The Current Approach to the Hepatocellular Carcinoma; A Mini Review of Etiology, Prognosis and Treatment

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Received: March 05, 2019
Published: March 11, 2019

Abstract

Hepatocellular carcinoma (HCC) is the most common liver malignancy worldwide and is one of the major causes of cancer-related deaths. HCC is reported to be the second most fatal malignancy. The major risk factors for HCC are well known; the known risk factors include hepatitis C virus (HCV) and hepatitis B virus (HBV). Major advances have been reported in the treatment of HCC. Success of early diagnosis increases when these risk factors are identified, and the cases are followed up. It is reported that in the treatment of early-diagnosed HCC cases, ethanol injection or radiofrequency ablation methods as well as surgical resection should be preferred, particularly in cases without liver cirrhosis and in cases where the tumor is restricted. Similarly, liver transplantation may be an option for patients that meet specific criteria.

Keywords: Hepatocellular Carcinoma; Liver Carcinoma; Liver; Cirrhosis

Introduction

Hepatocellular carcinoma (HCC) is one of the leading causes of cancer-related deaths in the world. HCC is reported to be the second most fatal malignancy after pancreatic carcinoma [1-3]. In the United States of America, 5-year survival of patients with HCC is reported to be approximately 9% [4]. In contrast to several other malignancies, the major risk factors for HCC are well known. These risk factors include hepatitis C (HCV) and hepatitis B virus (HBV). Major advances have been reported in the treatment of HCC. Success of early diagnosis increases when these risk factors are identified, and the cases are followed up [1]. A glance at the epidemiology of HCC reveals that it is more common in developing regions [5,6]. A 2008 study reported extremely high mortality rates for HCC [7]. Owing to the high mortality rate, HCC is one of the most common causes of cancer-related deaths. There are several predisposing factors in the etiology and pathogenesis of HCC. Following are the few examples of these factors: Hepatotropic viruses, HBV and HCV, are the most common cause [8]. Cirrhosis: Considering its annual incidence, approximately 3% of the patients with cirrhosis are expected to develop HCC [9]. Liver cell dysplasia (Large cell dysplasia and small cell dysplasia): Although both large and small cell dysplasia's are risk factors for HCC, some authors argue that the presence of small cell dysplasia is a more important risk factor for HCC [10,11]. Thorotrast: It has been reported to play a role in HCC development [12]. Alpha1 antitrypsin deficiency: It is reported that individuals born with this metabolic disorder have a predisposition for HCC [13]. Tyrosinemia: There is a high risk of HCC in individuals born with this metabolic problem [14]. Aflatoxin: Aflatoxin, derived from the metabolic wastes of the fungus Aspergillus flavus, is associated with HCC [15].

HCC can be radiographically diagnosed by computed tomography (CT) or magnetic resonance imaging (MRI). Moreover, dynamic imaging maintains contrast in the early arterial phase, which is then washed or released in the next portal phase. This imaging approach is 90% sensitive and 95% specific for
HCC [16]. Ultrasound (US) findings are similar to those of CT and MRI. However, in recent years, contrast-enhanced US is no longer accepted as a diagnostic tool owing to the fact that cholangiocarcinoma cases cannot be distinguished from HCC and further investigation methods are recommended. In terms of laboratory findings, alpha fetoprotein (AFP) elevation in patients with HCC has been known for more than 40 years. AFP elevation can also be detected in pregnancy, normal fetal yolk sac, and fetal liver tissues. Other than HCC, it is also detected in the malignancies of the biliary tract, pancreas, and gastrointestinal system, as well as in nonseminomatous tumors. AFP level is expected to decrease in infants at 300 days after birth, and any AFP elevation detected after this period is a cause to suspect malignancy [17]. However, it has been reported that approximately one-third of the diagnosed cases may have normal AFP levels at the time of diagnosis [18]. Despite the advances in imaging systems and the support provided by laboratory findings, final diagnosis of HCC is still made on the basis of histopathological examination. US or CT-guided cytological fine-needle aspiration biopsy (FNAB) or histological tru-cut biopsy might be preferred in cases suspected with HCC. It has been reported that the diagnostic accuracy of concurrent FNAB and tru-cut biopsy is higher than that of either procedure on its own, with a sensitivity of 96% and specificity of 95% [19]. In microscopic examination of HCC, neoplastic hepatocytes mimic normal liver tissue depending on the degree of differentiation. Well-differentiated tumors that are almost similar to normal tissue are generally difficult to histopathologically distinguish from differentiated liver adenoma tissues. Less differentiated anaplastic tumors can be identified using certain additional immunohistochemical or histochemical analyses because their similarity with normal tissue is reduced.

The most common histological patterns in microscopy are trabecular (sinusoidal), solid, and pseudoglandular (acinar) patterns [20]. Most HCC cases are immunohistochemically positive for AFP, epithelial membrane antigen, alpha1 antitrypsin, fibrinogen, IgG, ferritin, Heppar 1, MOC 31, glypican-3, and polyclonal CEA [21]. HCC staging is generally dependent on many criteria such as tumor size, number of tumor nodules, and the presence/absence of vascular invasion [21]. There are various treatment approaches available for HCC cases. Considering the HCC stage, the functional status of liver and the accompanying medical problems, the treatment decision for HCC should be made with a multidisciplinary team comprising a surgeon, oncologist, pathologist, radiologist, and hepatologist [1]. The most effective treatment in HCC cases is surgical resection and liver transplantation (Figures 1 & 2) [22]. Ablative therapies such as radiofrequency ablation (RF), microwave ablation, or percutaneous ethanol injection are among the treatments that should be primarily used for masses smaller than 2 cm. However, ablative therapies are also preferred in patients with advanced age and poor clinical condition, and in whom surgical resection or transplantation cannot be employed [23]. In cases of HCC, many factors such as the number of tumors, tumor size, presence of cirrhosis, and surgical experience should be taken into account before performing resection and transplantation procedures. In case of partial resection of cirrhotic livers, patient’s condition may deteriorate owing to impaired function and low regeneration capacity. For this reason, liver functions should be comprehensively evaluated in patients with cirrhosis, and then decision regarding surgery should be taken [22]. In patients in whom surgery cannot be performed, neoadjuvant therapies such as transarterial embolization (TAE), transarterial chemoembolization (TACE), RF ablation, and percutaneous acid injection can be employed. In addition to these techniques, the use of microwave therapy, transarterial radioembolization, and cryotherapy applications has also been reported. Furthermore, sorafenib, a tyrosine kinase inhibitor used for molecular therapy, can be preferred in patients with advanced stage HCC [24].
References


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