



Antimicrobial Reference Laboratory

GUIDELINE RANGES FOR TDM

2017

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Antimicrobial Reference Laboratory – Guideline Ranges 2017

Agent	Risk group	Expected levels (Guide-lines) (mg/L)	Re-assay interval* (days)
Aminoglycosides			
Gentamicin Tobramycin (Once-daily) ^a	All patients on 2nd-4th dose; earlier if changing renal function or other risk factors.	Pre <1 mg/L Post >10 mg/L or 8 h post (4.5 mg/kg) 1.5-6 mg/L or follow Hartford nomogram (but note this is for 7 mg/kg)	6-8
Gentamicin (Once-daily 5 mg/kg) ^b	Neonatal sepsis	Pre < 2mg/L BUT <1 mg/L after 3 rd dose Post >8mg/L	
Gentamicin Tobramycin (BD or TDS) ^{c,d}	All patients on 2nd-4th dose; earlier if changing renal function or other risk factors.	Gm- pneumonia Pre <2 mg/L Post >7 mg/L Infective endocarditis (IE) Pre <1mg/L Post 3-5 mg/L	3-7
Amikacin (Once-daily) ^a		Pre <5, Post >50	6-8
Amikacin (BD or TDS) ^c		Pre <10, Post >20	3-7
Streptomycin (7.5 mg/kg BD) ^{d,e}	All patients after 2nd-4th dose.	Infective endocarditis Pre <3.0 mg/L Post 10-25 mg/L	7-28
<p>* Assuming initial results are within the expected range</p> <p>^a Nicolau et al. 1995. Antimicrobial Agents & Chemotherapy 39:650-655.</p> <p>^b NICE Clinical Guideline 149, 2012.</p> <p>^c British National Formulary, Edition 67. 2014 section 5.1.4.</p> <p>^d Elliott et al. 2004. Journal of Antimicrobial Chemotherapy 54: 971-81.</p> <p>^e Note: these are different to the AHA Scientific Statement ranges. Baddour et al. 2015. Circulation 132:1435-86.</p>			

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Glycopeptides/Lipopeptides/Oxazolidinones			
Vancomycin ^{a-d}	All patients on >2-4 days therapy. Patients receiving other nephrotoxic drugs. Assay at 2nd-4th dose.	Pre dose 10-15 mg/L but 15-20 mg/L in complicated infection OR Steady state during continuous infusion 15-25 mg/L	6-8
Teicoplanin ^{e-f}	<i>Staph. aureus</i> a) Skin and soft tissue infection b) Bone and Joint infection d) Infective endocarditis e) OPAT on 25 mg/kg 3x per week	Pre 15-30 but <60 mg/L Pre 20-40 but <60 mg/L Pre 30-40 but <60 mg/L Pre 20-30 mg/L	6-8
Daptomycin ^g	Patients with CPK elevation, high dose therapy (>6 mg/kg) or renal impairment	Pre dose 5-20 mg/L Or Pre dose 10-20mg/L in severe sepsis	6-8
Linezolid (600mg BD) ^{h,i}	Patients on long-term therapy (>28d) or if on agents with potential drug interactions	Pre 3-8 mg/L	8-16
<p>* Assuming initial results are within the expected range ^aJeffres et al. 2006. Chest 130: 947-55. Lodise et al. 2008. Antimicrobial Agents & Chemotherapy 52: 1330-1336. ^bBritish National Formulary. 2008. Number 55. Rybak et al. 2009. Am J Health-Syst Pharm. 66:82-98. ^cIngram et al. 2008. Journal of Antimicrobial Chemotherapy 62: 168-171. ^dWysocki et al, 2001. Antimicrobial Agents and Chemotherapy 45: 2460-2467. ^eTeicoplanin: Summary of Product Characteristics. 2013. European Medicines Agency. Assessment report: Targocid and associated names. 2014. EMEA/H/A-30/1301. European Medicines Agency. ^fLamont et al, 2009. . Journal of Antimicrobial Chemotherapy 64: 181-187. ^gBhavnani et al. 2010. Clinical Infectious Diseases 50: 1568-74. Falcone et al. 2013. J. Infection Chemotherapy 19 :732-9, DiPaolo et al. 2013. Int J. Antimicrobial Agents 42 :250-5, Falcone et al. 2013. CID 57 :1568-76, Reiber et al. 2015 Therapeutic Drug Monitoring, 37 :634-40. . ^hPea et al. 2012. JAC 67:2034-42. Dong et al. 2014. Eur J. Clinical Microbiology & Infectious Diseases, Epub 12/02/14 ⁱMatsumoto et al. 2014. International Journal of Antimicrobial Agents 44:242-7.</p>			

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Antifungal agents			
Flucytosine ^a	Routine with 72h of starting therapy.	Pre 20-40 mg/L Post 50-100mg/L Pre dose concentrations <20 mg/L have been associated with treatment failure and emergence of resistance. Post dose concentrations >100 mg/L have been associated with toxicity.	4-8
Isavuconazole ^c	Not routinely monitored but may be useful in complex cases or in renal impairment	Pre 1-4 mg/L	4-8
Itraconazole ^{a-b}	Routine in 1 st week of therapy. Measure 4-7 days after starting therapy	By Chromatographic assay Prophylaxis pre 0.5-1.0 mg/L Therapy pre 1.0-2.0mg/L NB. These guidelines are different to those achieved by bio-assay.	4-8
Fluconazole ^a	Not routinely monitored but may be useful in complex cases or renal failure	AUC:MIC ratio of >100, call for advice on sampling.	4-8
Posaconazole ^{a-c}	Routine in majority of patients. Measure 3-8 days after starting therapy	Prophylaxis: Pre 0.7-1.5 mg/L Therapy: Pre 1.0-3.75 mg/L	4-8
Voriconazole ^{a,b,d}	Routinely within 5d of starting therapy	Prophylaxis and therapy Pre 1.0-4.5 mg/L	4-8
<ul style="list-style-type: none"> • Assuming initial results are within the expected range ^aVermes et al. 2000. Journal of Antimicrobial Chemotherapy 46: 171-179. Ashbee et al. 2014. J. Antimicrobial Chemotherapy 69:1162-1176. ^bAndes et al. 2009. Antimicrobial Agents and Chemotherapy 53: 24-34. Dolton et al. 2015. Current Opinion in Infectious Diseases 27:493-500. Chau et al. 2014 Intern Med J 44:1364-88. ^cDolton et al. 2012. Antimicrobial Agents and Chemotherapy 56: 2806-2813. Dekkers et al. 2016. Curr Fung Infect Rep 10:51-61 ^dPascual et al. 2012. Clinical Infectious Diseases 55:381-90. ^ePeixoto et al. 2014. JCM 52:1016-19. 			

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Agents used in Mycobacterial infection^a			
Streptomycin ^b (15 mg/kg OD)	All patients after 2nd-4th dose.	Pre <5 mg/L in <50y patients Pre <1 mg/L in >50y patients Post 15-40 mg/L	7-28d
Streptomycin ^c (25 mg/kg BIW)	All patients after 2nd-4th dose.	Pre <1 mg/L Post 65-80 mg/L	7-28d
Rifampicin ^c	Patients with poor clinical progression	Pre <0.5mg/L Post <4mg/L sub-therapeutic Post 4-8mg/L usually adequate Post 8-24mg/L ideal	Depending on levels & patient progression
Rifabutin ^d	Patients who fail to respond to treatment. Patients on agents with P450 interactions	Pre <0.1mg/L Post 0.45-0.9 mg/L	Depending on levels & patient progression
Levofloxacin ^d	Patients being treated for MDR TB.	Pre 0.5-2 mg/L Post 8-13 mg/L	Depending on levels & patient progression
Cycloserine ^d	All patients after 4th-6th dose.	Pre 10-20mg/L Post (3-4h) 20-35mg/L	10-30
Moxifloxacin ^d	Patients being treated for MDR TB.	Pre 0.3-0.7 mg/L, Post 3-5 mg/L	Depending on levels & patient progression
Linezolid ^e (600 mg OD oral)	Patients being treated for MDR TB.	Pre <5mg/L Post 12-26mg/L	Depending on levels & patient progression
<p>* Assuming initial results are within the expected range; BIW: twice a week ^a Assuming that patients are on standard (usually daily) therapy, for patients on intermittent therapy please call to discuss expected levels as these will vary depending on dosing regimen used. ^b British National Formulary, Edition 67. 2014 section 5.1.9. ^c Peloquin 2017. Microbiol Spectrum 5:1-8. Pasipanodya et al. 2013. J. Infectious Diseases 208:1464-73. ^d Holland et al. 2009. Pharmacotherapy 29:503-10. Srivastava et al. 2013. European Respiratory Journal, 42:1449-53. Ramachandran et al, 2015, Drug Safety, 38:253-69. Peloquin 2017. Microbiol Spectrum 5:1-8. Hwang et al.2013. Int J. Tuberc Lung Dis 17:1257-66. Park.et al. 2017. AACc 59:4429-4435 ^e Schechter et al. 2010. CID 50: 49-55; McGee et al. 2009. Antimicrobial Agents & Chemotherapy 53: 3981-3984. Dong et al. 2014. Eur J. Clinical Microbiology & Infectious Diseases, Epub 12/02/14</p>			

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Other agents			
Aciclovir and CMMG	Patients with renal impairment or on high dose therapy	There are too many dose regimens used to give single guideline ranges and interpretation of levels needs to be patient specific	6-8
Ganciclovir ^a	Young children, renally impairment or unstable renal function	Pre 0.5-1.0 mg/L Post 7-9 mg/L (ganciclovir) Post 5-7 mg/L (valganciclovir)	4-8
Chloramphenicol ^b	All patients but especially neonates.	Pre <10, Post (2h) 10-25	5-7
Co-trimoxazole (sulphamethoxazole + trimethoprim) ^c	High-dosage therapy (PCP) or renal impairment.	Sulphamethoxazole: Pre <100, Post 120-150 but <200 Trimethoprim: Pre 5-7, Post 5-10 but <20	6-8
Colistin ^d	Patients on IV treatment	Pre 2-4 mg/L	14-28
<ul style="list-style-type: none"> • Assuming initial results are within the expected range ^aLuck et al. 2011 International Journal of Antimicrobial Agents 37:445-448. ^b British National Formulary, Edition 67. 2014 section 5.1.7 ^c Joos et al. 1995. Antimicrobial Agents & Chemotherapy 39:2661-2666. ^d Nation et al. 2014. Lancet Infectious Diseases S1473-3099. 			

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