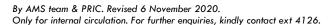
GENERAL CONSIDERATIONS

Optimal time to switch from IV to PO antibiotic: **48 to 96 hours** after IV antibiotic initiation

If patient **deteriorate** clinically after the conversion from IV to PO, IV therapy should be reinitiated

If patient fulfill the following criteria, MAKE a switch!

Manifestation of clinical improvement Afebrile (36 - 38°C) for 48 hours C-reactive protein (CRP) trending down White cell count (WCC) 4-12* x109/L or trending towards normal range No unexplained tachycardia (HR < 90 bpm) No unexplained hypotension (BP $\geq 90/60$ mmHg) and not vasopressor-dependent No tachypnoea (RR \leq 20 bpm) *Examine the patient's medications for potential cause of increase/sustained high WCC (eg. steroids) Able to tolerate oral therapy Not nil by mouth (NBM) Tolerating oral food or enteral feeding Oral absorption is not compromised by gastrointestinal conditions Good compliance, able to comply to oral medication at home Known indication for prolonged course of IV therapy? Conditions suitable for IV to PO switch: Conditions **not suitable** for IV to PO switch: **Endocarditis** Pneumonia × Central nervous system infections Skin & soft tissue infections (eg. meningitis, brain abscess) Urinary tract infection × MSSA bacteremia or fungemia Uncomplicated GNR bacteremia Intraabdominal infection without deep seated collections Conditions requiring approval from ID physician prior to IV to PO switch: Melioidosis Osteomyelitis Deep-seated infection Septic arthritis (eg. abscesses, empyema) Infected implant or prostheses Complicated orbital cellulitis Necrotising soft tissue infection





Equivalent oral antibiotic option?

Yes, available (Refer to page 2-3)

ANTIBIOTICS WITH **EQUIVALENT ORAL OPTIONS** (SEQUENTIAL THERAPY)

| Drug | For adults with normal renal function | | Bioavailability | |
|-----------------------------------|---|---|--|--|
| | IV dose | Equivalent oral dose | (BA) | Remarks |
| Amoxicillin/ Clavulanate | 1.2 g q8h | 625 mg TDS | Amoxicillin: 80% Clavulanate: 30-98% | - |
| Ampicillin/ Sulbactam | 1.5 g q8h 3 g q8h 3 g q6h | 375 mg BD 375-750 mg BD 750 mg BD | 80% | - |
| Azithromycin | 500 mg q24h | 500 mg OD | 34-52% | BA compensated by good tissue penetration |
| Cefuroxime | 750 mg q8h 1500 mg q8h | 250-500 mg BD 500 mg BD | 37-52% | 250 mg for sinusitis/pharyngitis, superficial SSTI and uncomplicated UTI |
| Ciprofloxacin* | 400 mg q12h 400 mg q8h | 500 mg BD 750 mg BD | 50-85% | - |
| Clindamycin | 600 mg q8h 600 mg q6h 900 mg q8h | 300 mg QID/ 600 mg TDS 600 mg QID 600 mg QID | ~ 90% | 300 mg QID for adult < 60 kg |
| Cloxacillin | 500 mg q6h 1000 mg q6h 2000 mg q6h | 250 mg QID 500 mg QID 1000 mg QID | ~50% | 250 mg for mild infections |
| Fluconazole | 200 mg q24h 400 mg q24h 800 mg q24h | 200 mg OD 400 mg OD 400 mg BD | > 90% | For opportunistic infections, dose may go up to 1200 mg/day |
| Levofloxacin* | 500 mg q24h 750 mg q24h | 500 mg OD 750 mg OD | ~ 99% | Hold tube feeds 1H before & 2H after levofloxacin given. |
| Linezolid | 600 mg q12h | 600 mg BD | ~100% | - |
| Metronidazole | 500 mg q12h 500 mg q8h 750 mg q8h | 400 mg BD 400 mg TDS 800 mg TDS | 100% | Abstain from alcohol to avoid disulfiram-like reaction |
| Trimethoprim/ sulfamethoxazole | 10-20 mg/kg/day (TMP) | 10-20 mg/kg/day (TMP) | 90-100% | - |

^{*} Space 2 hours before taking fluoroquinolones to avoid concomitant exposure to multivalent cations (Ca, Fe, Al, Mg, Zn) in dairy products, multivitamins and antacids. Cation chelate the drug and prevent absorption.



| For adu | | | |
|---|---|---|--|
| IV dose | Step-down oral dose | Bioavailability (BA) | |
| Benzylpenicillin 1-2 mega units q6h 3-4 mega units q4-6h | Phenoxymethylpenicillin 250 mg QID/500 mg BD 500 mg QID or Amoxicillin 500 mg TDS | Phenoxymethylpenicillin: 60-73% Amoxicillin: 80% | |
| Cefazolin 1 g q8h 2 g q8h | Cephalexin 500 mg QID 1000 mg QID | Cephalexin: 90% | |
| Cefepime | Amoxicillin/Clavulanate 625 mg TDS or Sultamicillin 750 mg BD | Amoxicillin: 80% Clavulanate: 30-98% Sultamicillin: 80% | |
| 2 g q8-12h | For definitive Pseudomonas infection: Ciprofloxacin* 500-750 mg BD | Ciprofloxacin: 50-85% | |
| Cefoperazone 1-2 g q12h | Amoxicillin/Clavulanate 625mg TDS or Sultamicillin 375-750 mg BD or Cefuroxime axetil 500 mg BD | Amoxicillin: 80% Clavulanate: 30-98% Sultamicillin: 80% Cefuroxime axetil: 37-52% | |
| | Amoxicillin/Clavulanate 625 mg TDS or Sultamicillin 750 mg BD | Amoxicillin: 80% Clavulanate: 30-98% Sultamicillin: 80% | |
| Ceftazidime | For definitive Pseudomonas infection: Ciprofloxacin* 500-750 mg BD | Ciprofloxacin: 50-85% | |
| 2 g q6-8h | For melioidosis: Trimethoprim/Sulphamethoxazole or Amoxicillin/Clavulanate Refer to HSgB Antibiotic Guideline 2019 for weight-based dosing & duration of antibiotic | TMP/SMZ: 90-100% Amoxicillin: 80% Clavulanate: 30-98% | |
| Amoxicillin/Clavulanate 625 mg TDS or Cefuroxime axetil 500 mg BD | | Amoxicillin: 80% Clavulanate: 30-98% Cefuroxime axetil: 37-52% | |
| Erythromycin Lactobionate 500 mg q6h 1000 mg q6h | Erythromycin Ethylsuccinate 400 mg QID 800 mg QID | Erythromycin: 18-45% | |
| Piperacillin/Tazobactam | Amoxicillin/Clavulanate 625 mg TDS or Sultamicillin 750 mg BD | Amoxicillin: 80% Clavulanate: 30-98% Sultamicillin: 80% | |
| 4.5 g q6-8h | For definitive Pseudomonas infection: Ciprofloxacin* 500-750 mg BD | Ciprofloxacin: 50-85% | |

^{*} Space 2 hours before taking fluoroquinolones to avoid concomitant exposure to multivalent cations (Ca, Fe, Al, Mg, Zn) in dairy products, multivitamins and antacids. Cation chelate the drug and prevent absorption.

References:

- Protocol on Antimicrobial Stewardship Program in Healthcare Facilities, MOH latest edition
- 2. Hospital Sungai Buloh Antibiotic Guideline 2019
- National Antibiotic Guideline 2019, MOH latest edition

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