HEPATOCELLULAR CARCINOMA IN PATIENTS UNDERGOING ORTHOTOPIC LIVER TRANSPLANTATION: radiological findings with anatomopathological correlation in Brazil

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ABSTRACT – Background - Hepatocellular carcinoma is one of the most common malignant tumors worldwide. Imaging techniques, specially computed tomography and ultrasound, are among the most useful diagnostic tools, although the accuracy of these methods may have a significant variability. Aims - To determine the prevalence of hepatocellular carcinoma in cirrhotic patients undergoing orthotopic liver transplantation at “Santa Casa de Misericórdia” of Porto Alegre, RS, Brazil; to estimate the sensitivity of computed tomography and ultrasound in pretransplantation detection of hepatocellular carcinoma in this population; to correlate the radiological characteristics with anatomopathological findings. Materials and Methods - Retrospective prevalence study. Population: adult, cirrhotic patients undergoing orthotopic liver transplantation from January 1990 to July 2003. Among the 292 transplanted patients, 31 cases of hepatocellular carcinoma were diagnosed, of which 29 were included in the study. Tumor characteristics in both ultrasound and computed tomography were compared to those observed in anatomopathological examination. Results - Prevalence of hepatitis C virus infection among patients with diagnosis of hepatocellular carcinoma was 93.5%, and the prevalence of hepatocellular carcinoma among transplanted patients was 10.6%. The overall sensitivity of the imaging techniques was 70.3% for computed tomography and 72% for ultrasound. Conclusion - The prevalence of hepatocellular carcinoma at our institution, as well as the sensitivity of both ultrasound and computed tomography to detect such tumors at pretransplantation screening were similar to those found by other authors, while the prevalence of hepatitis C virus infection, the most common etiological agent for liver disease in our patients, is one of the highest ever reported in literature. Factors influencing hepatocellular carcinoma detection rates were: time from examination to liver transplantation; acquisition of computed tomography images during arterial phase; lesion size. Arterial phase proved to be the most useful part of computed tomography examination in this study.


INTRODUCTION

Hepatocellular carcinoma (HCC) is one of the most common malignant tumors worldwide, and is closely associated with chronic liver disease. Approximately 80% to 90% of the cases occur in patients with cirrhosis, which is believed to be the most important risk factor for this neoplasm. Cirrhotic patients have a 5-year risk of developing HCC of about 20%⁷(13). Liver transplantation candidates, therefore, represent a very high risk population.

Imaging techniques, specially computed tomography (CT) and ultrasound (US), are among the most useful diagnostic tools for pretransplantation screening of HCC. However, the accuracy of these methods may vary due to both technical factors and tumor characteristics⁸(6).

The objective of this study was to undertake a retrospective analysis of pretransplantation radiological findings of all cases of HCC diagnosed in adult patients who were submitted to liver transplantation at “Santa Casa de
MATERIALS AND METHODS

Patients

We included in this study the adult cirrhotic patients who underwent orthotopic liver transplantation (OLT) at our institution from January of 1990 to July of 2003, and had post-transplantation anatomopathological diagnosis of HCC. Were performed 292 adult liver transplantsations (which represents more than 80% of adult OLT carried out at the city of Porto Alegre during this period) and diagnosed 31 cases of HCC through pathological analysis of the explanted liver. Of these patients 22 (71%) were males, and 9 (29%) were females; mean age was 55.2 years (29-71 years). Regarding Child-Pugh functional class, 3 patients were categorized as class A (9.7%), 17 as class B (58.4%), and 11 as class C (35.5%). The etiology of chronic liver disease in the population of patients undergoing liver transplantation is described in Table 1. Two patients were excluded from the study because their imaging exams were not available.

<table>
<thead>
<tr>
<th>Etiology</th>
<th>Number of patients</th>
<th>Number of cases of HCC</th>
</tr>
</thead>
<tbody>
<tr>
<td>HCV</td>
<td>153 (52.5%)</td>
<td>20 (64.5%)</td>
</tr>
<tr>
<td>HCV + alcohol</td>
<td>44 (15%)</td>
<td>9 (29%)</td>
</tr>
<tr>
<td>Alcohol</td>
<td>34 (11.6%)</td>
<td>0</td>
</tr>
<tr>
<td>HBV</td>
<td>17 (5.8%)</td>
<td>1 (3.5%)</td>
</tr>
<tr>
<td>Hemochromatosis + HBV</td>
<td>1 (0.3%)</td>
<td>1 (3.5%)</td>
</tr>
<tr>
<td>Other causes</td>
<td>43 (14.7%)</td>
<td>0</td>
</tr>
<tr>
<td>Total</td>
<td>292</td>
<td>31</td>
</tr>
</tbody>
</table>

Patients included in the transplantation waiting list were followed with medical examination, abdominal ultrasound and serum alpha-fetoprotein (AFP) at intervals of 3 to 6 months. In case of any abnormal findings, an abdominal CT was performed.

Pathological analysis

Following transplantation, the explanted liver was serially sectioned at 10-mm intervals for the anatomopathological examination. Nodules larger than 1.0 cm or with any differing characteristics, as compared to regenerative nodules, which would raise the suspicion of a malignant tumor, were examined microscopically. Malignant nodules were characterized as to their size, location and number. All cases were assessed by the same experienced pathologist blinded to the results of imaging.

Interpretation of imaging exams

All CT scans were retrospectively interpreted by one experienced radiologist who was unaware of any pathological characteristics of the tumor. The tumors seen on CT films were then characterized as to their size (diameter), total number of nodules, location (hepatic segment). Contrast-enhancement pattern was classified as hypodense, isodense, hyperdense or of mixed enhancement as compared to the surrounding hepatic parenchyma. CT scans showing one or more nodules, which were not suggestive of a benign process such as a cyst, hemangioma, or a regenerative nodule, were considered indicative of HCC and regarded as a positive exam. For the ultrasound exams, size, number, location, and ecogenicy (hypoechoic, hyperechoic, or of mixed ecogenicy) of each lesion was registered. Examinations showing solid tumors, which were not suggestive of any benign process, were considered positive for HCC.

In all cases in which imaging exams were considered positive for HCC, we verified whether the lesions seen on pathological analysis corresponded to the ones observed on radiological exams. In the case of US, information about size and location of the nodules was used, whereas in the case of CT scans, films were compared to photographic images of liver cross-section specimens.

Medical records were carefully reviewed and any previous intervention that could alter radiological findings of tumors, such as percutaneous ethanol injection or arterial chemoembolization, was registered. Time from imaging exam to transplantation was also registered. CT exams used in this study were obtained in different centers. For this reason, specification of type of equipment used as well as description of technical details of imaging acquisition was not possible. Besides imaging exams, serum AFP measures were also obtained.

RESULTS

A total of 292 OLT carried out at our institution were analyzed. The most frequent etiological agent causing chronic liver disease in these patients was hepatitis C virus (HCV), followed by a combination of alcohol and HCV (as shown in Table 1). The prevalence of HCV infection in our population of patients was 93.5%. Among these patients, 31 cases of HCC were seen at anatomopathological examination, of which 29 were included in the study. A total of 73 individual nodules were detected at pathological analysis, with sizes that ranged from 0.3 to 8 cm.

CT, US, and serum AFP levels were available in 27, 25 and 28 of the 29 patients, respectively. Arterial phase images were obtained in 17 of the 27 CT scans. Most imaging exams were acquired less than 6 months before liver transplantation.

Of the 27 patients examined by CT, 19 had at least one nodule detected, resulting in a sensitivity of 70.3 % (CI 95%; 51.3-85.1). Considering individual nodules, only 37% of the 64 lesions (CI 95%; 26.3-49.7) were detected. This proportion was 6.9% for nodules less than 1 cm, 55.2% for nodules from 1 to 3 cm, and 100% for tumors larger than 3 cm.

Sensitivity of CT was higher (82.3%) in the group of patients whose CT scans included arterial phase (Table 2). The contrast enhancement pattern during arterial, portal, and delayed phases is described in Table 3. In biphasic (arterial and venous phases) CT scans, the tumors most commonly appeared hyperdense during the arterial phase and isodense during de portal venous phase (35.3% of the cases), representing the most common enhancement pattern combination in our study (Figure 1).
Considering individual phases separately, hyperdense lesions were the most common ones during arterial phase (41.1%), while hypodense nodules predominated during portal venous phase (56.5%). In biphasic CT scans, a significant proportion of tumors (41.2%) was visible only during arterial phase, while a smaller percentage of lesions (11.8%) was identified only during portal venous phase. Seven patients underwent CT scans after lipiodol injection. Of these, HCC were detected in six, resulting in a sensitivity of 85.7% for this technique.

In our study, ultrasound showed at least one nodule in 72% of patients with HCC (CI 95%: 52.3-86.8). A total of 68 individual malignant nodules were diagnosed in this group of patients. Of these, 39.7% (CI 95%: 28.6-51.6) were detected – 12.1% of the lesions smaller than 1 cm, 37% of those with diameters from 1 to 3 cm, and 100% of nodules larger than 3 cm.

Both US and CT had their sensitivities affected by factors such as tumor size and time from examination to transplantation, as described in Tables 3 and 4. Of all nodules identified by these two imaging techniques, only two (one detected by US and one by CT) were not confirmed by histopathological examination to represent HCC.

### TABLE 2 – Detection of HCC according to type of imaging acquisition on CT scan

<table>
<thead>
<tr>
<th>Type of imaging</th>
<th>Number of patients included</th>
<th>Number of patients with detected nodules</th>
</tr>
</thead>
<tbody>
<tr>
<td>With arterial phase</td>
<td>17</td>
<td>14 (82.3%)</td>
</tr>
<tr>
<td>Without arterial phase</td>
<td>10</td>
<td>5 (50%)</td>
</tr>
<tr>
<td>Total</td>
<td>27</td>
<td>19 (70.3%)*</td>
</tr>
</tbody>
</table>

* IC 95%: 51.3%-85.1%

### TABLE 3 – HCC detection rate according to time from exam to transplantation* for each imaging modality

<table>
<thead>
<tr>
<th>Time interval</th>
<th>CT/AP*</th>
<th>US/AP*</th>
</tr>
</thead>
<tbody>
<tr>
<td>6 months or less</td>
<td>14/17 (82.3%)</td>
<td>16/20 (80%)</td>
</tr>
<tr>
<td>Over 6 months</td>
<td>5/10 (50%)</td>
<td>2/5 (40%)</td>
</tr>
</tbody>
</table>

*anatomopathological examination

### TABLE 4 – Sensitivity of imaging modalities according to lesion size

<table>
<thead>
<tr>
<th>Size</th>
<th>CT</th>
<th>US</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 1 cm</td>
<td>2/29 (6.9%)</td>
<td>4/33 (12.1%)</td>
</tr>
<tr>
<td>1 – 3 cm</td>
<td>16/29 (55.2%)</td>
<td>16/28 (37%)</td>
</tr>
<tr>
<td>&gt; 3 cm</td>
<td>6/6 (100%)</td>
<td>7/7 (100%)</td>
</tr>
<tr>
<td>Total</td>
<td>24/64 (37.5%)</td>
<td>27/68 (39.7%)</td>
</tr>
</tbody>
</table>

Overall sensitivity of AFP was 32.1% (IC 95%: 16.9-50.8) and 10.7% (IC 95%: 2.8-26.4) when serum levels above 20 ng/mL and 100 ng/mL, respectively, were considered a positive result. Sensitivity for combination of screening exams was 83.3% for US and AFP, 82.6% for US and CT, and 86.9% for all three methods combined.

Of the 73 nodules diagnosed by anatomopathological examination, 21 received some kind of pretransplantation treatment – 18 were treated with percutaneous ethanol injection (Figure 2) and 3 with chemoembolization. Excluding satellite lesions (<0.5 cm and adjacent to a larger lesion), the average observed tumoral necrosis was 74.3% for the treated group and only 8% for the nodules which were not treated (P<0.0001). Tumor size in both groups did not differ significantly (Table 5).

**DISCUSSION**

Twenty-nine cirrhotic patients with diagnosis of HCC were included in this study. A HCV infection prevalence of 67.3% was observed in the population of transplanted patients, while an even higher prevalence (93.5%) was seen in the group of transplanted patients with HCC. The literature reports diverge significantly on this issue. HCV infection is thought to be associated with approximately 70% of cases of HCC in Japan, 77% in Spain, and 55% in the United States,[11, 16] but these rates range from 7.6% to 81%[7, 21, 35]. The prevalence of HCV infection associated with HCC, in our study, therefore, is among the highest reported in literature.

The risk of developing HCC in cirrhotic patients is approximately 3% to 5% per year[5-9]. Patients in transplantation waiting lists constitute a high-risk population[12-13]. Early diagnosis of HCC
is of great importance, since several forms of therapy, such as percutaneous ethanol injection and arterial chemoembolization, are available and can alter the natural history of this tumor before transplantation can be carried out. OLT has been considered the treatment of choice for patients with tumors at early stages whose clinical conditions do not allow tumor resection. For these reasons, an adequate screening program is justified.

Table 5 – Comparison between treated and untreated patients in relation to tumor size and tumor necrosis

<table>
<thead>
<tr>
<th>Tumor features</th>
<th>Treated</th>
<th>Untreated</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean size</td>
<td>2.3 cm</td>
<td>2 cm</td>
</tr>
<tr>
<td>Mean % of necrosis*</td>
<td>74.3%</td>
<td>8%</td>
</tr>
<tr>
<td>Total number of nodules **</td>
<td>21</td>
<td>52</td>
</tr>
</tbody>
</table>

* P <0.0001; ** Three patients (3 nodules) treated with chemoembolization. All others were submitted to percutaneous alcohol injection.

Patient assessment includes medical examination, laboratory workup, and imaging exams. Clinical assessment, however, have a very low sensitivity and specificity in detecting HCC. AFP, a glycoprotein which has its gene expression reactivated during hepatocarcinogenesis, is the most useful laboratory exam. Serum levels above 20 ng/mL are considered abnormal, although such level might be related to active hepatitis, cirrhosis and other non-malignant conditions. AFP at this level, has a reported sensitivity of 55% to 80% (11, 34). Levels above 100 ng/mL are considered highly suggestive of HCC. In our study, AFP had a low sensitivity in detecting liver cancer.

The low sensitivity of AFP alone has led to the utilization of US as an additional screening technique. Overall US sensitivity reported in literature varies from 42% to 95% (5-6, 9). However, sensitivity for lesions smaller than 1 cm has been estimated to be about 37% (32), while for satellite nodules it may be as low as 16% (20). In this study, the observed sensitivity for US is similar to that found in most reports in which the entire explanted liver was examined at pathology. In studies in which surgical specimens were examined, however, reported sensitivity has usually been higher (6, 25, 26, 32).

For CT, HCC detection rates of 52% to 93.3% have been reported (8, 24, 25, 26), while sensitivity for individual nodules in patients undergoing OLT has ranged from 37% to 71% (14, 17, 24, 25, 26). In our study, despite the fact that many CT scans were not obtained using helical technique, the detection rate of HCC did not differ significantly from that reported by other authors.

Acquisition of images during arterial phase contributes significantly in increasing sensitivity for tumor detection by CT. The importance of arterial phase in CT is due to the fact that highly vascularized tumors, such as HCC, receive almost all their blood supply through the hepatic artery, resulting in early contrast enhancement. Normal hepatic parenchyma, on the other hand, receives only 25% of its blood supply through the hepatic artery, while the remaining 75% comes from the portal vein. As a result, tumor becomes conspicuous during arterial phase, since it becomes strongly impregnated by the contrast medium, as opposed to what happens to the surrounding liver parenchyma (10, 22). In cirrhotic livers, a greater proportion of blood supply comes from the hepatic artery, because it is less affected by the architectural changes secondary to fibrosis as compared to portal venous flow. For this reason, some authors believe that arterial phase assessment assumes an even greater importance in this group of patients (22). The use of arterial phase images can increase tumor detection rate in about 34% compared to CT scans with portal venous phase acquisitions alone. In 8% to 11% of patients, lesions can only be seen during arterial phase (41, 23). In our study, among patients with HCC at pathology whose CT scan included arterial phase images, the tumor detection rate was 82.3%, while the detection rate was only 50% for patients whose CT scans did not include arterial phase. In addition, 41.2% of the individual nodules assessed by biphasic CT were only seen during arterial phase.

The role of noncontrast CT in diagnosis of HCC remains controversial. Some authors report up to 25% of nodules being detected exclusively in noncontrast CT (3, 10). Other investigators question these findings, arguing that lesions which are visible
in non-contrast CT can usually be seen during delayed images as well\cite{27,33}. In our sample, no nodules were seen only in noncontrast CT.

A small number of patients underwent CT evaluation after arterial injection of iodized oil. Sensitivity for this technique, as reported in recent studies with transplanted patients, is approximately 60\%\cite{27,33}.

Several factors may interfere with accuracy of imaging techniques in the diagnosis of HCC. Time from surveillance CT to transplantation, for instance, is an important variable to be taken into consideration when assessing the exams sensitivity. A prospective study carried out in China has estimated the mean duplication period for HCC to be approximately 117 days\cite{30}. In fact, some studies have already demonstrated a correlation between interval from CT examination to transplantation and reported sensitivity\cite{27,33}. As described in Table 3, our study confirms this finding. Sensitivity of both US and CT were higher in the group of patients who were assessed less than 6 months before transplantation.

Nodule size is also a factor that may greatly affect sensibility of imaging exams. In our sample, a low sensitivity for small lesions (<1 cm), and a high sensitivity for large tumors (>3 cm) were observed. We must emphasize, however, that a large number of satellite tumors were detected at anatomopathological examination, and all of them were considered individual lesions when assessing sensitivities of imaging techniques. In studies in which only surgical specimens are examined, as is the case of most series that include pathological correlation in literature, anatomopathological identification of undetected nodules distant from the main tumor is not possible. As a result, a falsely increased sensitivity may be reported.

Of the 29 patients included in our study, 12 underwent percutaneous ethanol injection and 3 were treated with chemoembolization prior to transplantation. Anatomopathological examination is the gold standard for assessment of tumor response after these forms of treatment\cite{29,31}, which is expressed by percentage of tumoral mass that has undergone necrosis. Several authors report complete tumoral necrosis in 50\% to 73\% of tumors treated with ethanol injection\cite{15,28,29}. Of patients submitted to ethanol injection, in our study, 14 out of 18 nodules (77\%) underwent necrosis of at least 90\% of total tumoral mass. Complete necrosis, however, was observed in only 27\% of the cases.

CONCLUSION

The prevalence of HCC among population of adult, cirrhotic patients undergoing OLT at our institution was 10.6\%, and HCV infection was the etiological agent in 93.5\% of cases. Imaging exams used for screening of HCC in these patients had sensitivities which were similar to those reported in literature. Overall sensitivity of imaging techniques in detecting HCC in this population was 70.3\% for CT and 72\% for US. Sensitivity of both methods was lower for lesions smaller than 1 cm and high for tumors larger than 3 cm. Factors which influenced sensibility of imaging methods were: interval from CT assessment to transplantation; availability of arterial phase images in CT scans; lesion size. Of nodules detected by biphasic CT, 41.2\% were seen only during arterial phase, which was the most important phase in diagnosing HCC in our study. Finally, we observed significant tumoral necrosis in the group of patients treated with either percutaneous ethanol injection or chemoembolization prior to transplantation as compared to the group of patients who did not received any form of treatment.
Hepatocellular carcinoma in patients undergoing orthotopic liver transplantation: radiological findings with anatomo-pathological correlation in Brazil

REFERENCES


