## Antibiotics

#### Sources

- Keith Armitage, MD, MACP: https://www.youtube.com/watch?v=3XhBMg499\_w
- http://stritch.luc.edu/lumen/meded/therapy/pharm1\_blockiv\_2011.pdf

#### Key

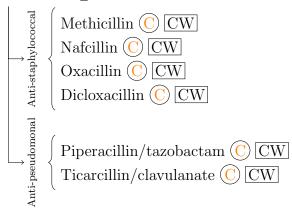
- \* Important to know (for non-ID docs)
- C Bactericidal
- S Bacteriostatic
- $\oplus$  Good coverage
- $\oslash$  Iffy coverage
- ⊖ Bad coverage
  - Mechanism of action:
    - CW Cell wall synthesis inhibitor
    - Protein synthesis inhibitor
    - NA Nucleic acid synthesis inhibitor
    - Met Metabolic inhibitor
  - Mechanism of resistance:
    - (Enz) Enzymatic degradation of drug
      - $\{\beta\}$   $\beta$ -lactamase
    - (PBP) Mutation of penicillin binding protein
    - (Mut) Mutation of the target
  - (Conc) Decrease in intracellular concentration (e.g., mutation in porins, development of efflux pumps)
    - (Seq) Sequestration of drug by proteins



### Penicillin C CW

- Still good against non-pneumococcal Strep (e.g., GAS, GCS) due to lack of resistance
- Resistance is mediated by PBPs, not  $\beta$ -lactamases, so resistance is overcome by increasing the dose, not by using  $\beta$ -lactamase inhibitors

#### Ampicillin (C) CW



# 

■ Not susceptible to  $\beta$ -lactamase

- ⊕ Staph (MSSA)
- $\oplus$  Strep
- $\oslash$  Gm- (E. coli, Proteus, Klebsiella)
- ⊖ Respiratory pathogens (Moraxella, H. influenzae, S. pneumoniae)
- → Anaerobes
- Bad CSF penetration

Generation 2 (C) (CW) (not important for non-ID docs) – cefuroxime, cefoxitin, cefotetan

- ⊕ Staph (MSSA)
- $\oplus$  Strep
- $\oplus$  Respiratory pathogens (Moraxella, H. influenzae, S. pneumoniae)
- Anaerobes
- ⊘ Gm−
- $\ominus$  Pseudomonas
- ⊖ Enterococcus
- Good CSF penetration

Generation 3 C CW – ceftriaxone\*

- ⊕ Staph (MSSA)
- $\oplus$  Strep
- Respiratory pathogens (Moraxella, H. influenzae, S. pneumoniae)
- Anaerobes
- ⊕ Gm-
- $\oslash$  Pseudomonas
- ⊖ Enterococcus
- Good CSF penetration

Generation 3.5 (C) (CW) – ceftazidime

- ⊕ Gm-
- $\oplus$  Pseudomonas
- $\ominus$  Gm+

Generation 4 (C) (CW) – cefepime\*

- ⊕ Gm−
- $\oplus$  Pseudomonas
- $\oplus$  Enterobacter
- ⊕ Gm+
- "Nosocomial cephalosporin"
- Neurotoxic (encephalopathy)

Generation 5 C CW – ceftaroline\*

- **•** MRSA
- Enterococcus
- $\bigcirc$  Pseudomonas
- → Healthcare-associated Gm-

## Penicillin Allergy

- $\blacksquare$  If immediate type I hypersensitivity (IgE)  $\to \underline{\rm not}$  safe to give cephalosporin
- Otherwise (other allergy or no allergy)  $\rightarrow$  safe

Carbapenems C CW Conc Enz PBP

• Not what they do cover, but what don't they cover?

Imipenem (C) CW

- $\oslash$  some Gm- rods
- ⊖ MRSA, VRE

Meropenem (C) CW • Safer than imipenem (esp., for people with seizure disorder) Doripenem (C) |CW| ■ Same as meropenem

Ertapenem (C) CW

- $\ominus$  Pseudomonas
- $\ominus$  Enterococcus
- Narrower than other carbapenems
- The once a day carbapenem

# Monobactams (C) $\overline{\text{CW}}$ $\{\beta\}$ (Conc)

• No immunologic cross-reactivity with  $\beta$ -lactams!

Aztreonam (C) CW

- ⊕ Aerobic Gm− rods
- ⊖ Gm+
- → Anaerobic

#### Aminoglycosides (C) Prot (Conc) (Enz) (Mut

- Most dangerous of the commonly used antibiotics; safe for a few days, not for >10-14
  - 1. Nephrotoxicity
  - 2a. Ototoxicity
  - 2b. Vestibular\* (the least reversible of the toxicities)
    - 3. Neuromuscular
- Toxicities usually temporary (if medication stopped early enough)
- Dosing: once per day
  - ▶ Except in neuropenic fever (controversial)
- ⊕ Gm− aerobes
- → Anaerobes

Gentamicin  $\bigcirc$  Prot, tobramycin  $\bigcirc$  Prot  $\rightarrow$  community-acquired infections

Amikacin (C) Prot  $\rightarrow$  nosocomial infections, resistant to aminoglycosidases

### Gm+ agents

Vancomycin (C) CW (Mut)

• Red man syndrome is an administration error, not an allergy

Linezolid (S) Prot (Mut)  Toxicities if used > 2 weeks
Daptomycin C CW Conc  Inactivated by surfactant (cannot be used for lung infections)
Tigecycline S Prot Conc Seq Enz  ⊖ Pseudomonas  ⊖ Proteus  Doxycycline S Prot Conc Seq Enz
Trimethoprim/sulfamethoxazole (TMP/SMX)
Rifampin C Prot Mut  Biofilm-associated infection  High resistance rates with monotherapy; should be paired with another drug  Revs up CYPs
Clindamycin $\bigcirc$ Prot $\bigcirc$ Mut $\bigcirc$ $\bigcirc$ Staph $\bigcirc$ Strep $\bigcirc$ Anaerobes $\bigcirc$ MRSA $\bigcirc$ Oral infections (high resistance)
Metronidazole (NA)

- Obligate anaerobes
- + Protozoa
- Can be paired with ceftriaxone (which covers facultative anaerobes)

## Quinolones $\bigcirc$ $\boxed{NA}$

Ciprofloxacin (C) NA

- ⊕ Gm− rods
- $\oplus$  Pseudomonas

Levofloxacin (C) NA
⊕ Gm− rods
$\oplus$ Pseudomonas
$\oplus$ Strep pneumoniae
Moxifloxacin C NA
⊕ Gm+
→ Gm-
Macrolides S Prot Conc Mut
Erythromycin $\textcircled{S}$ $\boxed{\operatorname{Prot}} \to \operatorname{GI}$ motility disorders
Clarithromycin (S) Prot
$\oplus$ Strep pneumoniae
Azithromycin (S) Prot
$\oplus$ Chlamydia (intracellular pathogen)
■ High intracellular concentration, low serum concentration
Anti-fungals
Azoles CW
Ketoconazole CW
Fluconazole CW
$\oplus \ \ Candida$
$\overline{\text{Itraconazole }\overline{\text{CW}}}$
Voriconazole CW
$\oplus \ Aspergillus$
→ Molds
Posaconazole CW
Isavuconazole CW

Amphoteric<br/>in B $\boxed{\text{CW}}$ 

 $\blacksquare$  For ID, BMT docs

Echinocandins  $\boxed{\text{CW}}$ 

 ${\rm Micafungin} \ \overline{\rm CW}$