



Modern approach to hepatocellular carcinoma treatment

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Abstract

Hepatocellular carcinoma (HCC) remains a leading cause of cancer-related mortality worldwide, necessitating innovative treatment strategies. Surgical resection and liver transplantation continue to be the gold standards for early-

stage HCC; however, advances in imaging and minimally invasive techniques have improved patient selection and outcomes. Additionally, the emergence of targeted therapies and immunotherapy has transformed the treatment landscape for advanced HCC. This review highlights the efficacy of agents such as tyrosine kinase inhibitors, alongside emerging options like immune checkpoint inhibitors, which have shown promise in clinical trials. Furthermore, the role of locoregional therapies, including ablation in the setting of combined treatment, transarterial chemoembolization and transarterial radioembolization with flow catheters, cone-beam computed tomography and 4D navigation guidance, is examined in the context of bridging therapies for patients awaiting surgical intervention. The integration of multidisciplinary care approaches and personalized treatment plans is crucial for optimizing outcomes. Future directions for HCC treatment are discussed, including the potential of novel biomarkers in prognosis and treatment response. This comprehensive overview aims to equip clinicians with the latest insights and foster collaborative efforts to improve HCC patient management and survival rates.

Key Words: Hepatocellular carcinoma; Robotic resection; Transplantation; Locoregional treatments; Radioembolization; Systemic treatment; Stereotactic body radiotherapy

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Core Tip: Hepatocellular carcinoma is the sixth most common cancer and is the fourth leading cause of cancer-related deaths worldwide. Its management can be challenging and relies on a multidisciplinary approach involving hepatobiliary surgeons, oncologists, hepatologists, endoscopists, interventional radiologists, radiotherapists and radiologists. This review investigates different treatments, with a special focus on most modern approaches.

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INTRODUCTION

Hepatocellular carcinoma (HCC) is the predominant form of primary liver malignancies, comprising approximately 90% of all cases[1,2]. This is reflected in a 5-year survival rate of less than 20% and an incidence-to-mortality ratio of around 1:1[3,4]. On a global scale, HCC is ranked as the sixth most frequently diagnosed malignancy and a major cause of cancer-related mortality, with a marked predominance in male patients[2,5,6].

Several established risk factors contribute to the development of HCC, including chronic viral hepatitis B and C, metabolic disorders (such as diabetes mellitus, non-alcoholic fatty liver disease and non-alcoholic steatohepatitis), toxic exposures (including excessive alcohol consumption, nicotine abuse, and aflatoxins), hereditary conditions (such as hemochromatosis, α 1-antitrypsin deficiency, tyrosinemia, and glycogen storage disease type 1a), and immune system-related disorders[7-9]. Regardless of the underlying cause, the progression to liver cirrhosis from chronic inflammatory damage is particularly concerning, as approximately one-third of these patients will develop HCC in their lifetime[1].

HCC's inherent resistance to conventional treatment modalities, such as chemotherapy and radiotherapy, complicates its management[2]. However, advances in imaging techniques, minimally invasive procedures, and precision oncology are paving the way for innovative therapeutic strategies, aiming for a meaningful extension of overall and recurrence-free survival rates[3,10].

The Barcelona Clinic Liver Cancer (BCLC) model has emerged as the most validated therapeutic algorithm for HCC management, guiding tumor staging, prognosis, and treatment allocation based on incorporated evidence and expertise [6,11].

ROBOTIC RESECTION AND TRANSPLANTATION

The role of surgical resection for HCC management has been heavily investigated in recent years. Several international organizations have developed comprehensive guidelines to standardize HCC treatment[12-14].

In summary, the current HCC resection guidelines highlight the importance of tailored treatment based on tumor characteristics. The most relevant factors included in the decision-making algorithm concern tumor characteristics (size and number of nodules, vascular invasion, and extra-hepatic spread), liver function [Child-Pugh, model for end liver disease (MELD)][15] and Eastern Cooperative Oncology Group performance status (ECOG PS)[16]. For major hepatectomies (≥ 3 segments), the volume and function of the future liver remnant (FLR) must also be analyzed[17]. The latter can be calculated by liver scintigraphy using ^{99m}Tc -labelled galactosyl human serum albumin[18] or ^{99m}Tc -labelled

mebrofenin[19]. Computed tomography (CT) or magnetic resonance imaging (MRI) can calculate the volume, and it is necessary to have about 30% FLR in non-cirrhotic patients and at least 40% in cirrhotic patients[17].

For cirrhotic patients with HCC, resection is indicated as an alternative to liver transplantation (LT) when there is a single nodule with preserved liver function, absence of extra-hepatic spread, and ECOG PS 0. However, several studies [20-22] have shown that minimally invasive liver surgery (MILS), either laparoscopic or robotic, reduces post-operative liver failure risk, thus expanding the pool of patients with HCC who can benefit from liver resection (LR). In particular, the indications of robotic LR (RLRs) have been expanded. Liu *et al*[23] demonstrated that the robotic system can be used in almost all types of LR, including minor hepatectomy, major hepatectomy, complex LR and donor hepatectomy. This is also due to the robotic system, which provides a 3D visualization of the surgery field, uses flexible robotic arms, and a tremor filter, breaking down some of the limitations of laparoscopic LR. However, lack of experience among practitioners and associated high costs limit the application of RLR[23]. However, the experience in RLR has been reported worldwide and its feasibility has gradually been accepted by clinical studies[24-26].

Unlike LR, LT can treat both HCC and underlying cirrhosis. However, to be a candidate for LT, HCC must fulfill certain criteria. The first criteria to be developed were the Milan criteria (one lesion ≤ 5 cm, or three lesions ≤ 3 cm and no macrovascular invasion)[27]; these were subsequently not widely used because they were too restrictive. To date, the most widely used are the Up-to-seven criteria[28], the University of California-San Francisco criteria[29], the total tumor volume criteria[30], the Metroticket 2.0 model[31], the 5-5-500 model[32] and the Toronto criteria[33](Figure 1).

In addition, if the patient does not fit within transplantability criteria, downstaging therapy can be performed, *e.g.*, performing a LR to bring the patient within the criteria[34,35]. The recent increase in indications for LT does not correspond to an increase in available donors, leading to HCC progression during the waiting list period; during active list maintenance checks, the patient may fall out of the transplantability criteria. For this reason, bridging therapy can be performed to reduce the risk of disease progression while the patient is on the waiting list[12].

However, approximately, one out of five patients are removed from the LT waitlist due to death or medical unsuitability[36] and this mortality continues to be partially driven by the discrepancy between organ supply and demand. The current model of organ allocation depends on the assessment of medical urgency for transplant and mortality on the waiting list using MELD-Na. Due to the paucity of organ supply, patients must often become very sick to receive a deceased donor organ. In the setting of scarce deceased donor organs, living donor LT (LDLT) represents an important alternative. Access to LDLT shortens the median waiting time and significantly decreases waitlist morbidity and mortality for all waitlisted patients[37,38]. However, the number of LDLTs has not increased much in recent years, and in the United States, they account for only 5% of LTs[36]. This is probably while donor risk is defined[39,40], there are still no adequately powered studies on the benefits of recipient survival. However in their case-control study, Jackson *et al*[41] demonstrated how LDLT recipients gained an additional 13 to 17 Life-years compared with patients who never received an LDLT.

When comparing LR and LT, there are no unequivocal reports in the literature. Indeed, some studies have shown better survival in patients undergoing LT[42], while others have found that LR is associated with better 5-year survival [43].

In conclusion, LT and LR, especially MILS, are curative surgical procedures. The difference lies in the fact that LR does not cure underlying liver disease, exposing the patient to a higher risk of recurrence. Conversely, LT is a more complex surgical procedure with significant operative and perioperative risks, in addition to the impact of immunosuppressive therapy. Although the criteria for inclusion on the waiting list for organ transplant are being broadened and new strategies are being developed to reduce waiting times such as bridging, downstaging and LDLT, the gap between patients on the waiting list for LT and available organs remains wide.

LOCOREGIONAL TREATMENTS

As reported by BCLC, locoregional therapies play a pivotal role in BCLC 0, A and B HCC[11]. In solitary HCC up to 2 cm, ablation should be considered as the first approach (evidence moderate; recommendation strong) if LT is not feasible because it is associated with similar survival outcomes to resection with fewer complications and similar life expectancy at a lower cost. Resection is favored beyond 2 cm, as ablation offers lower rate of complete responses and higher rate of local recurrences. Among ablation therapies, microwave ablation (MWA) is potentially the best option for HCC ≤ 4 cm given that it supplies more extensive tumor necrosis than radiofrequency ablation (RFA). Moreover, ablation can be used as bridging for LT candidates with expected waiting time > 6 months, as well as transarterial chemoembolization (TACE) and transarterial radioembolization (TARE)[11,44].

If a patient with BCLC 0 or BCLC A is not a candidate for any first line treatment options, TACE can be considered as a strategy, according to the concept of treatment stage migration, recently introduced by the 2022 BCLC update. TACE is the first option in patients with BCLC B that do not meet the extended LT criteria and have preserved portal flow and defined tumor burden[11]. According to BCLC guidelines, TACE should be performed in a selective manner (evidence high; recommendation strong). Both conventional-TACE (cTACE) and drug-eluting bead TACE (DEB-TACE) have similar outcomes, but the latter has a more favorable pharmacokinetic profile and leads to less post-procedural pain[44].

Similarly to TACE, TARE is a valid option in BCLC 0 and BCLC A for single HCC up to 8 cm based on the LEGACY trial results[45,46].

Cone beam computed tomography (CBCT) is an imaging modality integrated into the angiography suite. It provides 3D images in addition to conventional 2D imaging such as fluoroscopy and conventional digital subtraction angiography (DSA)[47] (Figure 2). For transarterial procedures, CBCT can improve detection of both tumor and associated arteries to

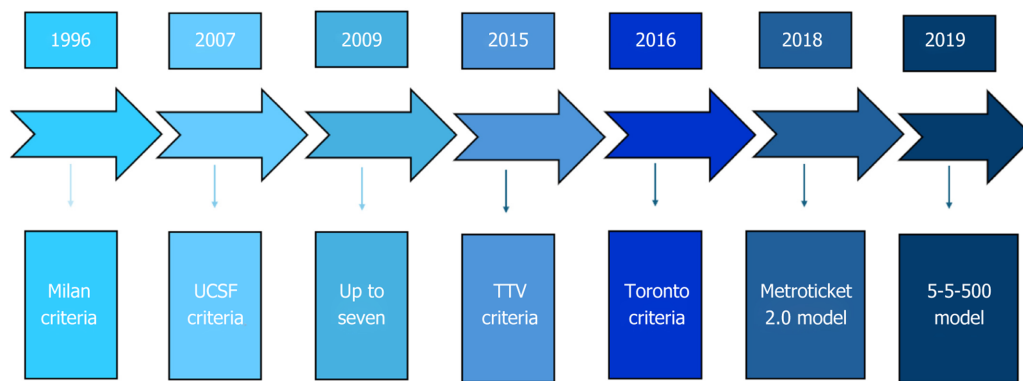


Figure 1 Timeline of eligibility criteria for liver transplantation.

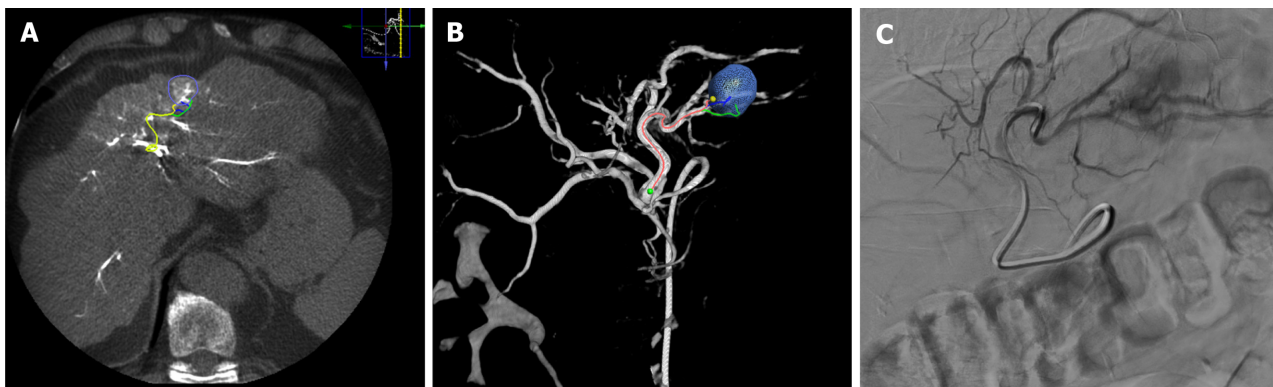


Figure 2 Transarterial chemoembolization performed with cone beam computed tomography. A: Cone beam computed tomography showing a nodule with contrast enhancement located in S4 with EmboGuide Software allowing the identification of the route (yellow line) from catheter tip to lesion; B: EmboGuide reconstruction; C: Angiography demonstrating catheterization of the lesion-feeder vessel.

obtain safer and more efficient procedures than DSA alone, especially for small and hypovascular HCCs[48]. Moreover, patients receiving TACE with CBCT guidance have higher overall and progression-free survival (PFS) than those receiving TACE under DSA guidance alone[49].

Because intra-arterial dissemination of radioisotopes may lead to potential life-threatening adverse events, a meticulous pre-procedural angiographic mapping is needed for TARE. CBCT provides relevant additional information about the perfusion of target tumor and extrahepatic nontarget tissues, not currently detectable by DSA or ^{99m}Tc MAA[50].

Thus, the use of CBCT is essential to optimize intra-arterial treatment and should be routinely used in the clinical practice of interventional radiology.

Curative intent of TARE

TARE, also known as selective internal radiotherapy (SIRT) or liver radioembolization (RE), has emerged as a significant endovascular treatment for HCC and has been endorsed by recent international guidelines[51,52] and supported by global practice patterns[53,54]. TARE is a minimally invasive procedure that involves the infusion of radioactive microspheres within the tumor *via* hepatic artery feeders and has been traditionally used as palliative treatment. However, recent advancements in TARE techniques, particularly ablative approaches (ablative TARE), have demonstrated its potential as a curative treatment modality[55,56]. Glass and resin microspheres containing yttrium-90 (Y-90) are the most extensively investigated radioactive microspheres for ablative TARE. However, clinical data on newer technologies, such as ^{166}Ho -labeled poly-L-lactic acid microspheres, are continuously expanding[55,57].

The concept of curative-intent TARE, also known as RE segmentectomy (Figure 3), was first introduced nearly 15 years ago[58] and involves catheterization of a segmental feeding artery to perform high-dose RE to selectively induce necrosis of a mass, including its relative liver segment (up to two Couinaud segments)[56,59]. The currently accepted threshold for ablative TARE using glass microspheres is ≥ 400 Gy and is based on recent pathological findings. Gabr *et al*[60] first suggested a higher threshold of ≥ 400 Gy for ablative targeted SIRT, as such doses demonstrated complete pathologic necrosis (CPN) without severe adverse events in 100% of the cases, compared to 65% complete necrosis using the previously proposed thresholds of 190 Gy. Subsequently, Montazeri *et al*[61] investigated the effect of treatment intensification on CPN in 75 HCC lesions following Y-90 glass microsphere radiation segmentectomy (RS). The authors reported that specific activity ≥ 327 Bq [area under the curve (AUC): 0.75, $P < 0.001$], dose ≥ 446 Gy (AUC: 0.69, $P = 0.005$), and treatment activity ≥ 2.55 Gbq (AUC: 0.71, $P = 0.002$) were predictive of CPN. Furthermore, a specific activity of ≥ 327 Bq was the sole independent predictor of CPN ($P = 0.013$).

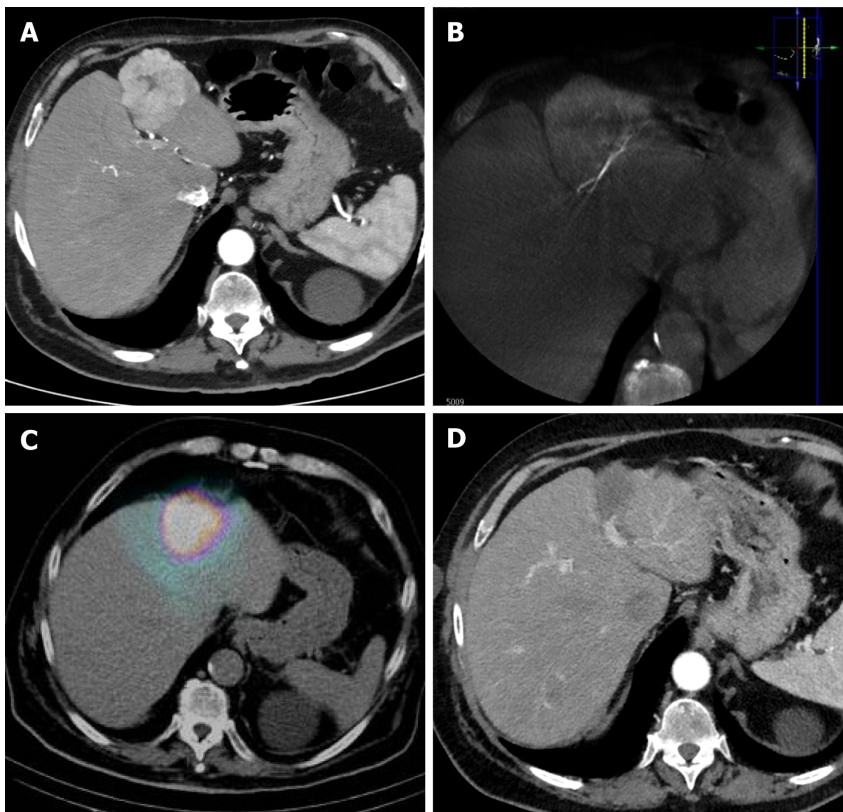


Figure 3 69-year-old man with hepatocellular carcinoma unfit for surgery due to comorbidities. A: Computed tomography (CT) showing hepatocellular carcinoma nodule (5-6 cm) located in S2; B: Cone beam CT demonstrating superselective catheterization of the segmental S2 artery with perfused nodule; C: Single-photon emission tomography combined with CT showing 99mTc-MAA uptake into S2; D: 12-months CT follow-up scan showing complete nodule necrosis with no contrast enhancement and atrophy of the treated liver lobe (radiation segmentectomy).

Curative-intent TARE can be used in patients with: (1) Unresectable tumors: Ablative RE can be used to completely ablate the tumor and the adjacent liver segment, simulating a R0 surgical resection. According to the updated BCLC guidelines, it is indicated for unresectable/unablatable solitary ≤ 8 cm HCC lesions. It can also be used for tumor downstaging, allowing for subsequent safe and efficient surgical resection. Left liver lobe RE lobectomy can be performed with curative intent; (2) Bridge to transplantation: Ablative RE can be used to control tumor growth and improve liver function in patients with > 6 months on the waiting list for LT; and (3) Limited remnant liver volume right hepatectomy: RE lobectomy can be an alternative to portal vein embolization, as it can simultaneously induce remnant liver hypertrophy and provide substantial tumor control while awaiting the hypertrophic response[11].

Based on updated evidence and expert opinions, the ideal candidates for TARE-segmentectomy are those presenting with ECOG 0-1, Child-Pugh score A cirrhosis without macrovascular invasion and an unresectable/unablatable solitary HCC tumor ≤ 8 cm in diameter. Using selective, liver-sparing approaches, ablative doses can be delivered only to the affected segment(s), while the surrounding normal liver parenchyma can be regenerated[62,63]. Moreover TARE-ablation can be performed in challenging or even inaccessible locations for percutaneous ablation, presenting high risk for incomplete ablation and recurrence. The relationship between lesion location to important structures is also less important, as TARE is not limited by the heat-sink effect and iatrogenic major vascular or biliary trauma. On the other hand, contraindications to avoid life-threatening adverse events include severe liver dysfunction with a special concern for abnormal bilirubin (> 2.0 mg/dL), albumin values (< 3 g/dL) and aspartate aminotransferase/alanine aminotransferase > 5 times the upper normal limit, gastrointestinal communications not amenable to protective coiling and significant lung shunting in which a radiation dose reduction would still be dangerous or ineffective. However a super-selective tumor ablative treatment should not greatly influence liver function[64].

Following the introduction of ablative SIRT in clinical practice, evidence indicating its safety and efficacy in selected HCC patients is constantly growing. In 2018, Lewandowski *et al*[65] published a retrospective series of 70 patients with early, solitary, ≤ 5 cm HCC who were treated with doses of > 190 Gy. The median overall survival (OS) was 6.7 years, with rates of 98%, 66% and 57% at 1-, 3- and 5-years, respectively.

Additional retrospective comparative findings suggested that ablative SIRT demonstrates equal safety and efficacy with established curative techniques such as MWA and open surgery. In a large retrospective analysis of 417 patients who underwent MWA and open surgery *vs* 235 patients who underwent TACE plus MWA for HCC lesions up to 3 cm, similar rates of complete response and OS were achieved[66]. Similarly, in a retrospective propensity score-matched (PSM) study comparing RS *vs* MWA (34 patients in each group; solitary HCC lesions < 4 cm), similar rates of liver toxicity, objective tumor response and OS were reported, while the targeted tumor mean PFS was superior in the TARE group (57.8 *vs* 38.6 months; $P = 0.005$)[67]. Impressively, in a retrospective multicenter study of 123 patients with solitary

HCC ≤ 8 cm, ECOG status of 0-1, and absence of macrovascular invasion or extrahepatic disease, RS outcomes (57 patients) were compared with those of open surgical resection (66 patients). After adjusting for confounders, OS was similar in both groups, while severe adverse events (grade ≥ 3 per the Clavien-Dindo classification), were significantly higher in the surgical group (20% *vs* 0%; $P < 0.001$) [68].

In 2021, the LEGACY multicenter (three United States centers), single-arm, retrospective study reported excellent outcomes of segmental delivery of doses > 400 Gy (median dose 410 Gy) for solitary HCC ≤ 8 cm in 162 patients with Child-Pugh A cirrhosis and ECOG status 0-1. Remarkably, no cases of radiation-induced liver disease (RILD) were reported despite the high dose protocol. OS was 86.6% at 3-years and further improved to 92.8% for patients who underwent subsequent transplantation or resection. Local recurrence and severe adverse events possibly related to the procedure were both 5.6%. The authors reported that dose > 400 Gy achieved complete pathological necrosis and should be considered as the threshold dose for ablative effect using glass Y90 microspheres [50].

Moreover, the DOSISPHERE multicenter, randomized, controlled trial published in 2021 demonstrated the superiority of “personalized dosimetry” using > 205 Gy dose in tumor delivery *vs* “standard dosimetry” with 120 Gy lobar delivery in patients with unresectable advanced HCC with at least one lesion ≥ 7 cm (OS 26.6 months *vs* 10.6 months) [69].

Finally, in 2023 the RASER trial (prospective single-center, single-arm), investigated TARE segmentectomy in 29 patients with very early and early stage HCC. Initial target lesion complete response was 83% and partial response was 17%. Initial objective response was 100% and sustained complete response at 24 months was 90% [70].

Ongoing protocols are evaluating personalized dosimetry protocols to maximize RS safety and efficacy [62]. The DOORwaY90 prospective, multicenter, open-label, single arm study was designed to evaluate the safety and effectiveness of a personalized Y90 resin microsphere dosimetry approach as first-line treatment in 100 patients with unresectable and unablatable HCC. The Partition model dosimetry will be used to determine the optimal dose, and the target mean dose to tumor will be ≥ 150 Gy [71].

TARE segmentectomy vs TACE

Recently, randomized data suggest a superior therapeutic effect using TARE (Table 1). In 2016, Salem *et al* [72] reported the results of the Premier randomized, phase 2, single-center study, of glass Y90 TARE with a median dose of 126 Gy *vs* cTACE. Although the study investigated doses below the currently suggested ablative threshold, TARE significantly prolonged time to progression. The 2022 TRACE phase II, single-center, prospective randomized, controlled trial, further supported the superiority in median OS of glass Y90 TARE using doses > 120 Gy *vs* doxorubicin DEB-TACE in selected patients with early or intermediate HCC, with similar rates of 30-day mortality and serious adverse events [73].

Further evidence on the superiority of RE segmentectomy over TACE was provided by two large retrospective PSM studies. The first, published in 2017 by Padia *et al* [74], investigated 104 patients with 132 tumors treated with 144 segmental RE procedures (target perfused tissue dose > 200 Gy) compared to 77 patients with 103 tumors treated with 138 chemoembolization procedures (76% DEB-TACE and 24% cTACE). Index and overall complete response rates were 92% and 84% for RE *vs* 74% and 58% for TACE ($P = 0.001$ and $P < 0.001$). Index tumor progression at 1 and 2 years was 8% and 15% for RE and 30% and 42% for TACE ($P < 0.001$). Median PFS and OS were 564 days and 1.198 days for RE *vs* 271 days and 1.043 days for TACE (PS-adjusted $P = 0.002$ and $P = 0.35$; censored by transplant PS-adjusted $P < 0.001$ and $P = 0.064$).

Subsequently, Biederman *et al* [75] reported the results of another retrospective PSM analysis of 112 patients who underwent RS *vs* 55 patients who underwent TACE for solitary HCC ≤ 3 cm without vascular invasion or metastasis. After PSM, the RS group showed better complete response and median time to secondary therapy, while OS was not significantly different.

Finally, although most studies investigating RS have used Y90 glass microspheres, recent retrospective data suggest that Y90 resins microspheres are equally effective. Villalobos *et al* [76] reported similar safety and efficacy outcomes in a retrospective study of 81 patients with HCC (20 in the resin cohort and 61 in the glass cohort). Notably, resin-based segmentectomy used lower mean tumor dose (TD) than glass-based segmentectomy ($308 \text{ Gy} \pm 210$ *vs* $794 \text{ Gy} \pm 523$, $P = 0.0002$), while mean TD thresholds able to predict the objective and complete responses were 176 Gy and 247 Gy for resin-based segmentectomy and 290 Gy and 481 Gy for glass-based segmentectomy.

Moreover, in 2024, Sarwar *et al* [77] published a retrospective single-center series of 67 patients with HCC who underwent TARE segmentectomy with 90-Y resin microspheres. Doses were calculated using the single-compartment medical internal radiation dosimetry model and the median tumor absorbed dose was 232 Gy (IQR: 163-405 Gy). Median posttreatment OS was 26 months, while per-lesion and overall duration of response ≥ 1 year was 88% and 72%, respectively. Notably, 2-year disease progression was lower, with no local progression through the end of follow-up, when tumors received ≥ 300 Gy (17% *vs* 61%; $P = 0.047$), in the former group. The 1-month grade 3 clinical adverse event rate was 1.5% (one patient with abdominal pain).

Novel technologies in intra-arterial therapies: 4D navigation and microcatheter balloon occluded TACE

Recent advances have significantly transformed the field of interventional radiology, enhancing precision, safety, and, consequently, patient outcomes. Among these innovations, the integration of 4D navigation systems with fusion imaging represents a cutting-edge platform designed to optimize precision and efficiency in interventional procedures. By merging 3D imaging with time-sequenced data, these systems provide dynamic, real-time visualizations of anatomical structures and procedural tools. This capability is invaluable for accurately guiding needles or catheters during complex interventions, such as tumor ablation or vascular procedures. For instance, they enable interventional radiologists to meticulously plan and monitor instrument trajectories, ensuring precise targeting and minimizing complication risks.

Table 1 Summary of the papers regarding the comparison between transarterial radioembolization and transarterial chemoembolization

Ref.	Main aim	Main findings
Salem <i>et al</i> [72]	Glass Y90 TARE (24 pt) <i>vs</i> cTACE (21 pt)	Time to progression: 26 <i>vs</i> 6.8 months ($P = 0.0012$)
TRACE[73]	Glass Y90 TARE (38 pt) <i>vs</i> DEB-TACE (34 pt)	Median OS: 30.2 months <i>vs</i> 15.6 months
Padia <i>et al</i> [74]	Segmental TARE (104 pt) <i>vs</i> TACE (138)	Overall complete response rate: 84% <i>vs</i> 58% ($P < 0.001$) Median PFS: 564 days <i>vs</i> 271 days ($P = 0.002$, adjusted $P > 0.001$) OS: 1198 days <i>vs</i> 1043 days ($P = 0.35$, adjusted $P = 0.064$)
Biederman <i>et al</i> [75]	Radiation segmentectomy (112 pt) <i>vs</i> TACE (55 pt)	Overall complete response rate: 92.1% <i>vs</i> 52.6% ($P = 0.005$) Median time to secondary therapy: 812 days <i>vs</i> 161 days ($P = 0.001$)

OS: Overall survival; PFS: Progression free survival; TACE: Transarterial chemoembolization; TARE: Transarterial radioembolization.

These novel techniques have a wide range of applications, including the treatment of HCC *via* both intra-arterial and percutaneous approaches. Specifically, the 4D navigation system offers a comprehensive one-room solution by integrating a CT scanner with a flat-panel angiography suite within the same interventional radiology room. The combination of real-time 2D imaging and CT-like images within the IR suite represents a significant advance in enhancing the reliability and precision of numerous procedures.

Several studies have demonstrated the utility of Angio-CT systems in image-guided procedures such as TACE. For instance, Piron *et al*[78] reported that radiation exposure during TACE varies significantly between high-end angiography suites, with lower total effective doses observed in Angio-CT suites compared to cone-beam CT suites. Creating a 3D roadmap from the celiac trunk at the beginning of the procedure provides a comprehensive visualization of liver vascularization, enabling faster tumor detection and facilitating super-selective catheterization. This approach reduces the number of required DSA runs and minimizes radiation exposure during fluoroscopy[78,79].

Emerging technologies include the integration of combined imaging modalities for the thermal ablation of hepatic tumors, offering effective solutions for treating lesions that are inaccessible with ultrasound (US)- or CT-guided approaches alone due to their localization or intrinsic characteristics. Ablation is a minimally invasive technique routinely performed as a standalone procedure or in combination with surgery[80]. For example, Hermida *et al*[81] proposed a method for the percutaneous thermal ablation of HCC located in the hepatic dome, a challenging region to access with US guidance, where lung interposition often compromises tumor visibility and accessibility. Their study demonstrated the efficacy and safety of CT-guided lesion ablation combined with CO₂-induced artificial pneumothorax. Additional applications of fusion imaging in the thermal ablation of liver lesions include "lipiodol tagging," which involves selective intra-arterial lipiodol injection into tumors, thereby enhancing the visibility of lesions that are difficult to detect using US guidance alone[82]. For hypovascular lesions, which appear as hypoattenuating nodules on CT during the portal phase, CT-portography offers a viable solution. CT arterial portography or CT hepatic arteriography enables multiple contrast medium injections through the superior mesenteric artery, proving highly effective for identifying hypovascular lesions in challenging locations. This technique facilitates precise, real-time intraprocedural needle placement, significantly improving the accuracy and safety of ablation procedures[81,83].

In summary, Angio-CT systems significantly enhance procedural precision, minimize radiation exposure, and optimize the management of complex cases, making them particularly beneficial for training and supporting young interventional radiologists.

Among the emerging technologies in intra-arterial therapies, Balloon-Occluded TACE (B-TACE) has emerged as a pivotal advancement.

TACE is well-established as one of the primary treatments for intermediate-stage HCC and is currently recommended for patients with HCC in BCLC stages 0, A, and B[11,84]. It is also highlighted in the recent European Association for the Study of the Liver guidelines, where it is described as a "disease control" strategy rather than a definitive treatment[51].

This limitation arises from the lack of standardization in the technique: Various chemotherapeutic agents (such as Doxorubicin, Idarubicin, Farmorubicin, *etc.*) can be used, and different procedural approaches are available, including cTACE or DEB-TACE[85]. To date, no clear evidence has demonstrated the superiority of one chemotherapeutic agent over another or one embolization technique over another. In terms of response rate, Veloso Gomes *et al*[86] analyzed 580 patients treated with DEB-TACE, reporting a response rate of 60.14% after one or more procedures. Similarly, in a large study by Peng *et al*[87] among 699 patients who underwent one or multiple sessions of cTACE or DEB-TACE, a complete response rate of 22.3% was observed.

To overcome these limitations, it would be advisable to standardize this technique across all its phases, including the execution method, product type, drug selection, and embolization approach.

In this context, B-TACE has emerged as a promising technique. Its encouraging results suggest the potential to achieve at least partial standardization of the TACE procedure. B-TACE is a variant of the TACE procedure that employs a balloon microcatheter for arterial occlusion. First described in 2013[88], it is not yet included in clinical guidelines[51] and its specific indications remain undefined. However, B-TACE is generally characterized by the infusion of a chemotherapeutic agent and lipiodol emulsion, followed by gelfoam administered under the occlusion of feeding arteries by a balloon.

The placement of the microballoon (*e.g.*, Occlusafe, Terumo®) proximal to all feeders maximizes therapeutic efficacy by temporarily occluding the feeding artery. This induces flow redistribution towards lower-resistance vascular territories and facilitates a pressure-gradient-driven chemoembolization. The drug-embolic mixture can be forcefully delivered into tumor vessels, ensuring higher drug accumulation, filling arteriportal microanastomoses, and achieving greater rates of complete tumor response. This approach has been particularly documented in cases where the balloon-occluded arterial stump pressure is 64 mmHg or lower and where significant collateral arteries are absent[89]. Additionally, the balloon minimizes the risk of leakage outside the target nodule, preventing backflow and non-target embolization[90].

B-TACE has demonstrated superior oncological outcomes compared to DEB-TACE in terms of response rate and longer time to progression, with no significant differences in adverse events, even in patients with larger tumors[91]. While there is no clear consensus on the optimal reference diameter for achieving the best outcomes with B-TACE, a 2021 study highlighted its superiority over cTACE for treating HCCs with diameters between 30-50 mm in a single session[92]. Similarly, in 2023, Chu *et al*[93] reported that B-TACE achieved superior local tumor progression control compared to cTACE in HCCs with diameters greater than 3 cm.

Recently, Lucatelli *et al*[94] performed a multicenter European study and confirmed the safety and efficacy of B-TACE, reporting a target complete response of 58.9% at 6 months with a mean lesion diameter of 37 mm. These findings are comparable to the response rate (60.14%) reported by Veloso Gomes *et al*[86].

Future directions include the potential standardization of this aspect of the procedure, which could pave the way for further studies aimed at standardizing additional procedural steps, ultimately ensuring greater consistency and reproducibility in interventional radiology practices for HCC management. Alongside B-TACE, advancements in interventional radiology, such as 4D navigation, fusion imaging, and Angio-CT, have significantly improved precision, safety, and oncological outcomes, heralding a promising future for minimally invasive therapies in HCC management.

Combined locoregional treatments

Although the combined locoregional treatments are not included in the BCLC guidelines[11], they are widely applied by interventional radiologists in clinical practice in HCC treatment.

The most widespread strategy is based on the combination of the percutaneous approach, such as RFA or MWA, and the intra-arterial approach, such as TACE, especially in the treatment of HCC larger than 3 cm[95-97].

Given that ablation therapies are curative for very early and early HCC and are negatively affected by the heat sink effect and because the tumor necrosis rate of TACE for intermediate HCC is low and the recurrence rate is high, combined therapies could be effective than single approaches[98,99].

A common combination is TACE + RFA, with two variants: TACE followed by RFA or RFA followed by TACE, each with different rationales. However, it is not clear which is superior[98]. Moreover, there is no clear consensus about time interval between TACE and RFA. Iezzi *et al*[98] recommended a single-step “combined” approach to amplify the synergistic effects of RFA and TACE, leading to reduction of hospitalization days, decrease in patient discomfort and cost. Jiang *et al*[95] recently confirmed that TACE combined with RFA was more effective for HCC than TACE alone. For patients with a tumor larger than 3 cm, the combined treatment also achieved better results than RFA alone.

According to two meta-analyses, including 1736[100] and 1799[101] patients, combination therapy with TACE and MWA is better than TACE alone in HCCs > 3 cm regarding OS, PFS and treatment response.

In 2020, Iezzi *et al*[102] first described a new combined single-step therapy with balloon-occluded MWA followed by B-TACE in patients with unresectable single HCC. Selective catheterization of the target vascular segment was performed by advancing a 0.014-in hydrophilic guidewire and a 2.8Fr microcatheter with an occlusion-balloon on the tip; the 10 mm long micro-balloon is made of compliant polyurethane and has a diameter of 1-4 mm according to inflation volume. Thereafter, the balloon was inflated to occlude the flow and obtain an arterial stump pressure drop. Ablation was then performed using a 14-G uncooled 2.450 MHz MWA system up to 40 W of applied power for 10 minutes with the balloon still inflated. After ablation, B-TACE was performed using 2 mL of 100-micron calibrated drug eluting-microspheres loaded with 50 mg doxorubicin mixed with 5 mL contrast medium and 3 mL distilled water. They treated five patients with good results. Technical success was obtained in all procedures and no residual tumor or local recurrence was registered at the 6-month CT follow-up. This combined approach has been strengthened by two multicenter studies, demonstrating a complete response, respectively, in 91, 3% at 1 month and in 85.7% at 3-6 months[103] and 79.4% at 1 month and 81.4% at 3-6 months[104]. The initial diameter of the lesion ≥ 5.0 cm has been proven to be as the sole independent prognostic factor for loss of complete response[104].

A recent and interesting meta-analysis demonstrated that for lesions > 5 cm, TACE + cryoablation (CRA) combination therapy has a significant decrease in alpha-fetoprotein (AFP) levels, a significant increase in CD4+ T cells and a significant decrease in CD8+ T cells compared to TACE alone, indicating that it is more beneficial for protecting or enhancing immune function. Moreover, TACE + CRA enhances the therapeutic efficacy and long-term survival rate, without increasing the risk of complications[105].

SYSTEMIC TREATMENT

Systemic treatment of HCC has seen significant advances over the past decade. The introduction of tyrosine kinase inhibitors, immune checkpoint inhibitors (ICIs), and combination therapies has improved OS and quality of life for patients with advanced HCC. Systemic therapy is advised for patients with intermediate-stage disease who are either not candidates for or have not responded to local treatments (BCLC B), as well as for patients with advanced HCC and preserved liver function (BCLC C).

The treatment for HCC is performed using immune therapies (designed to trigger an immune response against the tumor or enhance an ongoing immune reaction) or targeted therapies (which involve drugs that focus on specific molecular pathways crucial for HCC cell proliferation). The use of systemic cytotoxic drugs (which indiscriminately kill dividing cells) has not been beneficial in clinical trials. Immune checkpoints include co-inhibitory molecules expressed by effector lymphocytes that prevent excessive activation. ICIs block this effect and rejuvenate effector cells. ICIs used in randomized clinical trials (RCTs) target PD-1 (nivolumab, pembrolizumab, sintilimab) or its ligand PD-L1 (atezolizumab, durvalumab, tislelizumab) and cytotoxic T lymphocyte-associated antigen 4 (CTLA-4, ipilimumab, tremelimumab). Vascular endothelial growth factor (VEGF) is produced within the tumor microenvironment by tumor or stromal cells. It promotes tumor growth by stimulating angiogenesis and also has several immune-modulating effects, such as inhibiting dendritic cell activity or the creation of immunosuppressive myeloid-derived stromal cells and regulatory T cells, among other mechanisms. Anti-VEGF drugs may counteract these effects by binding to circulating VEGF (bevacizumab), or by specifically blocking its cellular receptors like VEGFR-2 (rivoceranib and ramucirumab). Targeted treatments used for HCC include multi-tyrosine kinase inhibitors that share an anti-angiogenic effect through inhibition of VEGF and platelet-derived growth factor receptors and vary in their inhibition of molecules involved in different molecular pathways (such as RAF, FGFR, RET, KIT, TIE2, or MET). Due to their multiple mechanisms of action, the effects of multi-TKIs on the immune system are not completely understood.

Combining immunotherapies and TKIs forms the cornerstone of systemic therapy in HCC. Their benefit has been confirmed in large-scale RCTs with control arms that have evolved over time. When making treatment choices, direct cross-trial comparisons might lead to inaccurate conclusions, so physicians should carefully consider the patient's profile, any relative contraindications, and the availability of treatments, always discussing these factors with the patient and, once again, in multidisciplinary team setting[51,106].

First line treatment

Based on the findings of the global IMbrave 150[107] and HIMALAYA trials[108], combination therapies involving PD-1 or PD-L1 inhibitors should be considered the first-line standard of care for patients without contraindications to ICIs (and bevacizumab). There is no evidence suggesting that one option is preferable over the other. Sorafenib and lenvatinib remain first-line choices for these patients, and single-agent durvalumab and tislelizumab can also be considered.

The IMbrave150 trial, which assessed the combination of atezolizumab and bevacizumab (A-B) in patients with unresectable HCC, was the first Phase III study to show survival benefit for any treatment over sorafenib. The median OS was 19.2 months for the (A-B) group, compared to 13.4 months with sorafenib (HR: 0.66). Combination therapy achieved an objective response rate (ORR) of 30%, while sorafenib had an ORR of 11%, with (A-B) showing an 8% complete response rate *vs* < 1% for sorafenib. Additionally, there was a significant delay in the decline of quality-of-life measures in the (A-B) group compared to the sorafenib group[107].

HIMALAYA was the first study to demonstrate the effectiveness of dual ICI therapy. It compared durvalumab-tremelimumab, durvalumab alone, and sorafenib alone. The study showed a better median OS with the single tremelimumab plus regular interval durvalumab (STRIDE) regimen compared to sorafenib alone (16.4 months *vs* 13.8 months; HR: 0.78)[108]. The 5-year OS rates were 19.6% for STRIDE and 14.4% for durvalumab monotherapy, compared to 9.4% for sorafenib[109]. TRAEs of any grade were less frequent with durvalumab compared to sorafenib.

The global Phase III CheckMate-9DW study assessed nivolumab-ipilimumab *vs* lenvatinib (administered to 90% of patients) or sorafenib. The results showed that OS was improved with nivolumab-ipilimumab compared to TKIs (median OS of 23.7 months *vs* 20.6 months; HR: 0.79). TRAEs of any grade occurred in 84% of patients receiving nivolumab-ipilimumab and 91% of patients receiving TKI[110].

The Phase III RATIONALE-301 trial compared tislelizumab, a monoclonal antibody with strong affinity and specificity for PD-1, to sorafenib. The primary non-inferiority endpoint was achieved, with a median OS of 15.9 months for tislelizumab *vs* 14.1 months for sorafenib (HR: 0.85). The ORR was 14.3% for tislelizumab. Tislelizumab was associated with a lower incidence of TRAEs, leading to discontinuation[111].

Lenvatinib was compared to sorafenib as first-line treatment in the open-label, global, Phase III REFLECT trial. The primary endpoint was achieved, showing non-inferiority for lenvatinib with a median OS of 13.6 months, compared to 12.3 months for sorafenib (HR: 0.92). However, the secondary endpoints favored lenvatinib, with a higher ORR of 24% *vs* 9% (by mRECIST) and improved PFS of 7.4 months *vs* 3.7 months[112]. Lenvatinib was also studied in combination with TACE in the open-label LAUNCH trial in China, (median OS 17.8 months for TACE-lenvatinib, *vs* 11.5 months for lenvatinib)[113].

Sorafenib was the first systemic therapy to show a survival benefit in advanced HCC in a placebo-controlled Phase III trial[114]. A subsequent confirmatory trial in Asia, which primarily included patients with a history of HBV infection, yielded a similar HR favoring sorafenib[115].

Second-line treatment

The Phase III RESORCE trial compared regorafenib to placebo in patients who had progressed on sorafenib but had tolerated a dose of ≥ 400 mg daily for at least 20 out of the 28 days prior to discontinuation. Regorafenib showed improved OS with a median of 10.6 months compared to 7.8 months for placebo (HR: 0.63)[116]. Cabozantinib was compared to placebo in the global Phase III CELESTIAL trial involving patients who had received one or two prior treatments, including sorafenib. The median OS was better in the cabozantinib group (10.2 months *vs* 8.0 months with placebo; HR: 0.63). The ORR for cabozantinib was 4% by RECIST v1.1[117]. Cabozantinib was later included in the COSMIC-312 trial, which compared first-line cabozantinib to sorafenib, with cabozantinib being assessed as a secondary endpoint[118]. The final analysis showed a median PFS of 5.8 months for cabozantinib *vs* 4.3 months for sorafenib. However, due to the lack of data showing non-inferior survival, first-line cabozantinib is not currently recommended

[119].

The Phase III REACH trial did not find that second-line ramucirumab was superior to placebo in advanced HCC, but a subgroup analysis indicated a potential benefit for patients with serum AFP ≥ 400 ng/mL [120]. The subsequent REACH-2 trial focused on patients with AFP ≥ 400 ng/mL. The results showed a significant improvement in OS with ramucirumab (median 8.5 months *vs* 7.3 months with placebo; HR: 0.71), although OS in both groups was relatively low, reflecting the poor prognosis associated with elevated AFP levels [121].

STEREOTACTIC BODY RADIOTHERAPY

Among the local treatment strategies, external beam conformal radiotherapy was historically considered a palliative treatment, mainly linked to the risk of RILD [122,123].

Stereotactic body radiotherapy (SBRT) is a targeted radiation therapy that delivers precise high radiation doses to specific targets in a limited number of fractions in a non-invasive manner, minimizing exposure to surrounding healthy tissue [124-126].

In the last decade, several studies have demonstrated that SBRT is endorsed as a safe and effective locoregional treatment, especially for patients with inoperable localized or recurrent HCC. It can also serve as bridging or down-staging therapy before LT. SBRT provides excellent local control (LC), survival benefits, reduced morbidity, and favorable cost-effectiveness [12,127,128].

The reported rates of LC at 1 year and 2 years after SBRT range from 87% to 97% in this patient population [129-131]. Biologically effective dose assuming an $\alpha/\beta = 10$ (BED10) ≥ 100 Gy is associated with significant LC [132]. A propensity score analysis of a retrospective single-institution cohort demonstrated durable LC rates of 97% at 1 year and 91% at 2 years for patients with one or two unresectable HCC tumors [133].

In the randomized multi-center phase 2 TRENDY trial, patients ineligible for RFA or additional surgery showed time to local recurrence of over 40 months following SBRT [134].

The RTOG 1112 study, a multi-center phase 3 randomized trial involving 177 patients with HCC who were unfit for transplantation, resection, RFA, or TACE, found that SBRT before Sorafenib resulted in median OS of 15.8 months, compared to 12.3 months for those who did not receive SBRT. PFS was significantly better with SBRT, showing 9.2 months *vs* 5.5 months. These patients had very advanced disease: 82% were classified as BCLC-C, 74% had MVI, 4% had metastases, and patients had at least five tumors [135].

Retrospective and prospective studies show that SBRT has low $> G3$ toxicity, ranging from 0% to 8% [133,134]. High quality-of-life scores were also reported, particularly regarding post-treatment improvements in pain [136].

An interesting topic could be the exploration of the immunomodulatory properties of SBRT combined with immunotherapy to improve local and systemic control of HCC. Preclinical studies are needed to understand the synergies between SBRT and immunotherapy, including optimal treatment volumes, dosages, predictive biomarkers, and the tumor microenvironment's role [137]. One example is Personalized Ultrahigh Fraction Adaptive Stereotactic Radiotherapy (PULSAR), which delivers high-dose radiation fractions weeks apart to allow for biological changes. When paired with immunotherapy, PULSAR may enhance anatomical adaptation and systemic response [138,139].

Deriving data about the use of SBRT as a bridge to transplant, recent studies have shown that SBRT could be considered in neoadjuvant treatment strategies for patients with locally advanced HCC [136,140]. A randomized controlled trial involving 164 patients with resectable HCC and tumor thrombus in the main trunk of the portal vein or its side branches compared two patient groups and found significant improvements in 12-month postoperative OS, HCC-related mortality, and rates of HCC recurrence [141].

The prospective phase 2 START-FIT study showed that a neoadjuvant regimen including TACE (day 1), followed by SBRT (day 28) and then Avelumab (14 days after SBRT), converted 55% of patients with initially unresectable, locally advanced HCC to curative treatment [142].

New therapeutic approaches are emerging in SBRT for HCC, such as stereotactic magnetic resonance-guided adaptive radiotherapy (SMART) and intensity-modulated stereotactic body proton therapy (SBPT).

SBRT targeting abdominal sites has limitations concerning the control of target motion and the daily reproducibility of organs at risk (OARs). Consequently, uncertainties during the planning and administration of SBRT restrict the safe delivery of ablative doses to tumors located near hepatobiliary and luminal OARs.

The SMART technique is useful in mitigating these uncertainties by employing on-table MRI from an MR linear accelerator for daily anatomical and positional adaptation, and movement management. This method enables a reduction in planning target volumes and allows for the assessment of dose escalation to improve LC [143]. Single-arm prospective and retrospective studies of SMART HCC, cholangiocarcinoma, and liver metastases have indicated a 2-year LC rate ranging from 73% to 100% and $G3$ toxicity ranging from 0% to 8% (no $G4$ toxicity was observed) [144].

Proton particles can help protect the uninvolved liver, reducing the risk of hepatic toxicity from radiotherapy and encouraging dose escalation [145]. Ongoing trials of SBPT for HCC (NCT04805788) are assessing the reduction of toxicity and the gain in oncologic outcomes.

Growing data on single-fraction SBRT of liver metastases reveal low toxicity and excellent control with doses of 18-40 Gy [146,147]. We await long-term results from larger series to ensure efficacy and safety.

FUTURE DIRECTIONS

Well-designed, multi-center RCTs are required to compare the safety and efficacy of ablative TARE with other curative-intent locoregional and surgical treatment options, as to provide solid data on its use in earlier-stage therapeutic lines. Additionally, defining the optimal dose for segmentectomy in various available TARE systems is imperative for further enhancing therapeutic efficacy[148]. Pressure-assisted microcatheters could also be used for more aggressive TACE/TARE but also as an occlusive tool to avoid non-selective delivery[94,149].

The combination of systemic and locoregional treatment modalities such as intraarterial treatments (TACE or TARE) plus immunotherapy strategies like checkpoint inhibitors or adoptive T-cell therapy[150], or between different locoregional treatments such as percutaneous ablation plus TACE or BTACE (B-TACE segmentectomy), have already provided optimistic initial results and are currently under further investigation[94,151].

The possibility of the abscopal effect following locoregional treatments is also a very promising concept under investigation. According to limited data, even subtherapeutic locoregional treatments for HCC could prime immune cells against tumor antigens to induce disease regression, but also enhance the effect of systemic immunotherapy against HCC. Such an effect could revolutionize treatment capabilities in poor prognosis patients with advanced and/or metastatic HCC[152-154]. More data are awaited soon regarding the synergistic effect of locoregional and systemic immunotherapy [155,156] or intra-lesional delivery of immunotherapy as sole treatment or combined with intraarterial therapy[157-159].

Personalized therapies are also proposed to improve outcomes. Recently, the molecular and immune classification of HCC distinguished two major molecular groups based on transcriptomic-based phenotypic classes (proliferation and non-proliferation class). The proliferation class is characterized by more aggressive tumors with poor histological differentiation, high vascular invasion and increased AFP[160]. Specific biomarkers such as circulating miRNAs and genomic profiling from ongoing translational studies could be used to guide personalized therapeutic protocols and select patients most likely to benefit from a given regimen. Tissue samples or biological fluids (blood, urine *etc.*) can undergo molecular analysis *via* omics-based methodologies to obtain molecular-based signatures for HCC recurrence. Omics signatures and clinical/pathological features could generate predictive systems and enable personalized treatment based on the molecular basis of recurrence[161].

More recently, AI deep learning models have been proposed to assist in patient selection, lesion characterization and to optimize personalized treatment protocols[162]. AI could be used in the near future to improve patient selection decision-trees and issues on optimal dosimetry and accurate pre- and post-procedural imaging.

Future research on synergistic therapies and personalized treatment protocols should address numerous issues including the nature, timing and sequence of immunotherapy, and locoregional treatments to warrant the best clinical outcomes, as well as the biological and genetic biomarkers to recognize patients likely to benefit from such combined approaches.

Technological advancements in fusion imaging and 4D-CT systems should also be used to improve numerous parameters that affect the outcomes of locoregional treatments such as target lesion(s) identification, intraprocedural guidance, treatment delivery, and immediate outcome assessment, and at the same time reduce radiation exposure and contrast use[163].

The reduction of radiation exposure and technical accuracy could also be enhanced with the use of novel robotic systems designed for peripheral endovascular procedures. Robotic endovascular treatments are a promising and appealing technology and is expected to transform clinical practice in the next decade[164].

Improved, personalized dosimetry is also a major investigational concern for TARE procedures. Dosimetry using intraprocedural cone-beam CT, SPET-CT dosimetry calculating the perfused -and not necrotic- mass volume, as well as 99mTc-MAA-based dosimetry using dedicated software and AI options are anticipated to further increase the safety and effectiveness of TARE[165-167].

Future improvements in dosimetry using MRI-based dosimetry are also awaited for Ho-166 TARE[168].

Additional data highlighting specific characteristic and advantages of Holmium (MRI visibility, 166Ho scout dose possibility) and more recently developed TARE platforms such as imageable RE microspheres (radiopaque Eye90 microspheres™), and Rhenium-188 [tungsten-188/rhenium-188 (W-188/Re-188) generator for onsite dose generation], will most likely increase their clinical penetration in the near future[169,170].

Finally, nanotechnology may offer new approaches to target HCC cells and can be employed for the optimization of intraarterial drug-delivery. Specifically, nanoparticles are currently under investigation as drug-carriers for local chemotherapy, or gene therapy agents. Nanoparticles functionalized with specific HCC-target ligands, could increase intratumoral drug accumulation. Moreover, nanoparticles allow the co-encapsulation of multiple drugs within a single nanoparticle and therefore enable a more homogeneous multi-drug DEB-TACE[171].

CONCLUSION

This review provides a comprehensive examination of current and emerging HCC treatment strategies and future directions in this intricate and multidisciplinary field. Emphasis is placed on the role of intraprocedural cross-sectional imaging-guided locoregional therapies, the effectiveness of targeted therapies, and the identification of novel biomarkers for both prognostic assessment and prediction of treatment response.

FOOTNOTES

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