment that is culturally appropriate and which enables respectful confidential discussion. This includes:

• The use of interpreter services where necessary,
• Advising consumers of their choice and ensuring informed consent is obtained,
• Providing care within scope of practice, meeting all legislative requirements and maintaining standards of professional conduct, and
• Documenting all care in accordance with mandatory and local requirements

This is a High Risk Medication
Use the term “units” (written in full) as the abbreviation of “U” can be misinterpreted as a “0”
An overdose can be rapidly fatal

Synonyms
Neutral insulin, soluble insulin, Actrapid®

Dose and Indications

Hyperkalaemia

Intravenous Injection: infuse over 15 minutes
0.1 unit/kg
Always prescribe with glucose to maintain glucose(g):insulin(units) ratio of 5:1
Reserved for the emergency treatment of cardiac arrhythmia due to hyperkalaemia

Continuous intravenous Infusion
0.1 to 0.2 units/kg/hour
Take care to avoid hypoglycaemia. Where plasma glucose levels are normal at baseline, consider the following to maintain glucose(g):insulin(units) ratio 2.5:1.
- Central access available: insulin 0.1unit/mL standard concentration in 25% glucose
- Central access not available: insulin 0.04unit/mL standard concentration in 10% glucose
If there is concern regarding high plasma glucose levels, consider alternative diluent. If glucose infusion is run concurrently (rather than as diluent above), ensure rate changes in insulin are adequately matched with glucose delivery to reduce the risk of hypoglycaemia.

Preparation and Administration

**Intravenous Injection**

**STEP ONE:**

Dilute 0.1mL of the 100units/mL soluble insulin with compatible fluid (e.g. sodium chloride 0.9%), to a total of 10mL. The solution now contains 1unit/mL.

<table>
<thead>
<tr>
<th>Dose</th>
<th>0.05units</th>
<th>0.1units</th>
<th>0.15units</th>
<th>0.2units</th>
<th>0.3units</th>
<th>0.4units</th>
<th>0.5units</th>
</tr>
</thead>
<tbody>
<tr>
<td>Volume</td>
<td>0.05mL</td>
<td>0.1mL</td>
<td>0.15mL</td>
<td>0.2mL</td>
<td>0.3mL</td>
<td>0.4mL</td>
<td>0.5mL</td>
</tr>
</tbody>
</table>

Draw up the prescribed dose of insulin (0.1unit/kg = 0.1mL/kg) for dilution in glucose solution (see STEP TWO)

Discard the remaining diluted insulin 1unit/mL solution.

**STEP TWO:**

- **Where central access available** - further dilute dose with 1mL/kg glucose 50% (glucose:insulin ratio of 5:1) and administer as a push over at least 15 minutes OR

- **Where central access is not available** – further dilute dose with 5mL/kg glucose 10% (glucose:insulin ratio of 5:1) and administer as a push over at least 15 minutes

**Continuous Intravenous Infusion**

Insulin adsorbs to PVC: new intravenous tubing should be flushed/primed with 20mL of a diluted insulin solution (use same strength as infused) prior to intravenous administration (no filter required).

Insulin Concentration Selection Tables can be found on the following pages of this guideline to assist prescribers to gauge which strength is best for the patient.

- The two standard concentrations to select from are:
  - Insulin 0.04unit/mL in 10% glucose
  - Insulin 0.1units/mL in 25% glucose

**Formulae**

To calculate infusion rate (mL/hr):

\[
\text{Rate (mL/hour)} = \frac{\text{dose (units/kg/hour)} \times \text{weight(kg)}}{\text{Infusion Strength (units/mL)}}
\]

To calculate the dose (units/kg/hour):

\[
\text{Dose (units/kg/hour)} = \frac{\text{Rate(mL/hr)} \times \text{Strength (units/mL)}}{\text{Weight (kg)}}
\]
Insulin Concentration Selection Table

**Insulin 0.04units/mL**

**Double dilution to make 50 mL syringe:**

**STEP ONE:** Dilute 0.5mL of 100unit/mL soluble insulin with 9.5mL of compatible fluid (total of 10mL). The resulting solution contains 5 unit/mL insulin.

**Step TWO:** Dilute 0.4mL insulin (5 units/mL) with 49.6mL of glucose 10% (or other compatible fluid) (total of 50mL)

Discard remaining 5unit/mL solution

Recommended for neonates who are limited to peripheral IV access

<table>
<thead>
<tr>
<th>Rate (mL/hr)</th>
<th>Weight (kg)</th>
<th>Approximate units/kg/hour</th>
<th>Weight (kg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.5</td>
<td>0.08 0.12 0.16 0.2 0.24 0.28 0.32 0.36 0.4</td>
<td>0.5</td>
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<tr>
<td>1</td>
<td>1</td>
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<tr>
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<tr>
<td>3</td>
<td>0.01 0.02 0.03 0.03 0.04 0.05 0.05 0.06 0.07</td>
<td>3</td>
<td></td>
</tr>
</tbody>
</table>

**Insulin 0.1units/mL**

**Double dilution to make 50 mL syringe:**

**STEP ONE:** Dilute 0.5mL of 100unit/mL soluble insulin with 9.5mL of compatible fluid (total of 10mL). The resulting solution contains 5 unit/mL insulin.

**Step TWO:** Dilute 1mL insulin (5 units/mL) with 49mL of 25% glucose (or other compatible fluid) (total of 50mL)

Discard remaining 5unit/mL solution

Recommended for neonates with central IV access

<table>
<thead>
<tr>
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<th>Approximate units/kg/hour</th>
<th>Weight (kg)</th>
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</thead>
<tbody>
<tr>
<td>0.5</td>
<td>0.1 0.2 0.3 0.4 0.5 0.6</td>
<td>0.5 1 1.5 2 2.5 3</td>
<td>0.5 1 1.5 2 2.5 3</td>
</tr>
<tr>
<td>1</td>
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<td>0.5 1 1.5 2 2.5 3</td>
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<tr>
<td>1.5</td>
<td>0.03 0.07 0.1 0.13 0.17 0.2</td>
<td>0.5 1 1.5 2 2.5 3</td>
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<tr>
<td>2</td>
<td>0.03 0.05 0.08 0.1 0.13 0.15</td>
<td>0.5 1 1.5 2 2.5 3</td>
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<td>2.5</td>
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<tr>
<td>3</td>
<td>0.02 0.03 0.05 0.07 0.08 0.1</td>
<td>0.5 1 1.5 2 2.5 3</td>
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</table>
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Compatible Fluids
Glucose 5%, glucose 10%, sodium chloride 0.9%
Glucose 25% (MUST be administered via central line)
Glucose 50% (MUST be administered via central line)

Adverse Effects
Hypoglycaemia, hypokalaemia

Monitoring
> Frequent blood and urine glucose levels as guided by the prescriber. Document in nursing care plan
> Electrolytes, particularly potassium

Practice Points
> The original vial of insulin may be reused for the same patient for up to 28 days
> Unopened vials to be stored in the fridge. Opened vials may be kept at room temperature for up to 28 days
> If ceasing insulin or changing the concentration, be careful to remove and replace the previous line and T-piece to avoid flushing through any insulin remaining in the tubing
> Insulin is incompatible with many medications (see Intravenous medication compatibility chart monograph or contact pharmacist for more information)
> Y-site compatible with parenteral nutrition, lipid emulsion and heparin

References
> Royal Children’s Hospital - Clinical Practice Guideline: Hyperkalaemia. Viewed 27 May 2020