<table>
<thead>
<tr>
<th>Alert</th>
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<tbody>
<tr>
<td>Indication</td>
<td>Treatment of hypotensive shock with or without myocardial dysfunction.</td>
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</tbody>
</table>
| Action | Catecholamine with alpha and beta adrenergic actions. Haemodynamic effects are dose dependent:  
  - At low doses of 0.01–0.1 microgram/kg/minute primarily stimulates cardiac and vascular beta 1- and beta 2-adrenoreceptors leading to increased inotropy, chronotropy, conduction velocity and peripheral vasodilation.  
  - At doses greater than 0.1 microgram/kg/minute adrenaline also stimulates vascular and cardiac alpha 1-receptors causing vasoconstriction and increased inotropy. The net effects are increases in blood pressure and systemic blood flow caused by the drug-induced increases in systemic vascular resistance (SVR) and cardiac output.¹ |
| Drug Type | Inotropic vasopressor. |
| Trade Name | Adrenaline 1:1,000 injection, Aspen Adrenaline 1: 10,000 injection. |
| Presentation | 1 mg/10 mL or 1:10,000 ampoule [100 microgram/mL]  
1 mg/mL or 1:1,000 ampoule [1000 microgram/mL] |
| Dosage / Interval | Low dose: 0.05–0.1 microgram/kg/minute  
High dose: 0.1–1 microgram/kg/minute |
| Route | Continuous IV infusion. |

### Preparation/Dilution

#### Preparation using **1:1,000 (1 mg/mL) ampoule**

#### LOW CONCENTRATION IV infusion

<table>
<thead>
<tr>
<th>Infusion dose</th>
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<tbody>
<tr>
<td>1 mL/hour = 0.05 microgram/kg/minute</td>
<td>150 microgram/kg adrenaline and make up to 50 mL</td>
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</table>

Draw up 150 microgram/kg [0.15 mL/kg] of 1:1000 adrenaline and add glucose 5%, glucose 10% or sodium chloride 0.9% to make a final volume of 50 mL with a concentration of 3 microgram/kg/mL. Infusing at a rate of 1 mL/hour = 0.05 microgram/kg/minute.

#### HIGH CONCENTRATION IV infusion

<table>
<thead>
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<th>Infusion dose</th>
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<tbody>
<tr>
<td>1 mL/hour = 0.2 microgram/kg/minute</td>
<td>600 microgram/kg adrenaline and make up to 50 mL</td>
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</table>

Draw up 600 microgram/kg [0.6 mL/kg] of 1:1000 adrenaline and add glucose 5%, glucose 10% or sodium chloride 0.9% to make a final volume of 50 mL with a concentration of 12 microgram/kg/mL. Infusing at a rate of 1 mL/hour = 0.2 microgram/kg/minute.

For infants requiring fluid restriction consider:  
**VERY HIGH CONCENTRATION IV infusion**

<table>
<thead>
<tr>
<th>Infusion dose</th>
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<tbody>
<tr>
<td>1 mL/hour = 0.4 microgram/kg/minute</td>
<td>1200 microgram/kg adrenaline and make up to 50 mL</td>
</tr>
</tbody>
</table>

Draw up 1200 microgram/kg [1.2 mL/kg] of 1:1000 adrenaline and add glucose 5% ONLY to make a final volume of 50 mL with a concentration of 24 microgram/kg/mL. Infusing at a rate of 1 mL/hour = 0.4 microgram/kg/minute.

*Stability data only available for 5% glucose for very high concentration.*

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¹ This RHW document is a modification of Neomed version. Dosage schedules remain the same. However, information on the commercial preparations not used at RHW is deleted. The risk rating is modified as per the local health district policy.
### Preparation using 1:10,000 (1 mg/10 mL) ampoule

#### LOW CONCENTRATION IV infusion

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Draw up 150 microgram/kg [1.5 mL/kg] of 1:10,000 adrenaline and add glucose 5%, glucose 10% or sodium chloride 0.9% to make a final volume of 50 mL with a concentration of 3 microgram/kg/mL. Infusing at a rate of 1 mL/hour = 0.05 microgram/kg/minute.

#### HIGH CONCENTRATION IV infusion

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Draw up 600 microgram/kg [6 mL/kg] of 1:10,000 adrenaline and add glucose 5%, glucose 10% or sodium chloride 0.9% to make a final volume of 50 mL with a concentration of 12 microgram/kg/mL. Infusing at a rate of 1 mL/hour = 0.2 microgram/kg/minute.

For infants requiring fluid restriction consider:

#### VERY HIGH CONCENTRATION IV infusion*

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Draw up 1200 microgram/kg [12 mL/kg] of 1:10,000 adrenaline and add glucose 5% ONLY to make a final volume of 50 mL with a concentration of 24 microgram/kg/mL. Infusing at a rate of 1 mL/hour = 0.4 microgram/kg/minute.

*Stability data only available for 5% glucose for very high concentration.

### Administration

Continuous intravenous infusion via a central line. Use with caution via a peripheral line.

### Monitoring

Continuous heart rate, ECG and blood pressure monitoring preferable. Assess urine output and peripheral perfusion frequently. Observe IV site closely for blanching and extravasation.

### Contraindications

Arrhythmia and tachyarrhythmia.
Cardiovascular disease resulting in arterial narrowing including cerebrovascular disease, coronary artery disease and digital ischaemia.
Phaeochromocytoma.
Thyrotoxicosis.
Glaucoma.
Known hypersensitivity to sympathomimetic amines.

### Precautions

Ensure adequate circulating blood volume prior to commencement.
Adrenaline is a potent chronotrope and vaspressor – may cause excessive tachycardia, severe hypertension and ventricular arrhythmias.
Adrenaline may cause lactic acidosis and hyperglycaemia.

### Drug Interactions

Hypotension may be observed with concurrent use of vasodilators such as glyceryl trinitrate, nitroprusside and calcium channel blockers.
Concurrent use of digitalis glycosides may increase the risk of cardiac arrhythmias.
Concurrent use of IV phenytoin with adrenaline may result in dose dependent, sudden hypotension and bradycardia.

### Adverse Reactions

Tachycardia and arrhythmia.
Systemic hypertension especially at higher doses.
May cause hypokalaemia.
Tissue necrosis at infusion site with extravasation.
Digital ischaemia.

Compatibility

Fluids: Glucose 5%, glucose 10%, Hartmann’s, sodium chloride 0.9%. Stability data only available for 5% glucose for very high concentration.

Y-site: Amino acid solutions. Amiodarone, anidulafungin, atracurium, bivalirudin, caspofungin, cisatracurium, dexametomidine, dobutamine, dopamine, ethanol, fentanyl, glyceryl trinitrate, heparin sodium, milrinone, morphine sulfate, pancuronium, potassium chloride, ranitidine, remifentanil, sodium nitroprusside, tigecycline, tirofiban, vecuronium.

Incompatibility

Fluids: Sodium bicarbonate.

Y-site: Aciclovir, aminophylline, ampicillin, atropine, azathioprine, calcium chloride, calcium gluconate, cefalotin, chloramphenicol, digoxin, ergometrine, ganciclovir, hyaluronidase, hydrocortisone sodium succinate, indomethacin, noradrenaline, phenobarbitone sodium, sodium bicarbonate, thiopentone, vancomycin.

Stability

Ampoule: Store below 30°C. Protect from light.
Diluted solution: Stable for 24 hours below 25°C.

Storage

Ampoule:
Store below 25°C.
Protect from light.
Discard remainder after use.

Special Comments

Ensure adrenaline has a “dedicated” line to avoid accidental bolus. Do not use as a side line with maintenance fluids.
Discard admixtures exhibiting colour change.

Evidence summary

Efficacy:
Treatment of hypotension in preterm infants: A single study of adrenaline 0.125–0.5 microgram/kg/minute versus dopamine 2.5–10 microgram/kg/minute reported they are equally effective at treating hypotension and increasing cerebral blood flow in very preterm infants. Adrenaline is associated with worse acid base status and increased hyperglycaemia. No difference in clinical outcomes was reported. [1–3] A single study of adrenaline 0.125, 0.250, 0.375, 0.5 microgram/kg/minute versus dopamine 5, 10, 15, 20 microgram/kg/minute reported dopamine reduced left ventricular output (LVO) 10% compared to a 14% increase in LVO with adrenaline. Dopamine and adrenaline caused significant increases in mean BP and pulmonary artery pressure. (LOE II, GOR C)

Infants and children with septic shock: Early administration of adrenaline 0.1–0.3 microgram/kg/minute was associated with increased survival compared to dopamine. [4] (LOE II, GOR B)

Vasopressors for hypotensive shock (newborns excluded): In treatment of hypotensive shock beyond the newborn period, there was no difference in mortality comparing adrenaline and other vasopressors (noradrenaline, noradrenaline and dobutamine, or noradrenaline and dopexamine). [5] (LOE I, GOR B)

Summary: Adrenaline may be used in hypotensive neonates with vasodilatory shock with or without myocardial dysfunction, particularly those with septic shock or unresponsive to other inotropes. (LOE II, GOR B)

Safety: Adrenaline may be associated with worse acid base status and increased hyperglycaemia. [3] Adrenaline is a potent vasoconstrictor. [6]

Pharmacokinetics: The onset of action is rapid and after intravenous infusion the half-life is approximately 5–10 minutes. [7] However, the half-life of intravenous adrenaline has not been reported in sick newborn infants. The plasma half-life of intratracheal adrenaline for newborn
Adrenaline (epinephrine) IV infusion

Resuscitation is likely to average approximately 50 minutes.[8]

References