



Yttrium-90 Selective Internal Radiation Therapy with Glass Microspheres for Hepatocellular Carcinoma: Current and Updated Literature Review

Edward Wolfgang Lee, MD, PhD^{1*}, Lourdes Alanis, MD, MPH^{1*}, Sung-Ki Cho, MD^{2*}, Sammy Saab, MD^{3*}

¹Division of Interventional Radiology, Department of Radiology, UCLA Medical Center, David Geffen School of Medicine at UCLA, Los Angeles, CA 90095, USA; ²Division of Interventional Radiology, Department of Radiology, Samsung Medical Center, Sungkyunkwan University School of Medicine, Seoul 06351, Korea; ³Division of Hepatology, Department of Medicine, Pflieger Liver Institute, University of California at Los Angeles, Los Angeles, CA 90024, USA

Hepatocellular carcinoma is the most common primary liver cancer and it represents the majority of cancer-related deaths in the world. More than 70% of patients present at an advanced stage, beyond potentially curative options. Yttrium-90 selective internal radiation therapy (Y90-SIRT) with glass microspheres is rapidly gaining acceptance as a potential therapy for intermediate and advanced stage primary hepatocellular carcinoma and liver metastases. The technique involves delivery of Y90 infused glass microspheres via the hepatic arterial blood flow to the appropriate tumor. The liver tumor receives a highly concentrated radiation dose while sparing the healthy liver parenchyma due to its preferential blood supply from portal venous blood. There are two commercially available devices: TheraSphere® and SIR-Spheres®. Although, Y90-SIRT with glass microspheres improves median survival in patients with intermediate and advanced hepatocellular carcinoma and has the potential to downstage hepatocellular carcinoma so that the selected candidates meet the transplantable criteria, it has not gained widespread acceptance due to the lack of large randomized controlled trials. Currently, there are various clinical trials investigating the use of Y90-SIRT with glass microspheres for treatment of hepatocellular carcinoma and the outcomes of these trials may result in the incorporation of Y90-SIRT with glass microspheres into the treatment guidelines as a standard therapy option for patients with intermediate and advanced stage hepatocellular carcinoma.

Index terms: *Yttrium 90; Selective internal radiation therapy; Hepatocellular carcinoma; Radioembolization*

INTRODUCTION

Hepatocellular carcinoma (HCC) is the most common

Received September 7, 2015; accepted after revision March 20, 2016.

*These authors contributed equally to this work.

Corresponding author: Edward Wolfgang Lee, MD, PhD, Division of Interventional Radiology, Department of Radiology, Ronald Reagan Medical Center at UCLA, David Geffen School of Medicine at UCLA, 757 Westwood Plaza, Suite 2125, Los Angeles, CA 90095-743730, USA.

• Tel: (1310) 267-8771 • Fax: (1310) 267-3631

• E-mail: EdwardLee@mednet.ucla.edu

This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (<http://creativecommons.org/licenses/by-nc/3.0>) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

primary liver cancer and is the second most common cause of death from cancer worldwide, with an estimated 745000 deaths in 2012 (1). It is the fifth most common cancer among men (554000 cases) and the ninth most common cancer in women (228000) (1). The age-adjusted worldwide incidence rates per 100000 for men and women are 15.3 and 5.3, respectively (1). In men, the incidence rates are highest in Eastern and South-Eastern Asia (31.9 and 22.22, respectively), intermediate in Southern Europe (9.5) and Northern America (9.3), and the lowest in Northern Europe (4.6) and South-Central Asia (3.7) (1). In women, the rates are generally lower than those in men with the highest rates in Eastern Asia and Western Africa (10.2 and 8.1, respectively) and the lowest in Northern Europe and Micronesia/Polynesia (1.9 and 1.4, respectively) (1). In the United States, an estimated 23000 deaths from liver cancer

occurred in 2014 (2). In this same year, an estimated 33190 new cases of liver cancer were expected (2). From 2006 to 2010, the rates of liver cancer increased by 3.7% per year in men and by 2.9% per year in women (2).

Hepatocellular carcinoma has a poor prognosis with an overall 5-year relative survival rate of 16% and survival decreases as patients are diagnosed with regional and distant stages of the disease (2). Only 41% of liver cancer patients are diagnosed at an early stage, while the majority of patients present with advanced disease. Curative treatments indicated for early HCC include resection, liver transplantation, and percutaneous ablation (3, 4). Patients with early HCC can achieve a survival rate of 50–70% at 5 years after undergoing resection, liver transplantation, or percutaneous ablation (5). Non-curative treatments to improve survival in patients with unresectable HCC include locoregional therapies such as transarterial chemoembolization (TACE) and selective internal radiation therapy (SIRT) for intermediate HCC and sorafenib for advanced HCC (3, 4).

Methods

A systematic literature search for yttrium-90 (Y90)-SIRT with glass microsphere studies (published, unpublished and ongoing) was carried out from August 19, 2014 to July 19, 2015, and it was updated on August 30, 2015. Two investigators developed and conducted the literature search. Studies were identified using database searches and citation searches of selected articles. The electronic database searched was PubMed (1965-present). Only articles in English were selected for the review and no date restrictions were applied to the search. The search was conducted using free-text terms and standardized subject terms appropriate for the specific database. Combinations of the following search terms were used: HCC, hepatocellular carcinoma, radioembolization, TARE, SIRT, Yttrium-90, Y90-SIRT, TheraSphere®, glass microspheres, therapy/treatment,

safety, efficacy, and Sorafenib (Table 1). Ongoing and completed clinical trials were identified through searches of clinical trial databases (6). Study eligibility was assessed by a single investigator in an un-blinded manner. Studies were retained if they met the following criteria: 1) TheraSphere® or glass microspheres use in the treatment for HCC, 2) effect of Y90-SIRT with glass microspheres on early, intermediate or advanced HCC, or 3) safety and/or efficacy of Y90-SIRT with glass microspheres or Sorafenib for HCC. All study designs were accepted. Studies were excluded if their aim was to evaluate: 1) SIR-Spheres® or 2) resin-based microspheres as our main focus in this literature review was the treatment of HCC with glass microspheres which is currently approved by the Food and Drug Administration (FDA) under an humanitarian device exemption (HDE).

Yttrium-90 Selective Internal Radiation Therapy (Y90-SIRT)

The technique uses Y90 impregnated glass or resin microspheres, which are delivered through a catheter directly into the hepatic arteries (7, 8). It takes advantage of the fact that primary and secondary hepatic tumors are vascularized mostly by arterial blood flow in comparison to healthy liver parenchyma which obtains its blood supply mainly from portal venous blood (7). The diameters of the glass and resin microspheres (20–30 microns versus 20–60 microns, respectively) allow them to become permanently embolized in the terminal arterioles of tumor (9). Y90 is a pure beta-emitter that disintegrates into stable zirconium-90 and has a half-life of 64.24 ± 0.30 hours (7, 8, 10). The average β -emission is 0.9367 MeV, with a mean tissue penetration of 2.5 mm and a maximum tissue penetration of 10 mm (7, 8). This allows delivery of high radiation doses to hepatic tumors while minimally affecting the healthy surrounding liver parenchyma unlike external radiation (3, 7, 11).

There are two commercially available microspheres:

Table 1. Summary of Search Strategy for PubMed

Search Strategy	Search Terms (Combined with AND/OR)
1	HCC AND (radioembolization OR TARE OR SIRT)
	(HCC therapy OR treatment) AND (TARE OR Y90 OR Yttrium-90)
	Hepatocellular carcinoma AND (TARE OR Y90 OR Y90-SIRT OR radioembolization)
2	HCC OR hepatocellular carcinoma
	HCC AND safety and/or efficacy of (Y90-SIRT OR Y90 OR TheraSphere® OR glass microspheres OR Sorafenib) (3, 6, 11, 24, 34, 43, 47, 48, 61-63, 66-69, 74)

HCC = hepatocellular carcinoma, TARE = transarterial radioembolization, Y90-SIRT = yttrium-90 selective internal radiation therapy

TheraSphere® (MDS Nordion, Ottawa, Canada) and SIR-Spheres® (Sirtex Medical Ltd., Lane Cove, Australia) (8). In 1999, the United States FDA approved the glass microspheres (TheraSphere®), under an HDE, for radiation treatment or as a neo-adjuvant to surgery or transplantation in patients with unresectable HCC who could have placement of appropriately positioned hepatic arterial catheters, and later it approved the use of TheraSphere® in patients with portal vein thrombosis (12). The FDA approved the use of resin microspheres (SIR-Spheres®) in 2002 via a premarket approval for its use in the treatment of unresectable metastatic colorectal cancer to the liver with adjuvant chemotherapy with floxuridine (13).

Pre-Treatment Evaluation

Prior to undergoing Y90-SIRT, patients must undergo pre-treatment evaluation to ensure that the therapy is successful (14, 15). This pre-treatment evaluation is performed 1–2 weeks prior to the procedure and it consists of the following studies: clinical studies, laboratory tests, and imaging (triphasic liver CT or MRI, angiography, and a technetium-99m labeled macroaggregated albumin [^{99m}Tc-MAA] scan) (14, 16).

Clinical Evaluation

The clinical evaluation is performed by a multidisciplinary team which determines a patient's candidacy for Y90-SIRT. The multidisciplinary team may consist of the following specialists depending on the local practices in the institution involved: surgical/medical oncologists, transplant surgeons, hepatologists, radiologists, and interventional radiologists (14). During the clinic visit, evaluation of a patient's performance status per the The Eastern Cooperative Oncology Group (ECOG) is completed (14). Patients with elevated baseline bilirubin (> 2 mg/dL), a Child-Pugh class C (score ≥ 10), an ECOG performance status ≤ 2, and an estimated radiation dose to the lungs > 30 Gy in a single treatment or 50 Gy in multiple treatments are not considered ideal candidates for Y90-SIRT (9, 14).

Laboratory Tests

Patients with HCC undergo laboratory tests that include liver function tests, tumor markers, serum bilirubin, serum albumin, and the prothrombin time (PT)/the international normalized ratio (14). Patients with cirrhosis are classified based on the commonly used Child-Pugh classification

which includes an assessment of encephalopathy and ascites in addition to the laboratory tests.

Imaging Evaluation

Triphasic Liver CT or MRI

The imaging evaluation consists of triphasic liver CT or MRI to evaluate variant vascular anatomy and to assess the extent and location of the hepatic tumor and its hypervascularity (7, 14). CT scans may identify several characteristics of primary HCC that are associated with a favorable response to Y90-SIRT such as well-defined tumor margins, central hypervascularity pattern, and hepatopulmonary fraction (17).

Angiography

Angiography is useful for arterial mapping as it provides the interventional radiologist with an assessment of the hepatic arterial anatomy, variant vasculature, and patency of the portal vein (14). It is also helpful for the occlusion of extrahepatic arteries, which supply the liver, with coil embolization (7, 14). Some extrahepatic arterial branches that may be coil embolized prior to therapy are: GDA, RGA, accessory left gastric artery, cystic artery, falciform ligament artery, phrenic arteries, inferior esophageal artery, supraduodenal artery, and retroduodenal artery (7, 14, 18, 19). These extrahepatic arterial branches are occluded prior to radioembolization to reduce extrahepatic diffusion of the microspheres which can result in significant side effects and/or to minimize delivery of only partial treatment during administration of the microspheres via the main hepatic vessel (7, 14). Between 17% to 30.8% of liver tumors and, in particular HCC, receive their blood supply from extrahepatic arterial branches (7). Development of extrahepatic branches can result from a history of chemoembolization, exophytic tumors, and occlusion of the main arterial pedicle and size of the tumor volume. In 63% of the tumors with development of extrahepatic branches, tumor size was larger than 6 cm (7).

Two commonly occluded extrahepatic arteries are the GDA and the RGA (7, 20). The GDA which branches off the common hepatic artery can be identified on CT and arteriography and is occluded by coiling (hydrocoils or metal coils) prior to the treatment as reflux carries a risk of pancreatitis and gastroduodenal ulceration (7). The study by Vesselle et al. (7) recommends that the GDA should be occluded as proximal as possible as extrahepatic

branches may arise very early and that collateral branches should also be occluded. However, occlusion of the GDA is contraindicated in the presence of a retrograde flow as there is no benefit and it may prove harmful to the patient (7). Another commonly occluded artery is the RGA, which arises from the main hepatic artery in 45–57% of patients (7). Catheterization of the RGA can prove to be technically difficult due to its narrow diameter, tight anatomical angle, and anatomical variants (7). In addition to using a microcatheter, a detachable hydrocoil or remodeling balloon can be used (7).

Technetium-99m Labeled Macroaggregated Albumin (^{99m}Tc-MAA) Scan

Prior to treatment, the patient must also undergo a ^{99m}Tc-MAA scan to assess pulmonary and splanchnic shunting and to confirm that there is no extrahepatic uptake (7, 14). A mixture of albumin particles similar to the size of the glass microspheres (25–35 microns) are bound to the gamma-emitting radioisotope ^{99m}Tc (21). The albumin particles are then imaged via single photon emission computed tomography gamma camera scintigraphy to detect shunting and determine the lung shunt fraction (LSF) (21). The LSF is used to calculate the appropriate dose that should be delivered to the lungs in an attempt to minimize the risk of radiation pneumonitis (9).

Dose Calculation for TheraSphere® (22-24)

To determine the liver volume for which the Y90 glass microspheres are delivered (i.e., volume of distribution) a 3-dimensional reconstruction of the target site is performed using CT or MR imaging. To calculate the mass of infused liver tissue (in kg), a conversion factor of 1.03 g/cm³ is used. The required activity for injection and the dose delivered to the target are calculated by the following formula:

$$A \text{ (in GBq)} = (D \text{ [in Gy]} \times M \text{ [in kg]}) / 50$$

Where A is the net TheraSphere® activity delivered to the liver, D is the dose administered to the target liver mass, and M is the target liver mass.

When the LSF and residual activity (R) in the vial after treatment are taken into account, the actual dose delivered to the target mass (Gy) is calculated by the following formula:

$$D \text{ (in Gy)} = \{A \text{ (in GBq)} \times 50 \times (1 - [\text{LSF} - R])\} / M \text{ (in kg)}$$

Procedure

After pre-treatment evaluation, the patient will undergo Y90-SIRT which is done on an outpatient basis (21). Under fluoroscopic guidance, glass microspheres (about 1–8 million glass spheres) are delivered to the liver via a catheter placed into the femoral artery, which is then guided into the hepatic artery and eventually positioned into the diseased lobe/segment of the liver by following the branch of the hepatic artery (21). Once the catheter is properly positioned, the microspheres are infused at a rate similar to that of the hepatic arterial flow over a period of 3–5 minutes (9). Delivery of the glass microspheres is dependent on the hepatic arterial flow distal to the catheter tip. It is necessary to ensure that the catheter does not occlude the vessel in which it is positioned in order to prevent reflux secondary to vessel spasm (9).

Post-Treatment Assessment

After patients receive their treatment, they undergo post-treatment imaging and laboratory evaluation to assess the response to treatment or lack thereof. Y90-SIRT is associated with low toxicity and patients who receive an administered activity of less than 3 GBq can be released without contact restrictions according to the Nuclear Regulatory Commission contact scenario (25). The time interval for obtaining post-treatment imaging and laboratory evaluation varies among authors and clinical institutions (20). The study by Riaz, Awais, and Salem obtained triphasic CT or MRI at 1 month following treatment and at 3 month intervals following the first post-treatment as it may take 3–6 months for the optimal response (size reduction) to occur (14). Laboratory studies obtained 1 month following Y90-SIRT include liver function studies, complete blood count and tumor markers (alpha-fetoprotein for HCC) (14).

Tumor Response Assessment Following Y90-SIRT

Tumor response assessment following Y90-SIRT can be performed with conventional cross-sectional imaging such as CT and MRI (9, 17). There is a lack of standardization of functional imaging in HCC and response assessment based on anatomical methods is considered as the standard (26, 27).

Four commonly used guidelines to assess tumor response

following therapy are: World Health Organization (WHO) (28) (bidimensional), Response Evaluation Criteria in Solid Tumors (RECIST) (29) (unidimensional), Modified Response Evaluation Criteria in Solid Tumors (mRECIST) (30) (unidimensional), and the European Association for the Study of the Liver (EASL) (31) (necrosis). The RECIST guideline was amended in 2010 to become the mRECIST (30). The mRECIST adopted the concept of viable tumor, which shows arterial enhancement on contrast-enhanced radiologic imaging techniques, thus enabling an evaluation of the tumor response after therapeutic strategy (30). The EASL guidelines address the limitations of the guidelines of the WHO and RECIST, which based their guidelines on systemic therapies resulting in limitations when applied to locoregional therapies such as Y90-SIRT (31).

The study by Rhee et al. (32) showed that diffusion-weighted imaging (DWI) functional MR may assist in early determination of the response or failure of Y90-SIRT in HCC. Unlike MR anatomic imaging studies which are able to assess tumor response until 3 months after treatment, DWI showed that imaging changes at 1 month preceded anatomic size changes seen at 3 months following Y90-SIRT (32). Earlier detection would allow for repeat treatment or alternative therapy such as TACE to be given sooner to patients rather than waiting 3 months to determine the treatment response (32).

In an effort to better identify the tumor response and therapy-related changes in serum markers after Y90-SIRT, a retrospective single-center study assessed the tumor response by using the RECIST 1.1, mRECIST, and Choi criteria (33). Using these three guidelines in addition to the model for end stage liver disease and the serum C-reactive protein, the study was able to create a prognostic model which could predict survival probability for individual patients as early as one month after treatment (33). However, this prognostic model needs to be validated further by performing larger prospective treatment studies.

Complications

Although studies have shown that Y90-SIRT can be considered a safe and efficacious treatment for HCC, complications associated with the therapy can occur (Table 2) (34). Some patients may experience a post-radioembolization syndrome (PRS) which consists of nausea, vomiting, fatigue, abdominal pain/discomfort, and/or cachexia (14). Hospitalization is rarely required and

the incidence of PRS ranges from 10% to 70% (14, 35). Extrahepatic microsphere delivery or radioactivity affecting surrounding structures can result in hepatic dysfunction, biliary system complications, radiation pneumonitis, gastrointestinal (GI) complications, acute pancreatitis, radiation dermatitis, and lymphopenia (14). Other complications include: thrombocytopenia, vascular injury, contrast-induced nephrotoxicity, and allergic reaction to iodinated contrast media (14).

A serious hepatic complication after Y90-SIRT is radiation-induced liver disease (RILD) (14). The incidence of RILD ranges from 0% to 4% and it occurs due to radiation exposure of healthy hepatic parenchyma (14, 36, 37). Clinically, patients develop hepatomegaly, ascites, jaundice and elevated serum transaminases, especially alkaline phosphatase (37). Treatment involves aggressive therapy to control symptoms of abdominal ascites (37). Sangro et al. (38) further described radioembolization-induced liver disease (REILD) in 20% of patients who had undergone chemotherapy either before or after radioembolization. Patients with REILD presented clinically with jaundice and ascites as late as 1 to 2 months post-radioembolization. Post-radioembolization biliary complications include radiation cholecystitis (may be prevented by identifying and coiling the cystic artery (18, 19)), radiation-induced cholangitis, and bilomas/abscess (14). The incidence of post-radioembolization biliary complications is less than 10% and patients with prior surgeries involving the ampulla of Vater have a higher risk (14, 39). Less than 1% of patients will present with radiation pneumonitis and it is recommended that delivery of Y90 to the lungs should be less than 30 Gy in one treatment or the cumulative dose should be less than 50 Gy in multiple treatments (40). Patients with GI complications may present with diarrhea or ulcers. GI ulcers occur in less than 5% of patients and prophylactic coil embolization of the GDA and the RGA may help prevent sequelae (7, 14, 41). Acute pancreatitis may present with severe epigastric pain and with elevated serum lipase and amylase levels (14). Although diffusion of microspheres in the extrahepatic arteries such as the falciform artery is rare, it may result in radiation dermatitis with periumbilical pain (7, 14). Patients may have a greater than 25% decrease in their lymphocyte count following Y90-SIRT without an increase in the incidence of opportunistic infections (11, 14, 34).

Table 2. Summary of Post-Y90 SIRT Complications (11, 14, 35-37, 39-41, 75)

Complications			Findings/Conclusions	Incidence	Prevent/Treatment
Hepatic	Hepatic dysfunction	RILD	Hepatomegaly, ascites, jaundice, elevated serum transaminases (esp. alkaline phosphatase)	0-4% (36, 37)	May require aggressive therapy to control symptoms from abdominal ascites
		REILD (36)	Jaundice and ascites 1 to 2 months after RE/observed in patient that had received chemotherapy pre- or post-RE	20% (36)	Systematically assess for liver damage 1 to 2 months after RE
Extrahepatic	GI	PRS	Nausea, vomiting, fatigue, abdominal pain/discomfort, and/or cachexia	10-70% (14, 35)	Antiemetics for nausea, vomiting; steroids
		GI ulcers		< 5% (41)	May prevent by coiling GDA and right gastric artery/prophylactic antacids; endoscopy to confirm
		RUQ pain or generalized abdominal pain			Over-the-counter analgesics
		Diarrhea			Antidiarrheal medications; fluids and electrolyte replacement
	Biliary	Radiation cholecystitis		< 10% (39)	May prevent by coiling cystic artery
		Radiation-induced cholangitis	Fever, jaundice, RUQ		May require antibiotics
		Bilomas/abscess			Conservative management/percutaneous drainage
	Pancreatic	Acute pancreatitis	Severe epigastric or periumbilical pain	Very rare (14)	Conservative treatment
	Pulmonary	Radiation pneumonitis	May see bat-wing appearance on chest CT	< 1% (40, 75)	Recommend delivery of Y90 to lungs < 30 Gy in one treatment or accumulative dose < 50 Gy in multiple treatments (11, 40)
		Atelectasis and/or pleural effusion			May require steroids
	Renal	Contrast-induced nephrotoxicity			May prevent by adequate hydration pre- and post-procedure and limited use of iodinated contrast
	Vascular	Vascular injury			May prevent by stopping blood thinners appropriately
		Radiation dermatitis and periumbilical pain	Diffusion of microspheres in falciform artery	Rare (14)	May prevent by coiling artery
	Hematology	Lymphopenia	Patients may have greater than 25% of their lymphocyte count decrease following Y90-SIRT		
Thrombocytopenia		Splenomegaly may be observed			
Immunology	Allergic reaction to iodinated contrast	Range from pruritic rash to anaphylactic shock		May require anti-histamine and/or steroids	

GDA = gastroduodenal artery, GI = gastrointestinal, PRS = post-radioembolization syndrome, RE = radioembolization, REILD = radioembolization-induced liver disease, RLD = radiation-induced liver disease, RUQ = right upper quadrant, Y90-SIRT = yttrium-90 selective internal radiation therapy

Y90-SIRT Outcomes

The majority of published studies on the treatment of HCC patients with Y90-SIRT are retrospective or small, non-controlled prospective studies, and therefore, they are only supported by level II-2 and II-3 evidence (42). There is a lack of large, randomized, controlled trials (Table 3). Studies have reported improved median survival in patients with intermediate- to advanced-stage HCC following Y90-SIRT (median 7–41.6 months) (24, 38, 43, 44). Objective response rates vary between 35–70% among studies depending on the guideline criteria used (24, 38, 44, 45). Approximately 20% of patients present with liver-associated toxicity and treatment-related deaths are estimated to occur in about 3% of patients (46).

Portal Venous Thrombosis (PVT)

A common indication for Y90-SIRT is HCC patients with portal venous thrombosis (PVT), which develops in approximately one third of all patients with unresectable HCC (24). Y90-SIRT causes minimal occlusion of the hepatic arteries, and it is therefore safe in the setting of PVT (16). Kulik et al. (24) conducted the first large-series analysis investigating the use of Y90-SIRT in the setting of PVT. The phase II study analyzed Y90-SIRT in 108 HCC patients with and without PVT (34% versus 66%, respectively) and reported partial response rates of 42.4% (size) and 70% (necrosis) (24). Patient survival varied according to the presence of cirrhosis and location of PVT. A phase II study conducted by Mazzaferro et al. (47), which assessed the efficacy of Y90-SIRT in patients with intermediate or advanced HCC, reported that the median time to progression (TTP) was 11 months with no significant difference between PVT versus no PVT (7 months vs. 13 months, respectively), median overall survival was 15 months with a nonsignificant trend in favor of patients without PVT (18 months) versus with PVT (13 months), objective response was 40.4%, and mortality at 30–90 days was 0–3.8%.

In 2010, the study by Salem et al. (46) confirmed the positive outcomes of Y90-SIRT in the treatment of 291 patients with HCC. This was a single-center, prospective, longitudinal cohort study that investigated long-term outcomes, response rate (size and necrosis), TTP and survival stratified by Child-Pugh, United Network for Organ Sharing (UNOS), and Barcelona Clinic Liver Cancer. Survival times differed between patients: Child-Pugh A (17.2 months), Child-Pugh B (7.7 months), and Child-Pugh B with

PVT (5.6 months). The overall TTP was 7.9 months. TTP was longer for Child-Pugh A and B without PVT (15.5 months versus 13 months, respectively) in comparison to those with PVT (5.6 months versus 5.9 months, respectively) (46). The 30-day mortality rate was 3% and the response rates were 57% and 42% based on EASL and WHO criteria, respectively (46). That same year, a European study by Hilgard et al. (43) analyzed Y90-SIRT in 108 patients with advanced HCC and confirmed its safety and efficacy. TTP was 10.0 months with an overall survival of 16.4 months. They observed complete and partial response by necrosis criteria in 3% and 37%, respectively.

Lobar or Segmental Biliary Tract Obstruction

Yttrium-90 Selective Internal Radiation Therapy has been reported to be safe in patients with lobar or segmental biliary tract obstruction and normal bilirubin levels (2 mg/dL or lower) (48). A retrospective study of 12 patients, with a median overall follow-up time of 22.9 months, showed no evidence of therapy-related progressive leukocytosis, bilirubin increase, or biliary complications (infection, sepsis, biliary necrosis/stricture, abscess, or biloma formation) after Y90-SIRT (48).

Novel Concepts

The use of Y90-SIRT has led to the discovery of several novel concepts which may help patients who are undergoing potentially curative resection (49, 50). One of them is the controversial downstaging as a bridge to liver resection or transplantation in selected candidates (16, 51–54). Kulik et al. (55) reported that following Y90-SIRT, 19 of 34 (56%) patients were successfully downstaged from UNOS T3 to T2, and of these patients, eight (23%) underwent liver transplantation. Lewandowski et al. (56) concluded that Y90-SIRT outperforms TACE in downstaging HCC in patients from UNOS T3 to T2 (58% versus 31%, respectively). Y90-SIRT has been shown to be a safe and effective treatment for patients with unresectable HCC and transjugular intrahepatic portosystemic shunt (TIPS) who are awaiting liver transplantation. In the study by Donahue et al. (57), six of 12 patients with existing TIPS underwent liver transplantation after Y90-SIRT. Ibrahim et al. (58) reported that Y90-SIRT appears to be a feasible, safe and effective treatment option for patients with unresectable caudate lobe HCC and had the potential to downstage. Eight out of 291 patients were downstaged to within transplantation criteria. Four patients (50%) were downstaged from UNOS

Table 3. Summary of Clinical Outcomes of Studies of Y90-SIRT (Glass Microspheres) for Treatment of HCC

Lead Author, Year	Study Design	n	Treatment	Prognostic Group	RR	Median TTP (Months)	Median Survival, Months (P)		
Comparative studies									
Moreno-Luna, 2013 (63)	Retrospective, non-randomized	116	TheraSphere vs. TACE				15.0 vs. 14.4 (0.47)		
			61	TheraSphere	CR 12% [‡] , PR 39% [‡]	NR	15.0		
					BCLC A	NR	NR	23.9 (0.04)	
					BCLC B	NR	NR	16.8 (0.16)	
					BCLC C	NR	NR	8.4 (0.47)	
				55	TACE		CR 4% [‡] , PR 47% [‡]	NR	14.4
					BCLC A	NR	NR	18.6	
					BCLC B	NR	NR	13	
					BCLC C	NR	NR	10.1	
		Salem, 2011 (45)	Retrospective, non-randomized	245	TheraSphere vs. TACE		49%* vs. 36%* (0.104)	13.3 vs. 8.4 (0.046)	20.5 vs. 17.4 (0.23)
123	TheraSphere				49%*	13.3	20.5		
					BCLC A	47%* (0.229)	25.1 (0.4)	27.3 (0.74)	
					BCLC B	51%* (0.581)	13.3 (0.047)	17.2 (0.42)	
					BCLC C	54%* (0.097)	13.8 (0.38)	22.1 (0.04)	
				122	TACE		36%*	8.4	17.4
					BCLC A	32%*	8.8	45.4	
					BCLC B	44%*	9.4	17.5	
					BCLC C	17%*	7.9	9.3	
Lance, 2011 (62)	Retrospective, non-randomized			73	TACE vs. SIR-Spheres or TheraSphere				8.0 vs. 10.3 (0.33)
		38	SIR-Spheres or TheraSphere		NR	NR	8.0		
		35	TACE		NR	NR	10.3		
El Fouly, 2015 (64)	Prospective, non-randomized	86	TheraSphere vs. TACE			13.3 vs. 6.8 (NS)	16.4 vs. 18 (NS)		
		44	TheraSphere BCLC B		CR 7% [‡] , PR 68% [‡]	13.3	16.4		
		42	TACE BCLC B		CR 5% [‡] , PR 45% [‡]	6.8	18		
Carr, 2010 (76)	Retrospective, non-randomized	790	TheraSphere vs. TACE				11.5 vs. 8.5 (< 0.05)		
			99	TheraSphere +PVT -PVT		CR 3%*, PR 33%*	NR	11.5 5 (< 0.05) 16 (NS)	
				691	TACE +PVT -PVT		CR 5%*, PR 55%*	NR	8.5 7 12
Lewandowski, 2009 (56)	Retrospective, non-randomized	86	TheraSphere vs. TACE				41.6 vs. 19.2 (0.008)		
		43	TheraSphere UNOS T3		61%*	33.3	41.6		
		43	TACE UNOS T3		37%*	18.2	19.2		

Table 3. Summary of Clinical Outcomes of Studies of Y90-SIRT (Glass Microspheres) for Treatment of HCC (Continued)

Lead Author, Year	Study Design	n	Treatment	Prognostic Group	RR	Median TTP (Months)	Median Survival, Months (P)		
Woodall, 2009 (77)	Prospective, non-randomized	52	TheraSphere		NR	NR	13.9 vs. 3.2 (0.01)		
		20		BCLC A-C -PVT	NR	NR	13.9		
		15		BCLC C +PVT	NR	NR	3.2 (0.26)		
		17		No treatment, screen failure	NR	NR	5.2		
Goin, 2004 (78)	Retrospective, non-randomized	63	TheraSphere vs. TACE		NR	NR	NR		
		34		TheraSphere + Okuda I	NR	NR	25.5		
				TheraSphere + Okuda II	NR	NR	10.9		
		29		TACE + Okuda I	NR	NR	11.3		
				TACE + Okuda II	NR	NR	11.7		
Non-comparative studies									
Dancey, 2000 (79)	Retrospective, non-randomized	20	TheraSphere	Okuda I/II	CP 5%, PR 15%	10.2	12.5		
Carr, 2004 (34)	Retrospective, non-randomized	65	TheraSphere		PR 38.4%	NR	21 vs. 10 (NS)		
		42		Okuda I	NR	NR	21		
		23		Okuda II	NR	NR	10		
Geschwind, 2004 (80)	Retrospective, non-randomized	80	TheraSphere		NR	NR	20.6 vs. 12.6 (0.02)		
		54		Okuda I	NR	NR	20.6		
		26		Okuda II	NR	NR	12.6		
Goin, 2005 (81)	Combined prospective and retrospective, non-randomized	121	TheraSphere		NR	NR	15.5 vs. 3.6 (< 0.0001)		
		88		Low risk	NR	NR	15.5		
		33		High risk	NR	NR	3.6		
Kulik, 2006 (55)	Retrospective, non-randomized	35	TheraSphere	UNOS T3	50%*	NR	26.3		
		Goin, 2005 (82)	Retrospective, non-randomized	88	TheraSphere		NR	NR	(< 0.001)
				26		CLIP 0	NR	NR	26.7
				41		CLIP 1-2	NR	NR	11.6
13		CLIP > 2	NR	NR	7.1				
Salem, 2004 (23)	Retrospective, non-randomized	15	TheraSphere	+Branch PVT	NR	NR	7.1		
Pressiani, 2013 (69)	Prospective, non-randomized	297	Sorafenib		NR	4.1	9.1		
		234		BCLC B/C CPA	NR	4.2	10.0 (< 0.001)		

Table 3. Summary of Clinical Outcomes of Studies of Y90-SIRT (Glass Microspheres) for Treatment of HCC (Continued)

Lead Author, Year	Study Design	n	Treatment	Prognostic Group	RR	Median TTP (Months)	Median Survival, Months (P)	
		63		BCLC B/C CPB	NR	3.8	3.8	
Bruix, 2012 (68)	Subgroup Study of Prospective Phase III	299	Sorafenib vs. Placebo			6.9 vs. 4.9	14.5 vs. 9.7	
		54		BCLC B	NR	6.9	14.5	
		245		BCLC C	NR	4.9	9.7	
Llovet, 2008 (66)	Prospective Phase III	602	Sorafenib vs. Placebo			5.5 vs. 2.8 (< 0.001)	10.7 vs. 7.9 (< 0.001)	
		299		Sorafenib BCLC B + C	PR 2% [‡]	5.5	10.7	
		303		Placebo BCLC B + C	PR 1% [‡]	2.8	7.9	
Cheng, 2009 (67)	Prospective Phase III	226	Sorafenib vs. Placebo			2.8 vs. 1.4 (0.005)	6.5 vs. 4.2 (0.014)	
		150		Sorafenib BCLC C, CPA	PR 5% [‡]	2.8	6.5	
		76		Placebo BCLC C, CPA	PR 1% [‡]	1.4	4.2	
Mazzaferro, 2013 (47)	Prospective Phase II	52	TheraSphere		OR 40.4%*	11	15	
		17		-PVT BCLC B CPA	OR 8%* OR 6%*	13 13	18 18	
		35		+PVT BCLC C CPA CPB	OR 13%* OR 10%* OR 3%*	7 6 NR	13 16 6	
Salem, 2010 (46)	Prospective, non-randomized	291	TheraSphere		42%*	7.9	NR	
				BCLC A	21%*	25.1	26.9	
				BCLC B	42%*	13.3	13.3	
				BCLC C, -EHD	40%*	6.0	7.3	
				BCLC C, +EHD	11%*	3.1	5.4	
Hilgard, 2010 (43)	Retrospective, non-randomized	108	TheraSphere		CR 3% ^{†¶} , PR 20% ^{†¶}	10.0	16.4	
		2		BCLC A	NR	NR	NR	
		51		BCLC B	NR	NR	16.4	
		55		BCLC C	NR	NR	NR	
				CPA	NR	NR	17.2	
				CPB	NR	NR	6	
				-PVT +PVT	NR NR	NR NR	16.4 10.0	
Kulik, 2008 (24)	Prospective Phase II	108	TheraSphere		PR 42.4%*, RR 70% [†]	NR	(0.0052)	
		71		BCLC C, -PVT	NR	NR	15.4	
		25		BCLC C, +Branch PVT	NR	NR	10.0	

Table 3. Summary of Clinical Outcomes of Studies of Y90-SIRT (Glass Microspheres) for Treatment of HCC (Continued)

Lead Author, Year	Study Design	n	Treatment	Prognostic Group	RR	Median TTP (Months)	Median Survival, Months (P)
		12		BCLC C, +Main PVT	NR	NR	4.4
Salem, 2005 (11)	Prospective Phase II	43	TheraSphere		PR 47%	NR	24.4 vs. 12.5 (< 0.001)
		21		Okuda I	NR	NR	24.4
		22		Okuda II	NR	NR	12.5

*WHO criteria, ¹EASL criteria, ²RECIST criteria, ³mRECIST criteria (30), ⁴76 out of 108 responses + necrosis after 30 days of treatment. BCLC = Barcelona Clinic Liver Cancer staging system, CLIP = Cancer of the Liver Italian Program scoring system, CP = Child-Pugh score, CR = complete response, -EHD/+EHD = without or with extrahepatic disease, HCC = hepatocellular carcinoma, NR = not recorded, NS = not statistically significant, OR = objective response, PR = partial response, PVT = portal vein thrombosis, RR = response rate, SIRT = selective internal radiation therapy, TACE = transarterial chemoembolization, TTP = time to progression, UNOS = United Network for Organ Sharing, Y90 = yttrium-90

T3 to T2, three patients underwent liver transplantation, and one of the patients despite being downstaged was unable to undergo transplantation, given the comorbid conditions. Moreover, downstaging can also be achieved in patients with PVT (59).

A second novel concept termed “radiation lobectomy” is observed in HCC patients whose right-lobe disease is treated with Y90-SIRT (16, 49, 50). After Y90-SIRT treatment, the irradiated lobe undergoes atrophy and the contralateral lobe undergoes hypertrophy, as opposed to portal vein embolization which can induce hypertrophy but does not treat HCC (16, 60).

The third novel concept is labeled “radiation segmentectomy” and it uses Y90-SIRT to obliterate small, single tumor-bearing liver segments that are contraindicated for ablation or resection secondary to location, insufficient liver reserve, and comorbidities (35, 61).

Y90-SIRT with Glass Microspheres versus TACE in Intermediate Stage HCC

Transarterial chemoembolization is the treatment of choice for patients with intermediate stage HCC, but studies have shown that Y90-SIRT may have a role in subgroups of these patients (16). It is difficult to perform randomized controlled trials comparing TACE versus Y90-SIRT.

In comparison to Y90-SIRT, TACE involves premedicating patients (i.e., antibiotics, antiemetics, and narcotics) and hospitalization ranging from 1 to 5 days following treatment for post-embolization syndrome (16). Moreover, prospective data comparing the efficacy with regard to response and/or survival between Y90-SIRT and TACE in patients with intermediate HCC are lacking.

In 2011 Salem et al. (45) published a large comparative

effectiveness study. The retrospective study of 245 patients who received either TACE (122 patients) or Y90-SIRT (123 patients), reported that although TTP was longer after Y90-SIRT than after TACE (13.3 months versus 8.4 months, respectively, $p = 0.046$), median survival times were not statistically different between the two treatment groups in patients with intermediate disease (17.2 months versus 17.5 months, respectively, $p = 0.42$) (45). Another retrospective study showed there was no significant difference in survival between Y90-SIRT and TACE (median 8 months versus 10.3 months, respectively, $p = 0.33$) (62). This study further showed that post-embolization syndrome was significantly more severe in patients who underwent TACE resulting in increased total hospitalization rates (62).

The retrospective case-control study by Moreno-Luna et al. (63) in 2013 showed that there was no significant difference in efficacy between Y90-SIRT and TACE. The median survival did not differ between Y90-SIRT (15.0 months) and TACE (14.4 months). The two-year survival was 30% for Y90-SIRT and 24% for TACE. Complete tumor response was more common after Y90-SIRT (12%) than after TACE (4%). However, Y90-SIRT patients reported more fatigue, had less fever and required less hospitalization than patients treated with TACE. The study by El Fouly et al. (64) in 2014 reported similar findings during the comparative analysis between TACE and Y90-SIRT in the treatment of 86 patients with intermediate stage HCC. Both treatments resulted in similar median overall survival rates (18 months for TACE versus 16.4 months for Y90-SIRT) and the TTP was not statistically different between treatments. However, the number of treatment sessions, total hospitalization time, and rate of adverse events were significantly higher in the TACE cohort. Future studies comparing Y90-SIRT and TACE

Table 4. Summary of Ongoing and Recruiting Clinical Trials for Y90-SIRT for HCC Treatment (6)

Clinical Trials.gov Identifier	Treatment Arm(s)	Patient Population	Primary Outcome Measure	Status	Sponsor
NCT01349075	TheraSphere (Yttrium-90)	Unresectable HCC	Response to treatment via diagnostic imaging	Recruiting	Thomas Jefferson University
NCT02072356	TheraSphere (Yttrium-90)	Unresectable HCC	Response to treatment; survival time, adverse experiences	Recruiting	Ohio State University Comprehensive Cancer Center
NCT00906984	TheraSphere (Yttrium-90)	Unresectable HCC	Response to treatment via diagnostic imaging	Recruiting	University of California, Irvine
NCT01176604	TheraSphere (Yttrium-90)	Unresectable HCC	Overall survival associated with treatment	Recruiting	M.D. Anderson Cancer Center
NCT00877136	TheraSphere (Yttrium-90)	Unresectable HCC	Evaluate patient quality of life and toxicities associated with treatment	Recruiting	St. Joseph Hospital of Orange
NCT01686880 [†]	SIR-Spheres (Yttrium-90)	HCC in cirrhotic liver	Peri-operative morbidity of SIRT prior to surgical resection or radiofrequency	Recruiting	Jules Bordet Institute
NCT00956930 [†]	Yttrium-90 glass microspheres vs. TACE (cisplatin, mitomycin, doxorubicin)	Unresectable HCC	Time to progression in patients treated with TACE vs. Y90 via diagnostic imaging	Recruiting	Northwestern University
NCT01381211 [†]	TheraSphere (Yttrium-90) vs. TACE-DEB (doxorubicin)	Intermediate HCC	Time to progression	Recruiting	University Hospital, Ghent
NCT00846131 [*]	TheraSphere (Yttrium-90) vs. Sorafenib + TheraSphere (Yttrium-90)	Pre-transplant HCC	Evaluate sorafenib as an adjunct to Y-90 for control of HCC as a bridge/downstage to transplant	Ongoing	Northwestern University
NCT01900002 [†]	TheraSphere (Yttrium-90) + Sorafenib	Advanced HCC	Toxicity of Sorafenib and Yttrium-90	Recruiting	M.D. Anderson Cancer Center
NCT01556490 [‡]	Sorafenib vs. TheraSphere (Yttrium-90) + Sorafenib	Unresectable HCC	Overall survival	Recruiting	BTG International Inc.
NCT01126645 [†]	RFA followed by sorafenib or placebo (local ablation group) or SIRT + sorafenib or sorafenib alone (palliative treatment group)	Unresectable HCC	Time to recurrence; overall survival; Primovist®-enhanced MRI is non-inferior or superior compared with contrast-enhanced multislice CT	Recruiting	University of Magdeburg
NCT01482442 [‡]	SIR-Spheres (Yttrium-90) vs. Sorafenib	Advanced HCC	Median overall survival time	Recruiting	Assistance Publique - Hôpitaux de Paris
NCT01135056 [‡]	SIR-Spheres (Yttrium-90) vs. Sorafenib	Locally advanced HCC	Overall survival	Recruiting	Singapore General Hospital
NCT02004210 [‡]	TACE vs. TARE	Advanced HCC	Overall survival	Recruiting	Seoul National University Hospital
NCT00530010	TheraSphere (Yttrium-90)	Unresectable HCC	Proportion of patients completing scheduled treatment plan	Recruiting	Northwestern University

Table 4. Summary of Ongoing and Recruiting Clinical Trials for Y90-SIRT for HCC Treatment (6) (Continued)

Clinical Trials.gov Identifier	Treatment Arm(s)	Patient Population	Primary Outcome Measure	Status	Sponsor
NCT02305459	TheraSphere (Yttrium-90) + QLQ-C30 with HCC module (Behavioral)	HCC	Change from Baseline in Quality of Life questionnaire QLQ-C30 with HCC Module	Recruiting	Cardiovascular and Interventional Radiological Society of Europe
NCT01775280 [†]	Yttrium-90 glass microspheres	Unresectable to borderline resectable HCC	Percentage of patients that can be downstaged to resectability	Recruiting	University of Zurich
NCT01798160 [‡]	TACE-DEB vs. SIR-Spheres (Yttrium-90)	HCC	Progression-free survival; overall survival	Ongoing	Johannes Gutenberg University Mainz
NCT01887717 [‡]	TheraSphere (Yttrium-90) vs. Sorafenib	Advanced HCC with PVT	Overall survival	Recruiting	BTG International Inc.

*Phase I trials, [†]Phase II trials, [‡]Phase III trials, [§]Phase IV trials. DEB = drug eluting beads, HCC = hepatocellular carcinoma, PVT = portal vein thrombosis, RFA = radiofrequency ablation, SIRT = selective internal radiation therapy, TACE = transarterial chemoembolization, TARE = transarterial radioembolization

should also assess their cost-effectiveness in therapy for intermediate stage HCC, given the increased hospital stay due to complications from repeated TACE treatments.

Although studies have not shown statistically significant differences in efficacy or survival between Y90-SIRT and TACE, Y90-SIRT seems to be better tolerated with significant differences in length of hospital stay and post-embolization symptoms. In addition, Y90-SIRT outperforms TACE with regard to downstaging and quality of life measures. As discussed previously, the ability to downstage from UNOS T3 to T2 as a bridge to liver transplantation was achieved more frequently with Y90-SIRT (58%) than with TACE (31%) (56). In 2013, the study by Salem et al. (65) demonstrated that Y90-SIRT outperformed TACE by validated quality-of-life (QoL) measures. The prospective study of 56 patients with HCC who underwent Y90-SIRT versus TACE (29 versus 27, respectively) showed that although patients who received Y90-SIRT had a larger tumor burden, they had higher QoL scores in comparison to patients who received TACE.

Y90-SIRT with Glass Microspheres versus Sorafenib in Advanced Stage HCC

The treatment of choice in patients with advanced stage HCC is sorafenib and it is associated with an overall survival of 6.5–14.5 months (43, 66–69). It is difficult to compare the efficacy of Y90-SIRT with sorafenib in advanced HCC through well-designed randomized controlled studies, given the high probability of crossover among treatment groups as HCC progresses. No study directly comparing sorafenib with Y90-SIRT with glass microspheres is currently available.

However, the observational cohort study by Hilgard et al. (43) showed that in comparison to the phase III trial which led to the approval of sorafenib (SHARP trial), the median overall survival in patients with advanced HCC treated with Y90-SIRT was slightly longer (16.4 months) than that in patients with advanced HCC treated with sorafenib (10.7 months) (66). In addition, unlike Y90-SIRT, sorafenib has been shown to have significant side effects, which result in treatment discontinuation (44%), dose reduction, or withdrawal (64%) (66, 69).

Future Prospects

There are currently 16 active clinical trials investigating the use of Y90-SIRT with glass microspheres for treatment of HCC (Table 4) (6). In the United States, there are studies investigating the use of Y90-SIRT for the treatment of unresectable HCC (NCT01349075, NCT02072356, NCT00906984, NCT01176604, and NCT00877136) and assessing the safety of Y90-SIRT compared to radiofrequency prior to surgical resection in patients with HCC (NCT01686880) (6). The PREMIERE trial in the United States is comparing Y90-SIRT with TACE in patients who are not candidates for radiofrequency ablation or in patients with unresectable HCC (NCT00956930) (6). The European TRACE trial is a multicenter randomized control study comparing TARE with TACE for the treatment of HCC (NCT01381211) (6).

The good toxicity profile of Y90-SIRT makes it an alternative or adjunct to treatment with sorafenib (70).

In the United States, a prospective randomized trial is studying the role of Y90-SIRT alone or in combination with sorafenib in the treatment of HCC patients who are awaiting liver transplantation (NCT00846131) and a phase II study is investigating the role of Y90-SIRT with sorafenib in advanced HCC (NCT01900002) (6). The STOP-HCC study, which is being conducted in the United States and Europe, is comparing sorafenib with and without Y90-SIRT (NCT01556490) (6). The outcomes of these trials may result in the incorporation of Y90-SIRT with glass microspheres into the treatment guidelines as a standard therapy option for patients with intermediate and advanced stage HCC.

CONCLUSION

Although the use of Y90-SIRT in treating liver malignancies dates back to the 1960s (71-73), it has recently begun to gain clinical acceptance as a promising treatment option for patients with intermediate and advanced HCC (8). Multiple studies have provided compelling data which suggest that Y90-SIRT in comparison to TACE, has a higher tumor response, less post-embolization symptoms, better downstaging outcomes, and provides a higher QoL. However, the EASL, the European Organization for Research and Treatment of Cancer, and the American Association for the Study of Liver Diseases do not currently recommend Y90-SIRT as a standard therapy for intermediate or advanced HCC outside clinical trials (70). This clinical lack of enthusiastic approval stems from the fact that there is a lack of research and evidence from large-scale randomized controlled trials. Nonetheless, with the growing body of level 2 and level 3 evidence, Y90-SIRT has found a place in the guidelines adopted by the European Society for Medical Oncology, the European Society of Digestive Oncology, and the National Comprehensive Cancer Network (21). Y90-SIRT may become widely accepted and may be incorporated into the treatment guidelines if further research shows reproducibility, multicenter implementation, and economic feasibility.

REFERENCES

1. Ferlay J, Soerjomataram I, Dikshit R, Eser S, Mathers C, Rebelo M, et al. Cancer incidence and mortality worldwide: sources, methods and major patterns in GLOBOCAN 2012. *Int J Cancer* 2015;136:E359-E386
2. Cancer Facts & Figures 2014. American Cancer Society. <http://www.cancer.org/research/cancerfactsstatistics/>
3. cancerfactsfigures2014/. Accessed December 13, 2015
3. European Association For The Study Of The Liver; European Organisation For Research And Treatment Of Cancer. EASL-EORTC clinical practice guidelines: management of hepatocellular carcinoma. *J Hepatol* 2012;56:908-943
4. Maida M, Orlando E, Cammà C, Cabibbo G. Staging systems of hepatocellular carcinoma: a review of literature. *World J Gastroenterol* 2014;20:4141-4150
5. Llovet JM, Bruix J. Systematic review of randomized trials for unresectable hepatocellular carcinoma: chemoembolization improves survival. *Hepatology* 2003;37:429-442
6. ClinicalTrials.gov. <http://clinicaltrials.gov/ct2/home>. Accessed December 13, 2015
7. Vesselle G, Petit I, Boucebc S, Rocher T, Velasco S, Tasu JP. Radioembolization with yttrium-90 microspheres work up: practical approach and literature review. *Diagn Interv Imaging* 2015;96:547-562
8. Salem R, Lewandowski RJ, Sato KT, Atassi B, Ryu RK, Ibrahim S, et al. Technical aspects of radioembolization with 90Y microspheres. *Tech Vasc Interv Radiol* 2007;10:12-29
9. Memon K, Lewandowski RJ, Riaz A, Salem R. Yttrium 90 microspheres for the treatment of hepatocellular carcinoma. *Recent Results Cancer Res* 2013;190:207-224
10. Volchok HL, Kulp JL. Half-life of yttrium-90. *Phys Rev* 1955;97:102
11. Salem R, Lewandowski RJ, Atassi B, Gordon SC, Gates VL, Barakat O, et al. Treatment of unresectable hepatocellular carcinoma with use of 90Y microspheres (TheraSphere): safety, tumor response, and survival. *J Vasc Interv Radiol* 2005;16:1627-1639
12. TheraSphere HDE Approval. US Food and Drug Administration. <http://www.fda.gov/ohrms/dockets/dailys/00/jan00/010500/aav0001.pdf>. Accessed December 13, 2015
13. Medical Devices SIR-Spheres. U.S. Food and Drug Administration. <http://www.fda.gov/MedicalDevices/ProductsandMedicalProcedures/DeviceApprovalsandClearances/Recently-ApprovedDevices/ucm083605.htm>. Accessed December 13, 2015
14. Riaz A, Awais R, Salem R. Side effects of yttrium-90 radioembolization. *Front Oncol* 2014;4:198
15. Lau WY, Kennedy AS, Kim YH, Lai HK, Lee RC, Leung TW, et al. Patient selection and activity planning guide for selective internal radiotherapy with yttrium-90 resin microspheres. *Int J Radiat Oncol Biol Phys* 2012;82:401-407
16. Salem R, Mazzaferro V, Sangro B. Yttrium 90 radioembolization for the treatment of hepatocellular carcinoma: biological lessons, current challenges, and clinical perspectives. *Hepatology* 2013;58:2188-2197
17. Salem ME, Jain N, Dyson G, Taylor S, El-Refai SM, Choi M, et al. Radiographic parameters in predicting outcome of patients with hepatocellular carcinoma treated with yttrium-90 microsphere radioembolization. *ISRN Oncol* 2013;2013:538376
18. Theysohn JM, Müller S, Schlaak JF, Ertle J, Schlosser TW, Bockisch A, et al. Selective internal radiotherapy (SIRT) of hepatic tumors: how to deal with the cystic artery. *Cardiovasc*

- Intervent Radiol* 2013;36:1015-1022
19. McWilliams JP, Kee ST, Loh CT, Lee EW, Liu DM. Prophylactic embolization of the cystic artery before radioembolization: feasibility, safety, and outcomes. *Cardiovasc Intervent Radiol* 2011;34:786-792
 20. Powerski MJ, Scheurig-Münkler C, Banzer J, Schnapauff D, Hamm B, Gebauer B. Clinical practice in radioembolization of hepatic malignancies: a survey among interventional centers in Europe. *Eur J Radiol* 2012;81:e804-e811
 21. Kennedy A. Radioembolization of hepatic tumors. *J Gastrointest Oncol* 2014;5:178-189
 22. Physicians Package Insert. TheraSphere. http://www.therasphere.com/physicians_us/package_insert.asp. Accessed December 13, 2015
 23. Salem R, Lewandowski R, Roberts C, Goin J, Thurston K, Abouljoud M, et al. Use of Yttrium-90 glass microspheres (TheraSphere) for the treatment of unresectable hepatocellular carcinoma in patients with portal vein thrombosis. *J Vasc Interv Radiol* 2004;15:335-345
 24. Kulik LM, Carr BI, Mulcahy MF, Lewandowski RJ, Atassi B, Ryu RK, et al. Safety and efficacy of 90Y radiotherapy for hepatocellular carcinoma with and without portal vein thrombosis. *Hepatology* 2008;47:71-81
 25. McCann JW, Larkin AM, Martino LJ, Eschelman DJ, Gonsalves CF, Brown DB. Radiation emission from patients treated with selective hepatic radioembolization using yttrium-90 microspheres: are contact restrictions necessary? *J Vasc Interv Radiol* 2012;23:661-667
 26. Riaz A, Memon K, Miller FH, Nikolaidis P, Kulik LM, Lewandowski RJ, et al. Role of the EASL, RECIST, and WHO response guidelines alone or in combination for hepatocellular carcinoma: radiologic-pathologic correlation. *J Hepatol* 2011;54:695-704
 27. Duke E, Deng J, Ibrahim SM, Lewandowski RJ, Ryu RK, Sato KT, et al. Agreement between competing imaging measures of response of hepatocellular carcinoma to yttrium-90 radioembolization. *J Vasc Interv Radiol* 2010;21:515-521
 28. World Health Organization. WHO Handbook for Reporting Results of Cancer Treatment. http://whqlibdoc.who.int/offset/WHO_OFFSET_48.pdf. Accessed December 13, 2015
 29. Therasse P, Arbuck SG, Eisenhauer EA, Wanders J, Kaplan RS, Rubinstein L, et al. New guidelines to evaluate the response to treatment in solid tumors. European Organization for Research and Treatment of Cancer, National Cancer Institute of the United States, National Cancer Institute of Canada. *J Natl Cancer Inst* 2000;92:205-216
 30. Lencioni R, Llovet JM. Modified RECIST (mRECIST) assessment for hepatocellular carcinoma. *Semin Liver Dis* 2010;30:52-60
 31. Bruix J, Sherman M, Llovet JM, Beaugrand M, Lencioni R, Burroughs AK, et al. Clinical management of hepatocellular carcinoma. Conclusions of the Barcelona-2000 EASL conference. European Association for the Study of the Liver. *J Hepatol* 2001;35:421-430
 32. Rhee TK, Naik NK, Deng J, Atassi B, Mulcahy MF, Kulik LM, et al. Tumor response after yttrium-90 radioembolization for hepatocellular carcinoma: comparison of diffusion-weighted functional MR imaging with anatomic MR imaging. *J Vasc Interv Radiol* 2008;19:1180-1186
 33. Weng Z, Ertle J, Zheng S, Lauenstein T, Mueller S, Bockisch A, et al. A new model to estimate prognosis in patients with hepatocellular carcinoma after Yttrium-90 radioembolization. *PLoS One* 2013;8:e82225
 34. Carr BI. Hepatic arterial 90Yttrium glass microspheres (Therasphere) for unresectable hepatocellular carcinoma: interim safety and survival data on 65 patients. *Liver Transpl* 2004;10(2 Suppl 1):S107-S110
 35. Padia SA, Kwan SW, Roudsari B, Monsky WL, Coveler A, Harris WP. Superselective yttrium-90 radioembolization for hepatocellular carcinoma yields high response rates with minimal toxicity. *J Vasc Interv Radiol* 2014;25:1067-1073
 36. Sangro B, Gil-Alzugaray B, Rodriguez J, Sola I, Martinez-Cuesta A, Viudez A, et al. Liver disease induced by radioembolization of liver tumors: description and possible risk factors. *Cancer* 2008;112:1538-1546
 37. Kennedy AS, McNeillie P, Dezarn WA, Nutting C, Sangro B, Wertman D, et al. Treatment parameters and outcome in 680 treatments of internal radiation with resin 90Y-microspheres for unresectable hepatic tumors. *Int J Radiat Oncol Biol Phys* 2009;74:1494-1500
 38. Sangro B, Bilbao JJ, Iñarrairaegui M, Rodriguez M, Garrastachu P, Martinez-Cuesta A. Treatment of hepatocellular carcinoma by radioembolization using 90Y microspheres. *Dig Dis* 2009;27:164-169
 39. Atassi B, Bangash AK, Lewandowski RJ, Ibrahim S, Kulik L, Mulcahy MF, et al. Biliary sequelae following radioembolization with Yttrium-90 microspheres. *J Vasc Interv Radiol* 2008;19:691-697
 40. Salem R, Parikh P, Atassi B, Lewandowski RJ, Ryu RK, Sato KT, et al. Incidence of radiation pneumonitis after hepatic intra-arterial radiotherapy with yttrium-90 microspheres assuming uniform lung distribution. *Am J Clin Oncol* 2008;31:431-438
 41. Murthy R, Brown DB, Salem R, Meranze SG, Coldwell DM, Krishnan S, et al. Gastrointestinal complications associated with hepatic arterial Yttrium-90 microsphere therapy. *J Vasc Interv Radiol* 2007;18:553-561; quiz 562
 42. U.S. Preventive Services Task Force Procedure Manual. AHRQ Publication No. 08-05118-EF. USPSTF. <http://www.uspreventiveservicestaskforce.org/Home/GetFile/6/7/procmanual/pdf>. Accessed December 13, 2015
 43. Hilgard P, Hamami M, Fouly AE, Scherag A, Müller S, Ertle J, et al. Radioembolization with yttrium-90 glass microspheres in hepatocellular carcinoma: European experience on safety and long-term survival. *Hepatology* 2010;52:1741-1749
 44. Sangro B, Salem R, Kennedy A, Coldwell D, Wasan H. Radioembolization for hepatocellular carcinoma: a review of the evidence and treatment recommendations. *Am J Clin Oncol* 2011;34:422-431
 45. Salem R, Lewandowski RJ, Kulik L, Wang E, Riaz A, Ryu RK, et al. Radioembolization results in longer time-to-progression and reduced toxicity compared with chemoembolization in

- patients with hepatocellular carcinoma. *Gastroenterology* 2011;140:497-507.e2
46. Salem R, Lewandowski RJ, Mulcahy MF, Riaz A, Ryu RK, Ibrahim S, et al. Radioembolization for hepatocellular carcinoma using Yttrium-90 microspheres: a comprehensive report of long-term outcomes. *Gastroenterology* 2010;138:52-64
 47. Mazzaferro V, Sposito C, Bhoori S, Romito R, Chiesa C, Morosi C, et al. Yttrium-90 radioembolization for intermediate-advanced hepatocellular carcinoma: a phase 2 study. *Hepatology* 2013;57:1826-1837
 48. Gaba RC, Riaz A, Lewandowski RJ, Ibrahim SM, Ryu RK, Sato KT, et al. Safety of yttrium-90 microsphere radioembolization in patients with biliary obstruction. *J Vasc Interv Radiol* 2010;21:1213-1218
 49. Theysohn JM, Ertle J, Müller S, Schlaak JF, Nensa F, Sipilae S, et al. Hepatic volume changes after lobar selective internal radiation therapy (SIRT) of hepatocellular carcinoma. *Clin Radiol* 2014;69:172-178
 50. Teo JY, Goh BK, Cheah FK, Allen JC, Lo RH, Ng DC, et al. Underlying liver disease influences volumetric changes in the spared hemiliver after selective internal radiation therapy with 90Y in patients with hepatocellular carcinoma. *J Dig Dis* 2014;15:444-450
 51. Lambert B, Sturm E, Mertens J, Oltenfreiter R, Smeets P, Troisi R, et al. Intra-arterial treatment with ⁹⁰Y microspheres for hepatocellular carcinoma: 4 years experience at the Ghent University Hospital. *Eur J Nucl Med Mol Imaging* 2011;38:2117-2124
 52. Khalaf H, Alsuhaibani H, Al-Sugair A, Al-Mana H, Al-Mutawa A, Al-Kadhi Y, et al. Use of yttrium-90 microsphere radioembolization of hepatocellular carcinoma as downstaging and bridge before liver transplantation: a case report. *Transplant Proc* 2010;42:994-998
 53. Luna LE, Kwo PY, Roberts LR, Mettler TA, Gansen DN, Andrews JC, et al. Liver transplantation after radioembolization in a patient with unresectable HCC. *Nat Rev Gastroenterol Hepatol* 2009;6:679-683
 54. Tohme S, Sukato D, Chen HW, Amesur N, Zajko AB, Humar A, et al. Yttrium-90 radioembolization as a bridge to liver transplantation: a single-institution experience. *J Vasc Interv Radiol* 2013;24:1632-1638
 55. Kulik LM, Atassi B, van Holsbeeck L, Souman T, Lewandowski RJ, Mulcahy MF, et al. Yttrium-90 microspheres (TheraSphere) treatment of unresectable hepatocellular carcinoma: downstaging to resection, RFA and bridge to transplantation. *J Surg Oncol* 2006;94:572-586
 56. Lewandowski RJ, Kulik LM, Riaz A, Senthilnathan S, Mulcahy MF, Ryu RK, et al. A comparative analysis of transarterial downstaging for hepatocellular carcinoma: chemoembolization versus radioembolization. *Am J Transplant* 2009;9:1920-1928
 57. Donahue LA, Kulik L, Baker T, Ganger DR, Gupta R, Memon K, et al. Yttrium-90 radioembolization for the treatment of unresectable hepatocellular carcinoma in patients with transjugular intrahepatic portosystemic shunts. *J Vasc Interv Radiol* 2013;24:74-80
 58. Ibrahim SM, Kulik L, Baker T, Ryu RK, Mulcahy MF, Abecassis M, et al. Treating and downstaging hepatocellular carcinoma in the caudate lobe with yttrium-90 radioembolization. *Cardiovasc Intervent Radiol* 2012;35:1094-1101
 59. Garin E, Lenoir L, Rolland Y, Edeline J, Mesbah H, Laffont S, et al. Dosimetry based on ^{99m}Tc-macroaggregated albumin SPECT/CT accurately predicts tumor response and survival in hepatocellular carcinoma patients treated with ⁹⁰Y-loaded glass microspheres: preliminary results. *J Nucl Med* 2012;53:255-263
 60. Edeline J, Lenoir L, Boudjema K, Rolland Y, Boulic A, Le Du F, et al. Volumetric changes after (⁹⁰)y radioembolization for hepatocellular carcinoma in cirrhosis: an option to portal vein embolization in a preoperative setting? *Ann Surg Oncol* 2013;20:2518-2525
 61. Riaz A, Gates VL, Atassi B, Lewandowski RJ, Mulcahy MF, Ryu RK, et al. Radiation segmentectomy: a novel approach to increase safety and efficacy of radioembolization. *Int J Radiat Oncol Biol Phys* 2011;79:163-171
 62. Lance C, McLennan G, Obuchowski N, Cheah G, Levitin A, Sands M, et al. Comparative analysis of the safety and efficacy of transcatheter arterial chemoembolization and yttrium-90 radioembolization in patients with unresectable hepatocellular carcinoma. *J Vasc Interv Radiol* 2011;22:1697-1705
 63. Moreno-Luna LE, Yang JD, Sanchez W, Paz-Fumagalli R, Harnois DM, Mettler TA, et al. Efficacy and safety of transarterial radioembolization versus chemoembolization in patients with hepatocellular carcinoma. *Cardiovasc Intervent Radiol* 2013;36:714-723
 64. El Fouly A, Ertle J, El Dorry A, Shaker MK, Dechène A, Abdella H, et al. In intermediate stage hepatocellular carcinoma: radioembolization with yttrium 90 or chemoembolization? *Liver Int* 2015;35:627-635
 65. Salem R, Gilbertsen M, Butt Z, Memon K, Vouche M, Hickey R, et al. Increased quality of life among hepatocellular carcinoma patients treated with radioembolization, compared with chemoembolization. *Clin Gastroenterol Hepatol* 2013;11:1358-1365.e1
 66. Llovet JM, Ricci S, Mazzaferro V, Hilgard P, Gane E, Blanc JF, et al. Sorafenib in advanced hepatocellular carcinoma. *N Engl J Med* 2008;359:378-390
 67. Cheng AL, Kang YK, Chen Z, Tsao CJ, Qin S, Kim JS, et al. Efficacy and safety of sorafenib in patients in the Asia-Pacific region with advanced hepatocellular carcinoma: a phase III randomised, double-blind, placebo-controlled trial. *Lancet Oncol* 2009;10:25-34
 68. Bruix J, Raoul JL, Sherman M, Mazzaferro V, Bolondi L, Craxi A, et al. Efficacy and safety of sorafenib in patients with advanced hepatocellular carcinoma: subanalyses of a phase III trial. *J Hepatol* 2012;57:821-829
 69. Pressiani T, Boni C, Rimassa L, Labianca R, Fagioli S, Salvagni S, et al. Sorafenib in patients with Child-Pugh class A and B advanced hepatocellular carcinoma: a prospective

- feasibility analysis. *Ann Oncol* 2013;24:406-411
70. Sangro B, Iñarrairaegui M, Bilbao JI. Radioembolization for hepatocellular carcinoma. *J Hepatol* 2012;56:464-473
 71. Grady ED. Internal radiation therapy of hepatic cancer. *Dis Colon Rectum* 1979;22:371-375
 72. Bal CS, Kumar A. Radionuclide therapy for hepatocellular carcinoma: indication, cost and efficacy. *Trop Gastroenterol* 2008;29:62-70
 73. Ariel IM. Treatment of inoperable primary pancreatic and liver cancer by the intra-arterial administration of radioactive isotopes (Y90 radiating microspheres). *Ann Surg* 1965;162:267-278
 74. Pons F, Varela M, Llovet JM. Staging systems in hepatocellular carcinoma. *HPB (Oxford)* 2005;7:35-41
 75. Leung TW, Lau WY, Ho SK, Ward SC, Chow JH, Chan MS, et al. Radiation pneumonitis after selective internal radiation treatment with intraarterial 90yttrium-microspheres for inoperable hepatic tumors. *Int J Radiat Oncol Biol Phys* 1995;33:919-924
 76. Carr BI, Kondragunta V, Buch SC, Branch RA. Therapeutic equivalence in survival for hepatic arterial chemoembolization and yttrium 90 microsphere treatments in unresectable hepatocellular carcinoma: a two-cohort study. *Cancer* 2010;116:1305-1314
 77. Woodall CE, Scoggins CR, Ellis SF, Tatum CM, Hahl MJ, Ravindra KV, et al. Is selective internal radioembolization safe and effective for patients with inoperable hepatocellular carcinoma and venous thrombosis? *J Am Coll Surg* 2009;208:375-382
 78. Goin J, Dancey JE, Roberts C, Sickles CJ, Leung DA, Soulen MC. Comparison of postembolization syndrome in the treatment of patients with unresectable hepatocellular carcinoma: trans-catheter chemo-embolization versus yttrium-90 glass microspheres. *World J Nucl Med* 2004;3:49-56
 79. Dancey JE, Shepherd FA, Paul K, Sniderman KW, Houle S, Gabrys J, et al. Treatment of nonresectable hepatocellular carcinoma with intrahepatic 90Y-microspheres. *J Nucl Med* 2000;41:1673-1681
 80. Geschwind JF, Salem R, Carr BI, Soulen MC, Thurston KG, Goin KA, et al. Yttrium-90 microspheres for the treatment of hepatocellular carcinoma. *Gastroenterology* 2004;127(5 Suppl 1):S194-S205
 81. Goin JE, Salem R, Carr BI, Dancey JE, Soulen MC, Geschwind JF, et al. Treatment of unresectable hepatocellular carcinoma with intrahepatic yttrium 90 microspheres: a risk-stratification analysis. *J Vasc Interv Radiol* 2005;16(2 Pt 1):195-203
 82. Goin JE, Salem R, Carr BI, Dancey JE, Soulen MC, Geschwind JF, et al. Treatment of unresectable hepatocellular carcinoma with intrahepatic yttrium 90 microspheres: factors associated with liver toxicities. *J Vasc Interv Radiol* 2005;16(2 Pt 1):205-213