

Appendix 2: Antibiotic dosing guidelines

*NB Antibiotic doses in this guideline are appropriate for empirical treatment or sensitive organisms only. For any organism categorised as 'I' (Susceptible – increased exposure), seek further advice or refer to local policy for appropriate dose selection' (see NHS GGC Clinical Guidelines Portal)*

**PIPERACILLIN/TAZOBACTAM (FORMERLY TAZOCIN®):**

<b>Dosing &amp; Scheduling:</b>	<ul style="list-style-type: none"> <li>• 90 mg/kg (max 4.5g) four times a day (dose banded as per pharmacy chart)</li> <li>• Vial size 2.25g, 4.5g</li> <li>• Give by IV bolus over 3-5 mins*                             <ul style="list-style-type: none"> <li>* In cases where isolate is reported as 'I' (increased exposure), prolonged infusion may be required</li> </ul> </li> <li>• Renal Impairment:                             <ul style="list-style-type: none"> <li>▪ Dose adjustments required for GFR 50ml/min or less. Refer to Renal Drug Database</li> </ul> </li> </ul>
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**GENTAMICIN:**

<b>Dosing &amp; Scheduling:</b>	<p>Gentamicin dosing in patients with normal renal function:</p> <ul style="list-style-type: none"> <li>▪ Gentamicin 7mg/kg/once daily (max 500mg/dose)</li> <li>▪ Overweight/obese patients: Dose as per ideal or ideal adjusted body weight (Consult Pharmacy for further advice)</li> <li>▪ Trisomy 21 patients (<b>use with caution, discuss before prescribing</b>):</li> <li>▪ In patients treated with platinum compounds or high dose Methotrexate regimens, use only when clinically indicated</li> <li>▪ Give by IV infusion in 50 – 100ml Sodium Chloride 0.9% over 30 minutes</li> <li>▪ Measure level after 1<sup>st</sup> dose                             <ul style="list-style-type: none"> <li>Trough: Plasma samples at 18-24 hrs post-dose</li> </ul> </li> <li>▪ If trough level is &gt;1mg/L, the dosing interval is normally increased by 12 hours. Please discuss further dosing with pharmacy/microbiology</li> <li>▪ If trough level &gt;2mg/L, patient is unsuitable for pulsed dosing regimen, and subsequent doses should be guided by levels</li> <li>▪ If there is no change in dosage regimen or renal function, repeat trough levels every 4 days only</li> </ul> <p>Renal Impairment:</p> <ul style="list-style-type: none"> <li>▪ Use with caution. Dose reduction required for GFR 70ml/min or less</li> <li>▪ Initial dose:                             <ul style="list-style-type: none"> <li>In AKI, give a single dose of 5mg/kg, await trough before prescribing any further doses</li> <li>In chronic renal failure, give a single dose of 2.5mg/kg. Consult renal/pharmacy for monitoring and further dosing advice.</li> </ul> </li> </ul>
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**MEROPENEM:**

<b>Dosing &amp; Scheduling:</b>	<ul style="list-style-type: none"> <li>• 20 mg/kg tds (max 1g/dose). Can increase to 40 mg/kg in severe infections (max 2g/dose).</li> <li>• Give as IV bolus over 5 minutes or infuse over 15-30 minutes (dilute 1g in at least 50 ml Sodium Chloride 0.9% or Glucose 5%)</li> <li>• <i>Renal Impairment:</i> Dose adjustments required for GFR 50ml/min or less. Refer to Renal Drug Database.</li> </ul>
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**CIPROFLOXACIN:**

<b>Dosing &amp; Scheduling:</b>	<ul style="list-style-type: none"> <li>• Ciprofloxacin may be used in children where the benefit is considered to outweigh any potential risk.</li> <li>• <b>NB: Committee on Safety of Medicines (CSM) warning: Quinolone antibiotics may lower seizure threshold &amp; may induce convulsions in patients with or without previous history. There are rare reports of aortic aneurysm/dissection. In patients with known heart valve disease, careful benefit/risk assessment is essential</b></li> <li>• Tendon damage is a rare side effect of quinolone antibiotics. This risk is increased by concomitant use of steroids. If tendinitis is suspected, discontinue immediately.</li> <li>• Ciprofloxacin can prolong the QT interval – consider additive effect when used alongside other supportive therapies</li> <li>• Dosing regimen for treatment:             <ul style="list-style-type: none"> <li>▪ Intravenous: 1month – 18 years 10mg/kg three times a day (maximum dose of 400mg)</li> <li>▪ Oral: 20mg/kg bd (max 750mg/dose)</li> </ul> </li> <li>• Available preparations:             <ul style="list-style-type: none"> <li>▪ Tablets: 100mg, 250mg, 500mg, 750mg tablets;</li> <li>▪ Oral Suspension: 250mg/5ml</li> <li>▪ Premixed solution for IV infusion: 2mg/ml (50ml &amp; 100ml bags available)</li> </ul> </li> <li>• Oral absorption is good but do not use with oral Magnesium, Calcium, or Iron supplements as these affect absorption. Nasogastric feeds should be stopped for 2 hours before and after each dose.</li> <li>• Infuse undiluted IV over 30-60 minutes - flush with Sodium Chloride 0.9%.</li> <li>• Renal Impairment:             <ul style="list-style-type: none"> <li>▪ Dose adjustments required for GFR 30ml/min or less. Refer to Renal Drug Database</li> </ul> </li> </ul>
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**VANCOMYCIN:**

<b>Dosing &amp; Scheduling:</b>	<p><b>Intermittent Infusion – use Paediatric Vancomycin Infusion Chart</b></p> <p><i>Loading dose: &lt;6mo 15mg/kg, 6mo-18y 20mg/kg</i></p> <p><i>Maintenance dose: 0-6mo 10mg/kg 6hrly</i></p> <p style="padding-left: 100px;"><i>&gt;6mo-≤12y 15mg/kg 6hrly</i></p> <p style="padding-left: 100px;"><i>12-18y 15mg/kg 8hrly</i></p> <p><i>Prescribe on inpatient drug administration chart as 'see chart for dosing'.</i></p> <p><i>Prescribe all doses as per guidance on Vancomycin chart.</i></p> <p><i>Trough before 3<sup>rd</sup> dose, thereafter every 2-3 days</i></p> <p><i>Infuse over 2-3 hours</i></p> <p><b>Continuous Infusion</b></p> <p><i>Loading dose as for Intermittent Infusion (if switching from intermittent infusion, give 15mg/kg when next scheduled dose is due)</i></p> <p><i>Maintenance dose: start at 50mg/kg/24hrs immediately after loading dose</i></p> <p><i>Prescribe on inpatient drug administration chart as 'see chart for dosing'.</i></p> <p><i>Represcribe each day on fluid administration chart according to levels.</i></p> <p><i>Trough 18-24 hours after starting maintenance infusion</i></p> <ul style="list-style-type: none"> <li>• Target trough: 15-20mg/L</li> <li>• Renal Impairment:             <ul style="list-style-type: none"> <li>▪ Use with caution. Dose reduction required for GFR 50ml/min or less</li> </ul> </li> </ul>
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**TEICOPLANIN:**

<b>Dosing &amp; Scheduling:</b>	<ul style="list-style-type: none"> <li>• Child &gt; 2 months: 10mg/kg every 12 hours for 3 doses then 10mg/kg daily. Child &lt; 2 months: 16mg/kg for 1 dose, then 8mg/kg daily 24 hours after loading dose.</li> <li>• For <i>empirical</i> use, no proven organism – Max dose 800mg</li> <li>• For treatment of proven sensitive organism – Max dose 1200mg</li> <li>• Administration: Slow IV bolus or infusion over 30 mins diluted in Sodium Chloride.</li> <li>• Renal impairment:             <ul style="list-style-type: none"> <li>▪ Dose adjustments required for GFR 80ml/min or less. Refer to Renal Drug Database</li> <li>▪ No monitoring routinely required unless requested by microbiology</li> </ul> </li> </ul>
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**AMBISOME MEROPENEM:**

<b>Dosing &amp; Scheduling:</b>	<ul style="list-style-type: none"> <li>• Dosing regimen: dependent on clinical situation.</li> <li>• 3 mg/kg/day with proven or highly suspicious fungal infection (doses of up to 5 mg/kg/day have been used for proven infections).</li> <li>• Infuse in Glucose 5% ONLY at a concentration of 0.2-2.0 mg/ml. If dose <math>\geq 5</math>mg/kg infuse over 2 hours otherwise 1 hour.</li> <li>• Renal impairment:             <ul style="list-style-type: none"> <li>▪ No dose adjustments regardless of degree of renal impairment. Due to the size of AmBisome liposomes, there is no renal elimination</li> </ul> </li> </ul>
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