


38 – HIV Drug Resistance


Speaker: Michael Saag, MD



HIV Drug Resistance

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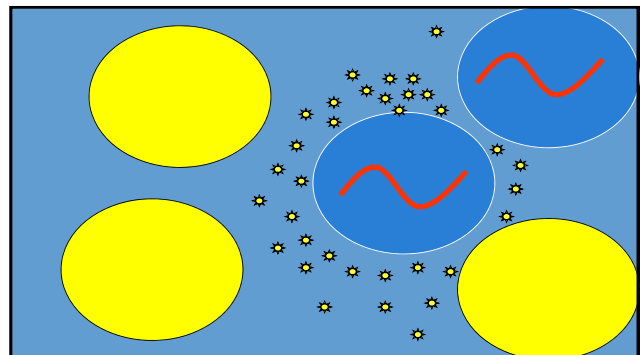
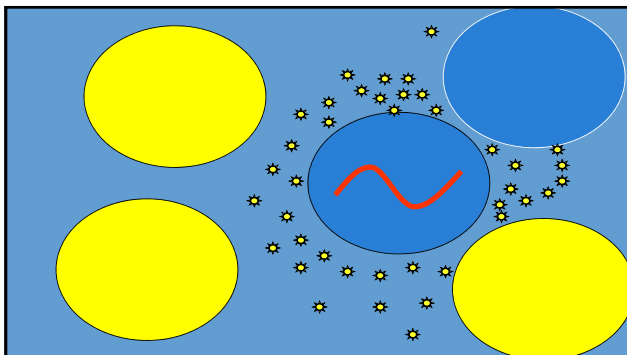
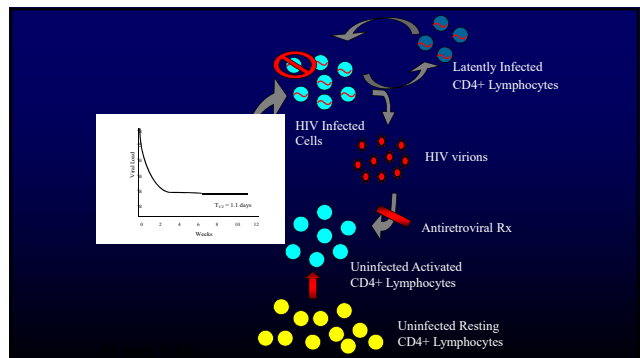
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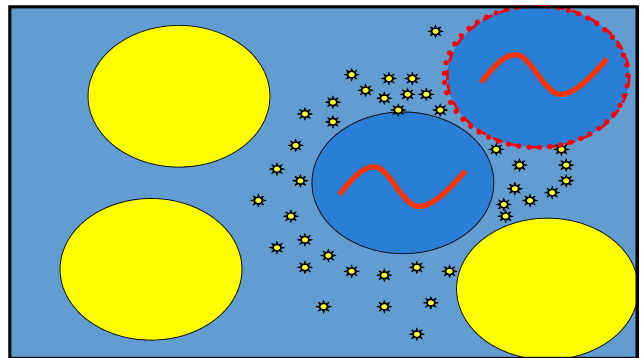
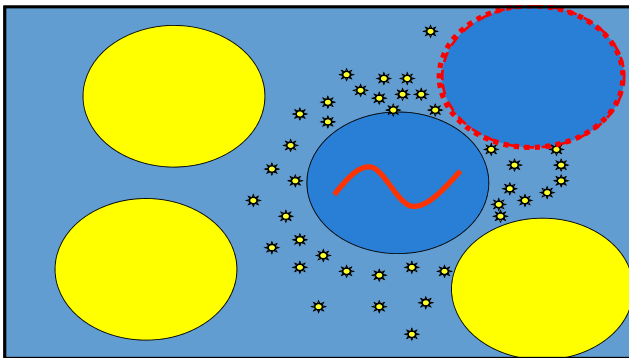
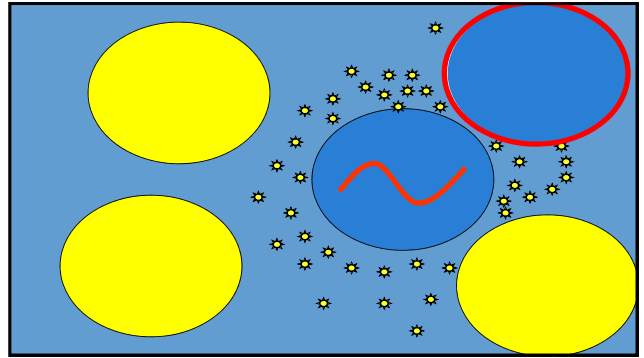
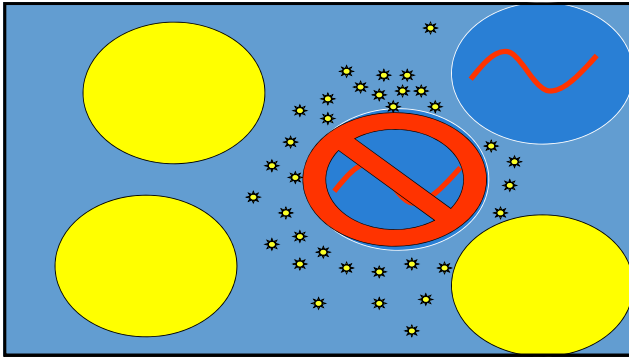
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How does resistance happen?



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Resistance Testing

- Genotypic resistance test
 - Perform test that gives mutations in viral genes
- Phenotypic resistance test
 - Perform test that describes growth of virus in the presence of anti-HIV drugs
- Limitations:
 - Cannot detect minority species (< 10% of viral population)

Key Issues in HIV Resistance

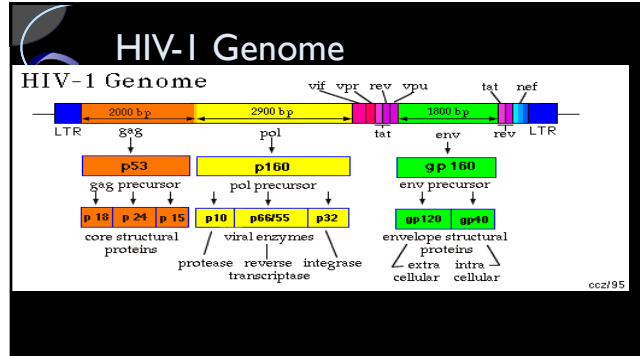
Easily Tested	Tough to Test
<ul style="list-style-type: none">• Specific Mutations• Cross – resistance• Prevalence of resistance at baseline	<ul style="list-style-type: none">• Definition of Phenotypes• Complex resistance patterns• Genetic Barrier• Nuances of Resistance• Relationship between Pk and Pd

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HIV Drug Resistance Testing

- Current guidelines recommend an **HIV genotype** as part of screening BEFORE ART is started.
- Following failure of 1st or 2nd regimens, **HIV genotype** is recommended to use with the history to choose the optimal next regimen.
- Following failure of 3rd and subsequent regimens, both **HIV genotype** AND **HIV phenotype** should be sent.
- If there is discordance between genotype and phenotype results, use the geno result (more sensitive).
- NOTE WELL:** Resistance mutations accrued from an earlier regimen **MAY NOT** be detected by tests obtained at the time of the current failing regimen



Mutation Nomenclature

Codon (position)
PR = 1-99 amino acids
RT = 1-560 amino acids

↓

M184V

Mutation Nomenclature

Codon (position)
PR = 1-99 amino acids
RT = 1-560 amino acids

↓

M184V

↙ Wild-type amino acid (consensus) ↘ Mutant amino acid

Alanine	A
Cysteine	C
Aspartate	D
Glutamate	E
Phenylalanine	F
Glycine	G
Histidine	H
Isoleucine	I
Lysine	K
Leucine	L
Methionine	M
Asparagine	N
Proline	P
Glutamine	Q
Arginine	R
Serine	S
Threonine	T
Valine	V
Tryptophan	W
Tyrosine	Y

Everything You Need to Know About Nucleoside Analog Resistance in One Slide!

Mutation	Selected by	Effects on other NRTIs
M184V	3TC, FTC	- Loss of susceptibility to 3TC, FTC - ↓ susceptibility to ABC, ddI (clinically insignificant) - Delayed TAMs and ↑ susceptibility to AZT, d4T, TDF
IABIs	AZT, d4T	- ↓ susceptibility to all NRTIs based on number of TAMs - More resistance with 41/210/215 than 67/70/219 pathway
Y115M, 69ins	AZT/ddI, ddI/d4T	- Resistance to all NRTIs - T69ins, TDF resistance
K65R	TDF, ABC, ddI	- Variable ↓ susceptibility to TDF, ABC, ddI (and 3TC, FTC) - ↑ susceptibility to AZT
74V	ABC, ddI	- ↓ susceptibility to ABC, ddI - ↓ susceptibility to AZT, TDF
44E, H81	AZT, d4T	- Increase NRTI resistance (with 41/210/215 pathway)

CASE 1

- 25 year old man presents with newly diagnosed HIV
- Had an episode c/w acute seroconversion syndrome 4 months ago
- Initial HIV RNA 40,000; CD4 443 cells/ul
- He wants to start ARV therapy

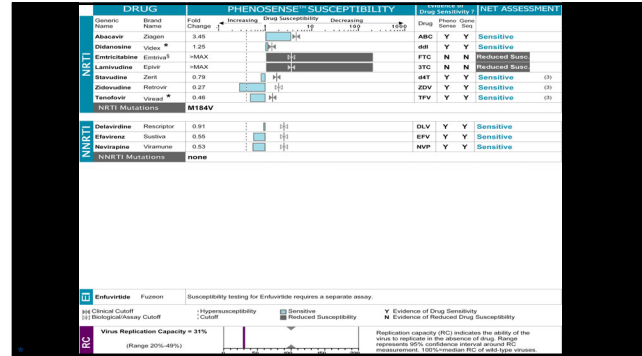
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Question #1

A baseline genotype is ordered that shows an M184V mutation. Which of the following drugs will have reduced susceptibility with this mutation?

- A. Efavirenz
- B. Zidovudine
- C. Tenofovir
- D. Etravirenz
- E. Emtricitabine



CASE 2

- 34 yo woman diagnosed with HIV 10 years ago
- Initially presented with PJP
- Initial Lab values
 - CD4 82 cells/uL
 - VL 106,000 c/mL
- Started on TDF / FTC / EFV (FDC)
- Did well for a while, then the regimen failed

Question #2

The genotype shows an M184V and K65R mutations. Which nRTI drugs would you include?

- A. ZDV
- B. TDF
- C. ddI
- D. ABC

DRUG	GENOTYPIC SUSCEPTIBILITY	PHENOTYPIC SUSCEPTIBILITY	NET ASSESSMENT
Abacavir	3.45	1.0	Reduced Susc
Didanosine	1.25	1.0	Reduced Susc
Emtricitabine	0.76	1.0	Sensitive
Lamivudine	0.76	1.0	Reduced Susc
Stavudine	0.27	1.0	Sensitive
Zalcitabine	0.45	1.0	Sensitive
Zidovudine	0.45	1.0	Sensitive
Tenofovir	0.45	1.0	Sensitive



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Non-nucleoside Reverse Transcriptase (NNRTI) Mutations

- **K103N** is the signature mutation for **efavirenz** (EFV).
- **Y181C** is the signature mutation for **nevirapine** (NVP).
- Older NNRTIs, efavirenz and nevirapine, have **low genetic barriers** (require only 1 mutation for resistance) and are **COMPLETELY** cross-resistant to one another.
- Newer NNRTIs, etravirine (ETR), rilpivirine (RPV), and doravirine (DOR) have higher barriers to resistance (require >1 mutation for resistance).
- **K103N** has no effect on etravirine susceptibility.
- **Rilpivirine** failure is associated with **E138K, K101E**, and/or **Y181C** and consequently, resistance to ALL NNRTIs.

CASE 3

- 34 yo woman diagnosed with HIV three years ago
- Initially presented with PJP
- Initial Lab values
 - CD4 82 cells/uL
 - VL 106,000 c/mL
- She was treated with TDF / FTC / ELV/ Cobi (FDC)
- The regimen failed after 12 months

Question #3

Which of the following mutations indicate high level resistance to elvitegravir ?

- Q148R
- L68I
- L68V
- K67N
- K65R

InSTI Resistance Mutations

Drug	66	67	70	74	105	107
Bictegravir TM	T	G	E	G	G	L
Cabotegravir TM	66	67	70	74	105	107
Dolutegravir TM	T	G	E	G	G	L
Elvitegravir TM	66	67	70	74	105	107
Raltegravir TM	74	89	97	121	140	142

Lenacapavir Resistance Mutations

MUTATIONS IN THE CAPSID GENE ASSOCIATED WITH RESISTANCE TO CAPSID INHIBITORS

Drug	L56	M66	Q67	K70	N74	A105	T107
Lenacapavir TM	I	I	H	S	S	R	R

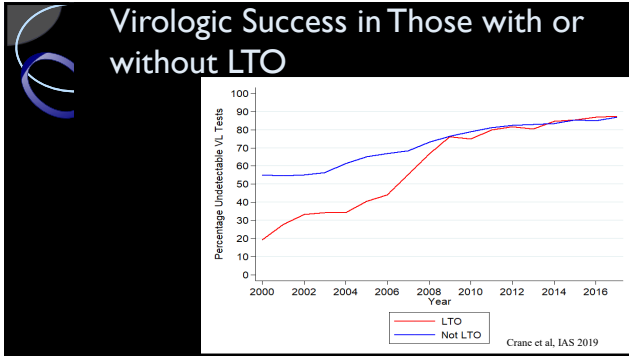
Case #4

PREVIEW QUESTION

- 34 yo MSM receiving CAB IM q 2 months for pre-exposure prophylaxis for last 6 months
- Asymptomatic
- HIV Ag/Ab test negative
- Routine screening: HIV RNA 6.1 c/ml

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Common Mutations To Memorize

• M184V/I	3TC and FTC
• M41L, D67N, K70R, L210W, T215Y, K219Q	"TAMS"
4 or more thymidine-analog mutations (TAMS) affect all approved nucleosides	
• K65R	tenofovir
• Q151M, 69SSS	multi-NRTI
• K103N	EFV (and NVP)
retains susceptibility to etravirine	
• Y181C	NVP and other NNRTI
• E138K, K101E	RPV and other NNRTI
• I50L	ATV
• N155H, Q148H/R/K	RAL and EVG
• Y143C	RAL
• R263K	DTG

- ### Summary
- High concern about resistance testing on Board Exams
 - Difficult to create test questions that do not require complex interpretation, have a single best answer, or are not 'multiple true-false'
 - Knowing common mutations and their role is a good way to prepare for the exam

