

VIRAL HEPATITIS CARE FORM

As cited in the Interim Guidelines on the Management of Patients with Hepatitis B and Hepatitis C infection, physicians and health care providers of hepatitis treatment facilities shall submit and report data to the Epidemiology Bureau and their regional, provincial and municipal counterparts. This form is to be filled-out on the initial and follow-up visit of the client. Please write in CAPITAL LETTERS and CHECK the appropriate boxes.

I. VISIT INFORMATION (Proceed to the Clinical Assessment Section if Case Report Form is recently accomplished)

Consult date: (mm/dd/yyyy) / /

Patient code:

Client visit type: ☐ Initial ☐ Follow-up

Unique Identifier Code: - - -

Facility name:

Tested positive for: ☐ Hepatitis B ☐ Hepatitis C

II. CLIENT DATA

Name (full name):

First Name Middle Name Last Name Suffix

Sex assigned at birth: ☐ Male ☐ Female

Age in years: Weight (in kg): Height (in cm):

If female, is she pregnant? ☐ Yes ☐ No

Nationality: ☐ Filipino ☐ Other:

III. CLINICAL ASSESSEMENT

Signs and symptoms	Result		Laboratory Tests	Date Done	Result	
Jaundice	<input type="checkbox"/> Yes	<input type="checkbox"/> No	HBsAg	/ /	<input type="checkbox"/> Positive	<input type="checkbox"/> Negative
Coagulopathy	<input type="checkbox"/> Yes	<input type="checkbox"/> No	HBeAg	/ /	<input type="checkbox"/> Positive	<input type="checkbox"/> Negative
Ascites	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Anti-HCV	/ /	<input type="checkbox"/> Reactive	<input type="checkbox"/> Non-reactive
Variceal Hemorrhage	<input type="checkbox"/> Yes	<input type="checkbox"/> No	HCV-RNA (quali)	/ /	<input type="checkbox"/> Positive	<input type="checkbox"/> Negative
Hepatic Encephalopathy	<input type="checkbox"/> Yes	<input type="checkbox"/> No	HCV-RNA (quant)	/ /		IU/mL
Hepatomegaly	<input type="checkbox"/> Yes	<input type="checkbox"/> No	HBV DNA	/ /		IU/mL
Splenomegaly	<input type="checkbox"/> Yes	<input type="checkbox"/> No	AST (SGOT)	/ /		IU/L
Pruritus	<input type="checkbox"/> Yes	<input type="checkbox"/> No	ALT (SGPT)	/ /		IU/L
Fatigue	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Total Bilirubin	/ /		mg/dL
Spider Angiomata	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Albumin	/ /		g/dL
Palmar Erythema	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Platelet Count	/ /		10 ⁹ /L
Other: (specify)			PT INR	/ /		sec

Non-invasive tests for detection of cirrhosis (use the most recent lab results)

Date done

Result

Aminotransferase/Platelet Ratio Index Score / /

☐ ≤ 2 ☐ > 2

Fibrosis 4 / FIB-4 Score / /

☐ ≤ 3.25 ☐ > 3.25

Transient Elastography / /

☐ < 12.5 kPa ☐ ≥ 12.5 kPa

Imaging (UTZ/CT/MRI) / /

☐ Cirrhosis ☐ No cirrhosis
☐ Mass ☐ No mass

Child-Turcotte-Pugh Class Score / /

☐ Class A ☐ Class B ☐ Class C

Did the client develop liver cirrhosis following treatment?

☐ Yes ☐ No

Did the client develop hepatocellular carcinoma following treatment?

☐ Yes ☐ No

Did the client acquire/develop any of the following compared from the last visit:

☐ Hepatitis B co-infection (if the client was initially diagnosed with Hepatitis C infection)

☐ HIV co-infection

☐ Hepatitis C co-infection (if the client was initially diagnosed with Hepatitis B infection)

☐ Other:

IV. TREATMENT

Is the client eligible for treatment?

☐ Yes, the client is eligible for treatment for Hepatitis B

☐ No, the client is not eligible for treatment

☐ Awaiting for laboratory results

☐ Yes, the client is eligible for treatment for Hepatitis C

and was advised to return for monitoring on:

(mm/dd/yyyy) / /

Treatment status:

☐ Enrolling this visit ☐ Continuing/refill ☐ Returning client ☐ Not on treatment (reason if not on treatment:)

Regimen:

☐ Sofosbuvir (400 mg)

☐ Tenofovir(TDF) (300 mg)

☐ Daclatasvir (60 mg)

☐ Sofosbuvir/Velpatasvir

☐ Ribavirin

☐ Other:

Date dispensed (mm/dd/yyyy)

pills missed (past 30 days)

of pills on hand

of pills dispensed

Expected date of refill (mm/dd/yyyy)

/ /

/ /

/ /

/ /

If the patient's treatment was discontinued , please completely fill out this section.

Date discontinued: / /

Reason: (D/C code) (For code 6 & 7, please specify)

Discontinuation codes (D/C):
1-Treatment Failure 2-Clinical progression 3-Patient Decision/Request 4-Compliance difficulties 5-Drug Interaction 6-Adverse Event (Specify) 7-Others (Specify) 8-Cured(for Hepatitis C) 9-Death