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INTRODUCTION

Hepatocellular carcinoma (HCC) is the third most common cause of cancer death in the world. It results in 598,000 deaths per year worldwide. Because of its poor prognosis, this number of deaths is almost the same as the number of cases being diagnosed each year (626,000) [1]. From a global perspective, the two most important risk factors for HCC are chronic hepatitis B and C infection. The geographic distribution patterns of HCC and hepatitis B virus (HBV) almost coincide with each other. The characteristics of patients with HCC are influenced by the etiology and the status of the underlying liver disease. The understanding of its natural history may influence the prognosis and choice of treatment.

HISTORY OF HCC

HCC was believed to be a rare disease in the early 1900s. In Germany, Eggel found only 163 cases of HCC autopsies when he surveyed the world literature for HCC. He proposed an anatomical classification that HCC was classified as nodular, massive and diffuse [2]. However, it was found that the disease was more prevalent outside Europe. In 1911, Yamagiwa suggested a classification based on the origin of the cancer cell, hepatocellular and cholangiocellular, with the terminology of “hepatoma” and “cholangioma” respectively. After Berman’s report of the extremely high HCC incidence among young Mozambican males in 1951, the interest in the studies of HCC increased. It has been shown that there is great regional difference in the pathology and epidemiology of the condition.

DIFFERENT HCC PATTERNS

HCCs in South Africa are mostly large sized and poorly...
HBV CARCINOGENESIS

As with many malignancies, the carcinogenesis of HCC is a multi-step process involving a number of different genetic alterations that ultimately lead to malignant transformation of the hepatocyte. It is postulated that HBV infection causes HCC via direct and indirect pathways. Continuous hepatocyte injury and regeneration in cirrhosis of the liver leads to increased liver cell turnover and hence accumulation of critical mutations in the host genome. This may result in genetic alterations, such as chromosomal rearrangements as well as activation of cellular oncoproteins or inactivation of tumour suppressor genes. However a higher rate of chromosomal abnormalities is found in HBV-related HCC than those linked to other risk factors. Alternative mechanisms in HBV carcinogenesis must also be involved. HBV belongs to the group of oncogenic viruses known as hepadna virus. It is able to integrate its DNA into the genome of the infected cell. Integrated HBV sequences have been observed in established hepatoma cell lines and in about 80% of human HBV related HCCs. It is postulated that the HBV DNA integration may confer a selective growth advantage on target cells and leads to the onset of tumor progression. The integration sites are frequently detected in cellular genes involving cell signaling or growth control. Host chromosomal instability is also enhanced by HBV DNA integration. Large inverted duplications, deletions, amplification, chromosomal translocation have all been observed to be associated with HBV integration. In addition, the regulatory proteins HBx and the PreS2 activators can exert a tumor promoter-like function, resulting in positive selection of cells producing a functional regulatory protein.

HCV CARCINOGENESIS

HCV infection is a more important factor than HBV in the development of HCC in western countries. Markers of HCV infection are found in a higher proportion of HCC patients than in most of the Asian countries; ranging from 44% to 66% in Italy, 27% to 58% in France, 60% to 75% in Spain. Japan, unlike other Asian countries, also has a high proportion of HCC caused by HCV infection accounting for 80% to 90% of all the cases.
HCC risk increases to 17-fold in HCV-infected patients compared with HCV-negative subjects\[11\]. The risk of HCC occurrence is different among all HCV patients. It is a function of the degree of liver fibrosis and the time of acquisition of the infection. The risk for cirrhotic HCV patients (F4) was the highest with 5.8% per year, compared to those who had less fibrosis (F1-3, 0.5% to 2.6%)\[12\].

Because of the absence of reverse transcription activity of the HCV RNA virus, its viral genome unlike HBV is not able to integrate into the genome of the infected cell. Therefore, HCV causes HCC via an indirect pathway by causing chronic inflammation, cell death, proliferation and cirrhosis. Accordingly, HCV-related HCCs are almost exclusively found in patients with cirrhosis. However, there are studies raising the possibility that HCV may also operate through other pathways in promoting malignant transformation of hepatocytes. HCCs without cirrhosis in HCV-infected patients, though rare, have been reported. The transforming potential of NS3 protein and core protein has been described.

**CLINICAL DIFFERENCES BETWEEN HBV- AND HCV-RELATED HCC**

Apart from the difference in the carcinogenic mechanisms, there are a number of clinical differences between HBV- and HCV-related HCC. For reasons yet to be known there is a larger proportion of male HBV infected patients with HCC than the HCV counterpart. The male-to-female ratio is higher in HBV-related HCC than in HCV-related HCC (6.7:1 and 3:1 respectively in one study)\[13\]. Studies consistently show that HCC develops 10 years earlier in HBV carriers than in HCV carriers, a difference possibly due to the earlier age at exposure to the causative agent\[14\].

Due to the different mechanisms involved in carcinogenesis, their rate of HCC development is also different. In a Japanese study, 795 patients with viral or alcoholic cirrhosis were observed from 1974 to 1989. Two hundred and twenty five patients (27.8%) developed HCC. The cumulative rates of HCC development in the 180 HBV cirrhotic subgroup at 3, 5, 10 and 15 years after follow up were 7.2%, 14.2%, 27.2% and 27.2% respectively. At 9 years after the follow up, the rate of HCC development began to decrease, and the cumulative rate remained the same thereafter. The corresponding rates in HCV-related cirrhosis were 10.4%, 21.5%, 53.2% and 75.2% respectively in the 349 HCV patients. The authors conclude that the cumulative rate of HCC occurrence has a linear relationship with time in HCV patients whereas it seems to plateau after 10 years for HBV patients\[15,16\]. However, according to an Italian study, the cumulative rates of HCC at 5, 10 and 15 years continue to increase with 6.5%, 23.4% and 31.9% for patients with HBV-related cirrhosis, and 4.6%, 24% and 56.2% for patients with HCV-related cirrhosis, respectively\[17\]. At the time of HCC diagnosis, a higher proportion of HCV infected patients than HBV infected patients has advanced liver histology and has a higher Child Pugh’s score\[18\].

The HCC tumor size and the growth pattern also tend to be different between HBV- and HCV-related HCCs. In most patients with HCV-related HCC, the tumors are more likely to be solitary, smaller sized and encapsulated whereas HBV-related HCC are more commonly infiltrative and multinodular\[19\]. Extensive hepatic involvement and portal vein invasion by the tumor were found in 44% and 52% respectively in 95 HBsAg-positive HCC patients compared to 25% and 28% respectively in 370 HBsAg-seronegative HCC patients\[19\].

**NATURAL HISTORY OF HCC**

The natural history of HCC depends on tumour growth characteristics as well as the underlying liver cirrhosis. The majority of hepatitis-related HCC develops in cirrhotic livers with a relatively poor liver reserve. Given a wide range of tumor doubling time, some patients may die from liver failure before the tumor grows to an advanced stage. Features of HCC shown to indicate more aggressive behaviour include poorly differentiated histology, lack of fibrous capsule, large tumour size, early vascular invasion and elevated serum levels of alpha fetoprotein (AFP). Tumor growth rates have a wide range of variability even among patients of the same region and regardless of disease stage. Doubling times ranges from 1 mo to 19 mo with a median of 4 to 6 mo\[20\].

Vascular supply of HCC is derived from the hepatic arterial network and the efferent vessels of portal origin have been described to create arterioporal or arteriovenous shunts which serve as a low resistance path for tumour thrombi fragments to survive and spread within the portal efferent network. In a large autopsy study from Sweden, 56% of HCC had evidence of vascular invasion but biliary tract involvement was observed in only 4%\[20\]. In another autopsy study in Japan, vascular involvement was noted up to 82% and the likelihood of vascular involvement was related not only to tumor size but also the macroscopic types\[20\]. The frequency of portal vein invasion is higher than hepatic vein invasion\[27\].

A retrospective analysis was performed in the United States of America on 347 HCC patients who received a metastatic workup including bone scans and computed tomography scans of the chest, abdomen, and pelvis. Clinical and tumor characteristics were evaluated as risk factors for metastasis by univariate and multivariate methods. One hundred forty-five patients had metastases: 72 had thoracic, 57 had abdominal, and 34 had bone metastases. Poor tumour differentiation, multinodular spread, and tumour size of 5 cm or more were the strongest predictors of metastatic disease by logistic regression analysis\[28\]. In a Japanese study, the relationship between HCC specimen staining for AFP, serum AFP levels and pathological findings were examined. The disease prognosis was studied with respect to the staining for AFP in excised tumors. The mean serum AFP level in patients with positive AFP staining. No significant relationship was found between AFP positivity and tumor size, tumor encapsulation, degree of vascular invasion, or the histological differentiation grade of the tumor. However, patients with AFP-positive
carcinomas had a poorer prognosis than those with AFP-negative carcinomas (5-year survival rate of AFP-positive and negative groups were 26.7% and 56.5%, respectively).

**PREDICTION OF SURVIVAL**

The prognosis of HCC has been dismal because patients with HCC usually present late and have already developed advanced liver cirrhosis. Curative resection was seldomly possible. Therefore, the life expectancies of patients with newly diagnosed HCC were classically measured in terms of weeks to months with mortality to incidence ratio close to 1. Due to the increased awareness of the disease and HCC surveillance programmes in some parts of the world, HCC can be diagnosed at its earlier phase permitting timely curative interventions. In a Hong Kong study, the clinical features and disease outcome of 306 HCC patients were analysed. The characteristics of those present with symptoms and those with the disease diagnosed by screening were compared. One hundred and forty two patients with HCC diagnosed by regular screening have a significantly lower serum AFP level, smaller tumour size, less bilobar disease, less portal vein infiltration and less distant metastasis compared with the 164 symptomatic subjects. As a result a higher proportion of patients could undergo curative resection (26.8% vs 7.9%).

Scores and models for predicting survival probability are invaluable to tailor suitable treatment options. The traditional TNM cancer staging system has limited applicability as it does not take into account the possible poor liver function in HCC patients. Eighty percent of HCC patients cannot be treated with surgery, so the tumour pathologic information needed in TNM cancer staging is not available. This further limits its prognostic use. Child-Pugh score on the other hand was designed to predict survival in liver cirrhosis without any component of HCC tumour morphology and consideration of disease extension. The first widely accepted staging system to predict survival for HCC which incorporates tumour biology and hepatic function was proposed by Okuda et al in 1985. Tumour size, serum bilirubin, albumin and the presence of ascites are included in the calculation of Okuda staging. In a Hong Kong study of 106 untreated unresectable HCC patients performed in 2001, the median survival was 5.1 mo in stage I, 2.7 mo in stage II and 1.0 mo in stage III. These figures were remarkably similar to the original report by Okuda et al, in which the median survival of stages I, II and III disease was 8.3, 2, and 0.7 mo respectively. Among other staging systems, only Okuda staging was independently correlated with prognosis in the final multivariate model. As more treatment options become available, newer prognostic models are introduced.

The Cancer of the Liver Italian Group Programme (CLIP) proposed a staging system which has been shown to be superior to that of the Okuda stage with a higher number of categories and greater discriminant ability. It reveals a subgroup of patients with an impressively more favourable prognosis who could be candidates for more aggressive therapeutic strategies. On the other hand, it also sorts out a subset of patients with a median survival long enough to be considered for clinical trials of palliative antineoplastic treatment. It includes Child-Pugh score, tumour size, presence of portal vein thrombosis, and serum AFP. Another survival model is the Barcelona Clinic Liver Group (BCLC) staging system which also effectively selects patients for aggressive treatments. It is constructed from several cohort studies and randomized controlled trials (RCTs) by the Barcelona group. This classification uses variables related to tumour stage, liver functional status, physical status, and cancer-related symptoms, and links the five stages described (0, A-D) with a treatment algorithm. The therapeutic approach configured into the BCLC staging system for patients with non-surgical HCC is based on the study by Llovet et al in which asymptomatic patients without evidence of disease extension (vascular invasion or metastasis) had significantly better survival compared to those with more advanced disease. Such patients are classified as having intermediate HCC and may be better candidates for tumour ablative interventions or chemoembolization.

**HBV VACCINATION**

HBV vaccination has been shown to be effective to protect against HBV infection. The vaccination is recommended to high risk population groups such as health care workers, patients with immunocompromised states or with chronic liver diseases or with the need for regular blood product transfusion or hemodialysis, and neonates of HBV carrier mothers. Vertical and early horizontal transmission is the major route of HBV transmission in high prevalent areas. The transmission rate was estimated to be as high as 90% in HBeAg positive mothers. The use of HBV vaccine and hepatitis B immunoglobulin (HBIG) can greatly reduce the infection rate of the neonates to 10% to 15%. The use of vaccination in all newborns can reduce the chance of infection acquired via other routes during early childhood and thus preventing the prolonged immunotolerant phase of this unique infection. By eliminating HBV seroprevalence in the population, HCC incidence should be greatly reduced in high prevalence areas. In Taiwan, a universal HBV vaccination programme was carried out in 1984. A study had shown that the HCC incidence in children was decreased since the introduction of the vaccination programme. In that study, the author showed that the annual incidence of childhood liver cancer was reduced from 0.70 per 100000 to 0.36 per 100000 in the children born between 1981 to 1986 and 1990 to 1994 respectively. Therefore a similar decrease in the adult HCC incidence is expected in the future. Likewise, vaccination programmes in high endemic areas like Gambia, intermediate endemic areas like Italy had shown impressive reduction of the HBV seroprevalence. This may transform to a reduction in HCC incidence similar to the Taiwan study. However, the development of an effective vaccine has been hampered by the high genetic variability of HCV.

**CONCLUSION**

HCC is a complex neoplasm often occurring in a preneoplastic cirrhotic liver, and thus, survival and treatment options depend on variables of both HCC and cirrhosis.
Among chronic viral hepatitis-related HCC, differences in carcinogenesis and clinical presentation exist between HBV and HCV patients. Survival models using tumour size, invasiveness, liver function, patient physical performance status can predict prognosis and guide treatments.

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