Intravenous Vancomycin use in Adults
Intermittent (Pulsed) Infusion

Background

This policy covers the use of intravenous vancomycin prescribed as an intermittent (pulsed) infusion. This can be used for treatment or prophylaxis. Evidence supporting this guidance is detailed below.

Vancomycin can also be given as a continuous infusion, when practical, for patients with severe or deep-seated infections (e.g. pneumonia, endocarditis, bone and joint infections) – refer to separate guidance.

This policy does not apply to the use of vancomycin in patients treated in Renal Units or receiving haemodialysis or haemofiltration.

Contra-indications and cautions

- Contra-indications to vancomycin therapy – hypersensitivity.
- Cautions for vancomycin therapy:
  - To avoid the risk of “red-neck/red-man syndrome”, pain or muscle spasm, ensure that the administration rate is not faster than 500 mg per hour.
  - Concurrent administration of neurotoxic and / or nephrotoxic agents increases the risk of vancomycin toxicity. Review therapy and consider amending or withholding nephrotoxic drugs during treatment with vancomycin. Where possible, avoid co-administration with the following:
    - amphotericin
    - potent diuretics
    - aminoglycosides
    - NSAIDs, and
    - ACE inhibitors.
  - The above list is not exhaustive – consult https://www.medicines.org.uk/emc/search?q=%22Vancomycin%22 for a full list
  - Patients with previous hearing loss due to potential ototoxicity.

Prescribing and documentation

- To ensure consistency, reduce risk and improve the prescribing of vancomycin, standardised charts (agreed nationally) should be used to document the prescription, administration and monitoring of intermittent vancomycin infusions. These should be used in conjunction with the existing inpatient prescription and administration record and the medical / nursing documentation.
- These charts contain a step-wise approach to safe and effective prescribing and key points of advice on monitoring, interpreting and re-prescribing.
- An Antimicrobial app and/or an online calculator is available in all boards and should be used to calculate the initial dose requirements.


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STEP 1: Prescribe the loading dose and maintenance dosage regimen

- To reduce the risk of mortality, commence vancomycin administration within 1 hour of recognising sepsis.
- If creatinine is known – use the online calculator (preferred method). The guidelines (below) in Table 1 (loading dose) and Table 2 (maintenance dose) can be used if the online calculator is not available. The dose amount and dosage interval are based on estimated creatinine clearance (Box 1) and actual body weight.
- If creatinine is not known – calculate and prescribe a loading dose based on actual body weight (Table 1). Calculate the maintenance dose once the creatinine is available.

Box 1: Estimation of creatinine clearance (CrCl)

The following ‘Cockcroft Gault’ equation can be used to estimate creatinine clearance (CrCl)

\[
\text{CrCl} = \frac{[140 - \text{age (years)}] \times \text{weight (kg)} \times 1.23 \text{ (male)} \text{ OR } 1.04 \text{ (female)}}{\text{serum creatinine (micromol/L)}}
\]

Cautions

- Use actual body weight or maximum body weight whichever is lower.
  - For maximum body weight table see https://www.sapg.scot/media/4471/maximum-body-weight-table.pdf
- In patients with low creatinine (< 60 micromol/L), use 60 micromol/L.
- Note: Use of estimated glomerular filtration rate (eGFR) is not recommended

Loading dose

Table 1: Initial vancomycin LOADING dose

<table>
<thead>
<tr>
<th>Actual body weight</th>
<th>Dose (mg)</th>
<th>Volume of sodium chloride (0.9%)*</th>
<th>Duration of infusion</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 40 kg</td>
<td>750 mg</td>
<td>250 mL</td>
<td>90 minutes</td>
</tr>
<tr>
<td>40 – 59 kg</td>
<td>1000 mg</td>
<td>250 mL</td>
<td>2 hours</td>
</tr>
<tr>
<td>60 – 90 kg</td>
<td>1500 mg</td>
<td>500 mL</td>
<td>3 hours</td>
</tr>
<tr>
<td>&gt; 90 kg</td>
<td>2000 mg</td>
<td>500 mL</td>
<td>4 hours</td>
</tr>
</tbody>
</table>

* Glucose 5% may be used in patients with sodium restriction.
Volumes used are for peripheral administration. More concentrated solutions (10mg/ml) must be given via a central line.

N.B. The loading dose is based on weight only so does not take account of renal function. When using the on-line calculator, on rare occasions a patient’s clearance of vancomycin may be so high that the maintenance dose is higher than the loading dose. In these circumstances, the loading dose should be the higher of the loading and maintenance doses i.e. if loading dose is calculated as lower than maintenance dose then give the maintenance dose as a loading dose instead.
Maintenance Dosage Regimen

- Give the first maintenance infusion 12, 24 or 48 hours after the loading infusion according to dose interval provided by the online calculator or Table 2 (below).

Table 2: Vancomycin MAINTENANCE dosage regimen

<table>
<thead>
<tr>
<th>CrCl (mL/min)</th>
<th>Dose amount</th>
<th>Volume of sodium chloride (0.9%)</th>
<th>Dose Interval</th>
<th>Time window for starting FIRST maintenance dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 20</td>
<td>500 mg over 1 hour</td>
<td>250 mL</td>
<td>48 hours</td>
<td>36-48 hours post loading dose</td>
</tr>
<tr>
<td>20 - 29</td>
<td>500 mg over 1 hour</td>
<td>250 mL</td>
<td>24 hours</td>
<td>18-24 hours post loading dose</td>
</tr>
<tr>
<td>30 - 39</td>
<td>750 mg over 1.5 hours</td>
<td>250 mL</td>
<td>24 hours</td>
<td>18-24 hours post loading dose</td>
</tr>
<tr>
<td>40 - 54</td>
<td>500 mg over 1 hour</td>
<td>250 mL</td>
<td>12 hours</td>
<td>6-12 hours post loading dose</td>
</tr>
<tr>
<td>55 - 74</td>
<td>750 mg over 1.5 hours</td>
<td>250 mL</td>
<td>12 hours</td>
<td>6-12 hours post loading dose</td>
</tr>
<tr>
<td>75 - 89</td>
<td>1000 mg over 2 hours</td>
<td>250 mL</td>
<td>12 hours</td>
<td>6-12 hours post loading dose</td>
</tr>
<tr>
<td>90 - 110</td>
<td>1250 mg over 2.5 hours</td>
<td>250 mL</td>
<td>12 hours</td>
<td>6-12 hours post loading dose</td>
</tr>
<tr>
<td>&gt;110</td>
<td>1500 mg over 3 hours</td>
<td>500 mL</td>
<td>12 hours</td>
<td>6-12 hours post loading dose</td>
</tr>
</tbody>
</table>

* Glucose 5% may be used in patients with sodium restriction.

Doses up to 2000 mg can be diluted in 500 mL fluid for peripheral administration. More concentrated solutions (10mg/ml) must be given via a central line.

The daily dose can be split into 3 equal doses and given 8 hourly. This approach is especially useful for patients who require high doses as it produces higher trough concentrations.

For example, 1500 mg 12 hourly (3000 mg per day) could be prescribed as 1000 mg 8 hourly and 750 mg 12 hourly (1500 mg per day) as 500 mg 8 hourly.

Note that patients who have unusual clinical characteristics, e.g. weight < 40 kg, weight >120 kg, age >90 years may require dose adjustments and require close monitoring. Contact pharmacy for advice.

STEP 2: Monitor the vancomycin concentration and reassess the dosage regimen

Concentrations are meaningless unless the dose & sample times are recorded accurately

- Due to wide variability in the handling of vancomycin, early analysis of a vancomycin concentration is required to ensure that the dosage regimen is appropriate.
- Take a trough sample (pre-dose) within 48 hours of starting therapy then every 2 - 3 days, or daily if the patient has unstable renal function.
- Monitor creatinine daily.
- Record the exact time of all vancomycin samples on the vancomycin prescribing chart AND on the
If renal function is stable, give the next dose before the trough result is available. If renal function is deteriorating, withhold until the result is available then follow the advice in Table 3.

**Target vancomycin concentrations**

- **Target trough concentration range:** 10 – 15 mg/L
- **If the patient is seriously ill (severe or deep-seated infections), the target range is 15 – 20 mg/L.**
  If the measured concentration is < 15 mg/L, consider increasing the dose amount or reducing the dosage interval (see 8 hourly dosing above).
- **If the patient is failing to respond, seek advice from microbiology or an infection specialist.**

**Adjustment of the vancomycin dosage regimen**

- Always check that dosage history and sampling time are appropriate before interpreting the result.
- Seek advice from pharmacy or microbiology if you need help to interpret the result. Dose adjustment tables may be useful within local guidelines to support changes in dose or dosing interval based on measured level.

If in doubt, take another sample before modifying the dosage regimen and / or contract pharmacy for advice

If the measured concentration is unexpectedly HIGH or LOW, consider the following:

- Were the dose and sample times recorded accurately?
- Was the correct dose administered?
- Was the sample taken from the line used to administer the drug?
- Was the sample taken during drug administration?
- Has renal function declined or improved?
- Does the patient have oedema or ascites?

**General points**

- Record the exact times of all measured concentrations on the vancomycin prescription chart.
- Undertake pre-prescribing checks (Box 2) to assess the risk of toxicity.
- Reassess the dose and continue or prescribe a dosage change.
- Document the action taken in the medical notes.
- Review the need for vancomycin daily.

**Box 2: Toxicity**

- Monitor creatinine daily. Seek advice if renal function is unstable (change in creatinine level)
- Signs of renal toxicity include increase in creatinine or decrease in urine output / oliguria
- Consider an alternative agent if creatinine is rising or the patient becomes oliguric.
- Vancomycin may increase the risk of aminoglycoside induced ototoxicity – use caution if co-prescribing.
Managing delays in vancomycin dose administration

This guidance has been developed by pharmacists in NHS Greater Glasgow and Clyde and applies to situations where a patient has stable renal function and a dose of vancomycin has been delayed unintentionally (for example due to loss of intravenous access).

Refer to the Table 3a below if dose unintentionally delayed by ≤50% of the dosing interval. Refer to Table 3b below if dose unintentionally delayed by >50% of the dosing interval.

Table 3a: Vancomycin dose - unintended delay of ≤50% of dosing interval

<table>
<thead>
<tr>
<th>Dose interval</th>
<th>Dose delay</th>
<th>Action</th>
</tr>
</thead>
<tbody>
<tr>
<td>12 hourly</td>
<td>≤6 hours</td>
<td>• Give the delayed dose immediately.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Record the date and exact time of administration on the chart and kardex with two nurse signatures.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Give the next dose at originally prescribed time.</td>
</tr>
<tr>
<td>24 hourly</td>
<td>≤12 hours</td>
<td></td>
</tr>
<tr>
<td>48 hourly</td>
<td>≤24 hours</td>
<td></td>
</tr>
</tbody>
</table>

Table 3b: Vancomycin dose - unintended delay >50% of dosing interval

<table>
<thead>
<tr>
<th>Dose interval</th>
<th>Dose delay</th>
<th>Action</th>
</tr>
</thead>
<tbody>
<tr>
<td>12 hourly</td>
<td>&gt;6 hours</td>
<td>• Give the delayed dose as soon as possible.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Record the date and exact time of administration on the chart and kardex with two nurse signatures.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Seek advice from pharmacy for further dosing.</td>
</tr>
<tr>
<td>24 hourly</td>
<td>&gt;12 hours</td>
<td></td>
</tr>
<tr>
<td>48 hourly</td>
<td>&gt;24 hours</td>
<td></td>
</tr>
</tbody>
</table>

This guidance below does not apply where the dose has been deliberately withheld (for example due to a high vancomycin trough level or deteriorating renal function). Contact pharmacy for advice in these cases. If you are unsure how to determine if the patient has deteriorating renal function, contact medical or pharmacy staff.