HEPATITIS B CARE

HEP B FOLLOW UP LABS

Every 6 months:

Liver function tests, HBsAq, and AFP

HBV DNA for those with previous HBV DNA above 1000 IU/ml or elevated ALT or AST, and those with hepatocellular carcinoma (HCC), family history of HCC, on antiviral therapy.

Yearly: HBV DNA, CBC

RUQ ULTRASOUND EVERY 6 MONTHS

Men ≥ age 40

Women ≥ age 50

Persons with AFP >10 (Note: if female under age of 50, screen for pregnancy first).

Family history of HCC

Hepatitis B Genotype F at any age

Previous HCC

Advanced liver fibrosis (Metavir F₃ or F₄ fibrosis)

WHO NEEDS HEPATITIS B TREATMENT - CONSULT LIVER DISEASE SPECIALIST FIRST

Those with HBV DNA > 20,000 and ALT > 2x ULN normal

If liver biopsy shows moderate or greater inflammation or fibrosis

Those with history of HCC with any detectable HBV DNA

Persons with moderate to advanced liver fibrosis (Metavir F2, F3, or F4)

HBV DNA > 2000 IU/ml plus elevated ALT plus FibroScan > 9kPa in persons who have not had a liver biopsy

Hepatitis B patients starting cancer chemotherapy or immunosuppressive therapy who are HBsAg +. Screen these patients for HBsAg and HBV DNA. Place those who are HBsAg+ on tenofovir or entecavir.

Note: <u>All persons</u> should be screened before starting cancer chemotherapy or immunosuppressive therapy by drawing HBsAg and HBcAb. If either of these are positive, then obtain HBV DNA.

HIV ab testing should be completed prior to starting any HBV antiviral medication

See Hepatitis B Medication Information on Next Page

HEP B TREATMENT MEDS (will be initiated by Hepatology provider do not discontinue w/o guidance). Note: There is a risk of severe acute exacerbation of Hepatitis B with discontinuation of Hep B meds!

	1	Adverse Effects	
Drug	Dosage	Auverse Effects	Monitoring labs and Management (in
			consultation w/Hepatology)
tenofovir disoproxil	300mg daily		Creatinine, phosphorus
<u> </u>	300ilig daily		
fumarate (TDF)			baseline, 2-8 wks after
(do not use if renal			starting, then q 3-6 mos
impaired)			A a i al a i al a
		New or worsening renal	Avoid use with other
		impairment	nephrotoxic drugs,
			monitor renal labs
			periodically and switch to
			TAF if worsening renal
			function
			Canaiday assassing DMD
		Bone loss	Consider assessing BMD
			and supplementing Ca+ and D.
		0. 5 1 1: 1	and D.
		• ~ 10% - Rash, diarrhea,	Comento matic
		headache, pain,	Symptomatic
		depression, asthenia,	management or switch to
		nausea	entecavir or TAF
entecavir	0.5 or 1mg		Creatinine at baseline & q
	daily 2 hours		6 mos. Dosage
	before or		adjustment if GFR <50.
	after a meal		
		Lactic acidosis and severe	If suspected, suspend
		hepatomegaly w/steatosis	treatment
			Cymptomatic
		• ≤3% - Headache, fatigue,	Symptomatic
		dizziness, nausea	management or switch to TDF or TAF
tenofovir alofenamide	arma onco		Creatinine, phosphorus,
(TAF)	25mg once daily w/food		est. cr.cl., urine glucose
(not recommended if est.	daily w/100d		, ,
Cr.Cl. <15)			and protein
C1.C1. <15)		Now or worsening renal	D/C med if significant
		 New or worsening renal impairment 	decrease in renal function
		ппрантненс 	or evidence of Fanconi
			syndrome
		• > r06 Hoodacha	Syndroine
		• > 5% - Headache,	Symptomatic
		abdominal pain, fatigue,	
		cough, nausea and back	management or switch to
		pain	TDF or entecavir

GOT A QUESTION? WHO TO CALL

Liver Disease & Hepatitis Program – 907-729-1560 or 800-655-4837 and ask for a provider or Mary Snowball, RN [907-729-1564, (T & Th)] or another liver disease RN when Mary is not available.