

Transarterial chemoembolization in patients with hepatocellular carcinoma: Predictors of survival

Summit Sawhney BSc¹, Aldo J Montano-Loza MD², Peter Salat MD FRCPC¹, Mairin McCarthy MBBCh FRCR MRCP¹, Norman Kneteman MD FRCPC³, Judith Meza-Junco MD⁴, Richard Owen MBBCh FRCR MRCP¹

S Sawhney, AJ Montano-Loza, P Salat, et al. Transarterial chemoembolization in patients with hepatocellular carcinoma: Predictors of survival. *Can J Gastroenterol* 2011;25(8):426-432.

BACKGROUND: Transarterial chemoembolization (TACE) is the mainstay of management for patients with hepatocellular carcinoma who are not suitable for curative treatments.

OBJECTIVE: To determine factors associated with mortality after the first TACE procedure.

METHODS: From January 2004 to May 2008, 60 patients underwent TACE as treatment for hepatocellular carcinoma. Clinical and biochemical parameters before TACE, and response after TACE, were evaluated with conventional classifications (WHO, Response Evaluation Criteria in Solid Tumors [RECIST], and European Association for the Study of the Liver [EASL] criteria) and with one-, two- and three-dimensional assessment.

RESULTS: Overall median survival after the first TACE procedure was 17.1±3.4 months. According to Cox regression analysis, having an alpha-fetoprotein level of greater than 200 ng/mL (HR 2.35 [P=0.02]) and a Model for End-stage Liver Disease (MELD) score of greater than 10 (HR 4.19 [P=0.001]) were associated with higher risk of mortality; whereas reduction in tumour size measured in one dimension (HR 0.96 [P=0.005]), two dimensions (HR 0.98 [P=0.004]) and three dimensions (HR 0.98 [P=0.001]) was associated with lower risk of mortality. Moreover, reduction in tumour size by 3% or more assessed in one, two or three dimensions was associated with lower risk of mortality (HR 0.48 [P=0.04]; HR 0.36 [P=0.01]; HR 0.31 [P=0.003], respectively). The three conventional classifications were not useful for predicting mortality (WHO: HR 1.07 [P=0.9]; RECIST: HR 0.94 [P=0.9]; EASL: HR 0.94 [P=0.9]).

CONCLUSIONS: Having an alpha-fetoprotein level of greater than 200 ng/mL and a MELD score of greater than 10 before undergoing TACE was associated with a greater risk of mortality. Conventional classifications of response were not useful for predicting mortality. Reduction in tumour size after the first TACE procedure was associated with better survival, primarily if patients had more than a 3% reduction in tumour size assessed by three-dimensional measurement.

Key Words: Hepatocellular carcinoma; Survival; Transarterial chemoembolization; Tumour size

Hepatocellular carcinoma (HCC) is the fifth most common cancer in the world, and the third most common cause of death (1). Prognosis in the majority of cases remains poor because of either advanced tumour staging or severe hepatic dysfunction. The best option for these patients is liver transplantation, with a five-year survival rate of up to 80% (2,3); however, for various reasons, most patients cannot receive a transplant. Surgical resection or percutaneous ablative techniques are considered to be curative treatments for HCC (4), but are only suitable for those with localized tumours or good underlying synthetic liver function. The majority of patients at

La chimioembolisation transartérielle chez les patients ayant un carcinome hépatocellulaire : les prédicteurs de la survie

HISTORIQUE : La chimioembolisation transartérielle (CETA) constitue le mode de prise en charge habituel des patients ayant un carcinome hépatocellulaire non admissibles à un traitement curatif.

OBJECTIF : Déterminer les facteurs associés à la mortalité après la première intervention de CETA.

MÉTHODOLOGIE : De janvier 2004 à mai 2008, 60 patients ont subi une CETA pour traiter un carcinome hépatocellulaire. Les chercheurs ont évalué les paramètres cliniques et biochimiques avant la CETA et la réponse après la CETA au moyen de classifications classiques (OMS, *Response Evaluation Criteria in Solid Tumors* [RECIST] et critères de l'*European Association for the Study of the Liver* [EASL]) et d'une évaluation unidimensionnelle, bidimensionnelle et tridimensionnelle.

RÉSULTATS : La survie médiane moyenne après la première intervention de CETA était de 17,1±3,4 mois. D'après l'analyse de régression de Cox, un taux de foetoprotéine alpha supérieur à 200 ng/mL (RR 2,35 [P=0,02]) et un indice de modèle de maladie hépatique en phase terminale (MMHT) supérieur à 10 (RR 4,19 [P=0,001]) s'associaient à un risque de mortalité plus élevé, tandis que la réduction de la dimension de la tumeur mesurée en une dimension (RR 0,96 [P=0,005]), en deux dimensions (RR 0,98 [P=0,004]) et en trois dimensions (RR 0,98 [P=0,001]) s'associait à un risque de mortalité plus faible. De plus, une réduction d'au moins 3 % de la dimension de la tumeur évaluée en une, deux ou trois dimensions s'associait à un plus faible risque de mortalité (RR 0,48 [P=0,04]; RR 0,36 [P=0,01]; RR 0,31 [P=0,003], respectivement). Les trois classifications classiques n'étaient pas utiles pour prédire la mortalité (OMS : RR 1,07 [P=0,9]; RECIST : RR 0,94 [P=0,9]; EASL : RR 0,94 [P=0,9]).

CONCLUSIONS : Un taux de foetoprotéine alpha supérieur à 200 ng/mL et un indice de MMHT supérieur à 10 avant la CETA s'associait à un plus fort taux de mortalité. Les classifications classiques de réponse n'étaient pas utiles pour prédire la mortalité. La réduction de la dimension de la tumeur après la première intervention de CETA s'associait à une meilleure survie, surtout si la réduction de la dimension de la tumeur des patients était supérieure à 3 %, d'après une mesure tridimensionnelle.

presentation are palliative, unless they can be downstaged to a curative option or cannot be treated with current techniques.

Until quite recently, there was no level 3 evidence of survival benefit in patients with advanced HCC treated with transarterial chemoembolization (TACE), and its use was controversial in patients who were not suitable for curative therapies. Two randomized clinical trials in 2002 (5,6) and a meta-analysis in 2003 (7) showed level 3 evidence of survival benefits of TACE compared with best support care.

The evaluation of response to TACE is transcendental because an objective response may become a surrogate marker for improved

¹Department of Radiology and Diagnostic Imaging; ²Division of Gastroenterology & Hepatology; ³Department of Surgery, University of Alberta Hospital;

⁴Department of Medical Oncology, Cross Cancer Institute, Edmonton, Alberta

Correspondence: Dr Richard Owen, Department of Radiology and Diagnostic Imaging, Walter Mackenzie Health Sciences Centre, University of Alberta, 8440 112th Street, Edmonton, Alberta T6G 2B7. Telephone 780-407-1210, fax 780-407-8628, e-mail drrichardowen@tbwifl.ca

Received for publication June 24, 2010. Accepted January 24, 2011

TABLE 1
Criteria for response assessment after transarterial chemoembolization in patients with hepatocellular carcinoma

Type of tumour response	WHO criteria	RECIST	EASL criteria
Complete	Disappearance of all target lesions	Disappearance of all target lesions	Absence of enhanced tumour areas, reflecting complete tumour necrosis
Partial	≥50% decrease	≥30% decrease	≥50% decrease of enhanced areas
Stable disease	≤50% decrease and ≤25% increase	≤30% decrease and ≤20% increase	≤50% decrease and ≤25% increase of enhanced areas
Progressive disease	≥25% increase or new lesions	≥20% increase or new lesions	≥25% increase of enhanced lesions or new enhanced lesions

EASL European Association for the Study of the Liver; RECIST Response Evaluation Criteria in Solid Tumors

survival. Criteria have been developed for response assessment, including WHO (8,9), the Response Evaluation Criteria in Solid Tumors (RECIST) (10,11) and the European Association for the Study of the Liver (EASL) criteria (12) (Table 1). However, these criteria have limitations; for example, WHO and RECIST evaluate only bidimensional and unidimensional tumour measurements, and disregard the extent of necrosis, whereas the EASL criteria evaluate response by measuring the extent of tumour necrosis; however, there are a lack of data supporting correlations with improving survival.

In the present retrospective analysis of prospectively acquired data, we analyzed factors associated with mortality after the first TACE procedure. We also determined the value of conventional classifications (ie, WHO, RECIST and EASL) to measure response and predict survival after TACE, and analyzed the clinical utility of the new values of response, using one-, two- and three-dimensional tumour measurements expressed as percentages to predict outcome.

METHODS

Study population

Between January 2004 and May 2008, 60 consecutive patients with HCC underwent TACE at the University of Alberta Hospital (Edmonton, Alberta). HCC was diagnosed using characteristic non-invasive imaging criteria (computerized axial tomography [CAT] scan or magnetic resonance imaging [MRI]) in 37 patients, and with liver biopsy in 23 patients. Each patient was followed after the first TACE procedure until specific end points were reached, which included death, liver transplant, resection or treatment failure (development of metastatic disease post-TACE and/or deterioration in liver function post-TACE). The present study was approved by the local ethics committee.

Chemoembolization criteria

Inclusion criteria: Preprocedure requirements: assumed to not be a candidate for 'curative treatment' (surgery or radiofrequency ablation, percutaneous ethanol injection) due to location, size, etc. Biopsy-proven or noninvasive criteria for HCC, preprocedure dynamic cross-sectional imaging (CAT scan or MRI), and tumours greater than 2 cm in diameter, bilirubin level lower than 50 µmol/L, Model for End-stage Liver Disease (MELD) score of less than 30, Child-Pugh class A or B, and Eastern Cooperative Oncology Group (ECOG) score of 0 to 1.

Exclusion criteria: Very advanced disease (multifocal tumour, large mass greater than 8 cm in diameter, etc), more than 50% liver volume, portal vein invasion, metastasis, ECOG score of greater than 1, life expectancy of less than three months, severe liver disease with MELD score of greater than 30, encephalopathy, or multigated acquisition (MUGA) scan ejection fraction of less than 60%.

Chemoembolization protocol

A MUGA scan was performed before the initial chemoembolization, after the fourth course of TACE, and after six or more courses of TACE to evaluate heart function. In patients with significant cardiac dysfunction, mitomycin or a combination of doxorubicin and mitomycin was used instead of doxorubicin alone. A triphasic abdominal CAT scan (liver mass protocol) was also performed before each first TACE, and subsequently after every second treatment to assess tumour

response. Prophylactic antibiotics were given to all patients: cefazolin 1 g intravenously (IV) every 8 h (three doses) and metronidazole 500 mg IV every 6 h (four doses). All patients also received diphenhydramine 50 mg IV, ondansetron 8 mg IV every 8 h (three doses) and dexamethasone 8 mg IV every 8 h (three doses). Fentanyl 50 µg to 150 µg IV or morphine 5 mg to 10 mg IV, and midazolam 1 mg to 5 mg IV were used to achieve moderate sedation.

A 5 Fr sheath was placed in the common femoral artery with selective catheterization of the superior mesenteric (SMA) and celiac axis arteries. An angiogram of the SMA was performed with delayed portal vein imaging. Further angiography of the celiac axis was performed to characterize the hepatic arterial supply. Following the initial arterial assessment, the catheter was advanced into the lobar or segmental hepatic artery supplying the tumour. If the initial 4 Fr or 5 Fr diagnostic catheter could be advanced into the optimal position, it was used for the TACE infusion and, in cases in which more selective catheterization was required, a 2.3 Fr or 2.5 Fr microcatheter was used (Rapid Transit, Cordis Endovascular Systems, USA, or Mira-Flex, Cook Medical, USA). In general, the TACE infusion point was chosen to enable selective tumour embolization. However, if there were separate arterial feeders, or a discrete blood supply could not be identified, a right or left hepatic arterial infusion was performed. If concern regarding hepatic synthetic function was present, a selective approach was chosen. Once the lesion and its blood supply were identified, an emulsion consisting of doxorubicin (50 mg in 10 mL of normal saline) and lipiodol (5 mL to 10 mL) was injected under fluoroscopic guidance into the arterial supply of the tumour. If the bilirubin level was elevated by more than 35 µmol/L, less than 10 mL of lipiodol was used. If there was continuing arterial supply to the tumour after administration of the lipiodol/doxorubicin emulsion, embolization of the feeding vessel to near stasis was achieved using polyvinyl alcohol particles 350 µm to 500 µm in size (Boston Scientific, USA).

Following TACE, pain medication was given as needed to maintain pain control. Second and subsequent treatments were scheduled at six-week intervals for the first four treatments, and at three-month intervals (up to a maximum of eight courses) for subsequent treatments. End points for treatment were as follows: contraindications to further TACE (deteriorating cardiac function, deteriorating hepatic function); progression of disease despite treatment; tumour regression or stabilization after a minimum of two treatments; development of metastasis; surgical resection; transplantation; or death. MUGA scans were repeated after the second, fourth, sixth and subsequent TACE procedures. Contraindications to TACE were the following: advanced liver disease (bilirubin level of greater than 50 µmol/L, super selective embolization only in cases for which the bilirubin level was greater than 30 µmol/L); Child-Pugh class C, MELD score of less than 30 and/or development of encephalopathy; renal failure (serum creatinine level of greater than 250 µmol/L); coagulation abnormality (international normalized ratio of greater than 1.5; platelet levels lower than $50 \times 10^9/L$); and evidence of extrahepatic spread of disease. Three qualified interventional radiologists performed the TACE procedures, with the majority completed by two interventional radiologists (28 and 30 cases each). On average, each patient underwent 2.5 TACE treatments (range one to eight).

TABLE 2
Clinical and laboratory features at accession in patients undergoing transarterial chemoembolization (n=60)

Clinical/biochemical	Mean ± SEM or n	Range or percentage
Features at presentation		
Age, years	60±2	24–91
Male:female ratio	50:10	82:18
Etiology of cirrhosis		
Alcohol	11	18
HCV	11	18
Alcohol + HCV	9	15
HBV	13	22
Cryptogenic	11	18
Others*	5	8
Child-Pugh class, A/B	39/21	65/35
MELD score, points	9±0.4	4–17
BCLC (A/B/C)	10/46/4	17/77/6
ALT (normal <50 U/L)	89±9	19–365
AST (normal <38 U/L)	94±9	11–292
AP (normal 30–130 U/L)	158±18	45–846
Bilirubin (<20 µmol/L)	23±2	8–105
International normalized ratio	1.2±0.5	0.9–3.0
Albumin (normal 35–50 g/L)	36±1	16–50
Creatinine (normal <125 µmol/L)	83±3	33–165
AFP (normal <10 ng/mL)	2664±1825	1–103,225
AFP >400 ng/mL	16	27
Total tumour size (1D), cm	6.9±0.6	1.3–20.4
Total tumour size (2D), cm ²	51.5±8.5	1.4–318.2
Total tumour size (3D), cm ³	293.9±71.8	1–3053.6
Tumours, n	2±0.2	1–10

*Cryptogenic primary biliary cirrhosis. 1D One dimension; 2D Two dimensions; 3D Three dimensions; AFP Alpha-fetoprotein; ALT Alanine aminotransferase; AP Alkaline phosphatase; AST Aspartate aminotransferase; BCLC Barcelona Clinic Liver Cancer staging classification; HBV Hepatitis B virus; HCV Hepatitis C virus; MELD Model for End-stage Liver Disease

Assessment of response following TACE

Tumour response was evaluated one to three months after the first TACE procedure. The evaluation of tumour response with conventional classifications, such as WHO, RECIST and EASL, was determined in 53 of 60 patients who underwent a CAT scan after the first TACE, and they were classified as complete response, partial response, stable disease or progressive disease. Then, the differences in the tumour size between the pre-TACE and post-TACE CAT scan were measured, assessed in one, two and three dimensions, with the differences expressed as a percentage. The three-dimensional assessment was performed as a surrogate of the total tumour volume, and was calculated as the sum of the volumes of all tumours [$4/3 \pi r^3$, where r is the maximum radius of each HCC], as described elsewhere (13). All of the imaging studies were evaluated by only one radiologist (RO).

Statistical analyses

Fisher's exact probability test was used to compare categorical variables, while the unpaired t test was used to compare differences in the means of continuous variables. Nonparametric variables in independent samples were compared using the Mann-Whitney U test. Survival was calculated using the Kaplan-Meier method, and they were compared using the log-rank (Mantel-Cox) test (14). Patients were followed after the first TACE procedure and were censored at the time of death, liver transplantation, surgical resection or at the time of their most recent visit. Prognostic factors for mortality were analyzed by Cox uni- and multivariate analysis (15). A ROC curve was plotted to measure how the response, via one-dimensional, two-dimensional and

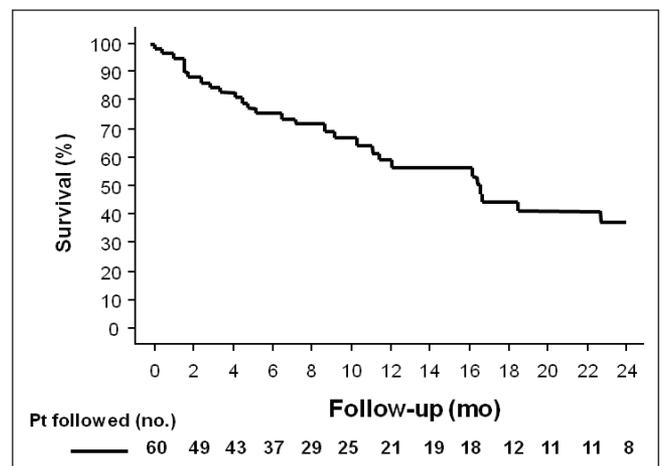


Figure 1 Overall probability of survival after the first transarterial chemoembolization procedure in patients (Pt) with hepatocellular carcinoma. The six-month and one-year probability of survival was 74% and 60%, respectively. mo Months; no Number

three-dimensional measurements after TACE (expressed as percentages) was able to discriminate mortality in patients with HCC. The validity of the model was measured by the area under the ROC curve (AUC) (16). Furthermore, ROC curves were used to determine the one-, two- and three-dimensional measurements expressed as percentage values that discriminated best between patients who died during follow-up after TACE and those who survived. Data are presented as the mean ± SEM in the tables and text.

RESULTS

Features at accession in patients receiving TACE

The mean age of the study population was 60±2 years (range 24 to 91 years); 50 patients (83%) were men. Table 2 summarizes the cohort characteristics. All patients were cirrhotic, the majority of which were secondary to a combination of chronic active hepatitis C and alcohol abuse (n=31 [52%]). Liver function was preserved in the majority: 39 patients (65%) were Child-Pugh class A, and mean MELD score before TACE was 9±0.4 points. According to the Barcelona Clinic Liver Cancer (BCLC) staging classification, 10 patients were in stage A (17%), 46 were in stage B (77%) and four were in stage C (6%). The basal size of the tumour, measured in one, two and three dimensions, and the number of tumours are shown in Table 2.

Survival after the first TACE procedure in patients with HCC

After the first TACE procedure, during a mean follow-up of 11±1 months (median eight months, range one to 38 months), 31 patients died, 14 patients received a liver transplant, 11 patients were alive, three patients were lost during follow-up and one patient underwent surgical resection. The overall median survival time after first TACE was 17.1±3.4 months (95% CI 10.5 to 23.7); the probability of survival at six months was 74%, and at one year was 60% (Figure 1).

Pre-TACE features associated with mortality according to univariate and multivariate Cox analysis

According to univariate Cox regression analysis of the pre-TACE variables, the alpha-fetoprotein (AFP) level (HR 1.01 [95% CI 1.00 to 1.02]; P=0.03), an AFP level of greater than 200 ng/mL (HR 2.35 [95% CI 1.14 to 4.87]; P=0.02) and a MELD score of greater than 10 (HR 4.19 [95% CI 1.79 to 9.78]; P=0.001) were significantly associated with a higher risk of mortality (Table 3). No other significant association was found with clinical and laboratory features before TACE and the risk of mortality, including age, sex, hepatitis viral infection, and the number and size of the tumour (Table 3).

TABLE 3
Pretransarterial chemoembolization features associated with mortality according to Cox regression analysis in patients with hepatocellular carcinoma

	Mortality (n=31)	Alive (n=29)	HR	95% CI	P
Features associated in the univariate analysis					
Age at diagnosis, years	61±2	58±2	0.41	0.98–1.04	0.4
Men:women, n:n	26:5	24:5	1.26	0.48–3.29	0.6
Viral hepatitis (B and C), n (%)	13 (42)	20 (69)	0.33	0.34–1.44	0.3
Alanine aminotransferase (normal <50 U/L)	80±14	79±11	0.99	0.99–1.003	0.4
Aspartate aminotransferase (normal <38 U/L)	95±14	93±12	1.00	0.99–1.006	0.9
Alkaline phosphatase (normal 30–130 U/L)	150±14	143±24	1.002	0.99–1.006	0.5
Bilirubin (normal <20 µmol/L)	23±3	24±3	1.02	0.99–1.05	0.1
International normalized ratio (normal 0.8–1.2)	1.2±0.02	1.2±0.08	0.58	0.17–1.93	0.4
Albumin (normal 35–50 g/L)	37±1	35±1	0.98	0.92–1.04	0.5
Creatinine (normal <125 mol/L)	89±5	76±3	1.01	0.99–1.03	0.08
Alpha-fetoprotein (normal <10 ng/mL)	3891±3315	925±626	1.01	1.00–1.02	0.03
Alpha fetoprotein >200 ng/mL	14 (45)	5 (17)	2.35	1.14–4.87	0.02
Model for End-stage Liver Disease score	8.9±0.6	8.2±0.6	1.11	0.98–1.26	0.09
Model for End-stage Liver Disease score >10, n (%)	10 (32)	5 (17)	4.19	1.79–9.78	0.001
Child-Turcotte-Pugh class, A:B, n	19:12	20:9	0.69	0.34–1.44	0.3
BCLC, A:B:C, n	2:26:3	8:20:1	2.58	0.95–7.03	0.06
Tumours, n	2.0±0.3	2.2±0.4	1.04	0.83–1.31	0.7
Total tumour size (1D), cm	7.9±0.9	5.6±0.7	1.07	0.98–1.16	0.1
Total tumour size (2D), cm ²	63±13	38±10	1.005	0.99–1.011	0.08
Total tumour size (3D), cm ³	381±115	194±76	1.001	1.00–1.001	0.06
Features associated in the multivariate analysis					
Alpha-fetoprotein (normal <10 ng/mL)	3891±3315	925±626	1.01	1.00–1.02	0.008
Model for End-stage Liver Disease score >10, n (%)	10 (32)	5 (17)	4.55	1.92–10.78	0.001

Data presented as mean ± SEM unless otherwise indicated. 1D One dimension; 2D Two dimensions; 3D Three dimensions; BCLC Barcelona Clinic Liver Cancer staging classification

In the multivariate Cox analysis, only the AFP level (HR 1.01 [95% CI 1.00 to 1.02]; P=0.008) and a MELD score of greater than 10 (HR 4.55 [95% CI 1.92 to 10.78]; P=0.001) were independently associated with a higher risk of mortality (Table 3).

Evaluation of tumour response after TACE

For the analysis of tumour response after the first TACE, seven patients were excluded because they did not undergo a subsequent CAT scan after TACE (three patients transplanted post-TACE [transplantation mean 101±41 days following TACE; median 100 days; range 31 to 173 days]; three died from treatment complications [mean 37±10 days following TACE; median 37 days; range 19 to 54 days]; and one patient was lost during follow-up).

Using the three conventional assessments of tumour response, none of the patients achieved a complete response classification (WHO, RECIST and EASL), and most patients exhibited stable disease (Table 4). According to WHO criteria, six patients achieved partial response, 45 stable disease and two progressive disease. According to the RECIST criteria, eight patients achieved partial response, 41 stable disease and four progressive disease. According to the EASL criteria, eight patients achieved partial response, 43 stable disease and two progressive disease (Table 4).

Performance of one-, two- and three-dimensional measurements of response after TACE

The best parameters to discriminate mortality in patients with HCC after TACE were the two- and three-dimensional reduction measurements in percentage (difference between pre-TACE CAT scan and post-TACE CAT scan), with an AUC of 0.69 (95% CI 0.54 to 0.84; P=0.02) for two-dimensional measurement, and an AUC of 0.72 (95% CI 0.58 to 0.86; P=0.009) for three-dimensional measurement (Figure 2). The one-dimensional response was less accurate for mortality discrimination (AUC 0.65 [95% CI 0.50 to 0.81]; P=0.07) (Figure 2). The best

TABLE 4
Tumour response assessment after transarterial chemoembolization in patients with hepatocellular carcinoma

Type of tumour response	Criteria		
	WHO	RECIST	EASL
Complete	0 (0)	0 (0)	0 (0)
Partial	8 (15)	6 (11)	8 (15)
Stable disease	41 (77)	45 (85)	43 (81)
Progressive disease	4 (8)	2 (4)	2 (4)

Data presented as n (%). EASL European Association for the Study of the Liver; RECIST Response Evaluation Criteria in Solid Tumors

cut-off value for one-, two- and three-dimensional assessment for discrimination of mortality after TACE was a reduction of 3% or more.

Post-TACE features associated with mortality according to univariate and multivariate Cox analysis

According to univariate Cox analysis of the post-TACE variables, reduction in the tumour size measured in one dimension (HR 0.96 [95% CI 0.93 to 0.99]; P=0.005), two dimensions (HR 0.98 [95% CI 0.97 to 0.99]; P=0.004) and three dimensions (HR 0.98 [95% CI 0.97 to 0.99]; P=0.001) were significantly associated with lower mortality after TACE (Table 5).

Moreover, if patients experienced a reduction in the size of the tumour of 3% or more assessed in one, two or three dimensions, they had a lower risk of mortality (HR 0.48 [95% CI 0.21 to 0.95]; P=0.04; HR 0.36 [95% CI 0.17 to 0.80]; P=0.01; and HR 0.31 [95% CI 0.14 to 0.67]; P=0.003, respectively) (Table 5).

Conventional classifications (ie, WHO, RECIST and EASL) were not useful in predicting mortality after the first TACE when patients

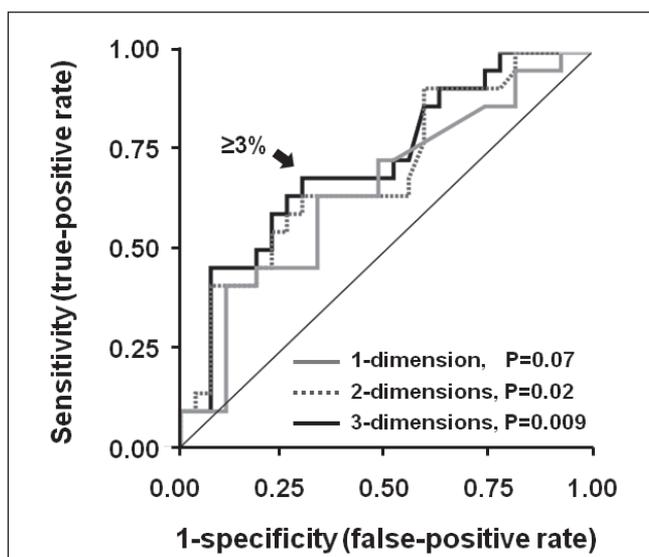


Figure 2) ROC curves for predicting mortality after transarterial chemoembolization in patients with hepatocellular carcinoma. The area under the curve (AUC) for two-dimensional tumour size reduction was 0.69 (95% CI 0.54 to 0.84; $P=0.02$), and for three-dimensional measurement, the AUC was 0.72 (95% CI 0.58 to 0.86; $P=0.009$). The one-dimensional measurement was less accurate for six-month mortality discrimination (AUC 0.65 [95% CI 0.50 to 0.81]; $P=0.07$)

with partial response and stable disease were compared with patients with progressive disease, or when comparing patients with partial response to patients with stable and progressive disease (Table 5).

In the multivariate Cox analysis, only the response measured in three dimensions was independently associated with a reduction in mortality risk after TACE (HR 0.94 [95% CI 0.89 to 0.99]; $P=0.03$) (Table 5).

Overall survival after TACE according to WHO, RECIST and EASL criteria

Using the WHO classification, median survival in partial response and stable disease ($n=49$) was 17 ± 3 months (95% CI 11 to 24), and in progressive disease ($n=4$) was 17 ± 12 months (95% CI 1 to 39; $P=0.9$

TABLE 5
Post-transarterial chemoembolization (TACE) features associated with mortality according to Cox regression analysis in patients with hepatocellular carcinoma

	Mortality (n=28)	Alive (n=25)	HR	95% CI	P
Features associated in the univariate analysis					
WHO: PR + SD versus PD	26:2	23:2	1.07	0.25–4.59	0.9
RECIST: PR + SD versus PD	27:1	24:1	0.94	0.13–7.04	0.9
EASL: PR + SD versus PD	27:1	24:1	0.94	0.13–7.04	0.9
WHO: PR versus SD + PD	4:24	4:21	1.36	0.47–3.94	0.6
RECIST: PR versus SD + PD	4:24	2:23	2.98	0.98–9.04	0.06
EASL: PR versus SD + PD	5:23	3:22	1.08	0.41–2.86	0.9
Response with 1D, cm	3.6 ± 3.4	15.7 ± 4.8	0.96	0.93–0.99	0.005
Response with 2D, cm ²	5.5 ± 5.9	28.5 ± 6.8	0.98	0.97–0.99	0.004
Response with 3D, cm ³	4.6 ± 8.3	42 ± 10	0.98	0.97–0.99	0.001
Reduction >3% with 1D	13 (46)	16 (64)	0.48	0.21–0.95	0.04
Reduction >3% with 2D	14 (50)	17 (68)	0.36	0.17–0.80	0.01
Reduction >3% with 3D	15 (54)	19 (76)	0.31	0.14–0.67	0.003
Features associated in the multivariate analysis					
Response with 3D, cm ³	4.6 ± 8.3	42 ± 10	0.94	0.89–0.99	0.03

Data presented as n:n, mean \pm SEM or n (%). Percentage of response measured as difference in size between the pre-TACE computed tomography scan and the first computed tomography scan post-TACE, expressed as percentage. 1D One dimension; 2D Two dimensions; 3D Three dimensions; EASL European Association for the Study of the Liver criteria; PD Progressive disease; PR Partial response; RECIST Response Evaluation Criteria in Solid Tumors; SD Stable disease

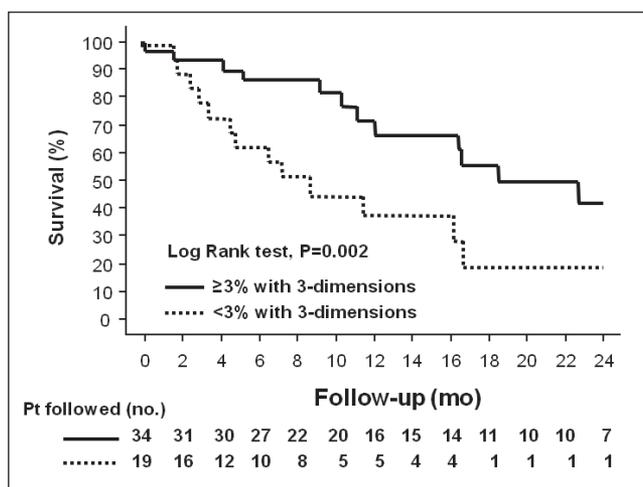


Figure 3) Kaplan-Meier survival curves of patients (Pt) with hepatocellular carcinoma after transarterial chemoembolization, separated into those who achieved a reduction in tumour size of 3% or more assessed in three dimensions compared with patients who did not ($P=0.002$ [log-rank test]). mo Months; no Number

[log-rank test]). Similar results were found using RECIST: partial response and stable disease ($n=51$) was 17 ± 3 months (95% CI 12 to 23), and in progressive disease ($n=2$) was 12 ± 7 months (95% CI 1 to 26; $P=0.9$ [log-rank test]); and EASL: partial response and stable disease ($n=51$) was 27 ± 3 months (95% CI 12 to 23 months), and in progressive disease ($n=2$) was 12 ± 7 months (95% CI 1 to 26; $P=0.9$ [log-rank test]).

Overall survival after TACE according to more than a 3% reduction in tumour size

Survival was significantly improved in patients who achieved a reduction in tumour size of 3% or more assessed in one, two and three dimensions, compared with patients who did not achieve this end point. This difference was greater if the tumour was assessed in three dimensions: median survival for patients who achieved a reduction in tumour size was 24 ± 5 months versus 8 ± 3 months ($P=0.002$ [log-rank test]) (Figure 3).

after TACE, we assumed HCC to be a spherical tumour, which in many cases, is not correct. Nevertheless, it seems that this assumption is more accurate than using one- and two-dimensional analysis in predicting survival, and more time efficient than computed tomography volumetry in assessing three-dimensional response. Finally, we only analyzed factors before and after the first TACE procedure, and most patients underwent more than one TACE (mean 2.5 ± 0.2 , range 1 to 8). However, despite these limitations, we believe that our study should encourage future studies to evaluate more effective ways to assess tumour response after locoregional treatments in patients with HCC.

SUMMARY

Mean survival after TACE in patients with noncurative HCC was approximately 17 months. Having an AFP level of greater than 200 ng/mL and a MELD score of greater than 10 before TACE was associated with a greater risk of mortality. Conventional classifications of response, such as WHO, RECIST and EASL, are not useful for predicting mortality. Reduction in tumour size after the first TACE procedure was associated with improved survival, primarily if patients experienced a reduction of more than 3% in tumour size assessed using three-dimensional measurement.

DISCLOSURE: This study was presented, in part, at the Annual Meeting of the American College of Gastroenterology in San Diego, California, on October 26, 2009.

REFERENCES

- Parkin DM, Bray F, Ferlay J, Pisani P. Estimating the world cancer burden: Globocan 2000. *Int J Cancer* 2001;94:153-6.
- Mazzaferro V, Regalia E, Doci R, et al. Liver transplantation for the treatment of small hepatocellular carcinomas in patients with cirrhosis. *N Engl J Med* 1996;334:693-9.
- Yao FY, Ferrell L, Bass NM, et al. Liver transplantation for hepatocellular carcinoma: Expansion of the tumor size limits does not adversely impact survival. *Hepatology* 2001;6:1394-403.
- Llovet JM, Burroughs A, Bruix J. Hepatocellular carcinoma. *Lancet* 2003;362:1907-17.
- Lo CM, Ngan H, Tso WK, et al. Randomized controlled trial of transarterial lipiodol chemoembolization for unresectable hepatocellular carcinoma. *Hepatology* 2002;35:1164-71.
- Llovet JM, Real ML, Montana X, et al. Arterial embolisation or chemoembolisation versus symptomatic treatment in patients with unresectable hepatocellular carcinoma: A randomized controlled trial. *Lancet* 2002;359:1734-9.
- Llovet JM, Bruix J. Systematic review of randomized trials for unresectable hepatocellular carcinoma: Chemoembolization improves survival. *Hepatology* 2003;37:429-42.
- Miller AB, Hoogstraten B, Staquet M, Winkler A. Reporting results of cancer treatment. *Cancer* 1981;47:207-14.
- World Health Organization. WHO Handbook for Reporting Results of Cancer Treatment. Offset Publication 48. Geneva: World Health Organization, 1979.
- Therasse P, Arbuck SG, Eisenhauer EA, 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;92:205-16.
- Therasse P, Eisenhauer EA, Verweij J. RECIST revisited: A review of validation studies on tumour assessment. *Eur J Cancer* 2006;42:1031-9.
- Bruix J, Sherman M, Llovet JM, 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;35:421-30.
- Toso C, Trotter J, Wei A, et al. Total tumor volume predicts risk of recurrence following liver transplantation in patients with hepatocellular carcinoma. *Liver Transpl* 2008;14:1107-15.
- Heinze G, Schemper M. Exact log-rank tests for unequal follow-up. *Biometrics* 2003;59:1151-7.
- Cox DR. Regression models and life tables (with discussion). *JR Stat Soc B* 1972;34:187-220.
- Hanley JA, McNeil BJ. The meaning and use of the area under a receiver operating characteristic (ROC) curve. *Radiology* 1982;143:29-36.
- Kasugai H, Kojima J, Tatsuta M, et al. Treatment of hepatocellular carcinoma by transcatheter arterial embolization combined with intraarterial infusion of a mixture of cisplatin and ethiodized oil. *Gastroenterology* 1989;97:965-71.
- Chang JM, Tzeng WS, Pan HB, et al. Transcatheter arterial embolization with or without cisplatin treatment of hepatocellular carcinoma. A randomized controlled study. *Cancer* 1994;74:2449-53.
- Bronowicki JP, Vetter D, Dumas F, et al. Transcatheter oily chemoembolization for hepatocellular carcinoma. A 4-year study of 127 French patients. *Cancer* 1994;74:16-24.
- Stefanini GF, Amorati P, Biselli M, et al. Efficacy of transarterial targeted treatments on survival of patients with hepatocellular carcinoma. An Italian experience. *Cancer* 1995;75:2427-34.
- Hatanaka Y, Yamashita Y, Takaha, et al. Unresectable hepatocellular carcinoma: Analysis of prognostic factors in transcatheter management. *Radiology* 1995;195:747-52.
- Chung JM, Park JH, Han JK, et al. Hepatic tumors: Predisposing factors for complications of transcatheter oily chemoembolization. *Radiology* 1996;198:33-40.
- Solomon B, Soulen MC, Baum RA, et al. Chemoembolization of hepatocellular carcinoma with cisplatin, doxorubicin, mitomycin-C, ethiodol, and polyvinyl alcohol: Prospective evaluation of response and survival in a US population. *J Vasc Interv Radiol* 1999;10:793-8.
- Takayasu K, Muramatsu Y, Maeda T, et al. Targeted transarterial oily chemoembolization for small foci of hepatocellular carcinoma using a unified helical CT and angiography system: Analysis of factors affecting local recurrence and survival rates. *Am J Roentgenol* 2001;176:681-8.
- Fornier A, Ayuso C, Varela M, et al. Evaluation of tumor response after locoregional therapies in hepatocellular carcinoma: Are response evaluation criteria in solid tumors reliable? *Cancer* 2009;115:616-23.
- Lencioni R, Llovet JM. Modified RECIST (mRECIST) assessment for hepatocellular carcinoma. *Semin Liver Dis* 2010;30:52-60.
- Toso C, Asthana S, Bigam DL, et al. Reassessing selection criteria prior to liver transplantation for hepatocellular carcinoma utilizing the Scientific Registry of Transplant Recipients database. *Hepatology* 2009;49:832-8.
- Riaz A, Ryu RK, Kulik LM, et al. Alpha-fetoprotein response after locoregional therapy for hepatocellular carcinoma: Oncologic marker of radiologic response, progression, and survival. *J Clin Oncol* 2009;27:5734-42.



Hindawi
Submit your manuscripts at
<http://www.hindawi.com>

