

### 1. Sulfonamides

**MOA: Structural analog of PABA → Inhibit dihydropteroate synthase → inhibit DNA synthesis**

Sulfamethoxazole - crystalluria

Sulfisoxazole

Sulfasalazine – poorly absorbed – used for GI/IBD

Silver Sulfadiazine – Topical TX for burns

**PK:** Acetylation and Glucuronidation by the liver – renal elimination

**Spectrum:**

- G(+) – Strep, Staph, Listeria, **Nocardia**
- G(-) – Enterobacteriaceae, H. Infla, Moraxella
- **Mycobacteria**
- **Parasites**
- **Fungi** – Pneumocystis – Pentadine for allergic Pts

**Resistance** – Alterantive metabolic pathway

**SE:** Crystalluria, Hypersensitivity, Hemolytic anemia

(G6PD), aplastic anemia, Jaundice, kernicterus, hyperkalemia

### 2. Dihydrofolate reductase inhibitors

**MOA: inhibit conversion of folate to tetrahydrofolate → Inhibit DNA synthesis**

Trimethoprim + sulfonamide

**PK:** Well distributed, crosses BBB/prostate, accumulates in acidic fluid (vaginal) – Renal elimination

**Spectrum:**

- G(+) – Strep, Staph, Listeria, **Nocardia**
- G(-) – Enterobacteriaceae, H. Infla, Moraxella
- **Fungi** – Pneumocystis

**Resistance** – Altered drug target

**SE:** N/V/D, folate deficiency,

### 3. Metronidazole

**MOA: Activated by anaerobic organisms → causes DNA fragmentation**

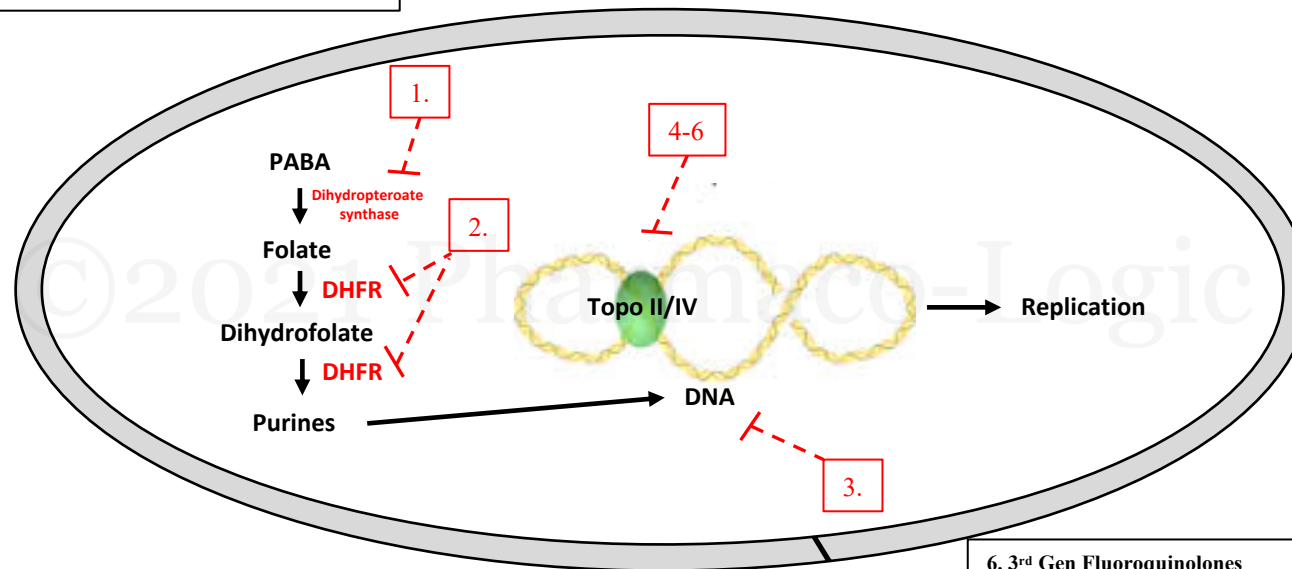
**PK:** Crosses the BBB, Renal – elimination

**Spectrum:**

- **Obligate anaerobes** – C. Difficile, B. fragilis
- Microaerophilic organisms – H. Pylori

**Resistance:** Obligate anaerobe - H. Pylori

**SE:** CNS effects, peripheral neuropathy, **Disulfiram effects**



### 4. 1<sup>st</sup> Gen Fluoroquinolones

**MOA: Inhibition of DNA gyrase (topoisomerase II/IV) → causes DNA breakage**

Norfloxacin

**PK:** renal excretion

**Spectrum** –

- Enterobacteriaceae
- Atypicals

### 5. 2<sup>nd</sup> Gen Fluoroquinolones

**MOA: Inhibition of DNA gyrase (topoisomerase II/IV) → causes DNA breakage**

Ciprofloxacin – CYP1A2 inhibitor

Ofloxacin

**PK:** Absorption decreased with cations, penetrates BBB and prostate – renal elimination

**Spectrum**

- G(+) – Bacillus anthracis, PRSP, CA-MRSA, enterococcus
- G(-) – Pseudomonas, Moraxella, enterobacteriaceae, H. influ., N. Meningitis, salmonella
- **Atypicals**
- **Mycobacteria** – MAC, TB, leprosy

**Resistance** – Decreased drug entry, Alterations in target enzymes

**SE:** Superinfections, CNS Effects, Photosensitivity, **cartilage deterioration** (quinolones in dem bones), Prolong QT, Hepatotoxicity

### 6. 3<sup>rd</sup> Gen Fluoroquinolones

**MOA: Inhibition of DNA gyrase (topoisomerase II/IV) → causes DNA breakage**

Levofloxacin

Moxifloxacin – Biliary Excretion

Gemifloxacin - Rash

**PK:** Absorption decreased with cations, penetrates BBB and prostate

**Spectrum:**

- G(+) – PRSP, S. aureus, enterococcus
- G(-) – Pseudomonas, Moraxella, enterobacteriaceae, H. influ., N. Meningitis
- **Atypicals**
- **Anaerobes**
- **Mycobacteria** – MDR-TB

**Resistance** – Drug target mutation

**SE:** Superinfections, CNS Effects, Photosensitivity, **cartilage deterioration** (quinolones in dem bones), Prolong QT, Hepatotoxicity