

Contents lists available at ScienceDirect

# Digestive and Liver Disease

journal homepage: www.elsevier.com/locate/dld



# Guidelines

Multidisciplinary Treatment of Hepatocellular Carcinoma in 2023: Italian practice Treatment Guidelines of the Italian Association for the Study of the Liver (AISF), Italian Association of Medical Oncology (AIOM), Italian Association of Hepato-Bilio-Pancreatic Surgery (AICEP), Italian Association of Hospital Gastroenterologists (AIGO), Italian Association of Radiology and Clinical Oncology (AIRO), Italian Society of Pathological Anatomy and Diagnostic Cytology (SIAPeC-IAP), Italian Society of Surgery (SIC), Italian Society of Gastroenterology (SIGE), Italian Society of Medical and Interventional Radiology (SIRM), Italian Organ Transplant Society (SITO), and Association of Patients with Hepatitis and Liver Disease (EpaC) – Part I – Surgical treatments

Giuseppe Cabibbo<sup>a,b,\*</sup>, Bruno Daniele<sup>c</sup>, Mauro Borzio<sup>d</sup>, Andrea Casadei-Gardini<sup>e</sup>, Umberto Cillo<sup>f</sup>, Agostino Colli<sup>g</sup>, Massimiliano Conforti<sup>h</sup>, Vincenzo Dadduzio<sup>i</sup>, Francesco Dionisi<sup>j</sup>, Fabio Farinati<sup>k,l</sup>, Ivan Gardini<sup>h</sup>, Edoardo Giovanni Giannini<sup>m</sup>, Rita Golfieri<sup>n,o</sup>, Maria Guido<sup>p</sup>, Andrea Mega<sup>q</sup>, Silvia Minozzi<sup>r</sup>, Fabio Piscaglia<sup>s,t</sup>, Lorenza Rimassa<sup>u,v</sup>, Laura Romanini<sup>w</sup>, Anna Pecorelli<sup>x</sup>, Rodolfo Sacco<sup>y</sup>, Marta Scorsetti<sup>z,aa</sup>, Luca Viganò<sup>ab,ac</sup>, Alessandro Vitale<sup>f</sup>, Franco Trevisani<sup>t,ad,\*\*</sup>

<sup>a</sup> Section of Gastroenterology and Hepatology, Department of Health Promotion, Mother and Child Care, Internal Medicine and Medical Specialties PROMISE, University of Palermo, Italy

- <sup>b</sup> Gastroenterology Unit, Azienda Ospedaliera Universitaria Policlinico "Paolo Giaccone", Palermo, Italy
- <sup>c</sup> Oncology Unit, Ospedale del Mare, ASL Napoli 1 Centro, Napoli, Italy
- <sup>d</sup> Centro Diagnostico Italiano (CDI), Milano, Italy
- <sup>e</sup> Department of Oncology, Vita-Salute San Raffaele University, IRCCS San Raffaele Scientific Institute Hospital, Milan, Italy
- <sup>f</sup> General Surgery 2-Hepato-Pancreato-Biliary Surgery and Liver Transplantation Unit, Padua University Hospital, 35128 Padua, Italy
- <sup>g</sup> Dipartimento di Medicina Trasfusionale ed Ematologia, Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico, Milano, Italy
- <sup>h</sup> EpaC Onlus, Italian Liver Patient Association, Turin, Italy
- <sup>i</sup> Medical Oncology Unit, "Mons. A.R.Dimiccoli" Hospital, Barletta, ASL BT, Italy
- <sup>j</sup> Department of Radiation Oncology, IRCCS Regina Elena National Cancer Institute Rome, Italy
- <sup>k</sup> Department of Surgery, Oncology and Gastroenterology, University of Padova, 35128 Padova, Italy
- <sup>1</sup>Gastroenterology Unit, Azienda Ospedale-Università di Padova, 35128 Padova, Italy
- m Gastroenterology Unit, Department of Internal Medicine, University of Genoa, IRCCS Ospedale Policlinico San Martino, Genoa, Italy
- <sup>n</sup> Alma Mater Studiorum" Bologna University, Bologna, Italy
- <sup>o</sup> Radiology Unit Madre Fortunata Toniolo Private Hospital, coordinator of Radiology centers Medipass Bologna, Bologna, Italy
- <sup>p</sup>Department of Medicine, University of Padova, Padova- Italy
- <sup>q</sup> Department of Gastronterology, Regional Hospital Bolzano, Italy
- <sup>r</sup> Oncology Department, Istituto di Ricerche Farmacologiche Mario Negri, IRCCS, Milano, Italy
- <sup>s</sup> Division of Internal Medicine, Hepatobiliary and Immunoallergic Diseases, IRCCS Azienda Ospedaliero-Universitaria di Bologna, Italy
- <sup>t</sup> Department of Medical and Surgical Sciences, University of Bologna, Italy
- <sup>u</sup> Department of Biomedical Sciences, Humanitas University, Via Rita Levi Montalcini 4, 20072 Pieve Emanuele, Milan, Italy

Social media: 🈏 (G. Cabibbo)

# https://doi.org/10.1016/j.dld.2023.10.029

1590-8658/© 2023 Editrice Gastroenterologica Italiana S.r.l. Published by Elsevier Ltd. All rights reserved.



<sup>\*</sup> Corresponding author at: Section of Gastroenterology and Hepatology, Department of Health Promotion, Mother and Child Care, Internal Medicine and Medical Specialties PROMISE, University of Palermo, Palermo, Piazza delle Cliniche n 2, 90127, Italy.

<sup>\*\*</sup> Corresponding author at: Unit of Semeiotics, Liver and Alcohol-related diseases, IRCCS Azienda Ospedaliero-Universitaria di Bologna, 40138, Bologna, Italy. E-mail addresses: giuseppe.cabibbo@unipa.it (G. Cabibbo), franco.trevisani@unibo.it (F. Trevisani).

<sup>v</sup> Medical Oncology and Hematology Unit, IRCCS Humanitas Research Hospital, Via Manzoni 56, 20089 Rozzano, Milan, Italy

<sup>w</sup> Radiology Unit, Ospedale di Cremona, ASST Cremona, Cremona, Italy

- <sup>x</sup> Department of Radiology, IRCCS Azienda Ospedaliero-Universitaria di Bologna, Bologna, Italy
- <sup>y</sup> Gastroenterology and Endoscopy Unit, Department of Surgical and Medical Sciences, University of Foggia, 71100 Foggia, Italy

<sup>2</sup> Department of Biomedical Sciences, Humanitas University, 20090 Pieve Emanuele, Milan, Italy

<sup>aa</sup> Department of Radiotherapy and Radiosurgery, Humanitas Research Hospital IRCCS, Via Manzoni 56, 20089, Rozzano, Milan, Italy

ab Hepatobiliary Unit, Department of Minimally Invasive General & Oncologic Surgery, Humanitas Gavazzeni University Hospital, Viale M. Gavazzeni 21,

24125 Bergamo, Italy

<sup>ac</sup> Department of Biomedical Sciences, Humanitas University, Viale Rita Levi Montalcini 4, 20090 Milan, Italy

<sup>ad</sup> Unit of Semeiotics, Liver and Alcohol-Related Diseases, IRCCS Azienda Ospedaliero-Universitaria di Bologna, 40138 Bologna, Italy

### ARTICLE INFO

Article history: Received 20 June 2023 Accepted 30 October 2023 Available online 28 November 2023

Keywords: Adjuvant therapy Cirrhosis Guideline GRADE Hepatocellular carcinoma Immunotherapy Liver transplant Liver cancer Multidisciplinary board PICO Recurrence Resection Surgery Survival Transplant benefit

# ABSTRACT

Worldwide, hepatocellular carcinoma (HCC) is the third most common cause of cancer-related death. The remarkable improvements in treating HCC achieved in the last years have increased the complexity of HCC management. Following the need to have updated guidelines on the multidisciplinary treatment management of HCC, the Italian Scientific Societies involved in the management of this cancer have promoted the drafting of a new dedicated document. This document was drawn up according to the GRADE methodology needed to produce guidelines based on evidence. Here is presented the first part of guidelines, focused on the multidisciplinary tumor board of experts and surgical treatments of HCC.

© 2023 Editrice Gastroenterologica Italiana S.r.l. Published by Elsevier Ltd. All rights reserved.

### 1. Introduction

This report summarizes the recommendations of Clinical Practice Guidelines regarding Surgical treatments of Hepatocellular Carcinoma (HCC) [1], drawn up according to the GRADE methodology [2] and promoted by the following scientific societies: Italian Association for the Study of the Liver (AISF), Italian Association of Medical Oncology (AIOM), Italian Association of Hepato-Bilio-Pancreatic Surgery (AICEP), Italian Association of Hospital Gastroenterologists (AIGO), Italian Association of Radiology and Clinical Oncology (SIAPeC-IAP), Italian Society of Surgery (SIC), Italian Society of Gastroenterology (SIGE), Italian Society of Medical and Interventional Radiology (SIRM), Italian Organ Transplant Society (SITO), and Association of Patients with Hepatitis and Liver Disease (EpaC).

Current knowledge on treatment of HCC is translated into relevant practical recommendations following the rules and the methodology indicated by the Centro Nazionale per l'Eccellenza delle Cure (CNEC) and the Istituto Superiore di Sanità (ISS).

The guideline developers, designated by the above-mentioned scientific societies, identified key questions that health care providers are frequently faced with in the management of patients with HCC.

#### 2. Background

HCC is a common cause of cancer-related mortality and morbidity worldwide [3,4] with variable, but on average still poor prognosis [5], that in the vast majority of cases occurs in patients with chronic liver disease, usually in the cirrhotic stage [6,7]. Early detection of HCC, increasing the percentage of early-stage tumors, expands the rate of patients amenable to curative treatments, favorably impacting overall survival [8].

In recent years, the therapeutic armamentarium of HCC has been remarkably enriched with new effective techniques and strategies, leading to the need of a management involving different specialists [9]. Indeed, prediction of outcome and treatment choice are particularly complex as they must consider the underlying liver disease and comorbidities, which condition treatment feasibility and have an inherent competing mortality risk.

### 3. Methods for developing the guideline

Twenty-two experts indicated by the above-mentioned scientific societies, plus 2 delegates of the EpaC patient association, selected by collegial discussion the key questions and draw up guidelines. This document was arranged according to the rules of the CNEC of the Italian Ministry of Health. The key questions were developed according to the Population, Intervention, Comparison, Outcomes (PICO) acronym. For each PICO question, the literature on MEDLINE/Pubmed, Embase and Cochrane Library databases was systematically searched with both Thesaurus terms and free text. A further hand-search was performed on the bibliography of articles and previously published guidelines.

Recommendations were formulated applying the GRADE approach [2] according to the CNEC manual [10]. All aspects concerning questions, assessment of evidence and conclusions were discussed among panel members and voted. Before voting, members declared their potential conflict of interest (COI) relevant to the PICO question, and only those without COI voted. The online GRADEpro GDT tool was used to develop questions, assess evidence, and make decisions [11]. The certainty of evidence was assessed applying the tool for Risk of Bias in randomized trials (RoB)

#### Table 1

PICO questions about Surgical treatment, Recommendations, Certainty of evidence, and Strength of recommendation of Clinical Practice Guidelines for the management of Hepatocellular Carcinoma (HCC).

PICO		Recommendation	Certainty of evidence	Strength of recommendation
1	Is management by a multidisciplinary team of experts versus management by a single expert indicated in patients with hepatocellular carcinoma (HCC)?	For patients with HCC, the panel recommends that the evaluation of the diagnostic and therapeutic workup be carried out by a multidisciplinary team of experts rather than by a single expert.	Moderate	Strong in favor of multidisciplinary management
2	In patients with Child-Pugh class A cirrhosis and single HCC, is hepatic resection indicated compared to the treatment with thermal ablation?	In patients with Child-Pugh class A cirrhosis and single HCC, the panel suggests preferring liver resection over thermal ablation, except for patients with HCC ≤2 cm, for whom the panel suggests thermal ablation.	Low	Conditional in favor of resection
3	In cirrhotic patients with good liver function and multinodular HCC, is liver resection indicated compared to transarterial chemoembolization (TACE)?	In well selected cirrhotic patients with good liver function and oligo-nodular (2–3 nodules) HCC, and after multidisciplinary board evaluation, the panel recommends liver resection over TACE.	Low	Strong in favor of resection
4	In cirrhotic patients with good liver function and HCC responsible for intrahepatic macrovascular invasion, is liver resection indicated compared to sorafenib-based systemic therapy?	In cirrhotic patients with good liver function and HCC responsible for intrahepatic macrovascular invasion, the panel suggests preferring liver resection versus sorafenib-based systemic therapy.	Very low	Conditional in favor of resection
5	In cirrhotic patients with HCC who are candidates for liver transplantation, is a selection according to the "transplant benefit" criterion indicated compared to other criteria?	Good Clinical Practice statement: The Panel believes that the criterion of "transplant benefit" instead of that of "transplant utility" should be used for the selection of candidates for transplantation, taking into account that the potential transplantability must consider overall health, comorbidity, nutritional status and age (although without a defined and universally accepted cut-off), in addition to the tumor burden.	Not applicable	Not applicable
6	In cirrhotic patients beyond Milan criteria, but without intrahepatic vascular invasion and extrahepatic tumor spread, is liver transplantation indicated compared to alternative treatments (liver resection, locoregional or systemic therapies)?	In potentially transplantable (comorbidities, nutritional status and age) cirrhotic patients with HCC beyond the Milan criteria, but without intrahepatic vascular (and biliary) invasion and extrahepatic tumor spread, the Panel recommends considering liver transplantation rather than alternative treatments (resection, locoregional treatments and systemic treatments).	Transplant vs. resection: moderate Transplant vs. alternative therapies: very low.	Strong in favor of transplantation
7	In patients with HCC single >2 cm or multifocal (within the centre's transplantability criteria) and treatable with any potentially radical therapy ("first-line" liver transplantation, resection or thermal ablation), is "salvage" transplantation (i.e. transplant performed at the time of cancer recurrence/ progression after resection or thermal ablation) indicated compared to the "first line" transplant?	In patients with HCC single >2 cm or multifocal (within the centre's transplantability criteria) and treatable with any potentially radical therapy, the panel suggests performing liver resection or thermal ablation followed, in case of cancer recurrence, by "salvage" transplantation.	Low	Conditional in favor of liver resection or thermal ablation followed by "rescue" transplant
8	In patients with HCC beyond the transplant criteria adopted by the center, is the "downstaging" procedure followed by transplantation indicated compared to treatments without subsequent transplantation?	In patients with HCC beyond the oncological transplant criteria adopted by the center, the panel recommends to perform the downstaging procedure aimed at bringing the patient back to the transplant criteria compared to all other therapies without transplantation.	Low	Strong in favor of transplantation

GRADE Working Group grades of evidence:.

High certainty: We are very confident that the true effect lies close to that of the estimate of the effect.

Moderate certainty: We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different.

Low certainty: Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect.

Very low certainty: We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect.

as suggested by Cochrane [12], and the Newcastle-Ottawa scale for non-randomized studies [13].

## 4. PICO questions and recommendations

Table 1 summarizes PICO questions about Surgical treatment, Recommendations, Certainty of evidence, and Strength of recommendation of Clinical Practice Guidelines for the management of Hepatocellular Carcinoma (HCC).

# 1. Is management by a multidisciplinary team of experts versus management by a single expert indicated in patients with hepatocellular carcinoma (HCC)?

The management of patients with HCC involves multiple professional specialists (Fig. 1), such as the hepatologist, radiologist,



Fig. 1. Multidisciplinary tumor board.

pathologist, surgeon, transplant surgeon, oncologist, radiation oncologist, as well as nurses and experts of palliative care [9,14,15]. The hepatologist represents the pivotal figure and the link between the other professional figures in all phases of the patient's journey [15], while the other specialists assume a fundamental role at different times, depending on the tumor stage and the therapeutic strategy adopted.

Patients with HCC have two peculiar features which necessitate of a multidisciplinary vision [9,16,17]: 1) HCC is associated with the presence of chronic liver disease in the vast majority of cases, most frequently in the cirrhotic stage (>90% of cases), which affects itself both the applicability of many therapies and the survival of patients, and whose management requires specialized knowledge; 2) numerous modalities of therapy are available, also including liver transplantation, a very peculiar case among solid tumors.

In this line, good liver function is generally defined as Child-Pugh class A (without ascites) and MELD score <10, also considering the grade of portal hypertension. However, it is important to consider that assessing liver function reserve before treatment and during follow-up is complex and multifaceted. So, also other scores are usually used as the model for end-stage liver disease (MELD) score, MELD-sodium score, albumin-bilirubin grade, and indocyanine green test.

The goal of a multidisciplinary approach is therefore to improve the patient's outcome by defining, in an individualized basis, the best diagnostic approach and the best therapeutic option applicable through a common unified discussion, where specialists examine together the individual case. Multidisciplinary teams are increasingly common, particularly in referral centers, replacing the referral of the patient to individual consultations with specific relevant specialists for the given stage of the tumor. Nevertheless, the available evidence demonstrating that the multidisciplinary approach improves the prognosis of patients with HCC remains relatively weak, resulting from retrospective studies and comparisons with historical control groups [18,19]. In particular, 7 retrospective studies with a control group and an analysis adjusted for confounders were found and scrutinized [20–26]. Of them, 3 included historical controls and 4 contemporary controls. When pooled, these studies reported a statistically significant reduction in mortality in patients managed by the multidisciplinary tumor board (MDT) compared to controls (HR 0.55, 95% CI 0.38–0.80).

Recommendation: For patients with HCC, the panel recommends that the evaluation of the diagnostic and therapeutic workup be carried out by a multidisciplinary team of experts rather than by a single expert.

**Certainty in evidence:** Moderate.

**Strength of recommendation:** Strong in favor of multidisciplinary management.

# 2. In patients with Child-Pugh class A cirrhosis and single HCC, is hepatic resection indicated compared to the treatment with thermal ablation?

Although liver transplantation remains the ideal treatment for all cirrhotic patients with HCC [8], the limited availability of grafts and the growing and improved efficacy of therapeutic alternatives to transplantation have led to consider resection and thermal ablation as first-line options for some of these patients [9,16,27–33].

Even if liver resection has long been the treatment of choice, radiofrequency or microwave thermal ablation have gained widespread use thanks to its less invasiveness and lower risk of complications, high reproducibility, adequate oncological efficacy and feasibility even in patients with a moderate liver dysfunction [34–36]. However, outcomes the feasibility of both resection and thermal ablation are affected by several features, such as underlying liver dysfunction, tumor diameter and position (superficial/deep, adjacent to hollow organs or vascular-biliary structures), general patient conditions (including comorbidity, frailty and Performance Status), that should be evaluated in a multidisciplinary context [37–44], and that are also considered in other Western HCC guidelines [16,27,28].

A total of 27 studies were included in the analysis, 6 randomized controlled trials (RCTs) [38–43] and 21 observational studies [44–64]. RCTs did not demonstrate a difference in overall survival between patients undergoing liver resection and those undergoing thermal ablation at 1, 3 and 5 years, while observational studies showed a survival advantage for surgery at 3 and 5 years. Observational studies also reported a higher recurrence free survival with resection with respect to thermal ablation. However, the long-term benefit of resection was not confirmed in the subgroup of patients with single HCC  $\leq 2$  cm. Therefore, as the two treatments were on-cologically equivalent in these patients, but thermal ablation has a lower risk of complications and a better cost-effectiveness, this choice should be preferred for treating HCC  $\leq 2$  cm whenever the tumor is clearly identifiable and adequately approachable [65,66].

Finally, in a multiparametric evaluation process, particularly when percutaneous thermal ablation procedures are considered unfeasible, it is important to consider the possibility of adopting a mini-invasive approach (laparoscopic or robotic).

Recommendation: In patients with Child-Pugh class A cirrhosis and single HCC, the panel suggests preferring liver resection over thermal ablation, except for patients with HCC  $\leq$ 2 cm, for whom the panel suggests thermal ablation.

Certainty in evidence: Low.

Strength of recommendation: Conditional in favor of resection.

# 3. In cirrhotic patients with good liver function and multinodular HCC, is liver resection indicated compared to transarterial chemoembolization (TACE)?

Approximately 35–40% of HCCs are multinodular at diagnosis [67,68]. The term multinodular includes extremely heterogeneous diseases, from oligo-nodular (2 or 3 nodules) to diffuse miliary disease, which require different treatments. Some proposals for the re-classification of multinodular HCC have been advanced [69–73] which, however, have not yet found correspondence with standard of care therapeutic indications.

The EASL guidelines [16] and AASLD guidance [28] recommend, as first-line treatment for multinodular HCC, transarterial chemoembolization (TACE) or, in oligo-nodular cases, percutaneous thermal ablation if liver transplantation is not feasible according to the selection criteria of the transplant center. This preclusion to resective surgery, especially for patients with 2–3 nodules, is not shared by both the Eastern [29,30] and Italian multisociety [31] guidelines and is not accepted by centers expert in liver surgery [9,67,68,33,74–76]. It should be noted that the BCLC update [77] and AASLD guidance [28], although excluding liver resection as first-line treatment for patients with intermediated stage (multinodular) HCC, consider not only TACE but also liver transplantation (LT) and systemic treatment.

The literature search identified 9 studies, consisting of 1 RCT [78] and 8 observational studies [67,79–85]. The RCT demonstrated a longer survival following liver resection than TACE at 1 year (76% vs 52%, RR 1.47, 95% CI 1.16–1.86), 3 years (64% vs 35%, RR 1.80, 95% CI 1.30–2.50) and 5 years (51% vs 18%, RR 2.90, 95% CI 1.75–4.79). In agreement, observational studies [67,79–85] demonstrated a better survival after liver resection than after TACE at 1year (5 studies, 2511 patients: 88% vs 79%, RR 1.12, 95% CI 1.08–1.16), 3 years (6 studies, 2775 patients: 54% vs 34%, RR 1.59, 95% CI 1.31–1.92) and 5 years (7 studies, 4875 patients: 53% vs 31%, RR 1.70, 95% CI 1.41–2.04).

The panel highlights that 90% of the patients included in the studies evaluated had 2–3 nodules: therefore, this represents the population in which liver resection may outperform TACE. Moreover, it is important to consider that the certainty of the evidence is highly conditioned by a series of limitations of the studies including heterogeneous inclusion criteria, high prevalence of hepatitis B virus etiology and heterogeneous residual liver function at baseline. Hence, due to the low level of evidence, the members of the Panel reiterate the importance of a multidisciplinary treatment evaluation on a case-by-case basis.

Recommendation: In well selected cirrhotic patients with good liver function and oligo-nodular (2–3 nodules) HCC, and after multidisciplinary board evaluation, the panel recommends liver resection over TACE.

**Certainty in evidence:** Low. **Strength of recommendation:** Strong in favor of resection.

### 4. In cirrhotic patients with good liver function and HCC responsible for intrahepatic macrovascular invasion, is liver resection indicated compared to sorafenib-based systemic therapy?

Approximately 10–15% of patients with HCC present with macroscopic vascular invasion (MVI) at diagnosis [68,86,87] with a median survival of 8–11 months [16,88]. The EASL guidelines [16] and AASLD guidance [28] consider MVI a contraindication to hepatic resection, proposing the systemic therapy as the unique option for these patients. Conversely, Eastern guidelines [29,30] consider the possibility of performing liver resection in selected patients, considering the results of numerous studies that have demonstrated the feasibility of the resection even in the presence of MVI, with acceptable postoperative mortality rates (3–6%) and survival at 3 and 5 years (17–49% and 10–39%, respectively) [75,87,89,90]. Similar position has been taken by the Italian multisocietal recommendations [31].

These patients are frequently candidates for a large hepatectomy to obtain surgical radicality [30,87], particularly in presence of MVI of large vessels and, therefore, require an accurate evaluation of the hepatic functional reserve and of the residual liver volume in order to minimize the risk of postoperative liver failure.

There is a clear association between the site of portal MVI and prognosis, and the prognosis is better for MVI of peripheral branches [86,91]. For this reason, portal invasion has been categorized into 4 classes [86,91]: Vp1, invasion of segmental or sectoral portal branches; Vp2, invasion of right or left portal branch; Vp3, invasion extending to the portal trunk; Vp4, invasion extended to the superior mesenteric vein. A survival advantage after surgery compared to nonsurgical treatment has been reported only in the presence of a MVI not extending to the portal trunk) [87,92–94]. The association between the site of invasion and prognosis has been reported even for MVI of hepatic veins [86,90]. Surgery may offer a survival benefit in patients with intrahepatic MVI, but not in those with the tumoral invasion of the inferior vena cava [90].

Four observational studies were identified [95–98]. They enrolled a total of 1143 patients, 618 of whom were treated with liver resection and 525 with systemic therapy. Among these studies, 3 had sorafenib as a control, while the fourth did not specify the systemic therapy used. These studies report a better survival after liver resection at 1 year (3 studies, 879 patients: 65% vs 41%, RR 1.60, 95% CI 1.12–2.29), 3 years (one study, 639 patients: 66% vs 18%, RR 3.82, 95% CI 2.92–5.00,) and 5 years (one study, 639 patients: 56% vs 13%, RR 4.35, 95% CI 3.14–6.03). The improved survival of surgery compared to sorafenib-based systemic therapy was demonstrated both in studies that included only patients with portal and hepatic vein invasion.

However, it is important to note that the overall certainty of the evidence was judged by the Panel to be very low, as it derives from observational studies, and with important limitations such as the risk of bias, imprecision, and poor generalizability. Therefore, their results should be interpreted with great caution particularly considering the impact in term of survival of the new systemic therapies (i.e., combinations based on immunotherapy) compared to so-rafenib [99–101]. Hence, new comparative studies are warranted.

Recommendation: In cirrhotic patients with good liver function and HCC responsible for intrahepatic macrovascular invasion, the panel suggests preferring liver resection versus sorafenib-based systemic therapy.

Certainty in evidence: Very low.

**Strength of recommendation**: Conditional in favor of resection.

# 5. In cirrhotic patients with HCC who are candidates for liver transplantation, is a selection according to the "transplant benefit" criterion indicated compared to other criteria?

In conditions of insufficient organ donation, the selection of patients to be included in the waiting list for liver transplantation (LT) and the intervention priority to assign to each patient ("prioritization") should follow the "transplant benefit" (TB) principle, which integrates the elements of urgency and utility [9,102–108].

The TB is calculated as the difference of the predicted survivals achievable with transplantation and alternative treatments. The major criticality inherent in the application of the TB concept derives from the lack of RCT comparing LT and alternative therapies, stratified by tumor stage, liver function [109,110] and downstaging therapies [111].

TB is also high for patients who suffer an early recurrence (within 2 years) of HCC after potentially radical treatments (resection or thermal ablation), especially if the tumor is multifocal and/or in the presence of deterioration of liver function, as this condition is burdened with an unfavorable prognosis [112].

Recently, Lai et al. [113] have created models of *Intention-to-treat* (ITT) survival with LT and with loco-regional therapies by a retrospective analysis of a large cohort of patients (2103 patients) waiting for LT. They showed that MELD score <13, response to locoregional therapy according to mRECIST criteria (either complete or absent with disease progression), alpha-fetoprotein levels >1000 ng/ml and T1-T2 stages were able to reduce the TB-ITT.

Although there is no broad international agreement, the Italian transplant community considers TB as the reference to select patients for LT and to calculate the priority for intervention [114–117]. In line with the principle of TB, recent cost-efficacy studies have shown that LT is cost-effective only for some categories of patients outside the Milan criteria and, hence, without effective therapeutic alternatives [103], but not for patients eligible for potentially radical alternative therapies such as resection or thermal ablation [118,119].

For all these reasons, the Panel agreed to formulate a Good Practice Statement [120,121] on the importance of adopting the principle of TB as a selection criterion for LT.

Indication of good clinical practice: The Panel believes that the criterion of "transplant benefit" instead of that of "transplant utility" should be used for the selection of candidates for transplantation, taking into account that the potential transplantability must consider overall health, comorbidity, nutritional status and age (although without a defined and universally accepted cut-off), in addition to the tumor burden.

6. In cirrhotic patients beyond the Milan criteria, but without intrahepatic vascular invasion and extrahepatic tumor spread, is liver transplantation indicated compared to alternative treatments (liver resection, locoregional or systemic therapies)?

It is well established that liver transplantation (LT) can provide excellent results also in patients who exceed the oncological limits established by the Milan criteria [122], provided they respect other validated "extended" criteria [123].

The "extended" criteria were established through observational studies that demonstrated that, adopting these criteria, post-transplant overall survival and cancer recurrence-free survival were comparable to those obtained using the Milan criteria [123]. Some extended criteria are purely morphological (largest tumor diameter, number of nodules, total tumor volume), such as the University of California San Francisco criteria [124,125], or the Asian crite-

ria [126,127]. Others combine morphological and biological (alphafetoprotein level or tumor grade) features, such as the Alphafetoprotein model [128,129], the Total Tumor Volume plus Alphafetoprotein criteria [130], the Metroticket 2.0 criteria [131], and the Padova-Toronto criteria [132,133]. All these extended criteria produced consistent "indirect" evidence that LT can guarantee excellent survival profiles (>70% at 5 years) even beyond the Milan criteria. Notably, these survival figures cannot be achieved with any therapy alternative to transplant [33].

Also, direct evidence about comparing LT and non-transplant therapies in patients beyond Milan criteria has been searched for. For the comparison between LT and liver resection, the Panel analyzed a systematic review, including 6 retrospective cohort studies, for evaluating the desired effects [134] and 2 studies for evaluating the undesirable effects [135,136].

Moreover, 3 cohort studies were considered for comparing LT and non-surgical therapies, one comparing LT vs. Sorafenib [137] and 2 comparing LT vs. transarterial chemoembolization (TACE) [138,139].

Observational studies [134] clearly showed a better overall survival (HR 0.83, from 0.68 to 1.01) and recurrence-free survival (0.45, from 0.37 to 0.56) after LT than after resection. Only slightly higher perioperative mortality and undesirable effects were detected after LT [135,136].

A significant long-term survival advantage in favor of LT was also confirmed by observational studies comparing LT with Sorafenib [137] or TACE [138,139].

Clinical recommendation: In potentially transplantable (comorbidities, nutritional status and age) cirrhotic patients with HCC beyond the Milan criteria, but without intrahepatic vascular (and biliary) invasion and extrahepatic tumor spread, the Panel recommends considering liver transplantation rather than alternative treatments (resection, locoregional treatments and systemic treatments).

**Certainty in evidence:** Transplant vs. resection: Moderate. Transplant vs. alternative therapies: Very low.

**Strength of recommendation:** Strong in favor of transplantation.

7. In patients with HCC single >2 cm or multifocal (within the centre's transplantability criteria) and treatable with any potentially radical therapy ("first-line" liver transplantation, resection, or thermal ablation), is "salvage" transplantation (i.e., transplant performed at the time of cancer recurrence/progression after resection or thermal ablation) indicated compared to the "first line" transplant?

The analysis of the role of the so-called "salvage liver transplant" (SLT), performed at the time of tumor recurrence after potentially radical treatments (resection or thermal ablation) is complex [140,141] due to the various factors that can influence the result, including: a) characteristics of the first tumor; b) type of recurrence (early or late, single or multifocal, local or intrahepatic distant); c) applicability of therapies alternative to LT; d) waiting times in list for SLT; e) availability of living transplant donors (LDLT). Due to this complexity, there are no randomized controlled trials comparing SLT with "front-line" transplantation.

Importantly, the option of SLT is in line with the principle of the "transplant benefit" aimed at providing grafts to the patients not amenable to therapeutic alternatives potentially radical [142–145]. Therefore, the possible loss of accessibility to SLT at the time of HCC recurrence represents the main drawback of this strategy.

Five systematic reviews were identified [146–150]. The most up-to-date and methodologically rigorous of them includes a to-tal of 9879 patients [149]. The studies reviewed by this review showed:

### Mortality:

- Mortality at an average follow-up of 1 year: moderately in favor of the SLT strategy (OR 0.86, 95% CI 0.75 0.98);
- Mortality at an average follow-up of 3 years: moderately in favor of the SLT strategy (OR 0.85, 95% CI 0.76 - 0.96);
- Mortality at an average follow-up of 5 years: moderately in favor of the SLT strategy (OR 0.85, 95% CI 0.76 - 0.96);
- Recurrence at a mean follow-up of 1 year: moderately in favor of the SLT strategy (OR 0.86, 95% CI 0.75 -0.99);
- Recurrence at a mean follow-up of 3 years: in favor of the SLT strategy (OR 0.56, 95% CI 0.39 - 0.81);
- Recurrence at a mean follow-up of 5 years: in favor of the SLT strategy (OR 0.75, 95% CI 0.66 - 0.86).

However, the results of this meta-analysis should be viewed with great caution as they do not consider patients who dropped out due to death or progression before receiving the SLT.

Adverse events:

- Biliary tract complications: slightly against SLT (OR 1.14, 95% CI 0.94 1.40);
- Sepsis: slightly against SLT (OR 1.14, 95% CI 0.63 2.06);
- Post-operative bleeding: against SLT (OR 1.32, 95% CI 1.03 1.71);
- Vascular complications: against SLT (OR 1.35, 95% CI 0.98 -1.85);
- Operative mortality: against SLT (OR 2.00, 95% CI 1.21 3.31).

Two cost-effectiveness studies were identified [151,152]. In the first one, SLT after partial hepatectomy led to a longer life expectancy and quality-adjusted life expectancy at a lower cost than front-line transplantation [151]. In the second study, first-line transplantation was superior and dominant over SLT performed after both liver resection and RFA [152].

Information regarding the average time spent on the transplant waiting list and the number of delisting before the intervention was also checked. The analysis of data from 10 studies [135,153–161] showed no significant differences between SLT and front-line transplant.

Clinical recommendation: In patients with HCC single >2 cm or multifocal (within the centre's transplantability criteria) and treatable with any potentially radical therapy, the panel suggests performing liver resection or thermal ablation followed, in case of cancer recurrence, by "salvage" transplantation.

Certainty in evidence: Low.

**Strength of recommendation:** Conditional in favor of liver resection or thermal ablation followed by "rescue" transplant.

# 8. In patients with HCC beyond the transplant criteria adopted by the center, is the "downstaging" procedure followed by transplantation indicated compared to treatments without subsequent transplantation?

A crucial aspect of the "downstaging" concerns the access criteria, i.e., the characteristics of the patients to whom downstaging is offered [162]. In fact, the access criteria can influence two important outcomes: a) the downstaging *failure rate* and the subsequent risk of dropout due to tumor progression while waiting for liver transplant (LT) [163]; b) the *outcomes of LT* and, in particular, the risk of death due to recurrence of HCC [162].

It is also worth noting that, if the outcome LT after downstaging is evaluated with the "intention to treat" method (i.e., the start of the follow-up coincides with the start of downstaging procedures), the overall survival is greatly reduced by the high percentage (21– 50%) of cases who do not complete the downstaging program due to lack of response to therapy - and consequent tumor progression - or worsening liver function or other causes [164]. Besides the entry criteria, the AFP value, and the Child-Pugh class [165,111] can affect the success rate of the downstaging.

An Italian consensus conference on the allocation criteria for LT [166] and a recent "position paper" of the Italian Association for the Study of the Liver [167] did not report criteria of eligibility for the downstaging. What was considered utmost important is the complete or at least partial response to loco-regional or systemic therapies (absolute or biological downstaging), after which patients can be allocated to the category with the highest priority for LT, given the relatively good prognosis after surgery and the temporariness of the results of locoregional treatments to contain the tumor [166].

The term "downstaging" refers to the attempt to reduce the tumor burden throughout liver resection, locoregional therapies such as transarterial chemoembolization (TACE) followed by resection or ablation [112,163,111,168–180] or transarterial radioembolization [181,182] or even systemic therapy, in order to bring back the patient within the transplant criteria adopted by the reference transplant center.

The downstaging can be distinguished in "relative" and "absolute". Relative downstaging has the purpose of bringing the patient back into the transplant criteria adopted by the center. Most of the studies which used this criterion (also defined as "morphological") included the patients in advanced or intermediate stage and without vascular or biliary invasion who were considered transplantable when downstaging reported the tumor within the Milan criteria (Milano-in) [168–171].

A seminal retrospective study by Otto et al. [179] indicated that patients with partial response to TACE had a 5-year survival after LT comparable to that of *ab initio* Milano-in controls, regardless of they met the Milan criteria. Subsequent studies confirmed that the response to locoregional therapies is a key factor to guide the selection of candidates for LT, being able to identify those patients with a favorable tumor biology which, in turn, leads to low post-transplant recurrence rates [112,111,180]. Therefore, a good radiological response to locoregional therapy *before* listing (downstaging) or *while waiting LT* (neoadjuvant therapy) detect the cases for which good medium- and long-term results can be expected. In fact, the good response to treatment is often associated with histopathological markers of good prognosis, such as the absence of micro-vascular invasion and a medium-low degree of tumor undifferentiation [170,182].

Furthermore, to better understand the degree of aggressiveness of the neoplasm, the majority of downstaging protocols adopt the so-called "time test", i.e., the presence of a "stability" of the result achieved with loco-regional therapies for a certain time (usually at least 3 months). The good response to therapy and its "stability" make downstaging (relative or absolute) a better selection system than morphological stage classification [112].

Moreover, a recent American study [172] has highlighted the prognostic role of alpha-fetoprotein (AFP) in patients undergoing downstaging, showing that the reduction of AFP levels below predefined values (for example, 500 or 100 ng/mL) heralds LT results similar to those obtained with *ab initio* Milano-in patients. Such an information underlines the importance of considering, besides the radiological response to therapy, the treatment-induced changes in biological indicators of tumor aggressiveness (18).

Of note, the unique randomized controlled study available on this topic [112] included 45 patients with HCC beyond the Milan criteria, no macrovascular invasion or extrahepatic tumor extension, good liver function (Child-Pugh class A-B7) and estimated post-transplant survival of at least 5 years, who had responded to downstaging with locoregional, surgical or systemic therapy and had received sorafenib for at least 3 months. The mean age was 57, 98.5% they were male. Patients were then randomized to receive LT or to continue to receive locoregional, surgical (liver resection, ablation, TACE and SIRT in various combinations) or systemic therapy The mortality (with a mean follow-up of 5 years) was remarkably lower in transplanted patients than in the counterpart (HR 0.32, 95% CI 0.11 - 0.92). Even the progression-free survival was clearly in favor of LT (HR 0.20, 95% CI 0.07 - 0.57).

A prospective observational study [174] conducted in China included 66 patients with intermediate HCC that met the following selection criteria for downstaging: single tumor <8 cm or 2- 3 tumors <5 cm and total diameter <8 cm, without vascular invasion and who had responded to locoregional downstaging therapy (TACE and/or thermal ablation). The patients underwent resection (n. 35) if they had cirrhosis and preserved liver function or LT (n. 31) if resection was not feasible for anatomical reasons and a live or deceased donor liver was available (n. 31). The mean age was 44 years; males were 60%. The baseline characteristics of patients were similar in the two groups. The study showed a mortality tendentially but not non significantly lower after LT compared to successful resection (HR 0.72, 95% CI 0.32 –1.62).

Therefore, although most of the evidence currently available regarding the downstaging comes from uncontrolled studies, they would indicate that this procedure, when succeeds in reducing tumor mass, is associated with a post-transplant survival similar to that obtained in patients who *ab initio* respect to the selection criteria for transplantation adopted by the center [168–180,183].

Even recurrence-free survival appears to be often superimposable, although a recent work reports a non-significant increase in recurrences in the downstaging group compared to controls at 5year (18). Similarly, an Italian single-center study including 43 patients who successfully underwent downstaging (transition from stage T3 to T2) reports a lower 5-year survival of down-staged patients (although not significantly) compared to patients initially in stage T2 (62% vs 76%) and a higher rate of tumor recurrence (20.9% vs 7.6%) [184].

Clinical recommendation: In patients with HCC beyond the oncological transplant criteria adopted by the center, the panel recommends to perform the downstaging procedure aimed at bringing the patient back to the adopted transplant criteria compared to all other therapies without transplantation.

Certainty in evidence: Low.

**Strength of recommendation**: Strong in favor of transplantation.

### 5. Future perspective

Future studies and up-dated treatment guidelines should: 1) evaluate the role of systemic therapies in conversion strategies [9] in a perspective of a forthcoming evolution in the management of advanced HCC; 2) evaluate the impact of adjuvant strategies [185–187]; 3) better define the role of pre-planned combined treatment strategies; 4) assess the surrogacy of intermediate radiological endpoints across different HCC stages and treatments; 5) include liver-related evolutionary events [187–191] to better understand competing risks with survival.

Given the complexity of the disease and the large number of potentially useful therapies, it is not surprising that the expertise of many physicians is required to provide optimal care to patients with HCC; so, patients diagnosed with liver cancer on cirrhosis should be referred to multidisciplinary teams. Finally, the role of *expert multidisciplinary tumor board*, able to adopt a personalized therapeutic approach tailored to the characteristics of each patient, should be further evaluated, and emphasized.

### **Conflict of interest**

The authors declare that there are no conflicts of interest.

### Acknowledgements

The authors thank Silvia Minozzi: Istituto di Ricerche Farmacologiche Mario Negri IRCCS - Milano Marta Monteforte: Istituto di Ricerche Farmacologiche Mario Negri IRCCS - Milano Veronica Andrea Fittipaldo: Istituto di Ricerche Farmacologiche Mario Negri IRCCS - Milano Gianluca Masi: Oncologia Medica 2 Universitaria, Azienda Ospedaliera Universitaria Pisana, Pisa Francesco Fiore: Radiologia Interventistica, Istituto Nazionale Tumori Fondazione Pascale, Napoli Luigi Maria Terracciano: Divisione di Anatomia Patologica, Humanitas University Hospital, Milano Cesare Guida: Radioterapia, Ospedale del Mare, Napoli Matteo Cescon: Chirurgia epatobiliare e dei trapianti dell'IRCCS Azienda Ospedaliera-Universitaria di Bologna Salvatore Gruttadauria: Dipartimento per la cura e lo studio delle patologie addominali IRCCS ISMETT UPMCI Massimo Alberto Iavarone: Gastroenterologia ed epatologia, Fondazione IRCCS Ca Granda, Ospedale Maggiore, Policlinico di Milano.

### References

- https://snlg.iss.it/wp-content/uploads/2023/02/LG97\_AISF-AIOM\_ Epatocarcinoma.pdf
- [2] Guyatt GH, Oxman AD, Vist GE, et al. GRADE Working Group. GRADE: an emerging consensus on rating quality of evidence and strength of recommendations. BMJ 2008;336:924–6.
- [3] Rumgay H, Arnold M, Ferlay J, et al. Global burden of primary liver cancer in 2020 and predictions to 2040. J Hepatol 2022;77(6):1598–606.
- [4] Garuti F, Neri A, Avanzato F, et al. The changing scenario of hepatocellular carcinoma in Italy: an update. Liver Int 2021;41(3):585–97.
- [5] Cabibbo G, Enea M, Attanasio M, et al. A meta-analysis of survival rates of untreated patients in randomized clinical trials of hepatocellular carcinoma. Hepatology 2010;51(4):1274–83.
- [6] Giannini EG, Farinati F, Ciccarese F, et al. Prognosis of untreated hepatocellular carcinoma. Hepatology 2015;61(1):184–90.
- [7] Cammà C, Cabibbo G. Prognostic scores for hepatocellular carcinoma: none is the winner. Liver Int 2009;29(4):478–80.
- [8] Singal AG, Zhang E, Narasimman M, et al. HCC surveillance improves early detection, curative treatment receipt, and survival in patients with cirrhosis: a meta-analysis. J Hepatol 2022;77(1):128–39.
- [9] Vitale A, Cabibbo G, lavarone M, et al. HCC Special Interest Group of the Italian Association for the Study of the Liver. Personalised management of patients with hepatocellular carcinoma: a multiparametric therapeutic hierarchy concept. Lancet Oncol 2023;24(7):e312–22.
- [10] CNEC Centro Nazionale per l'Eccellenza delle Cure Manuale metodologico per la produzione di linee guida di pratica clinica, Roma: ISS - Istituto Superiore di Sanità; 2020. Available at https://snlg.iss.it/wp-content/uploads/ 2019/04/MM\_v1.3.2\_apr\_2019.pdf.
- [11] GRADEpro gdt [Computer program] mcmaster university (developed by evidence prime) GRADEpro gdt, Hamilton (ON): McMaster University; 2021. Version accessed 5 November developed by Evidence PrimeAvailable at grade-pro.org.
- [12] Higgins JGS. Cochrane handbook for systematic reviews of interventions; 2011.
- [13] Wells GA, Shea B, O'Connell D, Peterson J, Welch V, Losos M, Tugwell P. The Newcastle-Ottawa Scale (NOS) for assessing the quality of nonrandomised studies in meta-analyses; 2000.
- [14] Cabibbo G, Latteri F, Antonucci M, et al. Multimodal approaches to the treatment of hepatocellular carcinoma. Nat Clin Pract Gastroenterol Hepatol 2009;6(3):159–69.
- [15] Cabibbo G, Aghemo A, Lai Q, et al. Optimizing systemic therapy for advanced hepatocellular carcinoma: the key role of liver function. Dig Liver Dis 2022;54(4):452–60.
- [16] European Association for the Study of the LiverEASL Clinical Practice Guidelines: management of hepatocellular carcinoma. J Hepatol 2018;69(1):182–236.
- [17] Cabibbo G, Petta S, Barbara M, Italian Liver Cancer (ITALICA) group, et al. Hepatic decompensation is the major driver of death in HCV-infected cirrhotic patients with successfully treated early hepatocellular carcinoma. J Hepatol 2017;67(1):65–71.
- [18] Byrd K, Alqahtani S, Yopp AC, et al. Role of multidisciplinary care in the management of hepatocellular carcinoma. Semin Liver Dis 2021;41(1):1–8.
- [19] Seif El Dahan K, Reczek A, Daher D, et al. Multidisciplinary care for patients with HCC: a systematic review and meta-analysis. Hepatol Commun 2023;7:e0143.
- [20] Agarwal PD, Phillips P, Hillman L, et al. Multidisciplinary management of hepatocellular carcinoma improves access to therapy and patient survival. J Clin Gastroenterol 2017;51(9):845–9.
- [21] Casadei Gardini A, Scarpi E, Foschi FG, et al. Impact of physician experience and multidisciplinary team on clinical outcome in patients receiving Sorafenib. Clin Res Hepatol Gastroenterol 2019;43(5):e76–8.

- [22] Duininck G, Lopez-Aguiar AG, Lee RM, et al. Optimizing cancer care for hepatocellular carcinoma at a safety-net hospital: the value of a multidisciplinary disease management team. J Surg Oncol 2019;120(8):1365–70.
- [23] Gaba RC, Kallwitz ER, Parvinian A, et al. Imaging surveillance and multidisciplinary review improves curative therapy access and survival in HCC patients. Ann Hepatol 2013;12(5):766–73.
- [24] Serper M, Taddei TH, Mehta R, et al. Association of provider specialty and multidisciplinary care with hepatocellular carcinoma treatment and mortality. Gastroenterology 2017;152(8):1954–64.
- [25] Sinn DH, Choi GS, Park HC, et al. Multidisciplinary approach is associated with improved survival of hepatocellular carcinoma patients. PLoS ONE 2019;14(1):e0210730.
- [26] Yopp AC, Mansour JC, Beg MS, et al. Establishment of a multidisciplinary hepatocellular carcinoma clinic is associated with improved clinical outcome. Ann Surg Oncol 2014;21(4):1287–95.
- [27] Marrero JA, Kulik LM, Sirlin CB, et al. Diagnosis, staging, and management of hepatocellular carcinoma: 2018 practice guidance by the American Association for the Study of Liver Diseases. Hepatology 2018;68:723–50.
  [28] Singal AG, Llovet JM, Yarchoan M, et al. AASLD practice guidance on pre-
- [28] Singal AG, Llovet JM, Yarchoan M, et al. AASLD practice guidance on prevention, diagnosis, and treatment of hepatocellular carcinoma. Hepatology 2023;78(6):1922–65.
- [29] Korean Liver Cancer Association; National Cancer Center2018 Korean Liver Cancer Association-National Cancer Center Korea Practice Guidelines for the Management of Hepatocellular Carcinoma. Gut Liver 2019;13:227–99.
- [30] Kokudo N, Hasegawa K, Akahane M, et al. Evidence-based clinical practice guidelines for hepatocellular carcinoma: the Japan society of hepatology 2013 update (3rd JSH-HCC Guidelines). Hepatol Res 2015:45.
- [31] Raccomandazioni Multisocietarie Italiane (AISF, AIOM, IT-IHPBA, SIC, SIRM, SITO) per la gestione clinica integrata del paziente con epatocarcinoma. https://www.webaisf.org/wpcontent/uploads/2019/02/position\_paper\_ hcc\_v30\_22.12.1.pdf
- [32] Vitale A, Peck-Radosavljevic M, Giannini EG, et al. Personalized treatment of patients with very early hepatocellular carcinoma. J Hepatol 2017;66(2):412–23.
- [33] Kawaguchi Y, Hasegawa K, Hagiwara Y, et al. Effect of diameter and number of hepatocellular carcinomas on survival after resection, transarterial chemoembolization, and ablation. Am J Gastroenterol 2021;116(8):1698–708.
- [34] Livraghi T, Meloni F, Di Stasi M, et al. Sustained complete response and complications rates after radiofrequency ablation of very early hepatocellular carcinoma in cirrhosis: is resection still the treatment of choice? Hepatology 2008;47:82–9.
- [35] Jia Z, Zhang H, Li N. Evaluation of clinical outcomes of radiofrequency ablation and surgical resection for hepatocellular carcinoma conforming to the Milan criteria: a systematic review and meta-analysis of recent randomized controlled trials. J Gastroenterol Hepatol 2021;36(7):1769–77.
- [36] Shin SW, Ahn KS, Kim SW, et al. Liver resection versus local ablation therapies for hepatocellular carcinoma within the Milan criteria: a systematic review and meta-analysis. Ann Surg 2021;273(4):656–66.
- [37] Viganò L, Laurenzi A, Solbiati L, et al. Open liver resection, laparoscopic liver resection, and percutaneous thermal ablation for patients with solitary small hepatocellular carcinoma (≤30mm): review of the literature and proposal for a therapeutic strategy. Dig Surg 2018;35(4):359–71.
- [38] Chen MS, Li JQ, Zheng Y, et al. A prospective randomized trial comparing percutaneous local ablative therapy and partial hepatectomy for small hepatocellular carcinoma. Ann Surg 2006;243(3):321–8.
- [39] Huang J, Yan L, Cheng Z, et al. A randomized trial comparing radiofrequency ablation and surgical resection for HCC conforming to the Milan criteria. Ann Surg 2010;252(6):903–12.
- [40] Feng K, Yan J, Li X, et al. A randomized controlled trial of radiofrequency ablation and surgical resection in the treatment of small hepatocellular carcinoma. J Hepatol 2012;57(4):794–802.
- [41] Fang Y, Chen W, Liang X, et al. Comparison of long-term effectiveness and complications of radiofrequency ablation with hepatectomy for small hepatocellular carcinoma. J Gastroenterol Hepatol 2014;29(1):193–200.
- [42] Ng K, Chok Chan A, et al. Randomized clinical trial of hepatic resection versus radiofrequency ablation for early-stage hepatocellular carcinoma. Br J Surg 2017;104(13):1775–84.
- [43] Lee H, Lee J, Yoon J, et al. A prospective randomized study comparing radiofrequency ablation and hepatic resection for hepatocellular carcinoma 2018. Ann Surg Treat Res 2018;94(2):74–82.
- [44] Lee S, Kang T, Cha D, et al. Radiofrequency ablation vs. surgery for perivascular hepatocellular carcinoma: propensity score analyses of long-term outcomes. J Hepatol 2018;69(1):70–8.
- [45] Chong C, Lee KF, Chu CM, et al. Microwave ablation provides better survival than liver resection for hepatocellular carcinoma in patients with borderline liver function: application of ALBI score to patient selection. HPB 2018;20(6):546–54.
- [46] Chong CC, Lee KF, Chu CM, et al. Laparoscopic hepatectomy (with or without Robotic Assistance) versus radiofrequency ablation as a minimally invasive treatment for very early-stage or early-stage hepatocellular carcinoma. Dig Surg 2020;37(1):65–71.
- [47] Chu HH, Kim JH, Kim PN, et al. Surgical resection versus radiofrequency ablation very early-stage HCC (≤ 2cm Single HCC): a propensity score analysis. Liver Int 2019;39(12):2397–407.

- [48] Conticchio M, Inchingolo R, Delvecchio A, et al. Radiofrequency ablation vs surgical resection in elderly patients with hepatocellular carcinoma in Milan criteria. World J Gastroenterol 2021;27(18):2205–18.
- [49] Di Sandro S, Benuzzi L, Lauterio A, et al. Single Hepatocellular Carcinoma approached by curative-intent treatment: a propensity score analysis comparing radiofrequency ablation and liver resection. Eur J Surg Oncol 2019;45(9):1691–9.
- [50] Harada N, Maeda T, Yoshizumi T, et al. Laparoscopic liver resection is a feasible treatment for patients with hepatocellular carcinoma and portal hypertension. Anticancer Res 2016;36(7):3489–97.
- [51] He W, Li B, Zheng Y, et al. Resection vs. ablation for alpha-fetoprotein positive hepatocellular carcinoma within the Milan criteria: a propensity score analysis. Liver Int 2016;36(11):1677–87.
- [52] Ito T, Tanaka S, Iwai S, et al. Outcomes of laparoscopic hepatic resection versus percutaneous radiofrequency ablation for hepatocellular carcinoma located at the liver surface: a case-control study with propensity score matching. Hepatol Res 2016;46(6):565–74.
- [53] Kaibori M, Yoshii K, Hasegawa K, et al. Treatment optimization for hepatocellular carcinoma in elderly patients in a japanese nationwide cohort. Ann Surg 2019;270(1):121–30.
- [54] Kato Y, Okamura Y, Omae K, et al. Propensity score-matched comparison of non-anatomical resection and radiofrequency ablation for hepatocellular carcinoma in patients with up to three tumours, each measuring up to 3cm in diameter. BJS Open 2018;2(4):213–19.
- [55] Kim T, Chang J, Um S, et al. Comparison of 2 curative treatment options for very early hepatocellular carcinoma: efficacy, recurrence pattern, and retreatment. Medicine 2019;98(26):e16279.
- [56] Li Y, Chen P, Yeh, et al. Clinical outcomes of surgical resection versus radiofrequency ablation in very-early-stage hepatocellular carcinoma: a propensity score matching analysis. BMC Gastroenterol 2021;21(1):418.
- [57] Liu P, Hsu Y, Hsia C, et al. Surgical resection versus radiofrequency ablation for single hepatocellular carcinoma ≤ 2cm in a propensity score model. Ann Sur 2016;263(3):538–45.
- [58] Miura JT, Johnston FM, Tsai S, et al. Surgical resection versus ablation for hepatocellular carcinoma  $\leq$  3 cm: a population-based analysis. HPB 2015;17(10):896–901.
- [59] Pan Y, Long Q, Yi M, et al. Radiofrequency ablation versus laparoscopic hepatectomy for hepatocellular carcinoma: a real world single center study. Eur J Surg Oncol 2020;46(4 Pt A):548–59.
- [60] Pompili M, Saviano A, De Matthaeis N, et al. Long-term effectiveness of resection and radiofrequency ablation for single hepatocellular carcinoma ≤3cm. Results of a multicenter Italian survey. J Hepatol 2013;59(1):89–97.
- [61] Ryu T, Takami Y, Wada Y, et al. Hepatic resection versus operative microwave ablation for single hepatocellular carcinoma ≤5 cm: a propensity scorematched analysis. Surgery 2019;166(3):254–62.
- [62] Song J, Yu Wang, Ma K, et al. Laparoscopic hepatectomy versus radiofrequency ablation for minimally invasive treatment of single, small hepatocellular carcinomas. Surg Endosc 2016;30(10):4249–57.
- [63] Takayasu K, Arii S, Sakamoto M, et al. Impact of resection and ablation for single hypovascular hepatocellular carcinoma ≤2cm analysed with propensity score weighting. Liver Int 2018;38(3):484–93.
- [64] Wang J, Houng W, Chi Chih, et al. Survival comparison between surgical resection and radiofrequency ablation for patients in BCLC very early/early-stage hepatocellular carcinoma. J Hepatol 2012;56(2):412–18.
- [65] Thein HH, Isaranuwuatchai I, Campitelli M, et al. Health care costs associated with hepatocellular carcinoma: a population-based study. Hepatology 2013;58(4):1375–84.
- [66] Cucchetti A, Piscaglia F, Cescon M, et al. Cost-effectiveness of hepatic resection versus percutaneous radiofrequency ablation for early hepatocellular carcinoma. J Hepatol 2013;59(2):300–7.
- [67] Fukami Y, Kaneoka Y, Maeda A, et al. Liver resection for multiple hepatocellular carcinomas: a japanese nationwide survey. Ann Surg 2020;272:145–54.
- [68] Roayaie S, Jibara G, Tabrizian P, et al. The role of hepatic resection in the treatment of hepatocellular cancer. Hepatology 2015;62:440–51.
- [69] Bolondi L, Burroughs A, Dufour JF, et al. Heterogeneity of patients with intermediate (BCLC B) hepatocellular carcinoma: proposal for a subclassification to facilitate treatment decisions. Semin Liver Dis 2012;32:348–59.
- [70] Ha Y, Shim JH, Kim SO, et al. Clinical appraisal of the recently proposed Barcelona Clinic Liver Cancer Stage B subclassification by survival analysis. J Gastroenterol Hepatol 2014;29:787–93.
- [71] Kim JH, Shim JH, Lee HC, et al. New intermediate-stage subclassification for patients with hepatocellular carcinoma treated with transarterial chemoembolization. Liver Int 2017;37:1861–8.
- [72] Yamakado K, Miyayama S, Hirota S, et al. Prognosis of patients with intermediate-stage hepatocellular carcinomas based on the Child-Pugh score: subclassifying the intermediate stage (Barcelona Clinic Liver Cancer stage B). Jpn J Radiol 2014;32:644–9.
- [73] Kudo M, Arizumi T, Ueshima K, et al. Subclassification of BCLC B stage hepatocellular carcinoma and treatment strategies: proposal of modified Bolondi's subclassification (Kinki criteria). Dig Dis 2015;33:751–8.
- [74] Ishizawa T, Hasegawa K, Aoki T, et al. Neither multiple tumors nor portal hypertension are surgical contraindications for hepatocellular carcinoma. Gastroenterology 2008;134:1908–16.

- [75] Torzilli G, Belghiti J, Kokudo N, et al. A snapshot of the effective indications and results of surgery for hepatocellular carcinoma in tertiary referral centers: is it adherent to the EASL/AASLD recommendations? An observational study of the HCC East-West study group. Ann Surg 2013;257:929–37.
- [76] Vitale A, Burra P, Frigo AC, et al. Survival benefit of liver resection for patients with hepatocellular carcinoma across different Barcelona Clinic Liver Cancer stages: a multicentre study. J Hepatol 2015;62:617–24.
- [77] Reig M, Forner A, Rimola J, Ferrer-Fàbrega J, Burrel M, Garcia-Criado Á, Kelley RK, Galle PR, Mazzaferro V, Salem R, Sangro B, Singal AG, Vogel A, Fuster J, Ayuso C, Bruix J. BCLC strategy for prognosis prediction and treatment recommendation: the 2022 update. J Hepatol 2022;76(3):681–93.
- [78] Yin L, Li H, Li AJ, et al. Partial hepatectomy vs. transcatheter arterial chemoembolization for resectable multiple hepatocellular carcinoma beyond Milan Criteria: a RCT. J Hepatol 2014;61(1):82–8.
- [79] Zhong JH, Ke Y, Gong WF, et al. Hepatic resection associated with good survival for selected patients with intermediate and advanced-stage hepatocellular carcinoma. Ann Surg 2014;260(2):329–40.
- [80] Tada T, Kumada T, Toyoda H, et al. Role of hepatic resection in patients with intermediate-stage hepatocellular carcinoma: a multicenter study from Japan. Cancer Sci 2017;108(7):1414–20.
- [81] Chen S, Jin H, Dai Z, et al. Liver resection versus transarterial chemoembolization for the treatment of intermediate- stage hepatocellular carcinoma. Cancer Med 2019;8(4):1530–9.
- [82] Lin CW, Chen YS, Lo GH, et al. Comparison of overall survival on surgical resection versus transarterial chemoembolization with or without radiofrequency ablation in intermediate stage hepatocellular carcinoma: a propensity score matching analysis. BMC Gastroenterol 2020;20(1):99.
- [83] Peng Y, Liu F, Xu H, et al. Is laparoscopic liver resection suitable for selected patients with BCLC stage B HCC? A propensity score-matched analysis. HPB (Oxford) 2020;22(4):595–602.
- [84] Oh JH, Sinn DH, Choi GS, et al. Comparison of outcome between liver resection, radiofrequency ablation, and transarterial therapy for multiple small hepatocellular carcinoma within the Milan criteria. Ann Surg Treat Res 2020;99(4):238–46.
- [85] Lu L, Zheng P, Wu Z, et al. Hepatic resection versus transarterial chemoembolization for intermediate-stage hepatocellular carcinoma: a Cohort Study. Front Oncol 2021;11:618937.
- [86] Ikai I, Arii S, Okazaki M, et al. Report of the 17th nationwide follow-up survey of primary liver cancer in Japan. Hepatol Res 2007;37:676–91.
- [87] Kokudo T, Hasegawa K, Matsuyama Y, et al. Liver Cancer Study Group of Japan. Survival benefit of liver resection for hepatocellular carcinoma associated with portal vein invasion. J Hepatol 2016;65:938–43.
- [88] Giannini EG, Bucci L, Garuti F, et al. Italian Liver Cancer (ITA.LI.CA) group. Patients with advanced hepatocellular carcinoma need a personalized management: a lesson from clinical practice. Hepatology 2018;67:1784–96.
- [89] Glantzounis GK, Paliouras A, Stylianidi MC, et al. The role of liver resection in the management of intermediate and advanced stage hepatocellular carcinoma. A systematic review. Eur J Surg Oncol 2018;44:195–208.
- [90] Kokudo T, Hasegawa K, Yamamoto S, et al. Surgical treatment of hepatocellular carcinoma associated with hepatic vein tumor thrombosis. J Hepatol 2014;61:583–8.
- [91] Shi J, Lai EC, Li N, et al. Surgical treatment of hepatocellular carcinoma with portal vein tumor thrombus. Ann Surg Oncol 2010;17:2073–80.
- [92] Guarino M, Cucchetti A, Pontillo G, et al. Pattern of macrovascular invasion in hepatocellular carcinoma. Eur J Clin Invest 2021;51(7):e13542.
- [93] Zheng N, Wei X, Zhang D, Chai W, et al. Hepatic resection or transarterial chemoembolization for hepatocellular carcinoma with portal vein tumor thrombus. Medicine 2016;95:e3959.
- [94] Wang K, Guo WX, Chen MS, et al. Multimodality Treatment for Hepatocellular Carcinoma with portal vein tumor thrombus: a large-scale, multicenter, propensity mathching score analysis. Medicine 2016;95:e3015.
- [95] Mei J, Li SH, Wang QX, et al. Resection vs. sorafenib for hepatocellular carcinoma with macroscopic vascular invasion: a real world, propensity score matched analytic study. Front Oncol 2020;10:573.
- [96] Famularo S, Donadon M, Cipriani F, et al. Hepatectomy versus sorafenib in advanced nonmetastatic hepatocellular carcinoma: a real-life multicentric weighted comparison. Ann Surg 2022;275(4):743–52.
- [97] Govalan R, Lauzon M, Luu M, et al. Comparison of surgical resection and systemic treatment for hepatocellular carcinoma with vascular invasion: national cancer database analysis. Liver Cancer 2021;10(5):407–18.
- [98] MaÃàhringer-Kunz A, Steinle V, Kloeckner R, et al. The impact of portal vein tumor thrombosis on survival in patients with hepatocellular carcinoma treated with different therapies: a cohort study. PLoS ONE 2021;16(5):e0249426.
- [99] Llovet JM, Ricci S, Mazzaferro V, et al. Sorafenib in advanced hepatocellular carcinoma. N Engl J Med 2008;359(4):378–90.
- [100] Finn RS, Qin S, Ikeda M, et al. Atezolizumab plus bevacizumab in unresectable hepatocellular carcinoma. N Engl J Med 2020;382(20):1894–905.
- [101] Abou-Alfa GK, Lau G, Kudo M, et al. Tremelimumab plus durvalumab in unresectable hepatocellular carcinoma. NEJM Evid 2022;1:EVIDoa2100070. doi:10.1056/EVIDoa2100070.
- [102] Persad G, Wertheimer A, Emanuel EJ. Principles for allocation of scarce medical interventions. Lancet 2009;373:423–31.

- [103] Vitale A, Farinati F, Burra P, et al. Italian Liver Cancer Group Utility-based criteria for selecting patients with hepatocellular carcinoma for liver transplantation: a multicenter cohort study using the alpha- fetoprotein model as
- a survival predictor. Liver Transpl 2015;21:1250–8.
  [104] Vitale A, Huo TL, Cucchetti A, et al. Survival benefit of liver transplantation versus resection for hepatocellular carcinoma: impact of MELD score. Ann Surg Oncol 2015;22:1901–7.
- [105] Schaubel DE, Guidinger MK, Biggins SW, et al. Survival benefit-based deceased-donor liver allocation. Am J Transplant 2009;9:970–81.
- [106] Berry K, Ioannou GN. Are patients with Child's A cirrhosis and hepatocellular carcinoma appropriate candidates for liver transplantation? Am J Transplant 2012;12:706–17.
- [107] Cillo U, Vitale A, Volk ML, et al. The survival benefit of liver transplantation in hepatocellular carcinoma patients. Dig Liver Dis 2010;42:642–9.
  [108] Vitale A, Morales RR, Zanus G, et al. Barcelona Clinic Liver Cancer stag-
- [108] Vitale A, Morales RR, Zanus G, et al. Barcelona Clinic Liver Cancer staging and transplant survival benefit for patients with hepatocellular carcinoma: a multicentre, cohort study; Italian Liver Cancer group. Lancet Oncol 2011;12:654–62.
- [109] Cillo U, Vitale A, Polacco M, et al. Liver transplantation for hepatocellular carcinoma through the lens of transplant benefit. Hepatology 2017;65:1741–8.
- [110] Takayasu K, Arii S, Kudo M, et al. Superselective transarterial chemoembolization for hepatocellular carcinoma. Validation of treatment algorithm proposed by Japanese guidelines. J Hepatol 2012;56:886–92.
- [111] Vitale A, Scolari F, Bertacco A, et al. Sustained complete response after biological downstaging in patients with hepatocellular carcinoma: XXL-Like prioritization for liver transplantation or "Wait and See" Strategy? Cancers 2021;13(10):2406.
- [112] Mazzaferro V, Citterio D, Bhoori S, et al. Liver transplantation in hepatocellular carcinoma after tumour downstaging (XXL): a randomised, controlled, phase 2b/3 trial. Lancet Oncol 2020;21:947–56.
- [113] Lai Q, Vitale A, Iesari S, et al. Intention-to-treat survival benefit of liver transplantation in patients with hepatocellular cancer. Hepatology 2017;66(1):910–1919.
- [114] Vitale A, Volk ML, De Feo TM, et al. A method for establishing allocation equity among patients with and without hepatocellular carcinoma on a common liver transplant waiting list. J Hepatol 2014;60(2):290–7.
- [115] Cillo U, Burra P, Mazzaferro V, et al. A multistep, consensus-based approach to organ allocation in liver transplantation: toward a "Blended Principle Model". Am J Transplant 2015;15:2552–61.
- [116] Mehta N, Bhangui P, Yao FY, et al. Liver transplantation for hepatocellular carcinoma. Working Group Report from the ILTS transplant oncology consensus conference. Transplantation 2020;104:1136–42.
- [117] Thorburn D, Taylor R, Whitney J, et al. Resuming liver transplantation amid the COVID-19 pandemic. Lancet Gastroenterol Hepatol 2021;6(1):12–13.
- [118] Spolverato G, Vitale A, Ejaz A, et al. The relative net health benefit of liver resection, ablation, and transplantation for early hepatocellular carcinoma. World J Surg 2015;39(6):1474–84.
- [119] Kim KC, Wang VW, Siddiqui FJ, et al. Cost-effectiveness analysis of liver resection versus transplantation for early hepatocellular carcinoma within the Milan criteria. Hepatology 2015;61(1):227–37.
- [120] Lofti T, Hajizadeh A, Moja L, et al. A taxonomy and framework for identifying and developing actionable statements in guidelines suggests avoiding informal recommendations. J Clin Epidemiol 2021;23 S0895- 4356(21)00314-0.
- [121] Guyatt GH, Alonso-Coello P, Schünemann HJ, et al. Guideline panels should seldom make good practice statements: guidance from the GRADE Working Group. J Clin Epidemiol 2016;80:3–7.
- [122] Mazzaferro V, Regalia E, Doci R, et al. Liver transplantation for the treatment of small hepatocellular carcinomas in patients with cirrhosis. N Engl J Med 1996;334(11):693–9.
- [123] Italian Association for the Study of the Liver (AISF), AISF Expert Panel. Position paper of the Italian Association for the Study of the Liver (AISF): the multidisciplinary clinical approach to hepatocellular carcinoma. Dig Liver Dis 2013;45(9):712–23.
- [124] Yao FY, Xiao L, Bass NM, et al. Liver transplantation for hepatocellular carcinoma: validation of the UCSF- expanded criteria based on preoperative imaging. Am J Transplant 2007;7:2587–96.
- [125] Mazzaferro V, Llovet JM, Miceli R, et al. Predicting survival after liver transplantation in patients with hepatocellular carcinoma beyond the Milan criteria: a retrospective, exploratory analysis. Lancet Oncol 2009;10:35–43.
- [126] Lee SG, Hwang S, Moon DB, et al. Expanded indication criteria of living donor liver transplantation for hepatocellular carcinoma at one large-volume center. Liver Transpl 2008;14:935–45.
- [127] Bonadio I, Colle I, Geerts A, et al. Liver transplantation for hepatocellular carcinoma comparing the Milan, UCSF, and Asan criteria: long-term follow-up of a Western single institutional experience. Clin Transplant 2015;29:425–33.
- [128] Duvoux C, Roudot-Thoraval F, Decaens T, et al. Liver transplantation for hepatocellular carcinoma: a model including alpha-fetoprotein improves the performance of Milan criteria. Gastroenterology 2012;143:986–94.
- [129] Notarpaolo A, Layese R, Magistri P, et al. Validation of the AFP model as a predictor of HCC recurrence in patients with viral hepatitis-related cirrhosis who had received a liver transplant for HCC. J Hepatol 2017;66(3):552–9.
- [130] Toso C, Meeberg G, Hernandez-Alejandro R, et al. Total tumor volume and alpha-fetoprotein for selection of transplant candidates with hepatocellular carcinoma: a prospective validation. Hepatology 2015;62:158–65.

- [131] Mazzaferro V, Sposito C, Zhou J, et al. Metroticket 2.0 model for analysis of competing risks of death after liver transplantation for hepatocellular carcinoma. Gastroenterology 2018;154(1):128–39.
- [132] Cillo U, Vitale A, Bassanello M, et al. Liver transplantation for the treatment of moderately or well-differentiated hepatocellular carcinoma. Ann Surg 2004;239(2):150–9.
- [133] Sapisochin G, Goldaracena N, Laurence JM, et al. The extended Toronto criteria for liver transplantation in patients with hepatocellular carcinoma: a prospective validation study. Hepatology 2016;64(6):2077–88.
- [134] Li C, Wen TF, Yan L, et al. Liver resection versus liver resection plus TACE for patients with hepatocellular carcinoma beyond Milan criteria. J Surg Res 2017;209(8–16).
- [135] Facciuto M, Koneru B, Rocca J, et al. Surgical treatment of hepatocellular carcinoma beyond Milan criteria. Results of liver resection, salvage transplantation, and primary liver transplantation. Ann Surl Oncol 2008;15(5):1383–91.
- [136] Lee KK, Kim DG, Moon I, et al. Liver transplantation versus liver resection for the treatment of hepatocellular carcinoma. J Surg Oncol 2010;101(1):47–53.
- [137] Cho Y, Lee JH, Lee DH, et al. Comparison of treatment outcome between living donor liver transplantation and Sorafenib for patients with hepatocellular carcinoma beyond the Milan criteria. Oncotarget 2017;8(29):47555–64.
- [138] Heinzow HS, Brockmann JG, Vahler M, et al. Liver transplantation versus supraselective transarterial chemoembolization in palliative patients with hepatocellular carcinoma exceeding the Milan Criteria - Is it time or a more individual approach? Ann Transplant 2013;18:515–24.
- [139] Kim J, Kwon C, Joh J, et al. Patients with unresectable hepatocellular carcinoma beyond Milan criteria: should we perform transarterial chemoembolization or liver transplantation? Transplant Proc 2010;42(3):821–4 20.Thein HH, Isaranuwatchai W, Campitelli MA, et al. Health Care Costs Associated with Hepatocellular Carcinoma: A Population-Based Study. Hepatology 2013;58(4):1375-84.
- [140] Gelli M, Sebagh M, Porcher R, et al. Liver Resection for early hepatocellular carcinoma: preoperative predictors of non-transplantable recurrence and implications for treatment allocation. Ann Surg 2020;272:820–6.
- [141] de Haas RJ, Lim C, Bhangui P, et al. Curative salvage liver transplantation in patients with cirrhosis and hepatocellular carcinoma: an intention-to-treat analysis. Hepatology 2018;67:204–15.
- [142] Cillo U, Vitale A, Volk ML, et al. Liver transplantation for T2 hepatocellular carcinoma during the COVID-19 pandemic: a novel model balancing individual benefit against healthcare resources. Cancers 2021;13(6):1416.
- [143] Vitale A, Volk M, Cillo U. Transplant benefit for patients with hepatocellular carcinoma. World J Gastroenterol 2013;19(48):9183–8.
- [144] Vitale A, Farinati F, Pawlik TM, et al. The concept of therapeutic hierarchy for patients with hepatocellular carcinoma: a multicenter cohort study. Liver Int 2019;39(8):1478–89.
- [145] Bhangui P, Allard MA, Vibert E, et al. Salvage versus primary liver transplantation for early hepatocellular carcinoma: do both strategies yield similar outcomes? Ann Surg 2016;264:155–63.
- [146] Li HY, Wei YG, Yan LN, et al. Salvage liver transplantation in the treatment of hepatocellular carcinoma: a Meta-analysis. World J Gastroenterol 2012;18(19):2415–22.
- [147] Murali AR, Patil S, Phillips KT, et al. Locoregional therapy with curative intent versus primary liver transplant for hepatocellular carcinoma: systematic review and meta-analysis. Transplantation 2017;101(8):e249–57.
- [148] Xiong Q, Geng TT, He L, et al. Harm and benefits of salvage transplantation for hepatocellular carcinoma: an updated meta-analysis. Transplant Proc 2019;48(10):3336–47.
- [149] Yadav DK, Chen W, Bai X, et al. Salvage liver transplant versus primary liver transplant for patients with hepatocellular carcinoma. Ann. Transplant. 2018;23:524–45.
- [150] Zhu Y, Dong J, Wang WL, et al. Short- and long-term outcomes after salvage liver transplantation versus primary liver transplantation for hepatocellular carcinoma: a meta-analysis. Transplant Proc 2013;45(9):3329–42.
- [151] Majno P, Sarasin F, Menth G, et al. Primary liver resection and salvage transplantation or primary liver transplantation in patients with single, small hepatocellular carcinoma and preserved liver function: an outcome- oriented decision analysis. Hepatology 2000;31(4):899–906.
- [152] Van Kleek EJ, Schwartz JM, Rayhill SC, et al. Liver transplantation for hepatocellular carcinoma: a survey of practices. J Clin Gastroenterol 2006;40(7):643–7.
- [153] Adam R, Azoulay D, Castaing D, et al. Liver resection as a bridge to transplantation for hepatocellular carcinoma on cirrhosis: a reasonable strategy? Ann Surg 2003;238(4):508–19.
- [154] Belghiti J, Cortes A, Abdalla EK, et al. Resection prior to liver transplantation for hepatocellular carcinoma. Ann Surg 2003;238(6):885–93.
- [155] Del Gaudio M, Ercolani G, Ravaioli M, et al. Liver transplantation for recurrent hepatocellular carcinoma on cirrhosis after liver resection: university of Bologna experience. Am J Transplant 2008;8:1177–85.
- [156] Liu F, Wei Y, Wang W, et al. Salvage liver transplantation for recurrent hepatocellular carcinoma within UCSF criteria after liver resection. PLoS ONE 2012(11).
- [157] Margarit C, Escartín A, Castells L, et al. Resection for hepatocellular carcinoma is a good option in Child-Turcotte-Pugh class a patients with cirrhosis who are eligible for liver transplantation. Liver Transplant. 2005;11(10):1242–51.

- [158] Sapisochin G, Bilbao I, Balsells J, et al. Optimization of liver transplantation as a treatment of intrahepatic hepatocellular carcinoma recurrence after partial liver resection: experience of a single European series. World J Surg 2010;34(9):2146–54.
- [159] Scatton O, Zalinski S, Terris B, et al. Hepatocellular carcinoma developed on compensated cirrhosis: resection as a selection tool for liver transplantation. Liver Transplant. 2008;14(6):779–88.
- [160] Vennarecci G, Antonini E, Santoro R, et al. First-line liver resection and salvage transplantation are increasing therapeutics strategies for patients with hepatocellular carcinoma and child A cirrhosis. Transplant Proc 2007;39(6):1857–60.
- [161] Cucchetti A, Cescon M, Trevisani F, et al. What is the probability of being too old for salvage transplantation after hepatocellular carcinoma resection? Dig Liver Dis 2012;44(6):523–9.
- [162] Lai Q, Vitale A, Halazun K, et al. Identification of an upper limit of tumor burden for downstaging in candidates with hepatocellular cancer waiting for liver transplantation: a West–East Collaborative Effort. Cancers (Basel) 2020;12:452.
- [163] Sinha J, Mehta N, Dodge JL, et al. Are there upper limits in tumor burden for down-staging of hepatocellular carcinoma to liver transplant? Anal. All-Comers Protocol Hepatol 2019;70:1185–96.
- [164] Murali AR, Romero-Marrero C, Miller C, et al. Predictors of successful downstaging of hepatocellular carcinoma outside Milan criteria. Transplantation 2016;100:2391–7.
- [165] Mehta N, Guy J, Frenette CT, et al. Excellent outcomes of Liver transplantation following down staging of hepatocellular carcinoma to within Milan criteria-a multi-center study. Clin Gastroenterol Hepatol 2018;16:955–64.
- [166] Cillo U, Burra P, Mazzaferro V, et al. I-BELT (Italian Board of Experts in the Field of Liver Transplantation). A multistep, consensus-based approach to organ allocation in liver transplantation: toward a "Blended Principle Model". Am J Transplant 2015;15:2552–61.
- [167] Burra P, Giannini EG, Caraceni P, et al. Specific issues concerning the management of patients on the waiting list and after liver transplantation. Liver Int 2018;38(8):1338–62.
- [168] Yao FY, Hirose R, LaBerge JM, et al. A prospective study on downstaging of hepatocellular carcinoma prior to liver transplantation. Liver Transpl 2005;11(12):1505–14.
- [169] Yao FY, Kerlan RK Jr, Hirose R, et al. Excellent outcome following down-staging of hepatocellular carcinoma prior to liver transplantation: an intentionto-treat analysis. Hepatology 2008;48(3):819–27.
- [170] Yao FY, Mehta N, Flemming J, et al. Downstaging of hepatocellular cancer before liver transplant: longterm outcome compared to tumors within Milan criteria. Hepatology 2015;61:1968–77.
- [171] Ravaioli M, Grazi GL, Piscaglia F, et al. Liver transplantation for hepatocellular carcinoma: results of down- staging in patients initially outside the Milan selection criteria. Am J Transplant 2008;8(12):2547–57.
- [172] Mehta N, Dodge JL, Roberts JP, et al. Alpha-fetoprotein decrease from >1,000 to < 500ng/mL in patients with hepatocellular carcinoma leads to improved posttransplant outcomes. Hepatology 2019;69:1193–205.</p>
- [173] Mehta N, Dodge JL, Grab JD, et al. National experience on down-staging of hepatocellular carcinoma before liver transplant: influence of tumor burden, alpha-fetoprotein, and wait time. Hepatology 2020;71(3):943–54.
- [174] Lei J, Wang W, Yan LJ. Downstaging advanced hepatocellular carcinoma to the Milan criteria may provide a comparable outcome to conventional Milan criteria. Gastrointest Surg 2013;17:1440–6.
- [175] Gordon-Weeks AN, Snaith A, Petrinic T, et al. Systematic review of outcome of downstaging hepatocellular cancer before liver transplantation in patients outside the Milan criteria. Br J Surg 2011;98(9):1201–8.
- [176] Chapman WC, Garcia-Aroz S, Vachharajani N, et al. Liver transplantation for advanced hepatocellular carcinoma after downstaging without up-front stage restrictions. J Am Coll Surg 2017;224(4):610–21.
- [177] Bova V, Miraglia R, Maruzzelli L, et al. Predictive factors of downstaging of hepatocellular carcinoma beyond the Milan criteria treated with intra-arterial therapies. Cardiovasc Intervent Radiol 2013;36:433–9.
- [178] Toso C, Meeberg G, Andres A, et al. Downstaging prior to liver transplantation for hepatocellular carcinoma: advisable but at the price of an increased risk of cancer recurrence – a retrospective study. Transpl Int 2019;32:163–72.
- [179] Otto G, Herber S, Heise M, et al. Response to transarterial chemoembolization as a biological selection criterion for liver transplantation in hepatocellular carcinoma. Liver Transpl 2006;12:1260–7.
- [180] Finkenstedt A, Vikoler A, Portenkirchner M, et al. Excellent post-transplant survival in patients with intermediate stage hepatocellular carcinoma responding to neoadjuvant therapy. Liver Int 2016;36:688–95.
- [181] Ettorre GM, Levi Sandri GB, Laurenzi A, et al. Yttrium -90 radioembolization for hepatocellular carcinoma prior to liver transplantation. World J Surg 2017;41:241–9.
- [182] Mazzaferro V, Bhoori S, Sposito C, et al. Milan criteria in liver transplantation for hepatocellular carcinoma: an evidence-based analysis of 15 years of experience. Liver Transpl 2011;17(Suppl 2):S44–57.
- [183] Lee S, Kim KW, Song G-W, et al. The real impact of bridging or downstaging on survival outcomes after liver transplantation for hepatocellular carcinoma. Liver Cancer 2020;9:721–33.
- [184] Ravaioli M, Odaldi F, Cucchetti A, et al. Long term results of down-staging and liver transplantation for patients with hepatocellular carcinoma beyond the conventional criteria. Sci Rep 2019;9:3781.

- [185] Qin S, Chen M, Cheng AL, et al. Atezolizumab plus bevacizumab versus active surveillance in patients with resected or ablated high-risk hepatocellular carcinoma (IMbrave050): a randomised, open-label, multicentre, phase 3 trial. Lancet 2023;402(10415):1835–47.
- [186] Singal AG, Rich NE, Mehta N, et al. Direct-acting antiviral therapy for hepatitis C virus infection is associated with increased survival in patients with a history of hepatocellular carcinoma. Gastroenterology 2019;157(5) 1253–63.e2.
- [187] Cabibbo G, Celsa C, Calvaruso V, Rete Sicilia Selezione Terapia HCV (RE-SIST-HCV) and Italian Liver Cancer (ITALICA) Group, et al. Direct-acting antivirals after successful treatment of early hepatocellular carcinoma improve survival in HCV-cirrhotic patients. J Hepatol 2019;71(2):265–73.
- [188] Reig M, Cabibbo G. Antiviral therapy in the palliative setting of HCC (BCLC-B and -C). J Hepatol 2021;74(5):1225–33.
- [189] Iavarone M, Nault JC, Cabibbo G, Torres F, Reig M. Indolent cancer and pattern of progression: two missing parameters in trial design for hepatology. Hepatology 2023. doi:10.1097/HEP.000000000000527.
   [190] Cabibbo G, Maida M, Genco C, Antonucci M, Cammà C. Causes of and preven-
- [190] Cabibbo G, Maida M, Genco C, Antonucci M, Cammà C. Causes of and prevention strategies for hepatocellular carcinoma. Semin Oncol 2012;39(4):374–83.
- [191] Pecorelli A, Lenzi B, Gramenzi A, et al. Curative therapies are superior to standard of care (transarterial chemoembolization) for intermediate stage hepatocellular carcinoma. Liver Int 2017;37(3):423–33.