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Risk calculators for hepatocellular carcinoma in patients affected with chronic hepatitis B in Asia

YangHI *et al.* HCC risk calculator for CHB

Hwai-I Yang, Mei-Hsuan Lee, Jessica Liu, Chien-Jen Chen

**Hwai-I Yang, Jessica Liu, Chien-Jen Chen,** Genomics Research Center, Academia Sinica, Taipei 115, Taiwan

**Hwai-I Yang,** Graduate Institute of Clinical Medical Science, China Medical University, Taichung 401, Taiwan

**Hwai-I Yang,** Molecular and Genomic Epidemiology Center, China Medical University Hospital, Taichung 401, Taiwan

**Mei-Hsuan Lee,** Institute of Clinical Medicine, National Yang-Ming University, Taipei 115, Taiwan

**Chien-Jen Chen,** Graduate Institute of Epidemiology and Preventive Medicine, College of Public Health, National Taiwan University, Taipei 115, Taiwan

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**Correspondence to: Hwai-I Yang, PhD,** Genomics Research Center, Academia Sinica, 128 Academia Road Section 2, Taipei 115, Taiwan. hiyang@gate.sinica.edu.tw

**Telephone: +**886-2-27871308 **Fax:** +886-2-27898811

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**Abstract**

Risk calculators are widely used in many clinical fields, and integrate several important risk factors through the conversion of a risk function into a single measure of risk. Several studies have been carried out to create risk calculators for the prediction of hepatocellular carcinoma (HCC) in patients with chronic hepatitis B (CHB). Most of them were hospital-based, with limited sample sizes and insufficient external validation. These study groups collaborated to establish the REACH-B risk score, which incorporated five clinical variables to predict HCC risk. This risk score was then validated in international clinical cohorts. Evidence suggests that quantitative serum HBsAg level provides additional predictability of HCC, especially in patients with low levels of HBV DNA. This novel marker was incorporated into a risk calculator and was internally validated. This tool will hopefully be externally validated in the near future. Risk calculators can be used to support clinical practice, and to establish preventive measures; several "off-label" extension usages have also been implemented. Albeit beneficial, several precautions and discussions should be noted in using the risk calculators. The future development of risk calculators for CHB patients can be extended by applying them to additional CHB-related outcomes, and by incorporating emerging risk parameters.

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**Key words:** Chronic hepatitis B; Hepatocellular carcinoma; Risk calculator

**Core tip:** The risk calculator is a useful tool in many fields of medicine, including hepatology. This paper reviews the history of the development and validation of risk calculators of hepatocellular carcinoma (HCC) for patients with chronic hepatitis B (CHB). The rationale for using HCC risk calculators is first given, followed by a description of the course and pathway towards deriving HCC risk estimation tools for treatment-free CHB patients. Examples of the application of HCC risk prediction tools in clinical and public health settings is also shown. The paper also discusses several issues raised by the application of HCC risk scores.

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**INTRODUCTION**

Hepatocellular carcinoma (HCC) is one of the most important adverse outcomes of chronic hepatitis B virus (HBV) infection, which affects more than 350 million people worldwide[1,2]. HCC causes poor quality of life and shortened survival, and is thus regarded as a major health challenge. The risk of chronic hepatitis B (CHB) progressing to HCC may be reduced by antiviral therapy[3], and surveillance with abdominal ultrasonography and serum alpha-fetoprotein tests can screen patients for early HCC treatment. Although the global number of individuals infected with chronic hepatitis B is extensive, especially in endemic areas such as Asian-Pacific and sub-Saharan African regions, only a small number of patients develop end-stage liver diseases. Therefore, the identification and triage of patients who are at high risk of HCC development is important. Several factors, such as gender, age, family history of HCC, presence of hepatic inflammation/fibrosis, alcohol consumption, elevated viral load, hepatitis B e antigen (HBeAg) positivity, and specific HBV genotypes (*e.g.,* genotype C), have been identified to be independently associated with elevated risk of HCC development (Figure 1)[4-8]. These factors, including patient, viral, and environmental factors, interact with one another and lead to HCC development in patients with chronic HBV infection. From the individualized medicine point of view, these factors should be used to reveal the future risk of HCC progression in patients with viral hepatitis so that preventive measures can be applied to those with high risk. Intuitively, a patient who has all of the risk factors for HCC can easily be categorized as very high risk (such as patient A in Table 1), whereas a patient without risk factors can be identified as very low risk (such as patient B in Table 1), despite not knowing the exact risk in these patients. However, the risk in patients with mixed risk profiles (patients C and D in Table 1) cannot easily be determined or compared. Thus, a tool that is evidence based and appropriately weights these risk predictors to provide a simple and accurate risk estimate for the interested outcome (*i.e.* HCC) would be very helpful. Consequently, some investigators have responded accordingly to this demand.

**RISK CALCULATORS FOR HEPATOCELLULAR CARCINOMA IN CHB PATIENTS WITHOUT ANTIVIRAL TREATMENT**

By 2011, several study groups had established prediction models that incorporated several clinical variables to estimate HCC risk for CHB patients (Table 2)[9-12]. Most of those were hospital-based in study design with limited sample sizes. These HCC prediction models had diverse sets of risk predictors, although some common parameters were used. The most important issue was the lack of external validation to a satisfactory extent, which was common among those tools. To solve this problem, these study groups collaborated to established an HCC risk score (REACH-B) incorporating gender, age, serum ALT concentration, HBeAg status, and serum HBV DNA level as the predicting parameters[13]. This study derived a 17-point risk model from 3,584 treatment- and cirrhosis-free CHB patients in a community-based Taiwanese cohort (REVEAL-HBV), and validated its use in a composite hospital-based cohort (*n* = 1505) from Hong Kong and Korea. This risk score could predict HCC with a wide range of risks, ranging from 0.0% to 23.6% at 3 years, 0.0% to 47.4% at 5 years, and 0.0% to 81.6% at 10 years for patients with the lowest through the highest scores. Although the derivation and validation cohorts were quite different in their distributions of sex, age, HBeAg serostatus, ALT concentration, HBV DNA level, and cirrhosis, the risk score developed from the derivation cohort accurately and reliably estimated the HCC risk at 3, 5, and 10 years of follow-up in the validation cohort. The area under the receiver operating characteristic curve (AUROC) and the corresponding 95% confidence intervals (CI) were 0.811 (0.790-0.831), 0.796 (0.775-0.816), and 0.769 (0.747-0.790), respectively, in predicting 3-, 5-, and 10-year HCC risk, which indicate a fair discriminatory capability. The performance of the risk score was improved when cirrhotic patients were excluded from the validation cohort[13]. Moreover, the predicted risks were similar to the observed risks as estimated by the Kaplan-Meier method, which indicate good calibration. This is the first study to provide firm external validation of the use of a HCC risk prediction tool in a group of clinical CHB patients.

Recently, several lines of evidence have suggested that quantitative serum HBsAg levels provide additional predictability of HCC, especially in patients with low levels of HBV DNA[14-16]. This novel marker features high reproducibility and relatively low cost. Thus, the original REVEAL nomograms were then upgraded by incorporating this novel risk predictor into the HCC risk prediction model[16]. In addition to HCC, this study also provided a prediction model for predicting the long-term development of cirrhosis as well. The risk prediction model for HCC included age, sex, family history of HCC, and a combined variable encompassing HBeAg serostatus, serum HBV DNA and ALT levels, quantitative serum HBsAg level, and HBV genotype as the predicting parameters (Table 3; all parameters were included in the REVEAL nomogram except for quantitative serum HBsAg level). The projected 5-, 10-, and 15-year HCC risk for each score was pre-calculated and depicted in a nomogram (Figure 2). This upgraded HCC risk calculator was internally validated using a third of the population from which the model was derived from, and showed excellent prediction accuracy and discriminatory ability. Taking the REACH-B score as a precedent, this upgraded version may hopefully be validated externally using clinical CHB patients to prove its accuracy, reliability, and added predictive capability over the original version in the near future. In addition to this, because testing serum HBV DNA levels is relatively costly compared to all other risk predictors in the risk calculator, a model/score might be generated in which quantitative serum HBsAg levels can be used in lieu of serum HBV DNA levels. This score/model, once generated and validated, can be used as a first-line risk prediction instrument by general practitioners and CHB patients who are interested in knowing their long-term risk, thereby largely extending the application of this tool into the community.

**IMPLICATIONS OF HCC RISK CALCULATORS FOR CLINICS AND PUBLIC HEALTH**

As a useful risk estimation tool, the HCC risk calculator can be used to support clinical practice as well as derive preventive measures. The risk score helps to create a gauge for HCC risk assessment that can be used for evidence-based decisions during clinical management of chronic HBV carriers. Based on patients’ personalized HCC risks, their follow-up intervals, surveillance patterns, and referral strategies can be tailored. Furthermore, as antiviral therapy has been shown to improve histology[17], timely antiviral therapy in high-HCC-risk patients may lead to improvement in quality of life and prolonged survival. The risk score also provides an appropriate platform for physician-patient communication, which may help to raise patients' awareness of HCC and assist in imparting knowledge to high-risk patients. Patients' willingness to receive antiviral therapies may hopefully be motivated by understanding their own risk. The risk prediction score might also complement clinical practice guidelines by functioning as a risk stratification system. From a public health viewpoint, the risk calculator can bridge the gap between personal risk profiles and population health impact resulting from HCC, and may help in proper allocation of health care resources. For example, by knowing the distributions of scores from community and clinical settings used in the REACH-B paper (shown in Figure 3),[13] one can try to estimate the impact of HCC according to different risk scores in specific time frames. Further analyses on its cost-effectiveness can be conducted in order to guide public health intervention plans.

A key question that has been raised about the usage of the risk calculator is, how will patients with diverse predicted risks be managed differently? An argument was also brought up that it may instead be more important to identify a group of patients with a zero risk for HCC within a specific time frame. AsFigure 2 shows, the HCC risk that can be predicted by the risk score is a risk continuum. The risk is never zero, even in patients with the lowest scores. The potential cut-off risk and corresponding management strategies is an issue. However, the risk calculator only deals with risk estimation of HCC. The management of CHB patients should take other information into consideration such as benefits, adverse effects, as well as the cost-effectiveness of management strategies. International guideline development committees may integrate all of these factors when setting up threshold scores for beginning various clinical interventions.

**EXTENDING THE USE OF HCC RISK SCORES IN TREATMENT RESEARCH**

The REACH-B scoring system has been used to classify anti-viral treatment eligibility of CHB patients according to the 2012 Asian Pacific Association for the Study of the Liver (APASL) treatment guidelines.[18] In this study, a total of 904 noncirrhotic CHB patients were enrolled, and their age, gender liver biochemistry, HBeAg status, and HBV DNA levels were recorded. This study showed that for patients to be eligible for anti-viral treatment, the minimal REACH-B score should be 7 and 6, respectively, for HBeAg-seropositive and –negative patients. Additionally, in HBeAg-seronegative patients, the REACH-B score could predict treatment eligibility, with an adjusted OR (95%CI) of 1.78 (1.61-1.98). In HBeAg-seropositive patients, however, this same score-dependent eligibility of treatment was not observed. In this study, the authors also showed that the REACH-B score was excellent in discriminating treatment eligibility for young (< 40 years) HBeAg-seropositive patients (AUC: 0.903) and in both young (< 45 years; AUC: 0.907) and older (≥ 45 years; AUC: 0.883) HBeAg-seronegative patients; but the discriminatory capability for older (≥ 40 years) HBeAg-seropositive patients was poor (AUC: 0.664). After an in depth investigation, they found that 46.4% of HBeAg-seropositive patients older than 40 years of age with high risk of HCC, as estimated by a REACH score ≥ 11, would be erroneously excluded from treatment, mainly because their ALT levels never exceeded 2xULN, even after frequent blood tests during follow up. In other words, these patients were excluded from treatment, despite the fact that all these patients were significantly viraemic. The authors suggested that the severity of liver fibrosis in these high-risk patients be evaluated through liver biopsy or complementary noninvasive tools such as liver stiffness measurements, rather than exclude them from treatment.

The accuracy of HCC risk scores that were derived from treatment-naïve CHB patients has been evaluated in 1531 patients treated with entecavir[19]. These patients were treated with entecavir 0.5 mg daily for at least 12 months, and were assessed once every 3-6 mo for symptoms, drug history, and adherence, in addition to collecting blood samples for biochemical analyses. Three HCC risk scores, including the CU-HCC, GAG-HCC, and REACH-B scores were used based on profiles at the start of treatment and 2 years later after treatment. The AUROCs for utilizing the CU-HCC, GAG-HCC, and REACH-B scores for predicting HCC were 0.80 (95%CI: 0.75-0.86), 0.76 (95%CI: 0.70-0.82), and 0.71 (95%CI: 0.62-0.81), respectively. This study also evaluated the on-treatment performance of HCC risk scores, as well as the change in HCC risk scores from baseline to 2 years after treatment with entecavir. Interestingly, the CU-HCC, GAG-HCC, and REACH-B scores, which were originally developed in treatment-naïve patients, can be applicable to predict subsequent development of HCC in patients treated with entecavir. However, because all of these scores were derived from and validated in treatment-free cohorts, and many potential predictors related to therapy and responses during and after treatment could not be collected, they may not be the optimal prediction tools for CHB patients under antiviral treatment. A risk calculator derived from treated patients, which includes pre-treatment, on-treatment, and post-treatment parameters, might best fit this purpose.

Recently, the REACH-B score has been used as a natural history counterpart to evaluate the long-term effect of antiviral therapy using tenofovir on HCC incidence over time.[20] The analysis included more than 600 patients from two multinational Phase 3, placebo-controlled tenofovir trials (one for HBeAg-seronegative patients and the other for HBeAg positive patients). Participants in both trials were randomly assigned to tenofovir treatment or placebo for 1 year, after which the patients could opt to continue on open-label tenofovir for 8 years. The REACH-B scores were estimated using patients' baseline data. The standardized incidence ratio (SIR) of observed versus expected number of HCC cases, which was estimated according to the REACH-B score, was calculated. The analysis showed that by year 6, the SIR (95%CI) was 0.50 (0.29-0.84), which meant a 50% decrease from the predicted number of HCC cases. The effect of tenofovir was noticeable in non-cirrhotic patients (75% of study patients), in which a SIR (95%CI) of 0.45 (0.23-0.91) could be observed at 6 years of therapy. The authors concluded that the incidence of HCC in patients on tenofovir in both trials was lower than what was predicted by the REACH-B model, and that potent antiviral therapy may reduce the risk of HCC. Of note, among patients with cirrhosis (25% of study patients), the observed numbers of HCC were closer to the curve of expected cases predicted by REACH-B. This result, however, does not necessarily rule out the long-term benefits of tenefovir, because the REACH-B score was generated from non-cirrhotic patients, which may have underestimated actual HCC risk in cirrhotic patients without treatment.

It has been stated in the conclusion of the National Institutes of Health Consensus Development’s statement on the management of hepatitis B[21] that the major goals of anti-HBV therapy are to prevent the development of progressive disease, specifically cirrhosis and liver failure, as well as HCC and subsequent death. However, to date, no randomized clinical trials (RCTs) of anti-HBV therapies have demonstrated a beneficial impact on overall mortality, liver-specific mortality, or development of HCC. Most published reports of hepatitis therapy use changes in short-term virologic, biochemical, and histologic parameters to infer the likelihood of long-term benefit. Approved therapies are associated with improvements in intermediate biomarkers, including suppressed HBV DNA levels, HBeAg loss or seroconversion, decreases in ALT levels, and improvement in liver histology. According to this statement[21], the most important research needs include representative prospective cohort studies to define the natural history of the disease, and large RCTs of monotherapy and combined therapies, including placebo-controlled trials, that measure the effects on clinical health outcomes.

The aforementioned study was a good example in using the risk score developed from a natural history cohort as a scale to quantify the effect of antiviral therapy. For the time being, it is no longer feasible to conduct placebo-controlled RCTs to prove the efficacy of antiviral drugs on long-term outcomes, because they need large populations and a very long time to observe enough patients with outcomes. In addition, when treatments began showing their initial benefits, not treating patients for the sake of an arm of a clinical trial was no longer ethical. In this case, comparing outcomes from treatment cohorts with an existing natural history cohort is much more feasible. In adopting this approach, a difficulty may be encountered - CHB patients who are eligible for antiviral treatment should typically have worse risk profiles. When comparing the long-term risk of disease progression between CHB treatment cohorts and CHB natural history cohorts, it is crucial to take patients’ risk profiles (such as gender, age, ALT, HBeAg, and HBV DNA level) into consideration because these are all important risk predictors of long-term clinical health outcomes. This issue can be overcome by introducing the natural history risk score, as shown by Kim e*t al*[20] which was used as a risk scale to be contrasted with treatment cohorts.

**PRECAUTION AND DISCUSSION OF USING THE RISK SCORES**

Although the risk calculators are easy-to-use and the REACH-B predictive score was externally validated to be an applicable tool for HCC risk estimation, several precautions and discussions should be taken. Because surveillance strategies derived from a Taiwanese population might not apply globally, further validation is still needed in patients of different ethnicities, geographical areas, ages at infection, genetic background, HBV genotypes or species, comorbidities, and exposures to environmental factors such as aflatoxin and alcohol[13,22,23].

Another key question for the REACH-B score is whether this particular risk calculator is suitable for cirrhotic patients. We know that cirrhosis occurs at a relatively late stage in the spectrum of liver disease, and studies indicate that the annual risk of developing HCC among these patients is extremely high[24]. However, to date, there is no HCC risk prediction tool for patients with severe fibrosis and cirrhosis. As the carcinogenesis of HCC is a multistage and multi-factorial process, the risk predictors for developing HCC in cirrhotic patients, in which CHB has already progressed to a relatively late stage, should be very different from those patients without cirrhosis. A universal risk prediction tool for the whole spectrum of patients is thus not reasonable. The predictability of the REACH-B risk score in cirrhotic patients is limited because the derivation cohort did not include CHB patients with this complication. The original REACH-B article showed that the accuracy and values of prediction diminished when applying the risk score to cirrhotic carriers[13]. However, for those in the early stages of liver disease, its accuracy was quite satisfactory. We should also bear in mind that cirrhosis itself is an important predictor for future HCC development. Therefore, patients with existing cirrhosis need close monitoring and timely initiation of antiviral therapy. Risk assessment in this group of patients, or the use of cirrhosis as a variable might therefore be pointless.

The concern has been raised on whether the risk score is suitable to be used for changing risk profiles during follow-up, either spontaneously or as a result of antiviral treatment. Because current HCC risk prediction tools were generated from a natural history cohort without history of antiviral therapy, the inference of predicted risks under circumstances of antiviral therapy should theoretically be inappropriate. Furthermore, current HCC prediction tools are based on one-shot baseline measurements; further validation studies are required to evaluate whether this risk score is applicable for changing risk profiles during follow-up, either changing spontaneously or through antiviral therapy. A risk calculator, which incorporates not only the baseline risk parameters but also the repeated measurements of patients’ risk profiles, would be expected to solve this problem, and thus deserves further efforts.

The risk calculator is a practical tool for managing various phases of a specific disease. Therefore, several other clinical outcomes and milestones of CHB, such as cirrhosis, liver-related mortality, as well as the seroclearance of HBeAg, HBsAg, and HBV DNA, can also be suitable for the development of risk or probability prediction tools. One example is a predictive scoring system for the spontaneous seroclearance of HBsAg in HBeAg-seronegative CHB patients[25]. Besides focusing on outcomes, some emerging clinical parameters might hopefully be incorporated into the current HCC risk score[26-29]. The incorporation of serum quantitative HBsAg titers is a good example; other factors such as host genetic and immunologic markers might also be candidates once their independent predictability are proved, even after taking known risk predictors into consideration. The realization of these prediction tools may benefit CHB patients by providing comprehensive and individualized management instruments, which have been developed based on scientific evidence of risk.

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**Table 1 Scenario for chronic hepatitis B patients with various risk profiles**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
|  | **Patient A** | **Patient B**  | **Patient C**  | **Patient D**  |
| Gender | Male | Female | Male | Female |
| Age (yr) | 70 | 30 | 55 | 40 |
| ALT (U/L) | 100 | 15 | 25  | 40 |
| HBeAg | Positive | Negative | Negative  | Negative |
| HBV DNA (copies/mL) | > 106 | < 300 | 104  | 105 |
| Risk of HCC  | Very high(but the exact risk is unknown) | Very low(but the exact risk is unknown) | Unknown | Unknown |

**Table 2 The hepatocellular carcinoma risk calculators for patients with chronic hepatitis B which have been developed before 2011**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
|  | **IPM[12]** | **GAG-HCC Risk Score[11]** | **CUHK Clinical Scoring System[10]** | **REVEAL Nomograms[9]** |
| Area | Korea | Hong Kong | Hong Kong | Taiwan |
| Origin of subjects | Hospital-based | Hospital-based | Hospital-based | Community-based |
| No. of subjects for derivation | 2020 | 820 | 1005 | 2435 |
| Feature of subjects | Not limited to CHB | CHB | CHB | CHB |
| No. of Model | 1 | 1 | 1 | 3 |
| Risk predictors | * Gender
* Age
* HCV infection
* HBV infection
* AFP level
* Chronic hepatitis
* Cirrhosis
* Heavy alcoholics
* ALT level
 | * Gender
* Age
* HBV DNA level
* CP mutations
* Cirrhosis
 | * Age
* Albumin
* Bilirubin
* HBV DNA level
* Cirrhosis
 | * Gender
* Age
* ALT level
* Family history of HCC
* Alcohol consumption
* HBeAg
* HBV DNA level
* HBV genotype
 |
| Risk function | Logistic regression | Cox regression | Cox regression | Cox regression |
| Predicted outcomes | HCC risk | 5-yr and 10-yr HCC risk | 5-yr and 10-yr HCC risk | 5-yr and 10-yr HCC risk |
| Validation (no. of subjects for validation) | 2-yr prospective validation (*n* = 833) | Leave-one-out cross-validation | External validation (*n* = 424) | External validation with data splitting (*n* = 1218) |

**Table 3 The most updated hepatocellular carcinoma risk score for chronic hepatitis B patients upgraded from the REVEAL nomogram**

|  |  |
| --- | --- |
| Baseline hepatocellular carcinoma predictor  | Risk score |
| Age (yr) |  |
|  30-34 | 0 |
|  35-39 | 1 |
|  40-44 | 2 |
|  45-49 | 3 |
|  50-54 | 4 |
|  55-59 | 5 |
|  60-65 | 6 |
| Sex  |  |
|  Female  | 0 |
|  Male  | 2 |
| Levels of ALT (IU/L)  |  |
|  < 15  | 0 |
|  15-44  | 1 |
|  ≥ 45  | 2 |
| Family history of hepatocellular carcinoma  |  |
|  No  | 0 |
|  Yes  | 2 |
| HBeAg/HBV DNA/HBsAg/Genotype  |  |
|  Negative/<104/<100/any type  | 0 |
|  Negative/<104/100-999/any type  | 2 |
|  Negative/<104/≥1000/any type  | 2 |
|  Negative/104-106/<100/any type  | 3 |
|  Negative/104-106/100-999/any type  | 3 |
|  Negative/104-106/≥1000/any type  | 4 |
|  Negative/≥106/any level/B or B+ C  | 5 |
|  Negative/≥106/any level/C  | 7 |
|  Positive/any level/any level /B or B+C  | 6 |
|  Positive/any level/any level /C  | 7 |

**Figure 1 Known risk factors for hepatocellular carcinoma disease progression for chronic hepatitis B.**

**Figure 2 The updated REVEAL nomogram for the prediction of hepatocellular carcinoma risk.**

**Figure 3 The distributions of score in the derivation cohort (from community) and the validation cohort (from clinical settings) used in the REACH-B study[13].**