				Н	IV TREATME	ENT: SINGL	E TABLET H	REGIMEN	S		
	l	Brand	NRTI Ba	ckbone		Anchor Antir	etroviral		HIVinfo		
		lames	1 <sup>st</sup> NRTI	2 <sup>nd</sup> NRTI	Integrase Inhibitor	N-NRTI PI PK B		PK Booster	Rating*	Considerations	Monitor
	9883	Biktarvy	Emtricitabine 200mg	<b>Tenofovir T<u>A</u>F</b> 25mg	<b>Bictegravir</b> 50mg				A1	<ul> <li>✓ w/ or w/o food. Take 2 hrs before or after Ca/cations</li> <li>✓ Good Lipid profile- consider for high cardiac risk</li> <li>✓ Not recommended in &lt; 30ml/min, severe hepatic impairment. Cl w/ dofetilde or rifampin</li> <li>✓ Severe acute exacerbation of Hep B upon d/c</li> </ul>	Renal function
	572 Tri	LamivudineAbacavir300 mg600 mgEmtricitabineTenofovir TDE			<b>Dolutegravir</b> 50 mg				A1	<ul> <li>✓ W or w/o food. Take 2 hrs before or 6 hrs after Ca</li> <li>✓ HLA-B*5701 has to be -ve before giving abacavir</li> <li>✓ No major CYP drug interactions ☺</li> <li>✓ Largest size tablet</li> <li>✓ Cl w/ dofetilde or rifampin</li> </ul>	HLA-B*5701
<u>&gt;</u>	1	Stribild	200 mg 300 mg		Elvitegravir 150 mg			<b>Cobicistat</b> 150 mg	B1	<ul> <li>✓ Take with food. Take 2 hrs before/after Ca/cations</li> <li>✓ TDF → Can use until 70 mL/min</li> <li>✓ TAF → Can use until 30 mL/min</li> </ul>	Renal Function BMD Lipids
ce Daily	510	Genvoya	Emtricitabine 200 mg	<b>Tenofovir T<u>A</u>F</b> 10 mg	Elvitegravir 150 mg			<b>Cobicistat</b> 150 mg	B1	<ul> <li>✓ Cobi inhibits renal tubular secretion of creatinine</li> <li>✓ Cobi has many drug inx via CYP3A4 inhibition (avoid w/ drugs highly dependent on CYP3A4 clearance</li> </ul>	Renal Function Lipids
	SV 137	Dovato	Lamivudine 300mg	-	<b>Dolutegravir</b> 50 mg				A1 (*NOT if VL>500,000 or HBV)	<ul> <li>✓ W or w/o food. Take 2 hrs before or 6 hrs after Ca</li> <li>✓ &lt; 50ml/min or Child-Pugh C not recommended</li> <li>✓ Cl w/ dofetilide</li> </ul>	Renal function
- OD	SV J3T	Juluca		-	<b>Dolutegravir</b> 50mg	<b>Rilpivirine</b> 25mg			A1	<ul> <li>Maintenance Therapy—for those already virologically suppressed and no known resistance. Take with a meal</li> <li>A/E: HSR, Hepatotoxicity. Monitor for ADE if CrCL &lt; 30ml/min</li> <li>C/I: Dofetilide, PPI</li> </ul>	Renal Function, Liver Function
lablet.		Cabenuva			<b>Cabotegravir</b> 30 mg (po), 600/400 mg IM	<b>Rilpivirine</b> 25 mg (po), 900/600 mg IM			A1	<ul> <li>Maintenance Therapy—for those already virologically suppressed and no known resistance</li> <li>Optional Lead-in (≥28 days): CAB 30 mg/RPV 25 mg with a meal. Take antacid/cation 2 hrs before/4hrs after oral CAB</li> <li>Initiation injection: CAB 600/RPV 900 mg IM</li> <li>Monthly maintenance: CAB 600/RPV 900 mg IM</li> <li>Q2month maintenance: CAB 600/RPV 900 mg IM</li> </ul>	Injection site reactions, pyrexia, fatigue, headache
	\$ 776	Delstrigo	Lamivudine 300mg	Tenofovir TDF 300mg		<b>Doravirine</b> 100mg			B1	<ul> <li>✓ Not recommended in CrCl&lt; 50ml/min</li> <li>✓ w/ or w/o food</li> <li>✓ May exacerbate hepatitis upon discontinuation</li> <li>✓ Avoid w/ strong CYP3A4 inducers (ie Rifampin)</li> </ul>	Renal Function
	123	Atripla	EmtricitabineTenofovir TDF200 mg300 mg			<b>Efavirenz</b> 600 mg			B1	<ul> <li>✓ Keep in mind CNS adverse effects of Efavirenz</li> <li>✓ Not recommended CrCL &lt;50ml/min</li> <li>✓ C/I: bepridil, elbasvir/grazoprevir</li> </ul>	Renal Function Lipids
	GSI	Complera	Emtricitabine 200 mg	<b>Tenofovir TDF</b> 300 mg		<b>Rilpivirine</b> 25 mg			B1 (TDF), B2 (TAF),	<ul> <li>✓ Take with meal (~ 350 kcal) for abs'n of RPV</li> <li>✓ Use if HIV RNA &lt; 100,000 &amp; CD4 &gt; 200</li> <li>✓ Avoid: Acid suppressing (PPI C/I)</li> <li>✓ PPI C/I)</li> </ul>	Renal Function BMD
	255	Odefsey	Emtricitabine 200 mg	<b>Tenofovir T<u>A</u>F</b> 25 mg		<b>Rilpivirine</b> 25 mg			if VL<100,000 and CD4>200	<ul> <li>✓ RPV fewer CNS s/e compared to Efavirenz</li> <li>✓ RPV fewer rash and dyslipidemia than Efavirenz</li> </ul>	Renal Function
	8121	Symtuza	<b>Emtricitabine</b> 200mg	<b>Tenofovir T<u>A</u>F</b> 10mg			<b>Darunavir</b> 800mg	<b>Cobicistat</b> 150mg	A1	<ul> <li>✓ Take with food</li> <li>✓ Not recommended in CrCL &lt;30ml/min or Severe hepatic impairment</li> <li>✓ C/I: Alfuzosin, Amiodarone, Bepridil</li> </ul>	Renal Function

\*Strength of Recommendation: A=strong, B=moderate, C=optional. Quality of Evidence: I=>1 randomized trials with clinical outcomes/validated lab endpoints, II=>1 non-randomized trials/observational cohort studies with long-term clinical outcomes, III=expert opinion

				HI	V PREVENT	ION: Pre-E	xposure Propl	hylaxis (PrE	EP)	
Class	Generic		Brand	Prepar	rations	Dosing	Side Effects	Drug Interactions	Indicated Populations	Comments
Nucleoside / Nucleotide Reverse Transcriptase Inhibitors	Emtricitabine, tenofovir alafenamide	FTC, TAF	Descovy	225	Emtricitabine 200 mg/TAF 10 or 25 mg	1 tablet daily	Mostly Well Tolerated • N/V/D/Gas	TAF- Substrate of P-gp and BCRP	<ul> <li>recommended in gbMSM and transgender women, and for people who are at risk via receptive vaginal sex</li> </ul>	<ul> <li>✓ only combo also effective against Hep B</li> <li>✓ TAF has ↓ rates of renal insufficiency and bone mineral density reduction vs TDF</li> <li>✓ Not recommended if Clcr&lt;15 and not on hemodialysis (HD)</li> </ul>
	Emtricitabine, tenofovir disoproxil fumarate	FTC, TDF	Truvada	GILEAD	Emtricitabine 200 mg/TDF 300 mg	Daily dosing: 1 tablet daily <u>On-demand ("2-1- 1") dosing</u> : 2 tabs between 2-24 hours before sex, then 1 tab every 24 hours until 2 days after last sexual encounter	Mostly Well Tolerated • N/V/D/Gas • Renal impairment • Reduced bone density	Monitor renal function with concomitant use of other nephrotoxic agents (incl. chronic high-dose NSAIDS)	<ul> <li><u>Daily dosing</u>: HIV-negative individuals at risk of acquiring HIV</li> <li><u>On-demand dosing</u>: HIV- negative gbMSM</li> <li>NOT indicated for those who are at risk via receptive vaginal sex or for those who inject drugs</li> </ul>	<ul> <li>✓ only combo also effective against Hep B</li> <li>✓ Renal dosing: 1 tablet q2days if Clcr 30-49 mL/minute; not recommended if &lt;30 mL/min or HD</li> </ul>
Integrase inhibitors	Cabotegravir	САВ	Apretude	Accude Research and the second	Cabotegravir 200 mg/mL IM injection	Oral lead in (optional): 30 mg QD for 28 days Initiation (3mL): 600 mg CAB IM q1month x 2 consecutive months Maintenance (3mL): 600 mg CAB IM q2month	Well Tolerated Injection site reactions, pyrexia, fatigue, headache, MSK pain, nausea, dizziness, sleep problems, rash (mild), diarrhea	No CYP3A4 inx UGT1A1, UGT1A9 (minor), P-gp, BCRP substrate ↓ [CAB] with: Inducers of UGT1A1/3A4	<ul> <li>✓ HIV-negative individuals weighing at least 35 kg at risk of sexually acquired HIV</li> <li>✓</li> </ul>	<ul> <li>✓ CAB is 1<sup>st</sup> long acting injectable indicated for PrEP</li> <li>✓ Optional oral CAB as lead-in dosing (≥28 days) to assess tolerability or for use as oral bridging therapy for missed Apretude injections</li> <li>✓ C/I: carbamazepine, oxcarbazepine, phenobarbital, phenytoin, rifampin, rifapentine.</li> </ul>
Capsid inhibitor	Lenacapavir	LEN	Yezługo	Microsoft Addition	Lenacapavir 300 mg tab 309 mg/mL (1.5 mL vials) Approved in US	Initiation: Day 1: 927 mg SC and 600 mg po Day 2: 600 mg po Maintenance: 927 mg q6mo (i.e q26 weeks +/- 2 weeks)	<ul> <li>Injection site reactions</li> <li>nausea</li> </ul>	Substrate of CYP3A4, P-gp, UGT1A1. Supplemental LEN dosing required with moderate/strong CYP3A4 inducers. Moderate CYP3A4 inhibitor.	<ul> <li>✓ HIV-negative individuals at risk of sexual acquisition of HIV</li> <li>✓ People who inject drugs and have sexual exposures</li> </ul>	✓ If discontinuing LEN for PrEP, alternative methods of HIV prevention are recommended beginning 6 months after last injection if exposures are ongoing

						HIV Anti	retroviral ( <i>i</i>	ART) Medicatio	ons	
	Class	Generic		Brand	Preparations	Combo Pill	Dosing	Side Effects	Drug Interactions	Comments
	Combined NRT	I Tablet Formula	ations							
E	AIDSinfo rating: paired with INSTI: Dolutegravir A1 Raltegravir B1	Emtricitabine, tenofovir alafenamide	FTC, TAF	Descovy	225	Emtricitabine 200 mg/TAF 10 or 25 mg	1 tablet daily	Mostly Well Tolerated • N/V/D/Gas	TAF- Substrate of P- gp and BCRP	<ul> <li>✓ only combo also effective against Hep B</li> <li>✓ Better viral suppression than Kivexa if VL &gt; 100,000</li> <li>✓ TAF has ↓ rates of renal insufficiency and bone mineral density reduction vs TDF</li> <li>✓ If on a booster, use 10 mg TAF instead of 25 mg</li> <li>✓ Not recommended if Clcr&lt;30 mL/minute or hemodialysis (HD)</li> </ul>
ors - NRTI	or a boosted PI: Darunavir <b>A1</b> Atazanavir <b>B1</b>	Emtricitabine, tenofovir disoproxil fumarate	FTC, TDF	Truvada	GILEAD	Emtricitabine 200 mg/TDF 300 mg	1 tablet daily	Mostly Well Tolerated • N/V/D/Gas • Renal impairment • Reduced bone density	↓ [atazanavir]; need to boost	<ul> <li>✓ only combo also effective against Hep B</li> <li>✓ Better viral suppression than Kivexa if VL &gt; 100,000</li> <li>✓ Renal dosing: 1 tablet q2days if Clcr 30-49 mL/minute; not recommended if &lt;30 mL/min or HD</li> </ul>
Inhibit	<b>paired</b> with: Darunavir <mark>B2</mark>	abacavir, lamivudine	ABC, 3TC	Kivexa	GS FC2	Abacavir 600 mg/lamivudine 300 mg	1 tablet daily	Mostly Well Tolerated • Headache/N//D/malaise • Hypersensitivity reaction		<ul> <li>Abacavir not ideal for those with CV risk factors</li> <li>HLA needs to be negative before giving abacavir</li> <li>Comments also apply to Triumeq</li> </ul>
se	Single Agent N	<b>RTI Formulation</b>	IS							
Transcriptase Inhibitors	MOA: Analogues of nucleo(t)side which replace a base during reverse	Tenofovir alafenamide Adenosine analogue Nucleo <u>tide</u> Reverse Transcriptase Inhibitor (NtRTI)	TAF	<b>Vemlidy</b> (for chronic HBV)	GSI 25 mg tab	Descovy <sup>1</sup> QD Genvoya <sup>1</sup> QD Odefsey <sup>1</sup> QD Biktarvy <sup>1</sup> QD Symtuza <sup>1</sup> QD	25 mg po QD (10 mg po QD if using with booster) <mark>Renal</mark>	Mostly Well Tolerated • N/V/D/Gas	TAF- Substrate of P- gp and BCRP	<ul> <li>✓ TAF = tenofovir alafenamide (targeted pro-drug), <i>less</i> bone &amp; renal issues</li> <li>✓ safe until renal function with CrCl of 30 mL/min</li> <li>✓ Preferred agent in cases of co-infection with HBV</li> </ul>
/ Nucleotide Reverse 1	transcription of viral RNA to DNA → chain termination <u>Resistance:</u> - "low genetic	Tenofovir disoproxil fumarate Adenosine analogue Nucleo <u>tide</u> Reverse Transcriptase Inhibitor (NtRTI)	TDF	Viread	GILEAD 4331 150, 200, 250, 300 mg tab 40 mg/g powder	Truvada <sup>1 QD</sup> Stribild <sup>1 QD</sup> Complera <sup>1 QD</sup> Delstrigo <sup>1 QD</sup> Atripla <sup>1 QD</sup>	300 mg po QD <mark>Renal</mark> avoid TDF in CKD	Mostly Well Tolerated • N/V/D/Gas • Renal impairment <sup>TDF</sup> • Reduced bone density <sup>TDF</sup>	↓[atazanavir] ↑[didanosine - ddi] Clinically not used with TDF anyways any longer	<ul> <li>✓ TDF = tenofovir disoproxil fumarate (pro-drug), efficacy of TDF = TAF</li> <li>✓ Renal: &lt; 10 mL/min not recommended, 10 - 29 mL/min give 300 mg po q72-96h, 30-49 mL/min give 300 mg po q48h, ≥ 50 mL/min no adjustment</li> <li>✓ Preferred agent in cases of co-infection with HBV</li> <li>✓ Favorable lipid profile</li> </ul>
ucleoti	barrier to resistance" - many mutations confer cross	Emtricitabine Cytidine analogue	FT <mark>C</mark>	Emtriva	200 mg cap	With TAF or TDF products above	200 mg po QD <sup>cap</sup> 240 mg po QD <sup>sol'n</sup> <mark>Renal</mark>	Well Tolerated • Headache <sup>common</sup> , dizziness • N/D • Rash, skin pig'n	Lamuvidine [X] → both <b>Cytosine</b> analogues (no point in using both)	<ul> <li>✓ Black Box: severe exacerbation of hep B on stopping drug in pts w Hep B</li> <li>✓ Only part of combos w Tenofovir in Canada</li> <li>✓ Rarely pts may experience bad diarrhea. Headache most common s/e.</li> </ul>
Nucleoside / N	resistance to others in the class Renal Dosing: Use with caution & check for renal dosing for each	Lamivudine Cytidine analogue	зт <mark>с</mark>	зтс	150, 300 mg tab	Kivexa <sup>1 QD</sup> Triumeq <sup>1 QD</sup> Dovato <sup>1 QD</sup> Delstrigo <sup>1 QD</sup> Combivir <sup>1 BID</sup> Trizivir <sup>1 BID</sup>	150 mg po BID 300 mg po QD <mark>Renal</mark>	Well Tolerated • Headache <sup>beginning</sup> • N/D/Abd pain <sup>transient</sup> • Insomnia <sup>uncommon</sup> Pancreatitis <sup>more peds</sup>	Emtricitabine [X] → both <b>Cytosine</b> analogues (no point in using both)	<ul> <li>Some people have headache in first few days, stick with it and use Tylenol and Advil if needed</li> <li>May exacerbate Hep B upon discontinuation</li> </ul>
	dosing for each agent	<b>Abacavir</b> Guanosine analogue	ABC	Ziagen	GX 623 300 mg tab	Kivexa <sup>1 QD</sup> Triumeq <sup>1 QD</sup> Trizivir <sup>1 BID</sup>	300 mg po BID 600 mg po QD can safely use in CKD	Common: • Headache, N/D, malaise Serious: • Hypersensitivity reaction (HSR)		<ul> <li>✓ Black Box: Only Rx for HLA-B*5701 negatives → Testing predicts HR in Caucasians. Rechallenge in HSR patients C/I → life threatening</li> <li>✓ Signs of HSR: fever, rash, tired, upset stomach, vomit, belly pain, flu-like sx, sore throat, cough. Occurs &lt; 6 wks after start (mean 11 days). Stop ASAP &amp; see MD.</li> <li>✓ Meta-analysis → no sign of ↑ MI → but if higher MI risk, ABC not best choice</li> <li>✓ Can cause hepatitis and lactic acidosis esp in women and obese</li> </ul>

	HIV Antiretroviral (ART) Medications											
Cla	ISS	Generic		Brand	Preparations	Combo Pill	Dosing	Side Effects	Drug Interactions	Comments		
longer recom first-lii	ovudine no r imended as ne therapy ost patients	<b>Zidovudine</b> Thymidine analogue	AZ <mark>T</mark>	Retrovir	100, 250 mg cap 10 mg/mL syrup 10 mg/mL inject	Trizivir <sup>1 Bid</sup> Combivir <sup>1 Bid</sup>	300 mg po BID Also <b>I.V.</b> form <mark>Renal</mark>	Not Well Tolerated • Headache <sup>62%</sup> • N <sup>50%</sup> / V <sup>17%</sup> / Anorexia <sup>20%</sup> • Insomnia • Nail pigmentation • Hematologic toxicity	stavudine [X] also a thymidine analogue	<ul> <li>Black Box: hematologic toxicity, myopathy, anemia, granulocytopenia, thrombocytopenia</li> <li>Often in subtherapeutic mono- and dual therapy regimens</li> <li>Resistance likely in Long term survivors</li> <li>Place for therapy: IV form and syrup still used in MTCT in <i>pregnancy and delivery</i> and infants with HIV</li> <li>No longer recommended**</li> </ul>		

Cla	SS	Generic	C	Brand	Preparations	Combo Pill	Dosing	Side Effects	Drug Interactions	Comments
	Integrase Strand Transfer Inhibitors tegravir Favorable lipid	Bic <u>tegravir</u>	BIC	-	9883 (Biktarvy)	Biktarvy <sup>1 QD</sup>	50mg po QD	Well Tolerated • Headache • Nausea/Diarrhea • Insomnia	CYP3A & UGT1A1 substrate (~50:50) Inhibits OCT2 & MATE1 • ^[Metformin]	<ul> <li>Only exists in combination</li> <li>Increase serum creatinine due to tubular inhibition without affecting glomerular function (increases usually in the first 4 weeks with median increase of 9.96umol/L after 48 weeks)</li> <li>May increase bilirubin</li> <li>Interacting classes: anticonvulsants, rifamycins, atazanavir</li> <li>C/I: Dofetilide, rifampin, St. John's wort</li> </ul>
Transfer Inhibitors - INSTI	profile as a class <u>Resistance</u> : Low genetic barrier to resistance with RAL and EVG. Higher with BIC, CAB, DTG	Cabo <u>tegravir</u>	САВ	Vocabria	SV CTV 200 mg/mL inj 30 mg tab	Cabe <mark>nuva</mark> IM injection	Oral: 30 mg QD (+25 mg RPV) Initiation (3mL): 600 mg CAB/900 mg RPV IM Maintenance: 400 mg CAB/600 mg RPV IM monthly <u>or</u> 600 mg CAB/900 mg RPV IM q2months	Well Tolerated Injection site reactions, pyrexia, fatigue, headache, MSK pain, nausea, dizziness, sleep problems, rash (mild), diarrhea	No CYP3A4 inx UGT1A1, UGT1A9 (minor), P-gp, BCRP substrate ↓ [CAB/RPV] with: Inducers of UGT1A1/3A4	<ul> <li>✓ CAB/RPV is 1<sup>st</sup> long acting injectable combination indicated as a switch regimen in virologically suppressed patients</li> <li>✓ Optional oral CAB as lead-in dosing (≥28 days) to assess tolerability or for use as oral bridging therapy for missed Cabenuva injections</li> <li>✓ NB: initiation injections: one month initiation if using q1month maintance injections. For q2month maintenance, start with two initiation injections one month apart.</li> <li>✓ Oral CAB C/I: carbamazepine, oxcarbazepine, phenobarbital, phenytoin, rifampin, rifapentine. Cabenuva C/I: as above plus rifabutin, systemic dexamethasone (&gt;1 dose), St. John's wort</li> </ul>
Strand	Class Interaction: Oral absorption is diminished when co-administered with polyvalent cations (Mg, Ca, Al, Fe). • BIC: take 2 hrs apart or together with food	Dolu <u>tegravir</u>	DTG	Tivicay	50 mg tab Pediatric: 10 mg, 25 mg tab 5 mg dispersible tabs	Triumeq <sup>1 QD</sup> Juluca <sup>1 QD</sup> Dovato <sup>1 QD</sup>	50 mg po QD 50 mg po BID*	Well Tolerated • Insomnia • Headache • ↑ SCr small (^~0.11mg/dL)	No CYP3A4 inx P-gp, UGT1A1, CY3A4 <sup>(10-</sup> 15%)substrate Inhibits OCT2 - Metformin (inc 2 fold [metformin]) - C/I Dofetolide	<ul> <li>✓ Take with/without food</li> <li>✓ Inhibits renal tubular secretion of creatinine, SCr "falsely" increases</li> <li>✓ May cause neural tube defects if taken at the time of conception</li> <li>✓ Higher barrier to resistance than EVG or RAL</li> <li>✓ *BID dosing if heavily tx-experienced, INSTI resistant, or given w enzyme inducers</li> <li>✓ High efficacy in those with baseline HIV RNA &gt; 100,000 copies/mL</li> <li>✓ C/I: Dofetilide, fampridine</li> </ul>
Integrase	<ul> <li>CAB: take 2 hrs before/4 hrs after ORAL CAB</li> <li>DTG: take 2 hrs before/6 hrs after or together with food</li> <li>EVG: take 2 hrs</li> </ul>	Elvi <u>tegravir</u>	EVG	Vitekta	85, 150 mg tab	Stribild Genvoya	85-150 mg po QD <sub>boosted</sub> w/ food	Well Tolerated • Hyperlipidemia • D/N • Headache	CYP3A4 substrate induces 2C9 (EVG) Inhibits CYP3A4, P-gp, BCRP, OATP1B1/3, OCT2, MATE1 (cobi)	<ul> <li>✓ Better absorption w food/snack</li> <li>✓ Coformulated with PK booster cobicistat</li> <li>✓ Cobicistat inhibits tubular secretion of creatinine w/o affecting glomerular function (if &gt;35.36umol/L need renal monitoring)</li> <li>✓ Lower genetic barrier to resistance than PIs or DTG</li> <li>✓ C/I: Eplereone, Lovastatin</li> </ul>
	apart • RAL: avoid (only Ca OK with Isentress; not HD)	Ral <u>tegravir</u>	RAL	lsentress & Isentress HD	227 400 mg tab 600mg tab (HD)	None	400 mg po BID 1200 mg po QD new study QDMRK	Well Tolerated • Rash • N/D, Headache • Insomnia ↑ LFTs, ↑ CK, rhabdo	No CYP3A4 inx UGT1A1 substrate	<ul> <li>✓ Take without regards to meals</li> <li>✓ 1<sup>st</sup> to market INSTI → Being studied: 1200 mg po QD (given as 2X 600mg)</li> <li>✓ Aluminum or Magnesium antacids reduce abs'n RAL (Can take Ca Antacids if on Isentress, NOT Isentress-HD)</li> <li>✓ Lower genetic barrier to resistance than PIs or DTG</li> <li>✓ Avoid strong inducers of UGT (ie carbamazepine)</li> </ul>

	Class	Generio	5	Brand	Preparations	Combo Pill	Dosing	Side Effects	Drug Interactions	Comments
itors - NNRTI	NNRTI vir MOA:	Dora <u>vir</u> ine	DOR	Pifeltro	700 100mg tab	Delstrigo <sup>TDF 1 QD</sup>	100mg po OD	Well tolerated Common SE • Headache • Diarrhea, Ab pain • Abnormal Dreams	Cyp3A4 Substrate	<ul> <li>✓ Take BID if using with rifabutin</li> <li>✓ Taken without regards to food</li> <li>✓ Favourable lipid profile – consider for high cardiac risk</li> <li>✓ Avoid use with Strong inducers of CYP3A4 (ie Carbamazepine, rifampin)</li> <li>✓ C/I: carbamazepine, oxcarbazepine, phenobarbital, phenytoin, enzalutamide, rifampin, rifapentine, mitotane, St.John's wort</li> </ul>
	NNRTIS bind allosterically in a pocket located near the catalytic site in the palm domain of the p66 subunit site of the Reverse Transcriptase (RT) enzyme Resistance: Low genetic barrier to	Efa <u>vir</u> enz	EFV	Sustiva	SUSTIVA 600 mg tab 50, 200 mg cap	Atripla <sup>TDF 1 QD</sup>	600 mg po QD avoid fatty meals on empty stomach (inc abs'n leading to s/e)	<ul> <li>CNS S/E <sup>52%</sup></li> <li>Dizziness, <u>vivid dreams</u></li> <li>Insomnia, somnolence</li> <li>Impaired concentration</li> <li>Hyperlipidemia</li> <li>Rash 26% (can treat through it mostly)</li> </ul>	CYP3A4 & 286 Substrate Potent inducer of CYP3A4,286, UGT1A1; inhibitor of CYP2C9/2C19 ↓ [conc] of: • Benzos (-olam are issues, - pams are ok) • most opioids	<ul> <li>✓ Let MD know if history of psych illness → should avoid this med</li> <li>✓ Vivid dreams bothersome to some, enjoyable to some other</li> <li>✓ CNS s/e worst after 1<sup>st</sup> or 2<sup>nd</sup> dose, often improve in 2-4 weeks</li> <li>✓ Methadone: monitor for symptoms of opioid withdrawal</li> <li>✓ May cause false +ve cannabinoid test</li> <li>✓ Pregnancy: birth defects reported in primate studies but no evidence of</li></ul>
oside RT Inhibitors	resistance with first generation (EFV,NVP), but second generation often still active depending upon genotype.	Etra <u>vir</u> ine	ETR	Intelence	(200) 100, 200 mg tab	None	200 mg po BID or 400 mg po QD <mark>w/ food</mark>	<ul> <li>Rash 9%</li> <li>Dyslipidemia</li> <li>Nausea</li> <li>Rhabdomyolysis uncommon</li> </ul>	CYP3A4, 2C9, 2C19 substrate Weak inducer of CYP2B6/ 3A4 Weak Inhibitor of 2C9/ 2C19	<ul> <li>✓ Tabs are large: dissolve readily in water for liquid dosing, however whole tablet is chalky, large and often difficult to swallow.</li> <li>✓ Severe rash reported</li> <li>✓ C/I: ombitasvir/paritprevir/ritonavir and dasabuvir regimens</li> </ul>
Non-nucleoside		Ne <u>vir</u> apine	NVP	Viramune	200 mg IR tab 400 mg SR tab	None	200 mg QD X 14 days then 200 mg po BID OR 400mg XR QD (more common)	<ul> <li>Rash 37%</li> <li>Hepatic failure</li> <li>Fever</li> <li>Nausea</li> </ul>	CYP3A4 substrate Potent inducer of CYP2B6/ 3A4	<ul> <li>✓ Black Box: severe rash &amp; hepatotoxicity. AVOID if CD4&gt;250 (women) or 400 cells/mm3 (male)</li> <li>✓ hypersensitivity → can treat through rash, but if with fever and elevated LFTs = sign of hypersensitivity, d/c</li> <li>✓ C/I: St. John's wort; avoid Strong inducers of CYP3A4 (Carbamazepine)</li> <li>✓ Lead-in phase to reduce rash, occurs in 1<sup>st</sup> 6 wks, more in women also drug is auto inducer (will reduce its own level)</li> </ul>
		Rilpi <u>vir</u> ine	RPV	Edurant	25 25 mg tab	Complera <sup>TDF 1 QD</sup> Odefsey <sup>TAF 1 QD</sup> Juluca <sup>1 QD</sup> Cabenuva <sup>IM q1-2</sup> months	25 mg po QD w/ food ++ q2 monthly IM injection (with cabotegravir/ Cabenuva)	<ul> <li>Rash 3%</li> <li>Headache 3%</li> <li>Insomnia</li> <li>Depression 8%</li> <li>Hyperlipidemia</li> <li>Hepatotoxicity</li> </ul>	CYP3A4 Substrate ↓ [Edurant] with: Inducers of CYP3A Drugs↑pH	<ul> <li>Among smallest HIV tablets</li> <li>Best absorbed with a good meal (350-500 calories)</li> <li>PPI contraindicated, H-2 blockers need dose reduction.</li> <li>Favorable lipid profile</li> <li>Lower virologic efficacy, not suggested for VL &gt; 100,000 &amp; CD4 &lt; 200</li> <li>Can exacerbate psych symptoms</li> <li>QTc prolongation (dose related)</li> <li>Available as long-acting q1-2 monthly injectable with cabotegravir (CAB): 900 mg IM initiation, then 600 mg IM monthly/900 mg IM q2months</li> </ul>

	Class	Gener	ic	Brand	Preparations	Combo Pill	Dosing	Side Effects	Drug Interactions	Comments
	Protease Inhibitor navir	Rito <u>navir</u> PK booster	RTV	Norvir	100 mg tab 80 mg/mL oral	None	100-200 po/day	<ul> <li>Bitter aftertaste</li> <li>Numbness around mouth at HIV doses</li> <li>N/V/D</li> <li>↑ LFTs, ↑ TG</li> <li>Hyperlipidemia</li> </ul>	<u>Inducer of:</u> • 1A2, 2B6, 2C9, 2C19, UGT <u>Inhibitor of:</u> • 3A4 <sup>strong</sup> 2D6, 2C8,	<ul> <li>✓ Black Box: many drug interactions→ life threatening</li> <li>✓ Extremely strong inhibition 3A4, P-GP and other transporters</li> <li>✓ HIV activity at higher doses but toxicity &amp; inx (not used for HIV treatment)</li> <li>✓ 100 mg per dose to boost (e.g. if using with BID drug, give 100 mg BID)</li> <li>✓ Fluorinated steroids (even inhaled, injected, topical) can lead to Cushing's syndrome</li> </ul>
itors - PI	Class S/E: Hyperlipidemia MOA: High genetic barrier to resistance when boosted	Daru <u>navir</u>	DRV	Prezista	Prezista:         600, 800 mg           tab         200, 800 mg           Prezcobix:         800 mg +           150 mg COB tab         150 mg	Prezcobix <sup>w</sup> cobicistat 1 QD Symtuza <sup>w</sup> cobicistat 1 QD	600 mg po BID or 800 mg po QD w/ food + RTV 100 mg QD- BID or cobicistat 150 mg QD	<ul> <li>Rash 10%</li> <li>Headache</li> <li>N/D</li> <li>↑ amylase</li> <li>Hepatotoxic</li> <li>Kidney stones?</li> </ul>	CYP3A4 Substrate/ Inhibitor CYP 2C9 inducer Failure of contraceptives	<ul> <li>✓ Currently highest prescribed PI: 2<sup>nd</sup> Gen PI</li> <li>✓ Works in those who are resistant to other PIs</li> <li>✓ Cobicistat will cause tubular creatinine reabsorption → SCr "pseudo" rise of 10-30 mmol/L from pts normal baseline</li> <li>✓ Needs RTV or COBI boosting</li> <li>✓ When boosted with RTV: 800 QD + 100 mg RTV for naïve, [600 mg + 100 RTV] BID for experienced</li> <li>✓ Contains <i>Sulfa</i> moiety</li> <li>✓ Avoid with use of drugs that depend on CYP3A4 metabolism and has narrow therapeutic window (ie Alfuzosin)</li> </ul>
Protease Inhibitors	1 <sup>st</sup> gen PIs not used usually: Fosamprenavir FPV (Telzir) Indinavir IDV (Crixivan) Nelfinavir NFV (Viracept) Saquinqvir SQV (Invirase)	Ataza <u>navir</u>	ATV	Reyataz	Revataz:         150, 200, 300mg tab           Evotaz:         300 mg + 150 mg COB tab	Evotaz <sup>w cobicistat</sup>	300 mg po QD boosted w RTV 100 mg or cobicistat 150 mg 400 mg po QD unboosted w/ food <sup>(&gt;390 cals)</sup>	<ul> <li>Kidney stone 10 fold inc</li> <li>Increased billi 60% (cosmetic, not harmful)</li> <li>D/N/Abd pain</li> <li>Headache <sup>6%</sup></li> <li>Rash <sup>20%</sup></li> </ul>	CYP3A4 substrate/ inhibitor inducers/inhibitors of 3A4 will interact Drugs inc pH	<ul> <li>✓ 2X150 mg (300 mg) + RTV 100 mg daily (TDF increases excretion of ATZ)</li> <li>✓ 2X200 mg (400 mg) unboosted with Kivexa (needs RTV boost w others)</li> <li>✓ Increased QTc, PR, more torsades</li> <li>✓ Jaundice as result of increased direct bilirubin → not harmful, pt may decide to switch for cosmetic reason</li> <li>✓ Absorption reduced when taken with H2Ra and PPI</li> <li>✓ H2RA: Unboosted → ATV≥2 hrs before or ≥ 10 hrs after Boosted → same time or &gt;10 hrs after H2RA</li> <li>✓ PPI: Unboosted → not recommended for co-administration, Boosted → ≥ 12 hrs after PPI</li> <li>✓ Consider avoiding in CKD</li> </ul>
	Tipranavir <b>TPV</b> ( <i>Aptivus</i> )	Lopi <u>navir</u> / RTV	LPV	Kaletra	200 mg + 50 mg RTV tab	Kaletra <sup>4</sup> QD or 2 BID	400 mg po BID 800 mg po QD	• Diarrhea <sup>24%</sup> • N • ↑ LFTs, billi, Lipids, MI	CYP3A4 Substrate/ Inhibitor Many ↑ [benzos] Fentanyl Phenytoin	<ul> <li>✓ Dangerous (deadly) interaction with fentanyl</li> <li>✓ Unpredictable interaction with phenytoin → RTV inhibitor, LPV inducer of CYP. Unpredictable pheny level (unpredictable)</li> <li>✓ +++ diarrhea, worse with q24h</li> <li>✓ May need higher doses if tx experienced or later in pregnancy</li> <li>✓ May have Cardiac risk</li> </ul>

Class	Generic		Brand	Preparations	Combo Pill	Dosing	Side Effects	Drug Interactions	Comments
CCR-5 Co Receptor Antagonists	Maraviroc	MVC	Celsentri	150, 300 mg tab	None	150-600 mg po BID Standard: 300mg BID with or without food	<ul> <li>cough <sup>13</sup></li> <li>Rash <sup>10%</sup>, Abdo pain</li> <li>Dizziness, myalgia</li> <li>Ortho hypo, syncope</li> <li>Upper resp infection</li> </ul>	CYP3A4, P-gp substrate inducers/inhibitors of 3A4 or P-gp will interact	<ul> <li>Black Box: hepatotoxicity, systemic allergic reaction</li> <li>Used later in tx only for CCR-5-tropic HIV virus, cannot use for CXCR-4-tropic virus which is seen more and more in advance dx</li> <li>Avoid: Rifapentine, Dasabuvir + Ombitasvir/Paritaprevir/RTV</li> </ul>
Fusion Inhibitor	Enfuvirtide	ENF	Fuzeon	Portuge Research	None	90 mg SC BID	<ul> <li>Inj site reaction<sup>~100% pt</sup></li> <li>Bacterial pneumonia</li> <li>Hypersensitivity<sup>&lt;1%</sup></li> </ul>	Neither inducer or inhibitor of CYP enzymes	<ul> <li>✓ Was historically used in era between 1<sup>st</sup> and 2<sup>nd</sup> generation PIs</li> <li>✓ Unstable drug, dose needs to be prepared before administering each dose</li> <li>✓ No cross resistance with other ARVs</li> </ul>
Entry Inhibitor	Ibalizumab- uiyk	IBA	Trogarzo	150mg/mL vial	None	2000mg IV single dose then, 800mg Q2W	• Dizziness • Diarrhea, Nausea • Skin Rash	Neither inducer or inhibitor of CYP enzymes	<ul> <li>Indication: Treatment of HIV with combination of other ARV in heavily experienced patients with multidrug resistant infection failing current therapy</li> <li>Infused over 15-30 minutes (Loading dose no less than 30 minutes)</li> <li>Each 2 mL vial delivers 1.33mL containing 200mg of IBA</li> <li>If maintenance dose missed (&gt;3 days) then loading dose needs to be given again</li> <li>No cross resistance with other ARVs</li> <li>Not Approved in Canada</li> </ul>
gp120 Attachment Inhibitor	Fostemsavir	FTR	Rukobia	600 mg tab	None	600 mg BID with or without food	<ul> <li>Headache</li> <li>Skin Rash</li> <li>Micturition Urgency</li> <li>N/V/D</li> <li>Fatigue</li> </ul>	CYP3A4 (Partial), P- gp, BCRP substrate Strong CYP3A4 inducers will interact; inhibits OATP1B1/3, BCRP	<ul> <li>Indication: Treatment of HIV in combination with other ARV in heavily treatment experienced HIV patients with multi-drug resistant HIV-1 failing current ARV due to resistance, intolerance or safety considerations</li> <li>Prodrug of small molecule Temsavir</li> <li>BRIGHTE study 96 wks (Ackerman et al. AIDS 2021;35:1061-72.)</li> <li>Contraindicated with strong CYP3A4 inducers (anticonvulsants, mitotane, enzalutamide, rifampin, St. John's wort)</li> </ul>

Class	Gener	ic	Brand	Preparations	Combo Pill	Dosing	Side Effects	Drug Interactions	Comments
Capsid inhibitor	Lenacapavir	LEN	Sunlenca	<b>62D</b> 300 mg tab 309 mg/mL (1.5 mL vials)	None	Initiation: Day 1 & 2: 600 mg po daily Day 8: 300 mg po Day 15: 927 mg SC Simplified initiation (approved in US): Day 1: 927 mg SC and 600 mg po Day 2: 600 mg po Day 2: 600 mg po	<ul> <li>Injection site reactions</li> <li>nausea</li> </ul>	Substrate of CYP3A4, P-gp, UGT1A1. Strong inducers of CYP3A4/P-gp/UGT1A1 are contraindicated; not recommended with moderate CYP3A4 and P-gp inducers, and not with strong inhibitors of CYP3A4/P-gp/UGT1A1 together. Moderate CYP3A4 inhibitor.	<ul> <li>✓ Indication: Treatment of HIV in combination with other ARV in adults with multi-drug resistant HIV-1 for whom it is otherwise not possible to construct a suppressive antiviral regimen</li> <li>✓ Contraindicated with strong CYP3A4/P-gp/UGT1A1 inducers (anticonvulsants, rifampin, St. John's wort)</li> </ul>

OBT = optimized background therapy