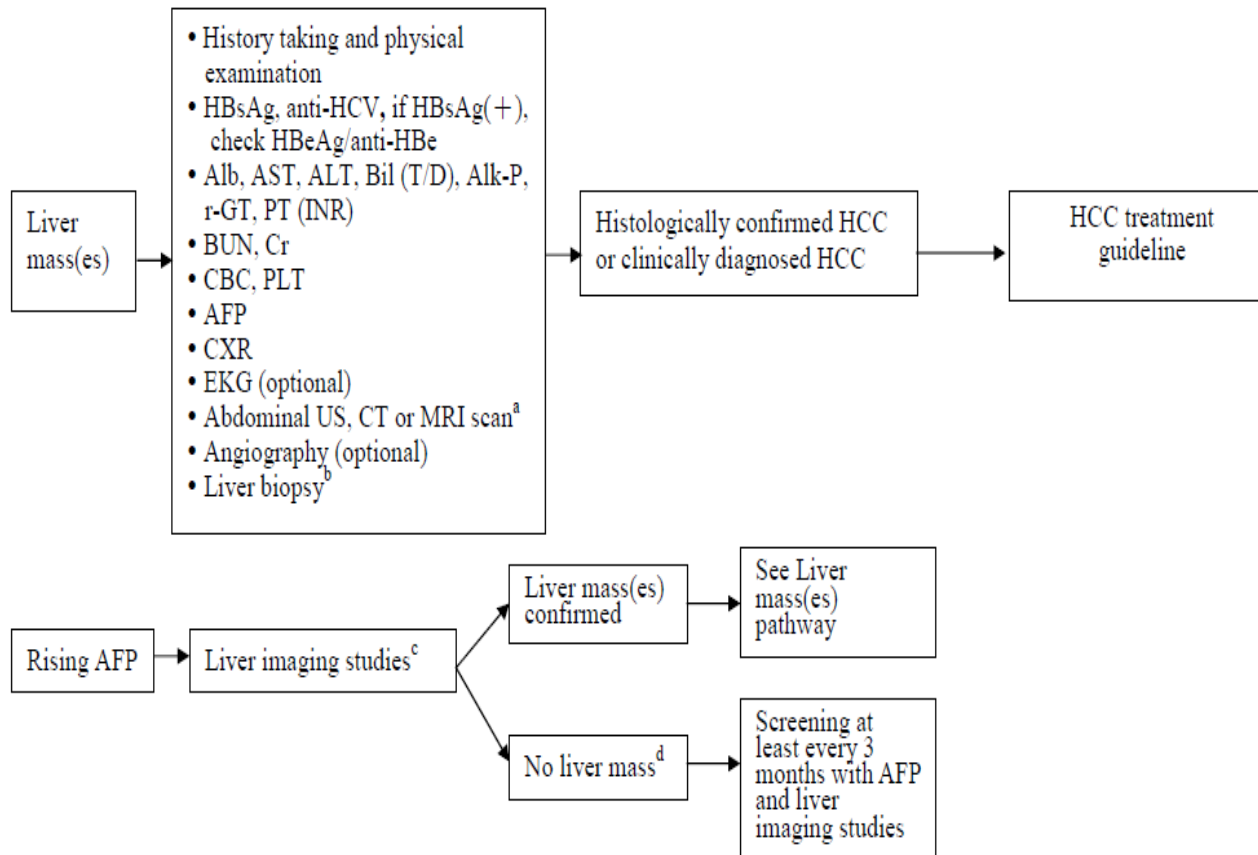


肝癌治療指引|肝癌治療指引

中國醫藥大學附設醫院

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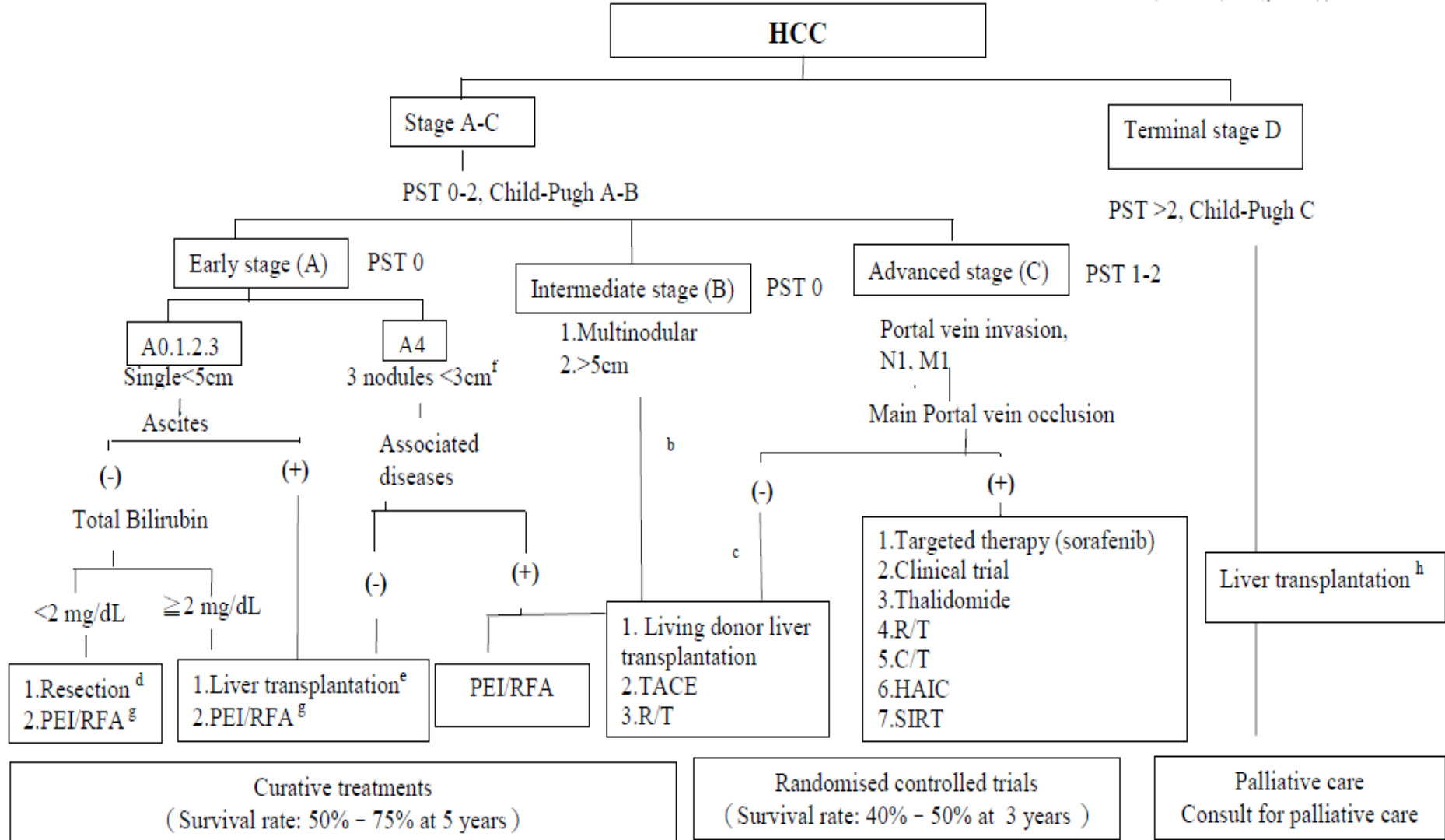
^a CT or MRI scan to define extent and number of primary lesions, vascular anatomy, involvement with tumor, and extrahepatic disease.

^b Liver biopsy is required to exclude metastatic hepatic tumor if less than 2 of the following

g 3 clinical diagnostic criteria is fitted: (1) HBsAg or anti-HCV(+) with liver cirrhosis; (2) At least one typical imaging study. Ultrasound is used as a screening tool. The lesions identified should be investigated further with either dynamic CT scan or MRI with contrast. At least one of the studies has to be typical of HCC; otherwise angiography is required. If the angiographic findings are not characteristic of HCC, biopsy should be performed. (3) AFP \geq 400 ng/mL.

^c If ultrasound negative, CT or MRI scan should be performed.

^dRule out germ cell tumor if clinically indicated.



^{a & b} If the number of nodules ≤ 3 , localized in one lobe or not centrally located, resection may be indicated

^c If the mass is in one lobe, Child-Pugh score is A and the tumor invasion is not beyond main portal vein, resection may be indicated

^d ICG 15'RR: <15% \rightarrow lobectomy, 15-30% \rightarrow subsegmentectomy, >30% \rightarrow wedge resection or enucleation

^e UCSF criteria for deceased donor liver transplantation: single tumor ≤ 6.5 cm, number ≤ 3 , each size ≤ 4.5 cm, total ≤ 8 cm

^f RFA + PEI may be indicated when single tumor with 3-5cm in size

^g PEI/RFA may be considered if patient rejects surgery or tumor < 3 cm

^h Expanding criteria for living donor liver transplantation: no evidence of extrahepatic metastasis or main portal vein thrombus

1. Criteria for partial hepatectomy (PH)
 - (1) Allowable liver function
 - (2) PVT or HVT can be removed
 - (3) Free section margin
 - (4) Usual indications:
 - Solitary tumor
 - Multiple tumors ≤ 3 , each size ≤ 5 cm (determined by the location)
2. Candidate for liver transplantation (LTx)
 - (1) Child B or C
 - (2) No contraindication for LTx
 - (3) No extrahepatic metastasis
 - (4) Tumor size ≤ 5 cm
number ≤ 3 , each size ≤ 3 cm
(Milan criteria for diseased donor liver transplantation)
 - (5) Single Tumor ≤ 6.5 cm
number ≤ 3 , each size ≤ 4.5 cm, total ≤ 8 cm
(UCSF criteria for living donor liver transplantation)
3. Criteria for PEI or RFA
 - (1) Child A or B, No or minimal amount of ascites
 - (2) Platelet $\geq 50,000/\text{mm}^3$, PT prolongation ≤ 5 seconds
 - (3) Ultrasound or CT identifiable and approachable lesions
 - (4) Patients can cooperate and hold breath adequately
 - (5) Liver tumor number ≤ 3 , each size ≤ 3 cm or single tumor ≤ 5 cm
 - (6) If tumor $> 2\text{-}3\text{cm}$, favor RFA
 - (7) IV General anesthesia indicated in RFA
4. Criteria for TACE
 - (1) Child A or B
 - (2) Patent main portal vein or main portal vein thrombosis with cavernous transformation
 - (3) Main portal vein obstruction but with peri-portal collateral circulation
5. Protocol for PEI or RFA
Pre-PEIT evaluation:
CBC/DC, WBC, PT, Alb, AST/ALT, Bil(T/D), BUN, Cr, AFP
Ultrasound

Dynamic CT scan
MRI (optional)
Biopsy (optional, recommended if normal AFP levels)

PEIT:

Total volume (ml) of ethanol needed: $4/3\pi(r + 0.5)^3$, r is the radius of the tumor in centimeter; preferably inject 2-10 ml of ethanol each session if the patient can tolerate
Schedule PEIT once to twice weekly

Post-PEI/RFA evaluation:

Clinical assessment of local side effects and bleeding complication
WBC/DC if infections cannot be ruled out
AFP every 1-3 months
Ultrasound and/or Dynamic CT scan/or MRI after completion of PEI/RFA
Schedule PEI/RFA again if dynamic CT scan shows viable tumors

6. Protocol for TACE

Pre-TACE evaluation:

CBC/DC, WBC, PT, Alb, AST/ALT, Bil(T/D), BUN, Cr, AFP
Ultrasound
Dynamic CT scan
MRI (optional)
Biopsy (optional, recommended if normal AFP levels)
Adequate hydration if borderline renal function

Post-TACE evaluation:

Clinical assessment of post-embolization syndrome
WBC/DC if infections cannot be ruled out
Alb/AST/ALT/Bil(T/D)/PT/Cr at 1-2 weeks after TACE (additional follow up if indicated)
AFP every 1-3 months

Ultrasound and/or Dynamic CT scan every 1-3 sessions of TACE

Schedule TACE every 1-3 months if dynamic CT scan shows viable tumors and liver function allows

7. Criteria for R/T to portal vein thrombosis or primary tumor

- (1) Absence of severe hepatoencephalopathy or uncontrolled ascites
- (2) Patient can cooperate
- (3) ECOG performance status ≤ 3
- (4) Total bilirubin ≤ 10 mg/dl

8. Protocol for R/T

Pre-R/T evaluation:

CBC, WBC & D/C, PLT, PT, Alb, AST/ALT, Bil(T/D), Alk-P, BUN, Cr, AFP

Dynamic CT scan

MRI (optional)

Post-R/T surveillance:

Alb/AST/ALT/Bil(T/D)/Alk-P at least every 2 weeks for 2 months, then every 1-2 months for 1 year, then every 3 months

CBC, WBC & D/C, PLT, PT, BUN, Cr 1 month later

AFP, if initially elevated, every 2 months in 1 year, then 3-6 months

Imaging study 1-2 months later, then every 3-6 months

9. Indication for Hepatic Arterial Infusional Chemotherapy (HAIC)

- (1) HCC: advanced HCC, diffuse invasion type, portal vein thrombosis
- (2) Liver metastasis from colorectal cancer
- (3) Liver metastasis from gastric cancer
- (4) Liver metastasis from breast cancer
- (5) Intrahepatic cholangiocarcinoma

10. Protocol for Hepatic arterial infusional chemotherapy:

- (1) Cisplatin 10mg/1hr, D1-5/ 4w, 5FU 250mg/5hr, D1-5/ 4w
- (2) Cisplatin 7mg/1hr, D1-5/ 4w, 5FU 170mg/5hr, D1-5/4w
- (3) Cisplatin 20mg/30min, 5FU 250mg/2hr, W1,2,3,4,5,7,9,11,13,15

11. Chemotherapy: There is no large randomized, controlled clinical trial to demonstrate the survival benefit of systemic chemotherapy.

Patients with good performance can be treated with doxorubicin-, epirubicin-, mitoxantron-, cisplatin-, 5-fluorouracil-, capecitabine-, etoposide- or gemcitabine-based chemotherapy after discussing with patient.

