

Optimizing Vancomycin Dosing & Monitoring

BOTTOM LINE: Use guideline based vancomycin dosing and monitoring to maximize treatment success and reduce unnecessary serum vancomycin levels and needless dosage changes.

Recommendations:

<p>Adult Dosing^{1,2}</p> <p>For more details and pediatric dosing, see Vancomycin Dosing</p> <p>Clcr = creatinine clearance</p>	<p>Loading dose¹⁻⁵:</p> <ul style="list-style-type: none"> Use a loading dose in: <ul style="list-style-type: none"> <u>serious infections</u> where rapid attainment of target trough level of 15-20 mg/L is desired, e.g. vertebral osteomyelitis, MRSA pneumonia, epidural abscess, septic shock <u>patients with significant renal dysfunction</u> in order to decrease the time required to attain target trough level 25-30 mg/kg (based on actual body weight; no maximum dose) IV single dose, followed by maintenance dose separated by recommended dosing interval <p>Maintenance dose: 15 mg/kg (based on actual body weight) IV/dose (maximum of 2 g/dose)</p> <ul style="list-style-type: none"> Doses greater than 500 mg - round to nearest 250 mg Doses less than 500 mg - round to nearest 50 mg <p>Dosing interval:</p> <table border="1" data-bbox="440 793 1435 1024"> <thead> <tr> <th>Calculated Clcr (mL/min)</th> <th>Dosing Interval for trough 10-20mg/L</th> <th>Dosing Interval for trough 15-20mg/L</th> </tr> </thead> <tbody> <tr> <td>80 or greater</td> <td>q12h</td> <td>q8h</td> </tr> <tr> <td>40 - 80</td> <td>q24h</td> <td>q12h</td> </tr> <tr> <td>20 - 40</td> <td>q36h</td> <td>q24h</td> </tr> <tr> <td>10 - 20</td> <td>q48h</td> <td>q48h</td> </tr> <tr> <td>Less than 10</td> <td colspan="2">Consider loading dose. Obtain pharmacist consult.</td> </tr> </tbody> </table>	Calculated Clcr (mL/min)	Dosing Interval for trough 10-20mg/L	Dosing Interval for trough 15-20mg/L	80 or greater	q12h	q8h	40 - 80	q24h	q12h	20 - 40	q36h	q24h	10 - 20	q48h	q48h	Less than 10	Consider loading dose. Obtain pharmacist consult.	
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<p>Levels^{1,2}</p> <p>For more details, see Vancomycin Monitoring</p> <table border="1" data-bbox="87 1178 355 1497"> <thead> <tr> <th>Infection</th> <th>Desired Trough Level (mg/L)</th> </tr> </thead> <tbody> <tr> <td>Osteomyelitis</td> <td rowspan="6">15-20</td> </tr> <tr> <td>Pneumonia</td> </tr> <tr> <td>CNS infections</td> </tr> <tr> <td>Endocarditis</td> </tr> <tr> <td>Bacteremia</td> </tr> <tr> <td>Serious MRSA infections</td> </tr> <tr> <td>Other infections</td> <td>10-20</td> </tr> </tbody> </table> <p>*Steady state (SS) occurs in 4 to 5 half lives and can be estimated for vancomycin by using the following equations: $k_e = Clcr * 0.00083 + 0.0044$ $t_{1/2} = 0.693 / k_e$ $SS = 4-5 * t_{1/2}$</p>	Infection	Desired Trough Level (mg/L)	Osteomyelitis	15-20	Pneumonia	CNS infections	Endocarditis	Bacteremia	Serious MRSA infections	Other infections	10-20	<ul style="list-style-type: none"> Peak (post) levels are NOT recommended. Trough (pre) levels (taken 30 minutes or less prior to next dose) are recommended in: <ul style="list-style-type: none"> patients with deteriorating/unstable renal function (increase in baseline Scr of 40 µmol/L or greater, or increase of 50% or more from baseline) morbidly obese patients [190% or greater of ideal body weight or BMI 40kg/m² or greater] patients with anticipated therapy greater than/equal to 7 days⁶ patients who are severely ill (i.e. sepsis) and/or require target trough of 15-20mg/L patients with altered volume of distribution or clearance of vancomycin (e.g. cystic fibrosis, pediatrics, elderly 60 years or older, cancer, burns more than 20% BSA) selected dialysis patients [e.g. high flux and continuous hemodialysis/filtration. First trough level should be taken at steady state* and after at least 2 maintenance doses (~ 30 hours if normal renal function: prior to 4th dose if q12h or prior to 5th dose if q8h). <ul style="list-style-type: none"> vancomycin clearance is enhanced in obesity. For morbidly obese patients, consider drawing first level sooner (e.g. before 2nd or 3rd dose). Subsequent trough levels: <ul style="list-style-type: none"> with dosage change: trough should be taken at new steady state* as described above once target trough achieved: trough should be taken every 7-10 days in hemodynamically stable patients; may need more frequently if hemodynamically unstable, renal function changing, or patient is on concurrent nephrotoxic drugs <p>NB: Do NOT hold next vancomycin dose while waiting for results of serum levels unless there is a specific order to do so, e.g. because of concerns of toxicity/adverse events and/or significant decline in kidney function.</p>							
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References:

- Rybak M, Lomaestro B, Rotschafer JC, et al. Therapeutic monitoring of vancomycin in adult patients: a consensus review of the ASHP, IDSA, and SIDP. Am J Health-Syst Pharm 2009;66:82-98.
- Blondel-Hill E, Fryters S. Bugs & Drugs website <http://bugsanddrugs.albertahealthservices.ca>. ©1998-2015 Alberta Health Services Accessed September 21, 2015.
- Mohammedi I, Descloux E, Argaud, et al. Loading dose of vancomycin in critically ill patients: 15 mg/kg is a better choice than 500 mg. Int J Antimicrob Agents 2006;27:259-62.
- Wang JT, Fang CT, Chen YC, et al. Necessity of a loading dose when using vancomycin in critically ill patients. J Antimicrob Chemother 2001;47:246.
- Matzke GR, McGory RW, Halstenon CE, et al. Pharmacokinetics of vancomycin in patients with various degrees of renal function. Antimicrob Agents Chemother 1984;25:433-7.
- Contreiras C, Legal M, Lau TTY, et al. Identification of risk factors for nephrotoxicity in patients receiving extended-duration, high-trough vancomycin therapy. CJHP 2014;67:126-32.

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