Small Steatotic HCC: A Radiological Variant Associated With Improved Outcome After Ablation

Margaux Hermida,1 Ancelin Preel,1 Eric Assenat,2 Lauranne Piron,1 Christophe Cassinotto,1 José Ursic-Bedoya,3 Chloé Guillot,1 Astrid Herrero,4 Fabrizio Panaro,4 Georges-Philippe Pageaux,3 and Boris Guiu1

Percutaneous thermal ablation is a validated treatment option for small hepatocellular carcinoma (HCC). Steatotic HCC can be reliably detected by magnetic resonance imaging. To determine the clinical relevance of this radiological variant, we included 235 patients (cirrhosis in 92.3%, classified Child-Pugh A in 97%) from a prospective database on percutaneous thermal ablation for <3 cm HCC. Among these patients, 52 (22.1%) had at least one steatotic HCC nodule. Nonalcoholic steatohepatitis was more frequent in patients with than without steatotic HCC (P = 0.057), whereas body mass index, diabetes mellitus, liver steatosis, and liver fat content did not differ between groups. Liver disease was less advanced in patients with than without steatotic HCC: lower total bilirubin (−2.1 µmol/L; P = 0.035), higher albumin (+0.8 g/L; P = 0.035), and lower Model for End-Stage Liver Disease score (−0.8; P = 0.014). Tumor phenotype was less aggressive in patients with steatotic HCC: lower alpha-fetoprotein (AFP) concentration (P = 0.019), less frequent AFP > 100 ng/mL (P = 0.045), and multifocality (P = 0.015). During the follow-up (median: 28.3 months), overall mortality (3.8% vs. 23.5%; P = 0.001) and HCC-specific mortality (0.0% vs. 14.2%; P = 0.002) rates were lower in patients with steatotic HCC. Early (<2 years) recurrence was also less frequent (32.7% vs. 49.2%; P = 0.041). The mean time to intrahepatic distant recurrence (16.4 vs. 9 months, P = 0.006) and the median time to recurrence and recurrence-free survival (32.4 vs. 18.6 months, P = 0.024 and 30.4 vs. 16.4 months, P = 0.018) were longer in patients with steatotic versus nonsteatotic HCC. The 3-year overall survival was 94.4% and 70.9% in steatotic and nonsteatotic HCC (P = 0.008). In multivariate analysis, steatotic HCC (hazard ratio = 0.12; P = 0.039) and AFP (HR=1.002; P < 0.001) independently predicted overall survival. Conclusion: Small steatotic HCC detected by magnetic resonance imaging is associated with a less aggressive tumor phenotype. In patients with such radiological variant, percutaneous thermal ablation results in improved outcome. (Hepatology Communications 2021;5:689-700).

Liver cancer is the sixth most common cancer and the third leading cause of death from cancer worldwide. Hepatocellular carcinoma (HCC) represents 80%-90% of all liver cancer types. For early-stage HCC, percutaneous thermal ablation (PTA) is a validated treatment option, together with surgical resection and liver transplantation. Specifically, PTA has become the first-line curative treatment of small (i.e., <3 cm) HCC due to its excellent tolerance, particularly in patients with portal hypertension or comorbidity. However, the rate of intrahepatic distant recurrence is high (i.e., 60%-80%)

**Abbreviations:** AFP, alpha-fetoprotein; ALBI, albumin-bilirubin; ASA, American Society of Anesthesiologists; BMI, body mass index; CI, confidence interval; CT, computed tomography; HCC, hepatocellular carcinoma; HR, hazard ratio; IDR, intrahepatic distant recurrence; IQR, interquartile range; ITT, intention-to-treat; MCI, Model for End-Stage Liver Disease; MRI, magnetic resonance imaging; NASH, nonalcoholic steatohepatitis; NLR, neutrophil-to-lymphocyte ratio; OS, overall survival; PLR, platelet-to-lymphocyte ratio; PTA, percutaneous thermal ablation; RFS, recurrence-free survival; SH-HCC, steatohepatitic HCC; SII, systemic immune-inflammation index; TTR, time to recurrence.

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conduct to provide much-needed data to (1) elucidate the mechanisms by which the outcome of patients with small steatotic HCC might be improved and (2) determine whether the detection of such radiological variant might be helpful in treatment decision making.

Therefore, the objective of this study was to explore the baseline characteristics of patients with small steatotic HCC detected by MRI, their outcome, and tumor recurrence after PTA compared with patients with nonsteatotic HCC.

**Materials and Methods**

This retrospective analysis was performed using data from a prospective database of patients who underwent PTA for HCC at our institution. This study was approved by our institutional review board (NCT03428321 [www.clinicaltrials.gov]), and written informed consent for the procedure and the prospective anonymized data collection was obtained from all patients.

Inclusion criteria were HCC diagnosed by histopathology or by imaging criteria, no extrahepatic metastasis or macrovascular invasion, tumor size ≤30 mm, World Health Organization performance status 0 or 1, prothrombin time ratio >50%, and platelet count higher than 50 G/L.

Exclusion criteria were follow-up <3 months, Child-Pugh class ≥B8, HCC nodule adjacent to the hepatic hilum, history of biliary-digestive anastomosis or endoscopic sphincterotomy, combined treatment with embolization or chemoembolization, and no baseline MRI.

Treatment was validated at a multidisciplinary meeting that included interventional radiologists, liver surgeons, oncologists, hepatologists, and radiation oncologists. PTA was considered as first-line treatment.
treatment for patients with Barcelona Clinic Liver Cancer (BCLC) 0-A HCC.(2)

All patients underwent contrast-enhanced dynamic MRI including chemical-shift gradient-echo imaging, within 1 month before PTA.

PATIENT AND TUMOR DATA

The following patient data were collected: age, sex, American Society of Anesthesiologists physical status score (assessed by the anesthesiologist), body mass index (BMI), diabetes mellitus, liver disease (cirrhosis was defined as typical hepatic dysmorphia on imaging, or by noninvasive evaluation of fibrosis, or by histological analysis of liver biopsy; patients were considered noncirrhotic based on liver biopsy or noninvasive evaluation of fibrosis; the others were considered undetermined), steatosis (defined as signal intensity loss on opposed-phase gradient-echo sequences at baseline MRI), liver fat content,(19) and Child-Pugh, Model for End-Stage Liver Disease (MELD), albumin-bilirubin (ALBI), and AFP scores.

Steatotic HCC was defined by two radiologists (5 and 15 years of experience in liver imaging, respectively) when signal intensity loss was noted on opposed-phase compared with in-phase gradient-echo images for at least one HCC nodule. HCC location also was recorded(18): dome, subcapsular, adjacent to large vessels, or adjacent to at-risk organs. The following inflammation-based scores were calculated: platelet-to-lymphocyte ratio (PLR), neutrophil-to-lymphocyte ratio (NLR), lymphocyte-to-monocyte ratio (LMR), and systemic immune-inflammation index (SII).(20)

PERCUTANEOUS THERMAL ABLATION

All procedures were done under general anesthesia with endotracheal intubation in a multimodality interventional suite. PTA was performed by four interventional radiologists (5-15 years of expertise in liver PTA) using a radiofrequency or a microwave device, depending on the operator’s choice. Ultrasonography was the first-line guidance modality, but computed tomography (CT) guidance was used after tumor tagging whenever necessary.(18) Contrast-enhanced CT (portal phase) was performed immediately after the procedure to evaluate the ablation zone (i.e., the area of low attenuation) and to detect postprocedural complications. In the case of incomplete ablation or insufficient margins, the ablation needle(s) was reinserted during the same procedure to achieve complete ablation.

FOLLOW-UP AND OUTCOME

Complications were recorded and classified according to the Society of Interventional Radiology (SIR) guidelines.(21) Technical success was defined as complete ablation of the target tumor(s)(21) on the immediate post-PTA CT images.

Patients were monitored by MRI (including dynamic acquisitions) at week 4-6 after PTA, and then every 3 months, and by chest CT scan every 6 months.

Complete ablation observed on the first follow-up MRI was considered as the primary treatment success. Secondary treatment success was treatment success observed only after a second PTA performed within 8 weeks after the first one.

Local tumor progression (LTP) was defined as any growing or enhanced tumor focus within or at the edge (direct contact) of the ablation zone, after complete ablation documented by at least one MRI. (22) Intrahepatic distant recurrence (IDR) was defined as any new HCC nodule in the liver, defined according to EASL criteria.(22)

STATISTICAL ANALYSIS

Normally distributed continuous variables were described using means ± SD, and nonnormally distributed continuous variables using medians and interquartile range (IQR). Categorical variables were compared with the Fischer’s exact test, and continuous variables with the two-sided t-test or Kruskal-Wallis test, as appropriate.

Time to recurrence (TTR) was defined as the interval between PTA and death, recurrence, or last follow-up. Recurrence-free survival (RFS) was defined as the time from PTA until the first recurrence, death, or last follow-up. OS was defined as the interval between PTA and death (any cause) or last follow-up. For TTR and survival analyses, patients who underwent liver transplantation were censored at transplantation date.
The median (and 95% confidence interval [CI]) follow-up was calculated using the reverse Kaplan-Meier method. Survival curves were estimated using the Kaplan-Meier method and compared with the log-rank test. Univariate and multivariate Cox proportional-hazards models of all potential baseline predictors were built to compute the hazard ratios (HRs) with their 95% CI. A robust variance estimator was used systematically. Log linearity was checked using fractional polynomials.

All analyses were performed with the Stata software version 16.1 (Stata Corporation, College Station, TX). A P value < 0.05 was considered significant.

Results

STUDY POPULATION

Between January 2015 and November 2019, 279 consecutive patients underwent PTA for HCC at our institution. Forty-four patients were excluded due to (1) absence of baseline liver MRI (n = 27), (2) combined treatment with embolization (n = 9), (3) metastatic progression discovered at PTA day (n = 1), (4) HCC > 30 mm (n = 4), and (5) follow-up < 2 months (n = 3). Finally, 235 patients (median age of 65 years [IQR: 58-72 years]; 79.2% [186 of 235] men) who underwent PTA of 419 small HCC nodules were included in this study. Cirrhosis was detected in 92.3% (217 of 235) of patients, primarily of alcoholic origin (55.7%). Nevertheless, liver function was well-preserved, because 97% of patients with cirrhosis were classified as Child-Pugh A (A5 = 83% and A6 = 14%) with a median MELD score of 8.8 (95% CI: 6-13) and ALBI grade 1 or 2 in 65.9% and 34.1% of patients, respectively. All patients had an AFP score ≤2. Liver biopsy was obtained in 19.2% patients (Fig. 1).

BASELINE CHARACTERISTICS OF PATIENTS WITH STEATOTIC AND NONSTEATOTIC HCC

Both radiologists independently classified (100% interobserver agreement) 52 patients (22.1%) as having at least one steatotic HCC (Fig. 2), and 183 (77.9%) as without. Among the 6 patients with multifocal steatotic HCC, 3 (50%) presented fatty change in all HCC nodules. Nonalcoholic steatohepatitis (NASH) was more frequent in patients with than without steatotic HCC (21.2% vs. 10.4%; \( P = 0.057 \)). Except for BMI, which did not differ between patients with and without steatotic HCC, metabolic conditions tended to be more frequent in the group with steatotic HCC group (diabetes mellitus: 46.2% vs. 38.8%, and liver steatosis: 40.4% vs. 32.8%, respectively; not significant).

![Study flowchart](image-url)
Liver fat content was also higher in patients with steatotic HCC (6% vs. 4.6%), but the difference was not significant (Table 1).

Liver disease was slightly less advanced in patients with than without steatotic HCC: Baseline total bilirubin concentration was lower (−2.1 µmol/L; \( P = 0.035 \)); albumin concentration was higher (+0.8 g/L; \( P = 0.035 \)); and MELD score was lower (−0.8; \( P = 0.014 \)) in patients with steatotic HCC. These data also significantly differed in a per-tumor comparison (data not shown).

Tumor phenotype differed significantly between groups. AFP concentration was lower (\( P = 0.019 \)), and AFP > 100 ng/mL (\( P = 0.045 \)) and multifocality were less frequent (\( P = 0.015 \)) in patients with steatotic HCC, whereas tumor size did not differ between groups. Similar results were obtained in a per-tumor comparison (data not shown). Finally, PTA was more frequently performed under ultrasonography guidance (vs. CT, \( P = 0.01 \)) and using radiofrequency (vs. microwave ablation, \( P = 0.02 \)) in patients with steatotic HCCs.

None of the inflammation-based scores (NLR, PLR, LMR, and SII) differed significantly in patients with and without steatotic HCCs (Table 2).

**FOLLOW-UP AND OUTCOME (TABLE 3)**

Technical success was 100% in both groups. Primary treatment success was obtained in 100% of patients with steatotic and 98.9% of patients with nonsteatotic HCC (\( P = 0.9 \)), whereas secondary treatment success was 100% in both groups.

Complications (SIR grade B/C) were observed after 3% and 3.7% of PTA in patients with and without steatotic HCC, respectively (\( P = 1 \)). No PTA-related death, needle track seeding, or liver abscess was recorded.

The median follow-up of the whole cohort was 28.3 months (95% CI: 25.6-33.4) and did not differ between patients with and without steatotic HCC (median: 27.5 and 29.8 months; \( P = 0.59 \)). Twenty-six (11.1%) patients underwent liver transplantation. In 12 (46.2%) patients, PTA was performed in a context of bridge to transplant. The number of transplanted patients did not differ between groups (\( P = 0.9 \)).

During the follow-up, 3.8% (2 of 52) and 23.5% (43 of 183) of patients with and without steatotic HCC died (\( P = 0.001 \)). HCC-specific mortality also was much lower in patients with steatotic HCC (0 vs. 14.2% [26 of 183]; \( P = 0.002 \)).

Overall, recurrence was less frequent in patients with than without steatotic HCC (42.3 vs. 54.6%), but the difference was not significant (\( P = 0.12 \)). Conversely, early (i.e., <2 years) recurrences were significantly less frequent in patients with steatotic HCC (32.7% vs. 49.2%; \( P = 0.041 \)).

LTP and IDR tended to be less frequent in patients with than without steatotic HCC (43 of 183 of patients with and without steatotic HCC died (\( P = 0.001 \)). HCC-specific mortality also was much lower in patients with steatotic HCC (0 vs. 14.2% [26 of 183]; \( P = 0.002 \)).

The first distant recurrence was beyond the Milan criteria in 5.8% of patients with steatotic HCC and in 8.5% of patients without steatotic HCC (\( P = 0.77 \)). In these patients, the median time to recurrence was
### TABLE 1. CHARACTERISTICS OF PATIENTS WITH STEATOTIC AND NONSTEATOTIC SMALL HCC NODULES TREATED BY PTA

| Characteristic                                    | Steatotic HCC | Nonsteatotic HCC | PValue |
|--------------------------------------------------|---------------|------------------|--------|
| Patients                                         | n = 52        | n = 183          |        |
| Age, years (mean ± SD)                           | 65.3 ± 9.1    | 64.7 ± 9.9       | 0.69   |
| Sex (n [%])                                      |               |                  |        |
| Male                                             | 41 (78.9%)    | 145 (79.2%)      | 1      |
| Female                                           | 11 (21.1%)    | 38 (20.8%)       |        |
| ASA score (n [%])                                |               |                  |        |
| 1-2                                              | 31 (59.6%)    | 87 (47.5%)       | 0.12   |
| 3-4                                              | 21 (40.4%)    | 96 (52.5%)       |        |
| Diabetes (n [%])                                 |               |                  |        |
| No                                               | 28 (53.8%)    | 112 (61.2%)      | 0.34   |
| Yes                                              | 24 (46.2%)    | 71 (38.8%)       |        |
| Metformin treatment (n [%])                      |               |                  |        |
| No                                               | 10 (19.2%)    | 32 (17.5%)       | 0.77   |
| Yes                                              | 9 (17.3%)     | 29 (15.8%)       |        |
| Statin treatment (n [%])                         |               |                  |        |
| No                                               | 9 (17.3%)     | 29 (15.8%)       | 0.83   |
| Yes                                              | 10 (19.2%)    | 32 (17.5%)       |        |
| BMI, kg/m² (mean ± SD)                           | 27.8 ± 5.3    | 27.4 ± 5         | 0.56   |
| Prior treatment for HCC (n [%])                  |               |                  |        |
| No                                               | 26 (50%)      | 86 (47%)         | 0.7    |
| Yes                                              | 26 (50%)      | 97 (53%)         |        |
| Liver disease                                    |               |                  |        |
| Cirrhosis (n [%])                                |               |                  |        |
| No                                               | 3 (5.8%)      | 15 (8.2%)        | 0.56   |
| Yes                                              | 49 (94.2%)    | 168 (91.8%)      |        |
| Causes of liver disease (n [%])                  |               |                  |        |
| Alcohol (with or without viral hepatitis)        | 28 (53.8%)    | 103 (56.3%)      | 0.24   |
| Viral hepatitis B or C                           | 10 (19.2%)    | 51 (27.9%)       |        |
| Hemochromatosis                                  | 3 (5.8%)      | 8 (4.4%)         |        |
| NASH                                             | 11 (21.2%)    | 19 (10.4%)       |        |
| Unknown                                          | 0             | 2 (0.9%)         |        |
| Steatosis (n [%])                                |               |                  |        |
| Absent                                           | 31 (59.6%)    | 121 (67.2%)      | 0.31   |
| Present                                          | 21 (40.4%)    | 59 (32.8%)       |        |
| MRI quantification (%)                           | 6% ± 5.6      | 4.6% ± 4.3       | 0.12   |
| Child-Pugh class                                 |               |                  |        |
| A5                                               | 46 (88.5%)    | 149 (81.4%)      | 0.3    |
| A6                                               | 4 (7.7%)      | 29 (15.9%)       |        |
| B7                                               | 2 (3.8%)      | 5 (2.7%)         |        |
| MELD score (mean ± SD)                           | 8.2 ± 1.9     | 9 ± 2.3          | 0.014  |
| Laboratory data (mean ± SD)                      |               |                  |        |
| AFP (ng/mL)                                      | 9.4 ± 13.5    | 44.7 ± 186.6     | 0.019  |
| AFP > 100 ng/mL (n [%])                          | 0             | 13 (8.2%)        | 0.044  |
| Total bilirubin (µmol/L)                         | 11.9 ± 5.6    | 14 ± 9.1         | 0.035  |
| Albumin (g/L)                                    | 41.7 ± 5      | 39.9 ± 4.7       | 0.035  |
| Prothrombin activity (%)                         | 87.3 ± 11.1   | 84.4 ± 13.1      | 0.11   |
| AST (UI/mL)                                      | 41 ± 38       | 38 ± 23          | 0.61   |
| ALT (UI/mL)                                      | 34 ± 32       | 33 ± 31          | 0.77   |
| GGT (UI/mL)                                      | 189 ± 204     | 155 ± 171        | 0.28   |
| Platelet count (×10³/mm³)                        | 139 ± 66      | 136 ± 74         | 0.76   |
| Neutrophils (×10³/mm³)                           | 3.65 ± 1.48   | 3.31 ± 1.32      | 0.15   |
| Lymphocytes (×10³/mm³)                           | 1.58 ± 0.77   | 1.62 ± 1.21      | 0.78   |
10.5 months (IQR: 8.1-32.4) and 5.1 months (IQR: 3.5-12.2), respectively. At first IDR, the AFP score was >2 in 1.9% and 3.3% of patients with and without steatotic HCC (P = 1), respectively.

IDR was treated by a new curative approach in 72.7% and 56.7% of patients with and without steatotic HCC (P = 0.23). During the follow-up, extrahepatic metastases were detected in 1 (1.9%) and 17 (9.3%) patients with and without steatotic HCC (P = 0.12) (Table 3).

### TABLE 1. Continued

| Characteristic                      | Steatotic HCC | Nonsteatotic HCC | PValue |
|-------------------------------------|---------------|------------------|--------|
| Monocytes (x10^3/mm^3)              | 0.51 ± 0.22   | 0.64 ± 0.22      | 0.34   |
| Creatinine (µmol/L)                 | 79.8 ± 24.9   | 83 ± 28.7        | 0.44   |
| ALBI score                          |               |                  |        |
| 1                                   | 34 (69.4%)    | 113 (64.9%)      | 0.56   |
| 2                                   | 15 (30.6%)    | 61 (35.1%)       |        |
| HCC Tumor size (mean ± SD)          |               |                  |        |
| <20 mm                              | 44 (84.6%)    | 149 (81.4%)      | 0.6    |
| >20 mm                              | 8 (15.4%)     | 34 (18.6%)       |        |
| No. of nodules per patient (n [%])  |               |                  |        |
| 1                                   | 46 (88.5%)    | 132 (72.1%)      | 0.017  |
| 1>1                                 | 6 (11.5%)     | 51 (27.9%)       |        |
| Dome location (n [%])               | 10 (19.2%)    | 47 (25.7%)       | 0.34   |
| Subcapsular location (n [%])        | 15 (28.9%)    | 72 (39.3%)       | 0.17   |
| Near large vessel (n [%])           | 16 (30.8%)    | 37 (20.2%)       | 0.11   |
| Near surrounding organ (n [%])      | 7 (10.6%)     | 27 (7.7%)        | 0.43   |
| PTA PTA modality (n [%])            |               |                  |        |
| Radiofrequency                      | 26 (50%)      | 77 (42%)         | 0.31   |
| Microwave                           | 26 (50%)      | 106 (57.9%)      |        |
| Imaging guidance (n [%])            |               |                  |        |
| Ultrasonography guidance            | 33 (63.5%)    | 93 (50.8%)       | 0.11   |
| CT guidance                         | 19 (36.5%)    | 90 (49.2%)       |        |

Note: Unless otherwise indicated, results are presented as numbers (percentages). Significant P values are indicated in bold.

Abbreviations: ALT, alanine aminotransferase; ASA, American Society of Anesthesiologists; AST, aspartate aminotransferase; GGT, gamma-glutamyltransferase.

### TABLE 2. INFLAMMATION-BASED SCORES IN PATIENTS WITH STEATOTIC AND NONSTEATOTIC HCC

| Scores                  | Steatotic HCC (n = 52 patients) | Nonsteatotic HCC (n = 183 patients) | PValue |
|-------------------------|---------------------------------|-------------------------------------|--------|
| NLR                     | 2.91 ± 1.85                     | 2.53 ± 1.27                        | 0.18   |
| PLR                     | 102 ± 50.2                      | 99.3 ± 58.2                        | 0.74   |
| LMR                     | 3.25 ± 1.53                     | 3.18 ± 2.63                        | 0.82   |
| SII                     | 364.7 ± 223.3                   | 329.5 ± 251.9                      | 0.34   |

Significant P values are indicated in bold.

### TTR, RFS, AND OS

The 3-year cumulative recurrence rate was 40.4% and 49.7% in patients with and without steatotic HCC. TTR was longer in patients with steatotic (median: 32.4 months; 95% CI: 17.1-not reached) than without steatotic HCC (median: 18.6 months; 95% CI: 12-23.6)) (HR = 0.6 [95% CI: 0.39-0.94], P = 0.024), but in multivariate analysis, steatotic HCC was not independently associated with TTR.

The 3-year RFS was 44.6% and 30.2% in patients with and without steatotic HCC. RFS was longer in patients with steatotic (median: 30.4 months [95% CI: 14.6-not reached) than without steatotic HCC (median: 16.4 months [95% CI: 11.4-21.1]) (HR = 0.6, P = 0.018), but in multivariate analysis, steatotic HCC was not independently associated with RFS.

The 3-year OS was 94.4% (95% CI: 79.2%-98.6%) and 70.9% (95% CI: 61.3%-78.6%) in patients with and without steatotic HCC. OS was longer in patients with than without steatotic HCC (HR = 0.14 [95%
CI: 0.03–0.61], P = 0.008). In multivariate analysis, only steatotic HCC (HR = 0.12; P = 0.039) and AFP (HR = 1.002; P < 0.001) independently predicted OS (Fig. 3 and Table 4).

Discussion

This study compares the baseline characteristics of patients with small steatotic versus nonsteatotic HCCs, and their outcome after PTA. We confirm that steatotic HCC is an independent predictor of improved OS in patients treated by PTA for early-stage tumors. In our Western population, liver disease was slightly less advanced in patients with steatotic HCC (mean difference between groups: +0.8 g/L [albumin], −2.1 µmol/L [bilirubin], −0.8 point [MELD score]). Although these differences were significant, it is very unlikely that the underlying liver disease explains the survival benefit. Indeed, PTA was performed in patients with BCLC 0–A HCC and well-compensated cirrhosis in nearly all cases (97% were Child–Pugh A), and the number of deaths from liver cause (outside HCC) was similar in both groups (3.8% in steatotic vs. 4.4% in nonsteatotic HCC). On the other hand, despite a comparable follow-up, no death from HCC was reported in the group with steatotic HCC, whereas it concerned 14.2% of patients with nonsteatotic HCC. Both TTR and RFS were also longer in patients with steatotic HCC (median of 32.4 vs. 18.6 months and 30.4 vs. 16.4 months, respectively).

Early (<2 years) tumor relapse is primarily related to metastatic spread.(7) The lower rate of early recurrence (32.7% vs. 49.2%; P = 0.041) and the longer time to IDR (16.4 vs. 9 months; P = 0.006) in patients with steatotic HCC strongly pleads for slower metastatic escape in this group.

Limited data are available on steatotic HCC.(12,23) In these studies, fatty change was detected by pathology analysis in approximately 20% of patients with HCC treated by surgical resection. We observed a similar rate (22.1%) in our series based on noninvasive MR diagnosis. The perfect interobserver agreement is not surprising, as intratumoral fat detection can be carried out by chemical-shift MRI with 100% specificity,(16,17) due to the difference in resonance frequency between water and fat protons. In a large series (516 patients), Chan et al.(12) showed that steatotic HCC was associated with diabetes mellitus (36.7% vs. 22.1% in patients without steatotic HCC). Our European population with high prevalence of alcohol-related cirrhosis and high rate

| TABLE 3. FOLLOW-UP AND OUTCOME AFTER PTA IN 235 PATIENTS WITH STEATOTIC (n = 52) AND NONSTEATOTIC HCCS (n = 183) SMALL HCC NODULES |
|---------------------------------------------------------------|
| Follow-up (median, 95% CI) | 27.5 months (20.6-34.8) | 29.1 months (25.2-33.6) | 0.54 |
| Liver transplant (n [%]) | 6 (11.5%) | 20 (10.9%) | 0.9 |
| Death, all causes (n [%]) | 2 (3.8%) | 43 (23.5%) | 0.001 |
| HCC-related death (n [%]) | 0 | 26 (14.2%) | 0.002 |
| Death from liver cause, except HCC (n [%]) | 2 (3.8%) | 8 (4.4%) | 0.62 |
| Overall recurrence (n [%]) | 22 (42.3%) | 100 (54.6%) | 0.16 |
| Early (<2 years) recurrence (n [%]) | 17 (32.7%) | 90 (49.2%) | 0.041 |
| First distant recurrence beyond Milan criteria (n [%]) | 3 (5.8%) | 17 (8.5%) | 0.58 |
| LTP, per tumor |
| Occurrence (n [%]) | 5 (7.6%) | 55 (15.5%) | 0.12 |
| Time to LTP (mean ± SD) | 18.9 ± 12 months | 12.1 ± 8.7 months | 0.18 |
| IDR, per patient |
| Occurrence (%) | 21 (40.4%) | 90 (49.2%) | 0.27 |
| Time to IDR (mean ± SD) | 16.4 ± 13.1 months | 9 ± 8.7 months | 0.006 |
| Largest HCC nodule (mean ± SD) | 18 ± 7 mm | 17 ± 12 mm | 0.81 |
| AFP score > 2 at first IDR (n [%]) | 1 (1.9%) | 6 (3.3%) | 1 |
| Curative treatment of IDR | 16 (72.7%) | 51 (56.7%) | 0.23 |
| Extrahepatic recurrence (n [%]) | 1 (1.9%) | 17 (9.3%) | 0.13 |

Note: Unless otherwise indicated, data are presented per patient. Significant P values are indicated in bold.

CI: 0.03–0.61], P = 0.008). In multivariate analysis, only steatotic HCC (HR = 0.12; P = 0.039) and AFP (HR = 1.002; P < 0.001) independently predicted OS (Fig. 3 and Table 4).
of diabetes mellitus strongly differs from the Chinese patients with high hepatitis B virus infection rate of this previous study.\(^{12}\) In addition, our study only included small HCCs, whereas Chan et al. retrospectively collected data on patients undergoing curative surgery basically for larger tumors.

More data are available on SH-HCC, which was described by Salamao et al. in 2010.\(^{11}\) SH-HCC should be regarded as a subset (57.8% in Chan et al.\(^{12}\)) of steatotic HCC, in which—besides intratumoral fat—ballooning changes, Mallory-Denk bodies, and intratumoral inflammatory infiltrate are present. In most reports, SH-HCC has been associated with underlying fatty liver, steatohepatitis, diabetes mellitus, and generally with metabolic syndrome risk.\(^{11-14,24}\) The association between SH-HCC and metabolic syndrome risk factors or liver steatosis\(^{11-14,24-26}\) led us to consider that in HCC, the presence of steatohepatitic features is due to a global liver response (seen in both benign and malignant tissue) to the metabolic syndrome.\(^{27}\) However, this hypothesis has been called into question, because HCC can also develop

![Graphs A, B, C, D, E, F](image_url)

**FIG. 3.** TTR (A), RFS (B), and OS (C) after PTA in patients with small steatotic and nonsteatotic HCCs.
steatohepatitic morphology outside the setting of fatty liver disease or metabolic syndrome, possibly related to genetic alterations inherent to HCC itself. In agreement, recurrent loss of chromosome 9q12-q31 has been reported in a subset of SH-HCC, but not in NASH-associated HCC. Although none of the metabolic factors (BMI, diabetes mellitus, metformin consumption, liver steatosis) investigated in our series was significantly associated with steatotic HCC, most of them were more frequent in this subgroup, including NASH (21.2% vs. 10.4%, \( P = 0.057 \)). Inflammation is involved in NASH and SH-HCC, and is recognized as a cancer hallmark. The host inflammatory response is associated with cancer progression and patient survival. The NLR, PLR, and SII inflammation-based prognostic scores are predictors of early HCC recurrence after surgical resection. SII also independently predicts post-recurrence survival. In our study, these inflammation-based scores were comparable in patients with and without non-steatotic

### Table 4. Univariate and Multivariate Cox Regression Models to Predict OS (Per Patient Analysis)

| Variables | Univariate Analysis | Multivariate Analysis |
|-----------|---------------------|----------------------|
|           | HR (95% CI)         | \( P \) Value        | HR (95% CI)         | \( P \) Value |
| Patients  |                     |                      |                     |              |
| Age       |                     |                      |                     |              |
| Sex female vs. male | 0.70 (0.32-1.54) | 0.38 | 1.92 (0.88-4.17) | 0.1 |
| ASA (>2 vs. ≤2) | 2.13 (1.17-3.9) | 0.014 |                     |              |
| Diabetes  | 1.36 (0.76-2.42) | 0.3 |                     |              |
| Metformin treatment | 0.99 (0.43-2.3) | 0.98 |                     |              |
| Statin treatment | 0.51 (0.18-1.42) | 0.19 |                     |              |
| Treatment-naive patient | 0.59 (0.31-1.09) | 0.09 |                     |              |
| Local recurrence | 0.94 (0.49-1.81) | 0.85 |                     |              |
| IDR       | 1.83 (0.96-3.47) | 0.07 |                     |              |
| Cirrhosis | 0.8 (0.33-1.94) | 0.62 |                     |              |
| Child-Pugh (B vs. A) | 0.72 (0.08-6.63) | 0.78 |                     |              |
| Cause of liver disease (vs. alcohol) |                     | |                     |              |
| Viral hepatitis B or C | 0.91 (0.46-1.8) | 0.79 |                     |              |
| Hemochromatosis | 0.38 (0.05-2.6) | 0.32 |                     |              |
| Others (including NASH) | 0.56 (0.2-1.52) | 0.25 |                     |              |
| Steatosis | 0.76 (0.41-1.4) | 0.38 |                     |              |
| AFP ≥ 100 vs. <100 ng/mL | 4.71 (2.03-10.9) | <0.001 |                     |              |
| AFP (per unit) | 1.002 (1.001-1.003) | <0.001 | 1.002 (1.001-1.003) | <0.001 |
| MELD (>9 vs. ≤9) | 2.59 (1.44-4.67) | 0.001 | 2.01 (0.95-4.26) | 0.07 |
| Tumor size (per mm) | 1.01 (0.96-1.06) | 0.73 |                     |              |
| Tumor size ≤20 mm | 1.48 (0.65-3.33) | 0.35 |                     |              |
| Nb. of HCC (1 vs. >1) | 1.52 (0.78-2.96) | 0.22 |                     |              |
| Steatotic HCC | 0.14 (0.03-0.61) | 0.008 | 0.12 (0.02-0.9) | 0.039 |
| Dome tumor | 0.79 (0.39-1.63) | 0.53 |                     |              |
| Subcapsular | 1.58 (0.89-2.81) | 0.12 |                     |              |
| Near large vessel | 0.95 (0.49-1.84) | 0.88 |                     |              |
| Near surrounding organ | 0.53 (0.17-1.63) | 0.27 |                     |              |
| PTA modality: MWA vs. RF | 1.54 (0.84-2.8) | 0.16 |                     |              |
| US vs. CT guidance | 0.76 (0.42-1.37) | 0.36 |                     |              |

Note: Otherwise indicated, numeric predictors are investigated per unit. Significant \( P \) values are indicated in bold. Abbreviations: MWA, microwave ablation; RF, radiofrequency; US, ultrasonography.
HCC, suggesting that inflammation does not play a major role in the occurrence of steatotic HCC and in their improved outcome.

In surgical studies, steatotic HCCs are more frequently well-differentiated, associated with lower serum AFP level, earlier stage, and lower frequency of major vessel and microvascular invasion. Similarly, in our series, tumor phenotype was less aggressive in patients with small steatotic HCC: Serum AFP concentration was significantly lower (no patient with AFP > 100 ng/mL in this group), and multifocality was less frequent. In studies on SH-HCC outcome after resection, no difference was noted in terms of OS, development of metastatic disease or local recurrence, except in one that found improved disease-free survival for typical SH-HCC. Conversely, in our series of small tumors treated by PTA, steatotic HCC was associated with improved outcome. Studies on PTA for early-stage HCC usually report LTP rates between 10% and 30%, and 3-year cumulative recurrence, RFS, and OS rates of 51%-57%, 29%-40%, and 60%-83%, respectively. In our study, patients with small steatotic HCC showed excellent oncological outcome (LTP rate = 7.6%, and 3-year cumulative recurrence, RFS, and OS = 40.4%, 44.6%, and 94.4%, respectively). These results suggest that focus should be redirected toward steatotic HCC rather than SH-HCC, especially because steatotic HCC can be noninvasively diagnosed by MRI in routine practice.

It remains quite uncommon to discover a predictive factor of improved outcome. Indeed, the commonly used predictors (macrotobecular-massive histological subtype, AFP, and multifocality) are associated with poor prognosis. Outside liver transplantation, in which some of them are used to exclude poor candidates, they are of limited interest in clinical practice due to the absence of any validated neo-/adjuvant treatment. On the other hand, the radiological detection of steatotic HCC as a predictor of good outcome could help in treatment decision making. In early-stage steatotic HCC, PTA is certainly the right treatment option, given the improved outcome of patients, its low complication rate, and improved cost-effectiveness compared with resection. In these patients with small steatotic HCC and well-compensated cirrhosis, liver transplant might be unnecessary or could be kept as a salvage therapy. Indeed, at first recurrence, the risk of recurrence outside the Milan criteria was very low in steatotic HCC (5.8%), compared with the 12.6% risk commonly reported in < 3 cm HCC. Additionally, at first recurrence, the AFP score was > 2 only in 1.9% of patients with steatotic HCC.

Several limitations to our study must be acknowledged. First, this is a retrospective, monocentric study, although data came from a prospectively maintained database with a strict imaging follow-up policy. Second, the median follow-up was relatively short (28.3 months). Finally, like in most HCC studies, the very low percentage of patients with liver biopsy did not allow exploring histological/molecular/genetic features of steatotic HCC. In this context, the SH-HCC subtype is associated with overexpression of C-reactive protein, and lack of satellite nodules and vascular invasion. These tumors belong to the G4 subclass with more favorable prognosis. Genetically, SH-HCCs very rarely harbor activated Wnt/beta-catenin pathway, and they lack CTNNB1, TP53, and TERT promoter mutations. It is certainly worth exploring molecular and genetic pathways in the wider subgroup of steatotic HCCs.

In conclusion, small steatotic HCC detected by MRI is associated with a less aggressive tumor phenotype. In patients with such radiological variant, PTA results in improved outcome and might be the treatment of choice.

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