



LEGACY
HEALTH PARTNERS

Cirrhosis, MASLD, Hep B, Hep C Clinical Collaboration Guide

Right patient, right service, right time

Last Reviewed: April 2025 – **FINAL**

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Cirrhosis

Who to suspect may have cirrhosis:

- F4 suggested on fibroscan, US elastography
- Ultrasound showing evidence of portal hypertension (i.e. ascites, splenomegaly, reversed portal venous flow on doppler)
- Physical exam - spider angiomas, ascites, asterixis (for hepatic encephalopathy), palmer erythema
- Laboratory – elevated INR, thrombocytopenia, elevated bilirubin, low albumin, abnormal AST/ALT

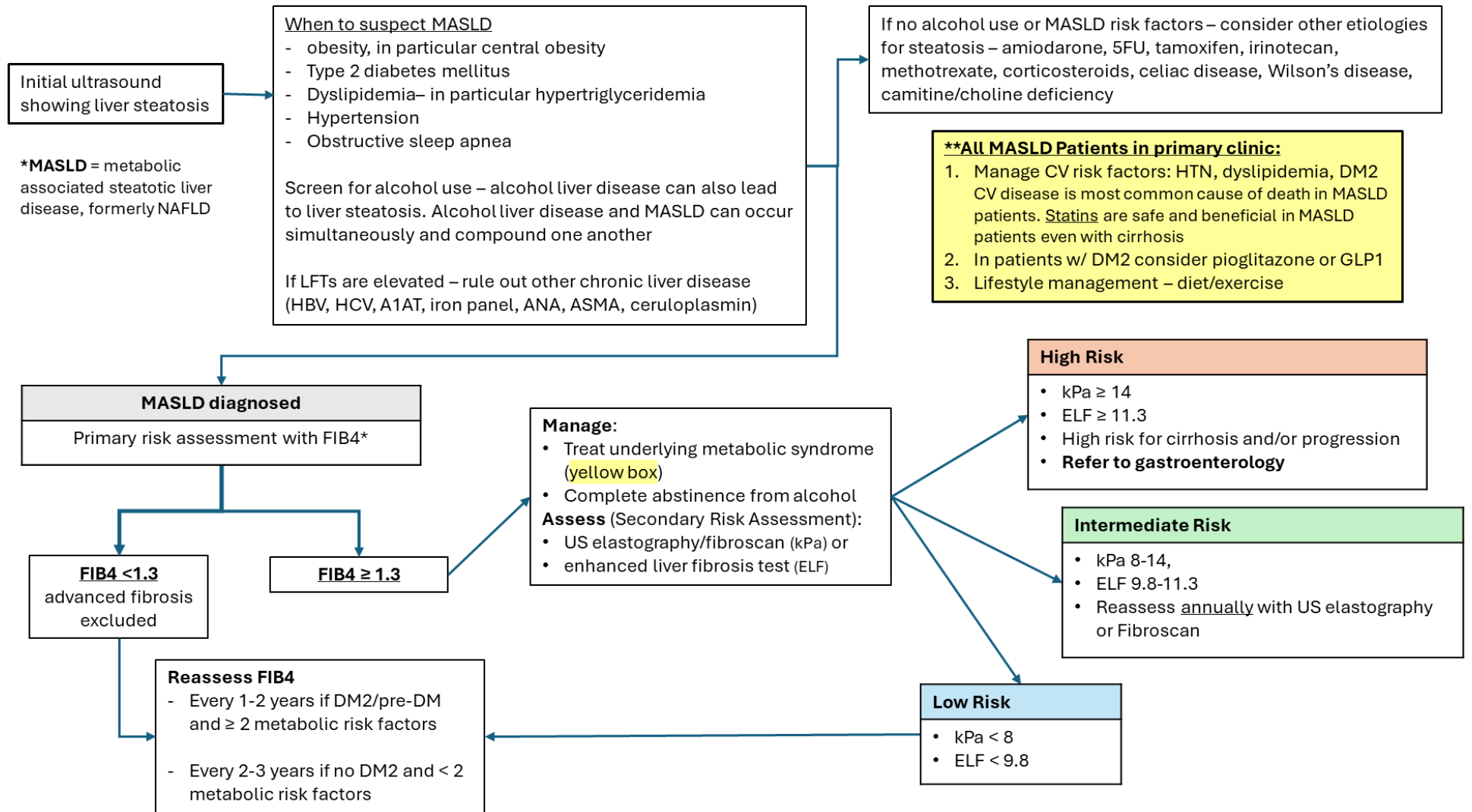
*If patients have cirrhosis: referral to GI recommended

Who to screen for HCC:

- All patients with cirrhosis
- Chronic hepatitis B without cirrhosis:
 - males from endemic country > 40 yo
 - females from endemic country > 50 yo
 - Person from Africa at earlier age (third decade of life)
 - Family history of HCC
 - PAGE-B score ≥ 10
- *F3 MASLD patients (off guidelines, but overall, a lot of GI docs do so out of a lot of caution; though higher healthcare costs)*

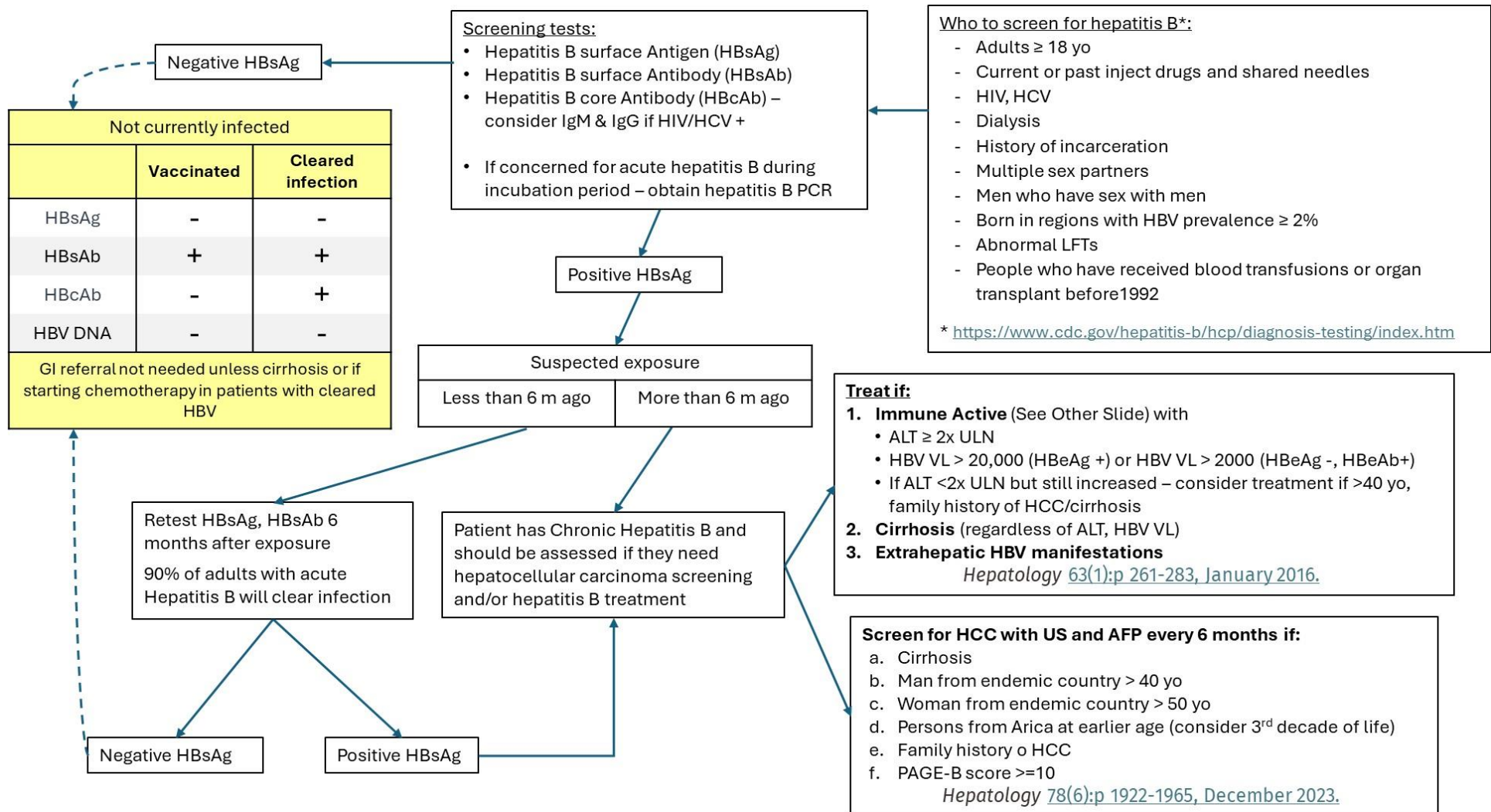
HCC screening = RUQ Ultrasound & AFP lab every 6 months
Hepatology 78(6):p 1922-1965, December 2023.

MASLD



*FIB4 calculator: <https://www.mdcalc.com/calc/2200/fibrosis-4-fib-4-index-liver-fibrosis>

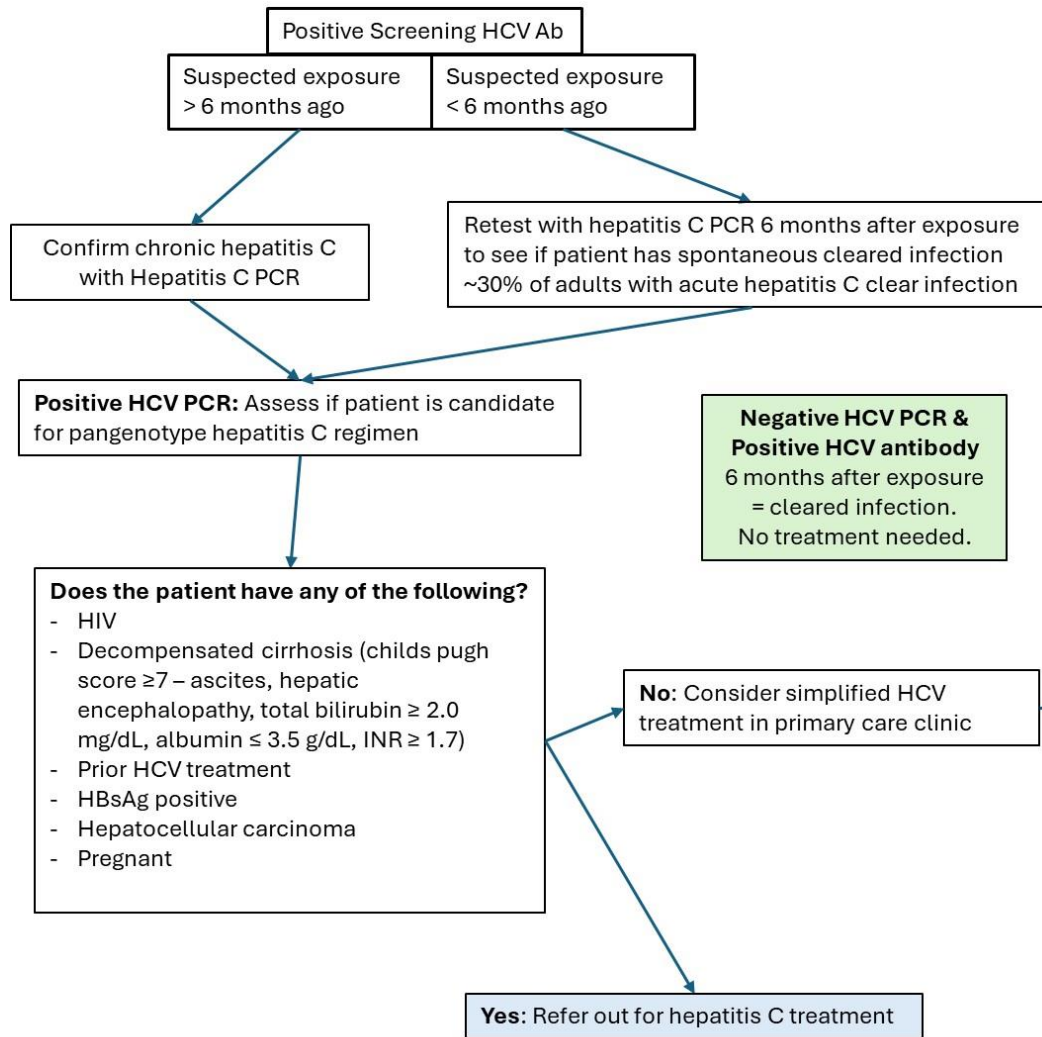
Hepatitis B



Chronic Hepatitis B

	IMMUNE TOLERANT	IMMUNE ACTIVE HBeAg +	INACTIVE	IMMUNE-ACTIVE HBeAg -	OCCULT
HBsAg	+	+	+	+	-
HBsAb	-	-	-	-	-
HBeAg	+	+	-	-	-
HBeAb	-	-	+	+	+/-
HBc IgM/IgG	+	+	+	+	+/-
HBV DNA	+++	+++	+	++	+/- (in serum) + (in liver)
ALT	Normal	Increased	Normal	Increased	Normal

Hepatitis C



Who to screen for hepatitis C with HCV Ab:

- Adults ≥ 18 yo
- Patients who currently or in the past inject drugs and shared needles
- People with HIV or who routinely undergo hemodialysis
- Abnormal LFTs
- People who have received blood transfusions or organ transplant prior to 1992

www.cdc.gov/hepatitis-c/hcp/diagnosis-testing/index.html

Pangenotype HCV Treatment : www.hcvguidelines.org/

Pre-treatment testing:

- CBC, CMP, INR, HCV PCR, HIV, HBsAg, HBcAb
- Plus RUQ US if cirrhosis– to rule out hepatocellular carcinoma
- Assess for drug-drug interactions with pharmacy

No Cirrhosis

- Glecaprevir/pibrentasvir x 8 weeks
- Sofosbuvir/velpatasvir x 12 weeks

Compensated cirrhosis (Childs Pugh A)

- FIB4 >3.25, elastography with >12.5kPa, positive FibroSure or Enhanced Liver Fibrosis Test, imaging showing liver nodularity or splenomegaly
- Glecaprevir/pibrentasvir x 8 weeks
- Sofosbuvir/velpatasvir x 12 weeks for genotypes 1,2,4,5,6
 - Genotype 3 – baseline NS5A resistance substitution(RAS) testing

Post-treatment testing:

- 12 weeks after treatment end - CMP, HCV PCR → HCV PCR is negative, cure achieved!
- Patients with cured HCV still need HCC surveillance

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Positive Screening HCV Ab

Suspected exposure
> 6 months ago

Suspected exposure
< 6 months ago

Negative HCV PCR & Positive HCV antibody 6 months after exposure = cleared infection. No treatment needed.

Confirm chronic hepatitis C with Hepatitis C PCR

Retest with hepatitis C PCR 6 months after exposure to see if patient has spontaneous cleared infection
~30% of adults with acute hepatitis C clear infection

Positive HCV PCR: Assess if patient is candidate for pangenotype hepatitis C regimen

No: Consider simplified HCV treatment in primary care clinic

Yes: Refer out for hepatitis C treatment

Does the patient have any of the following?

- HIV
- Decompensated cirrhosis (Childs pugh score ≥ 7 – ascites, hepatic encephalopathy, total bilirubin ≥ 2.0 mg/dL, albumin ≤ 3.5 g/dL, INR ≥ 1.7)
- Prior HCV treatment
- HBsAg positive
- Hepatocellular carcinoma
- Pregnant

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Disclaimer: No guideline can anticipate all the unique circumstances of patient care, and as such, there are times when good clinical judgement will result in, and will require deviation from this guideline. In those settings, the reason for such deviation from this guideline should be documented in the medical record.

Contact: If you have questions or comments about this guide, or are interested in the development of future collaboration guides, please email LHP medical director Albert Chaffin, M.D., at achaffin@lhs.org.