

**AMILORIDE**

<b>Trade Name</b>	Amiloride (Biomed)
<b>Class</b>	Potassium sparing diuretic
<b>Mechanism of Action</b>	<p>Amiloride inhibits sodium resorption and induces natriuresis by direct inhibition of the sodium channel in the apical plasma membrane of the distal nephron. Excretion of potassium, hydrogen, calcium and magnesium ions is reduced.</p> <p>The potassium sparing effects of amiloride are most likely secondary to hyperpolarization of the apical plasma membrane and are not related to any effect on aldosterone.</p>
<b>Indications</b>	Control of fluid overload from heart failure
<b>Contraindications</b>	Hyperkalaemia, severe renal or hepatic impairment, anuria, Addison's Disease.
<b>Supplied As</b>	Amiloride Suspension 1mg/mL
<b>Dilution</b>	N/A
<b>Dosage</b>	100 – 200 microgram/kg/dose
<b>Interval</b>	12 hourly
<b>Administration</b>	Oral – give with food to reduce risk of stomach upset.
<b>Compatible With</b>	N/A
<b>Incompatible With</b>	N/A
<b>Interactions</b>	<p>Risk of <b>hyperkalaemia</b> if used in combination with:  Potassium supplements, ACE inhibitors (captopril)  Aldosterone antagonists (spironolactone)</p> <p>Risk of electrolyte disturbance when used with trimethoprim + sulphamethoxazole</p> <p>NSAIDS (indomethacin) may antagonise the diuretic effects and increase risk of acute renal injury and hyperkalaemia.</p> <p>May increase risk of toxicity if used in combination with digoxin</p>
<b>Monitoring</b>	Monitor renal function & electrolytes (especially sodium and potassium) prior to and during treatment.
<b>Stability</b>	As per date on the bottle, no preservative. Discard 7 days after opening
<b>Storage</b>	Refrigerate, shake before use
<b>Adverse Reactions</b>	Stomach upset, dry mouth, rash, cough, shortness of breath, confusion, muscle cramps, postural hypotension, hyperkalaemia, hyponatraemia, metabolic acidosis polyuria.

<b>Metabolism</b>	Amiloride is not metabolised in the liver. Approximately 50% is excreted as unchanged drug in urine and the other 50% is excreted in faeces.								
<b>Comments</b>	Amiloride liquid is an unregistered medicine in NZ and is subject to regulations of Section 29, please notify ward pharmacist for help with supply if required to be continued after discharge from NICU.								
<b>References</b>	<ol style="list-style-type: none"> <li>1. <a href="http://www.micromedex.hcn.net.au">www.micromedex.hcn.net.au</a></li> <li>2. BNF for Children 2009</li> <li>3. <a href="http://www.medsafe.govt.nz">www.medsafe.govt.nz</a></li> <li>4. New Zealand Formulary for Children (NZFC) <a href="http://www.nzf.org.nz">www.nzf.org.nz</a></li> </ol>								
<b>Updated By</b>	<table> <tr> <td>A Lynn, B Robertshawe</td> <td>December 2009</td> </tr> <tr> <td>A Lynn, B Robertshawe</td> <td>June 2012 (re-order profiles)</td> </tr> <tr> <td>A Lynn, M Wallenstein B Robertshawe, A Evison</td> <td>May 2020 (review&amp; update)</td> </tr> <tr> <td>A Lynn, B Robertshawe</td> <td>May 2023 (routine review)</td> </tr> </table>	A Lynn, B Robertshawe	December 2009	A Lynn, B Robertshawe	June 2012 (re-order profiles)	A Lynn, M Wallenstein B Robertshawe, A Evison	May 2020 (review& update)	A Lynn, B Robertshawe	May 2023 (routine review)
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