

# Fluoroquinolone Study Cheat Sheet

## Mechanisms of Action

1. Targets DNA gyrase (topoisomerase II), causing excess supercoiling that arrests DNA replication
2. Targets topoisomerase IV, interfering with separation of daughter DNA molecules

**Killing:** Generally considered to have concentration-dependent (e.g., AUC:MIC) bactericidal activity, however pharmacodynamics can vary from organism to organism

**NOTE:** Fluoroquinolones have different affinity for these targets. For example ciprofloxacin has more DNA gyrase activity, making it more targeted towards Gram negative organisms.

	<b>Ciprofloxacin</b>	<b>Levofloxacin</b>	<b>Moxifloxacin</b>	<b>Delafloxacin</b>
Brand name	Cipro <sup>®</sup>	Levaquin <sup>®</sup>	Avelox <sup>®</sup>	Baxdela <sup>®</sup>
FDA approval	1994	1996	1999	2017
Available IV	Yes	Yes	Yes	Yes
Available PO	Yes	Yes	Yes	Yes
Oral bioavailability	80%	100%	100%	67%
Typical adult dose	500mg PO BID 400mg IV BID	250-750mg PO or IV daily	400mg PO or IV daily	450mg PO BID or 300mg IV BID
Adjust doses for renal function	Yes	Yes	No	Yes
<i>Streptococcus pneumoniae</i> coverage	+	+++	+++	+++
<i>Pseudomonas aeruginosa</i> coverage	+++	++	+	+++
Anaerobic coverage	-	-	++	++
Atypical coverage	+++	+++	+++	+++

### Respiratory FQs

1. Levofloxacin
2. Moxifloxacin
3. Gemifloxacin

Named such for their activity versus the common respiratory pathogen *Streptococcus pneumoniae*

### Resistance Genes

GyrA  
GyrB  
ParC

### FDA Boxed Warnings

1. Tendonitis
2. Tendon rupture
3. CNS effects
4. Peripheral neuropathy
5. Worsening of myasthenia gravis

### Other Possible Side Effects

- QT interval prolongation
- Photosensitivity
- Hypoglycemia
- Altered mental health
- Diarrhea

- The base molecule for FQs is naladixic acid
- FQ use is linked to MRSA infection and FQ resistance in Gram negatives. Per the FDA, they are not 1<sup>st</sup>-line drugs for acute bacterial sinusitis, acute bacterial exacerbation of chronic bronchitis, or uncomplicated UTI.
- Generally avoid FQ monotherapy for Enterococci and Staphylococci. Delafloxacin may be an exception, having a role versus MRSA.
- In general avoid moxifloxacin for UTI due to low urine concentrations compared to other FQs
- FQs are widely considered the DOC for prostatitis due to excellent penetration into the prostate
- Beware *E. coli* resistance to fluoroquinolones
- FQs can have a role for treating Mycobacterium tuberculosis and non-tuberculosis mycobacteria
- FQs bind to cations (e.g., calcium, magnesium, aluminum) when given orally at the same time. Separate by taking 2 hours before or 6 hours after cation-containing medications or foods.
- Beware drug-drug interaction with warfarin
- FQs have a variety of ophthalmic and otic preparations

**Abbreviations:** ABX = antibiotics, DOC = drug of choice, FQ = fluoroquinolone, UTI = urinary tract infection