Preclinical and clinical studies on the effectiveness of nucleoside analogues in suppressing HBV replication and in ameliorating HBV-related HCC recurrence have been reported to be an independent risk factor for recurrent HBV-related HCC.1-4 Factors that have been shown to be associated with a lower risk of recurrence include serum α-fetoprotein level, cirrhosis, hepatitis B e antigen (HBeAg) status, and HBV viral load.5,6 The association between nucleoside analogue use and risk of HCC recurrence among patients with HBV-related HCC has been investigatedparatively and statistically.7,8 Patients taking nucleoside analogues (treated cohort, n=518) had a lower risk of recurrence (n=106 [20.5%] vs n=1765 [43.6%]; P<.001), and lower overall death (n=55 [10.6%] vs n=1145 [28.3%]; P<.001). After adjusting for competing mortality, the treated cohort had a significantly lower 6-year HCC recurrence rate (45.6% vs 54.6%; 95% CI, 36.5%-54.6% vs untreated, 54.6%; 95% CI, 52.5%-56.6%; P<.001). Six-year overall mortality for treated cohorts was 29.0% (95% CI, 20.0%-38.0%) and for untreated 42.4% (95% CI, 40.0%-44.7%; P<.001). On modified Cox regression analysis, nucleoside analogue use (HR, 0.67; 95% CI, 0.55-0.81; P<.001), statin use (HR, 0.68; 95% CI, 0.53-0.87; P=.002), and nonsteroidal anti-inflammatory drugs or aspirin use (HR, 0.80; 95% CI, 0.73-0.88; P<.001) were independently associated with a reduced risk of HCC recurrence. Multivariable stratified analyses verified the association in all subgroups of patients, including those who were noncirrhotic (HR, 0.56; 95% CI, 0.42-0.76) and diabetic (HR, 0.52; 95% CI, 0.31-0.89).

Conclusion Nucleoside analogue use was associated with a lower risk of HCC recurrence among patients with HBV-related HCC after liver resection.

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For editorial comment see p 1922.

Surgery is considered the standard curative treatment option for hepatocellular carcinoma (HCC). However, the rate of long-term disease-free survival after liver resection remains unsatisfactory due to persistent high incidences of HCC recurrence.1 Many factors affect HCC recurrence risk after liver resection, including tumor size and stage, serum α-fetoprotein level, cirrhosis, hepatitis B e antigen (HBeAg) status, and HBV viral load.2,3 Among these factors, HBV viral load is the most clinically controllable.

Higher HBV viral load has been reported to be an independent risk factor for HCC recurrence in patients with HBV-related HCC.4 Nucleoside analogues are effective in suppressing HBV replication and in ameliorating HBV-related liver disease.5,6 They have been shown to be associated with a lower risk of HCC and other cirrhosis-related complications in those with chronic hepatitis.7,8 However, studies on the effectiveness of nucleoside analogues for HCC recurrence are limited.

Association Between Nucleoside Analogues and Risk of Hepatitis B Virus–Related Hepatocellular Carcinoma Recurrence Following Liver Resection

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Ming-Shiang Wu, MD, PhD
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Context Tumor recurrence is a major issue for patients with hepatocellular carcinoma (HCC) following curative liver resection.

Objective To investigate the association between nucleoside analogue use and risk of tumor recurrence in patients with hepatitis B virus (HBV)–related HCC after curative surgery.

Design, Setting, and Participants A nationwide cohort study between October 2003 and September 2010. Data from the Taiwan National Health Insurance Research Database. Among 100 938 newly diagnosed HCC patients, we identified 4569 HBV-related HCC patients who received curative liver resection for HCC between October 2003 and September 2010.

Main Outcome Measures The risk of first tumor recurrence was compared between patients not taking nucleoside analogues (untreated cohort, n=4051) and patients taking nucleoside analogues (treated cohort, n=518). Cumulative incidences and hazard ratios (HRs) were calculated after adjusting for competing mortality.

Results The treated cohort had a higher prevalence of liver cirrhosis when compared with the untreated cohort (48.6% vs 38.7%; P<.001), but lower risk of HCC recurrence (n=106 [20.5%] vs n=1765 [43.6%]; P<.001), and lower overall death (n=55 [10.6%] vs n=1145 [28.3%]; P<.001). After adjusting for competing mortality, the treated cohort had a significantly lower 6-year HCC recurrence rate (45.6% vs 54.6%; 95% CI, 36.5%-54.6% vs untreated, 54.6%; 95% CI, 52.5%-56.6%; P<.001). Six-year overall mortalities for treated cohorts were 29.0% (95% CI, 20.0%-38.0%) and for untreated 42.4% (95% CI, 40.0%-44.7%; P<.001). On modified Cox regression analysis, nucleoside analogue use (HR, 0.67; 95% CI, 0.55-0.81; P<.001), statin use (HR, 0.68; 95% CI, 0.53-0.87; P=.002), and nonsteroidal anti-inflammatory drugs or aspirin use (HR, 0.80; 95% CI, 0.73-0.88; P<.001) were independently associated with a reduced risk of HCC recurrence. Multivariable stratified analyses verified the association in all subgroups of patients, including those who were noncirrhotic (HR, 0.56; 95% CI, 0.42-0.76) and diabetic (HR, 0.52; 95% CI, 0.31-0.89).

Conclusion Nucleoside analogue use was associated with a lower risk of HCC recurrence among patients with HBV-related HCC after liver resection.

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HBV, NUCLEOSIDE ANALOGUES, AND HCC RECURRENCE

side analogue use in HCC recurrence have been relatively limited and have yielded conflicting results.12-14 In Taiwan, under the National Health Insurance (NHI) program, reimbursement for nucleoside analogues for HBV patients meeting certain criteria began on October 1, 2003 (eTable 1 and eTable 2 available at http://www.jama.com).

The purpose of this study was to examine the association between use of nucleoside analogues and risk of HCC recurrence among patients with HBV-related HCC after curative liver resection. We also examined this association among different subpopulations and calculated the number needed to be treated (NNT) for 1 less HCC recurrence.

METHODS
Study Design
We conducted a nationwide cohort study by retrieving all patients receiving curative liver resection for HCC from Taiwan’s National Health Insurance Research Database (NHIRD). The NHIRD has been described in detail in previous studies.15-17 In brief, it consists of detailed health care data from more than 25 million enrollees, representing more than 99% of Taiwan’s entire population. The accuracy of diagnosis of major diseases in the NHIRD, such as stroke and acute coronary syndrome, has been validated.17,18 This study has been approved by the ethical review board of the National Health Research Institutes, Taiwan.

Study Population
We identified all hospitalized patients who were admitted with a primary diagnosis of HCC (International Classification of Diseases, Ninth Revision [ICD-9] codes 155.0, 155.2) for the first time and who received curative liver resection between October 1, 2003, and September 30, 2010. The diagnostic accuracy of HCC was confirmed by both specific admission ICD-9 codes and inclusion in the Registry for Catastrophic Illness Patient Database (RCIPD), a subpart of the NHIRD.15,16 Surgical pathological confirmation or typical image presentation of HCC is required for patients to be registered in the RCIPD.

Only HCC patients with HBV infection (ICD-9 codes 070.2, 070.3, and V02.61) were included in our study cohorts. Patients were excluded if diagnosed with hepatitis C (ICD-9 codes 070.41, 070.44, 070.51, 070.54, and V02.62), other viral hepatitis (ICD-9 code V02.69), malignant tumor (ICD-9 codes 140-208), or if they received antiviral treatments for more than 3 months before the index admission. Patients were also excluded if they received liver resection, transarterial chemoembolization, percutaneous ethanol injection, radiofrequency ablation, or liver transplantation before the index hospitalization.

Main Outcome Measurements
Hepatocellular carcinoma recurrence was defined as rehospitalization with a primary diagnosis of HCC after the index admission date and a treatment modality for HCC recurrence, such as surgery, transarterial chemoembolization, percutaneous ethanol injection, radiofrequency ablation, or liver transplantation during the study period. Patients with HCC recurrence in the first 3 months after the index hospitalization for liver resection were excluded. We used the incident user design with follow-up for each patient beginning on the date of first prescription of nucleoside analogue in the treated cohort.19,20 The follow-up for the untreated cohort began on the first day after the index hospitalization for liver resection. Both cohorts were followed up until the date of HCC recurrence, death, or the end of 2010. Death was defined as withdrawal of the patient from the NHI program. Causes of death were defined according to the primary diagnosis of hospitalization in the 3 months preceding death.

Covariate Assessment
Propensity score was calculated using logistic regression as proposed by Rosenbaum and Rubin21,22 to estimate the probabilities of assigning a patient to the treated cohort given the background variables including age, sex, extent of resection, liver cirrhosis, other comorbidities listed in TABLE 1, and use of statins, nonsteroidal anti-inflammatory drugs (NSAIDs), and metformin. The mean and median propensity scores were compared between the 2 cohorts.

Since the chance of HCC recurrence can be confounded by competing risk of mortality, we identified comorbidities that may be associated with mortality based on diagnostic codes from both outpatient and inpatient datasets prior to the outcome of interest. All diseases included in the Charlson Comorbidity Index were analyzed except for human immunodeficiency virus, metastatic solid tumor, and cancer, because patients with these conditions were excluded from the present study.23 Comorbidities, listed by ICD-9 code, included acute coronary syndrome (410-414), cerebrovascular accident (430-438), chronic obstructive pulmonary disease (490-496), diabetes mellitus (250), liver cirrhosis (571.5), liver failure (570), renal failure (584-586), hypertension (401-405), hyperlipidemia (272.0-272.2), and peptic ulcer disease (531-534).
Certain drugs, including statins, aspirin, NSAIDs, and metformin, which might alter the risk of recurrent HCC, were analyzed. Exposure to these drugs was defined as frequency of use of more than 1 tablet per month during the observation period. Liver resection modalities for treatment of the original HCC, including major resection (extensive and partial hepatic lobectomy with at least 3 segmental resections of liver parenchyma) and minor resection (extensive and partial hepatectomy with 2 or fewer segmental resections of liver), were also analyzed.

Table 1. Study Cohorts Following Liver Resection

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Untreatedb (n = 4051)</th>
<th>Treatedb (n = 518)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, mean (SD), y</td>
<td>54.6 (12.5)</td>
<td>54.4 (11.8)</td>
<td>.80</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Women</td>
<td>716 (17.7)</td>
<td>83 (16.0)</td>
<td>.39</td>
</tr>
<tr>
<td>Men</td>
<td>3335 (82.3)</td>
<td>435 (84.0)</td>
<td></td>
</tr>
<tr>
<td>Resectionc</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Major</td>
<td>1431 (35.3)</td>
<td>139 (26.8)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Minor</td>
<td>2620 (64.7)</td>
<td>379 (73.2)</td>
<td></td>
</tr>
<tr>
<td>Follow-up, y</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean (SD)</td>
<td>2.18 (1.77)</td>
<td>2.64 (1.74)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>1.57 (0.77-3.15)</td>
<td>2.18 (1.21-3.69)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Interval to start therapy, y</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean (SD)</td>
<td>1.19 (1.38)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>0.66 (0.14-1.84)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Survival duration, y, Mean (SD)</td>
<td>-</td>
<td>1.45 (1.38)</td>
<td></td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>0.95 (0.48-1.94)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hospital visits</td>
<td></td>
<td></td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Mean (SD)</td>
<td>56.54 (57.77)</td>
<td>74.37 (68.17)</td>
<td></td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>37 (19-73)</td>
<td>60 (32-99.75)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Drug userd</td>
<td>Statins</td>
<td>158 (3.9)</td>
<td>.55</td>
</tr>
<tr>
<td></td>
<td>17 (3.3)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NSAIDs or aspirin</td>
<td>2181 (53.8)</td>
<td>271 (52.3)</td>
<td>.51</td>
</tr>
<tr>
<td>Metformin</td>
<td>498 (12.3)</td>
<td>60 (11.6)</td>
<td>.72</td>
</tr>
<tr>
<td>Major coexisting diseases</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cirrhosis</td>
<td>1569 (38.7)</td>
<td>252 (48.6)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Hypertension</td>
<td>978 (24.1)</td>
<td>109 (21.0)</td>
<td>.13</td>
</tr>
<tr>
<td>Diabetes</td>
<td>663 (16.4)</td>
<td>72 (13.9)</td>
<td>.16</td>
</tr>
<tr>
<td>Peptic ulcer diseases</td>
<td>603 (14.9)</td>
<td>78 (15.1)</td>
<td>.90</td>
</tr>
<tr>
<td>COPD</td>
<td>206 (5.1)</td>
<td>19 (3.7)</td>
<td>.20</td>
</tr>
<tr>
<td>Acute coronary syndrome</td>
<td>178 (4.4)</td>
<td>14 (2.7)</td>
<td>.08</td>
</tr>
<tr>
<td>Cerebral vascular disease</td>
<td>147 (3.6)</td>
<td>9 (1.7)</td>
<td>.03</td>
</tr>
<tr>
<td>Renal failure</td>
<td>83 (2.0)</td>
<td>5 (1.0)</td>
<td>.12</td>
</tr>
<tr>
<td>Hypercholesteremia</td>
<td>35 (0.9)</td>
<td>2 (0.4)</td>
<td>.43</td>
</tr>
<tr>
<td>Propensity score</td>
<td>Mean (SD)</td>
<td>0.11 (0.03)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td></td>
<td>0.12 (0.03)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>0.11 (0.08-0.14)</td>
<td>0.12 (0.10-0.15)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Charlson scoree</td>
<td>Mean (SD)</td>
<td>1.20 (1.68)</td>
<td>.84</td>
</tr>
<tr>
<td></td>
<td>1.22 (1.53)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Median (IQR)</td>
<td>1 (0-1)</td>
<td>1 (0-2)</td>
<td>.16</td>
</tr>
</tbody>
</table>

Abbreviations: COPD, chronic obstructive pulmonary disease; IQR, interquartile range; NSAIDs, nonsteroidal anti-inflammatory drugs.

Statistical Analysis

Death prior to tumor recurrence was considered a competing risk event. The death-adjusted cumulative incidences of HCC recurrence were calculated using a 2-step process and tested for equality among the study cohorts. Calculation and comparison of cumulative incidences in competing risk data ratios were conducted using modified Kaplan-Meier method and Gray method. We tested differences in the full time-to-event distributions between the study groups using log-rank test.

To determine the independent risk factors for HCC recurrence, multivariable analyses and stratified analyses using hazard ratios (HRs) were carried out with modified Cox proportional hazards models in the presence of competing risk event after adjusting for age, sex, resection modality, liver cirrhosis, diabetes, propensity score, and use of statins, NSAIDs or aspirin, and metformin. To assess the dose-dependent association of recurrence with nucleoside analogue use, we further conducted multivariable analysis with nucleoside analogue use as a continuous variable. On multivariable stratified analyses, the association between nucleoside analogue use and the risk of HCC recurrence was examined in different subgroups. All subgroup comparisons were preplanned to control for potential confounding factors reported in previous studies.

The NNT represented the number of patients who needed to be treated for 1 less HCC recurrence or mortality; it was calculated by the inverse of the absolute risk reduction. All results in the present study originated from a nationwide registry database. Hepatocellular carcinoma is defined as a major disease and HCC patients can apply for a catastrophic illness certificate, which grants exemption from co-payment. It is nearly impossible for these HCC patients to withdraw from the NHI program before death. Therefore, there were no missing data or loss of follow-up in our study population.

All data management was performed using SAS 9.2 software (SAS Institute Inc.). Calculations of cumulative incidences and Cox models in the competing risk analyses were carried out using the “cmprsk” package.
package of R (http://cran.r-project.org/web/packages/cmprsk/index.html). Calculated results were expressed as the estimated number together with the 95% confidence interval. Based on statistical power at 0.9, type I error rate at .05, and our case numbers in both cohorts, the detectable risk difference was estimated to be 0.02.

RESULTS
Demographic Characteristics of the HCC Cohort
We first identified 100,938 potentially eligible HCC patients admitted for the first time and registered in the RCIPD. We excluded 78,948 patients who did not receive liver resection and 8173 patients who received liver resection before October 1, 2003. In addition, 7870 patients without HBV infection or coinfection with hepatitis C or other hepatitis were excluded. Those using antiviral agents 3 months before liver resection or for less than 3 months were also not enrolled. There were 1019 patients who received other liver therapy or with another type of tumor before liver resection who were excluded. Another 359 patients with follow-up for less than 3 months were not included in the study. Therefore, 4569 patients were enrolled into the study cohorts (untreated, 4051 patients; treated, 518 patients) (Figure 1). In the treated cohort, 487 patients received only 1 nucleoside analogue, including 159 patients who received lamivudine, 292 patients who received entecavir, and 36 patients who received telbivudine. The remaining patients received more than 1 nucleoside analogue.

Demographic characteristics, confounding drugs, comorbidities, and follow-up durations of the study cohorts are presented in Table 1. The follow-up durations for the untreated cohort were a mean (SD) of 2.18 (1.77) years and a median (IQR) of 1.57 (0.77-3.15) years, and for the treated cohort, a mean (SD) of 2.64 (1.74) years and a median (IQR) of 2.18 (1.21-3.69) years. The mean (SD) interval to start of antiviral therapy after liver resection was 1.19 (1.38) years and the median (IQR) was 0.66 (0.14-1.84) years. The mean (SD) duration of nucleoside analogue use in treated patients was 1.45 (1.38) years and the median (IQR) was 0.95 (0.48-1.94) years. The mean (SD) propensity score for the untreated cohort was 0.11 (0.03) and the median (IQR) was 0.11 (0.09-0.14), and for the treated cohort, the mean (SD) was 0.12 (0.03) and the median (IQR) was 0.12 (0.10-0.15). The treated cohort had a significantly higher prevalence of liver cirrhosis (48.6%) when compared with the untreated cohort (38.7%) (P < .001).

Six-Year Cumulative Incidences of HCC Recurrence and Overall Mortality
Among the 359 patients excluded with follow-up of less than 3 months there were 226 HCC recurrences and 93 deaths. During the observation period, 1765 patients in the untreated cohort (43.6%) and 106 patients in the treated cohort (20.5%) developed HCC recurrence. Death before the recurrence of HCC was defined as competing mortality. Compared with the untreated cohort (451 deaths; 11.1%), only 26 patients (5.0%) died before HCC recurrence in the treated cohort. The major identifiable causes of competing mortality in the untreated cohort were HCC or HCC treatment–related mortality (314 patients), liver cirrhosis (14), pneumonia (12), sepsis (9), and gastrointestinal bleeding (7). The identifiable causes of competing mortality in the treated cohort included HCC or HCC treatment–related mortality (20 patients), pneumonia (2), liver cirrhosis (1), and sepsis (1). During the follow-up period, overall deaths for the untreated and treated cohorts were 1145 (28.3%) and 55 (10.6%), respectively. For overall mortality, the major identifiable causes of death in the untreated cohort were HCC or HCC treatment–related mortality (882 patients, liver cirrhosis (47), sepsis (28), pneumonia (22), and gastrointestinal bleeding (21). The major identifiable causes of overall mortality in the treated cohort were HCC or HCC treatment–related mortality (38 patients), liver cirrhosis (5), pneumonia (5), and sepsis (3).

In Figure 2, cumulative incidences of HCC recurrence after adjustment for competing mortality are shown. The risk of HCC recurrence was significantly lower for patients in the treated cohort (6-year cumulative incidence, 45.6%; 95% CI, 36.5%-54.6%) than for patients in the untreated cohort (54.6%; 95% CI, 52.5%-56.6%) (P < .001). The difference in 6-year cumulative incidence was 9%. The unadjusted NNT associated with 1 less HCC recurrence within 6 years was 12 (95% CI, 7.4-22.6). This implies that use of nucleoside analogues in 12 HCC patients af-
ter liver resection is associated with 1 less HCC recurrence within 6 years.

In the eFigure, we stratified patients by liver cirrhosis and NSAID use. We found that nucleoside analogue use was associated with lower risk of HCC recurrence in noncirrhotic patients, but not in cirrhotic patients. For NSAID users and nonusers, use of nucleoside analogues was associated with reduced risk of HCC recurrence.

Likewise, the risk of overall mortality was significantly lower in patients in the treated cohort (6-year cumulative incidence, 29.0%; 95% CI, 20.0%-38.0%) than in patients in the untreated cohort (42.4%; 95% CI, 40.0%-44.7%) (P < .001) (Figure 2). These observations further confirmed the association between nucleoside analogue use and attenuated risk of HCC recurrence in HBV-related HCC patients after liver resection.

**Multivariable Analysis**

Compared with the untreated cohort, the treated cohort was associated with a significantly lower risk of HCC recurrence (HR, 0.67; 95% CI, 0.55-0.81; P < .001). Use of statins (HR, 0.68; 95% CI, 0.53-0.87; P = .002) and use of NSAIDs or aspirin (HR, 0.80; 95% CI, 0.73-0.88; P < .001) were significantly associated with lower risk of tumor recurrence. Liver cirrhosis was found to be an independent risk factor for HCC recurrence (HR, 1.23; 95% CI, 1.12-1.35; P < .001) (Table 2). Each incremental year of use of nucleoside analogues was associated with reduced risk of HCC recurrence (HR, 0.59; 95% CI, 0.51-0.68; P < .001) (eTable 3).

**Multivariable Stratified Analysis**

The treated cohort was found to be associated with a reduced risk of HCC recurrence on all stratified analyses, including for noncirrhotic patients (HR, 0.56; 95% CI, 0.42-0.76) and diabetic patients (HR, 0.52; 95% CI, 0.31-0.89) (Figure 3). These observations further confirmed the association between nucleoside analogue use and attenuated risk of HCC recurrence in HBV-related HCC patients after liver resection.

**COMMENT**

The roles of HBV in HCC recurrence have been widely investigated. Kubo et al. first reported that high viral load is associated with lower risk of HCC recurrence after liver resection. The association between serum HBV loads and risk of HCC recurrence after liver resection or transcatheter arterial embolization has been confirmed in previous studies.

In the present study, we did not have data regarding a patient’s HBV viral load or liver function. However, Taiwan’s NHI program has strict regulations regarding reimbursement for nucleoside analogues. Reimbursement is granted only to patients in high-risk populations (eTable 1). Under such regulations, patients in the treated cohort should have a higher baseline HBV viral load, higher ALT level, or higher prevalence of liver decompensation than those in the untreated cohort to be eligible for reimbursement. In the present study, the treated cohort had higher prevalence of liver cirrhosis. If higher HBV viral load is associated with higher risk of HCC recurrence, higher baseline HBV viral load in the treated patients may have led to a more conservative estimation of the association in the present study.

Although it is generally accepted that HBV viral load plays an important role...
in HCC recurrence, studies regarding the effectiveness of nucleoside analogues in 
HCC recurrence have been very limited and have produced conflicting re-

csults.28 Based on a study of 14 patients 
receiving lamivudine and 10 control par-
cipants, Kubo et al13 found a lower 
5-year HCC disease-free survival rate af-
	er surgery in the lamivudine-treated pa-
tients. Chan et al14 reported that the 
5-year tumor free survival rates in the 
lamivudine or entecavir-treated group 
(42 participants; 51.4%) were signifi-
cantly higher than in the control group 
(94 participants; 33.8%). In contrast, Li 

The present study, we con-
ermed the association between 
nucleoside analogue use and reduced 
risk of HCC recurrence in HBV pa-
tients after liver resection based on a na-
tionwide database. However, reimburse-
ment for oral antiviral agents for HBV 
fection is strictly limited to specified 
indications in Taiwan, and HCC does 
not qualify as an indication. Therefore, 
most of the HBV-infected HCC pa-
tients in the present study did not ful-
fill the NHI criteria for oral antiviral 
therapy. 

In previous studies, HCC with cir-
rhosis has been associated with de-
creased overall survival compared with 
HCC without cirrhosis after curative 

liver resection.2 However, the out-
come of patients with HBV-related com-

pensated cirrhosis has not been shown to 
be worse than that of noncirrhotic patients.4 In the present study, nucleo-
side analogue use was not only associ-
ed with improved disease-free sur-
ival, but also with improved overall 

Survival in HBV-related HCC after liver 

resection. The higher prevalence of cir-
rhosis in the treated cohort was the 
main reason that the absolute differ-
ence in HCC recurrence rates (54.6% 
vs 45.6%) was much smaller than the 
HR (0.67) observed on multivariable 

analysis. 

On multivariable analysis, we found that statin and NSAID or aspirin use 

were associated with a lower risk of 

HCC recurrence. The protective role of 

statins in HBV-infected HCC has been 
reported in a recent study.30 The po-

tential mechanisms involve AMP-

activated protein kinase, p21 expres-
sion, endoplasmic reticulum stress, and 
autoaphagy.31 The association between 

the use of NSAIDs or aspirin and a lower 

risk of HCC recurrence is a novel find-
ing. Aspirin has been reported to in-

duce cell cycle arrest and apoptosis, me-
diated by increased metabolic and 

oxidative stress.32 An in vivo study 

showed that aspirin results in tumor 

growth inhibition.33 More recently, Si-
tia et al demonstrated that aspirin di-

minishes the number of intrahepatic 

HBV-specific CD8(+) T cells and HBV-
nonspecific inflammatory cells, the se-

verity of liver fibrosis, and the de-

velopment of HCC in an HBV transgenic 

mouse model.34 Based on prior studies 

and our results, we postulated that 

NSAIDs, including aspirin, may be ben-

eficial for reducing tumor recurrence 

in HCC patients. Future randomized 

controlled trial studies are necessary to 

clarify this issue. 

There are several limitations to 
the present study. First, a causal associa-

| Table 2. Risk of HCC Recurrence After Adjusting for Competing Mortality
<table>
<thead>
<tr>
<th>No.</th>
<th>HCC Recurrence No.</th>
<th>HR (95% CI)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Treated vs untreated</td>
<td>Untreated</td>
<td>4051</td>
<td>1765</td>
</tr>
<tr>
<td></td>
<td>Treated</td>
<td>518</td>
<td>106</td>
</tr>
<tr>
<td>Age</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;50 y</td>
<td>1568</td>
<td>655</td>
<td>1 [Reference]</td>
</tr>
<tr>
<td>50-59 y</td>
<td>1466</td>
<td>590</td>
<td>0.96 (0.83-1.10)</td>
</tr>
<tr>
<td>≥60 y</td>
<td>1536</td>
<td>626</td>
<td>1.01 (0.90-1.13)</td>
</tr>
<tr>
<td>Sex</td>
<td>Women</td>
<td>799</td>
<td>306</td>
</tr>
<tr>
<td></td>
<td>Men</td>
<td>3770</td>
<td>1565</td>
</tr>
<tr>
<td>Resection</td>
<td>Minor</td>
<td>2999</td>
<td>1228</td>
</tr>
<tr>
<td></td>
<td>Major</td>
<td>1570</td>
<td>643</td>
</tr>
<tr>
<td>Liver cirrhosis</td>
<td>Yes</td>
<td>2748</td>
<td>1046</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>1821</td>
<td>625</td>
</tr>
<tr>
<td>Diabetes</td>
<td>Yes</td>
<td>735</td>
<td>317</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>3834</td>
<td>1554</td>
</tr>
<tr>
<td>Statin use</td>
<td>Yes</td>
<td>4394</td>
<td>1814</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>175</td>
<td>57</td>
</tr>
<tr>
<td>NSAID or aspirin</td>
<td>Yes</td>
<td>2452</td>
<td>939</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>2117</td>
<td>932</td>
</tr>
<tr>
<td>Metformin use</td>
<td>Yes</td>
<td>2452</td>
<td>939</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>4011</td>
<td>1634</td>
</tr>
<tr>
<td>Propensity score</td>
<td>Yes</td>
<td>558</td>
<td>237</td>
</tr>
<tr>
<td></td>
<td>No</td>
<td>4569</td>
<td>1871</td>
</tr>
</tbody>
</table>

Abbreviations: HCC, hepatocellular carcinoma; HR, hazard ratio; NSAID, nonsteroidal anti-inflammatory drug.

a Adjusted for covariate factors, including age, sex, resection extent, liver cirrhosis, diabetes, and use of statins, NSAIDs, aspirin, and metformin.
b Treated and untreated categories indicate patients with hepatitis B virus who are receiving nucleoside analogues, and those who are not, respectively.
c Major resection indicates extensive and partial hepatic lobectomy with at least 3 segmental resections of liver paren-
hyma; minor resection indicates extensive and partial hepatectomy with 2 or fewer segmental resections of liver.
d Drug users indicate patients using a drug at least 1 day per month on average.
tion between a drug of interest and risk of HCC recurrence cannot be inferred based on an observational study. Confounding by indication may exist and account for differences in outcomes. The patients in the study cohorts may differ in many measured and unmeasured ways. We did not have personal information for our patients such as lifestyle, family history of malignant diseases, body mass index, or laboratory parameters including HBV DNA viral load, which may contribute to tumor recurrence risk. To avoid these biases, we selected only patients receiving curative liver resection because resectable patients are comparable in terms of disease extent and remnant liver function. We analyzed propensity scores and Charlson scores to examine the comparability of these 2 cohorts. Multivariable analysis was performed to adjust for potential confounders. Furthermore, we conducted multivariable stratified analysis to examine the risk of HCC recurrence after liver resection for the study cohorts in different strata. Although unmeasured confounders may still exist, we believe the methodology used in the present study is solid and robust.

Second, coding error is possible in a database. We were unable to check the accuracy of nucleoside analogue use in the NHIRD. However, the information regarding insurance-paid nucleoside analogues was accurate because every prescription is strictly regulated and only patients fitting specific criteria are eligible to receive reimbursement.

Third, some patients may have used self-paid nucleoside analogues and thus may have been inappropriately classified into the untreated cohort. Conversely, nucleoside analogue users in the treated cohort may have poor compliance. These potential misclassifications may have led to an underestimation of the association.

Fourth, HCC recurrence was defined as rehospitalization with a primary diagnosis of HCC and further HCC therapy. It is possible that some incidence of HCC recurrence might have been missed, for example if a patient did not receive therapy for the recurrence. Since all enrolled participants were able to undergo curative resection, the majority of them were unlikely to give up therapy for a recurrence. Furthermore, such misclassification would have underestimated the incidence of recurrent HCC in both cohorts and biased the results toward null difference.

Fifth, information about adverse events of nucleoside analogues was not available from the NHIRD. However,
there is evidence of excellent short-term safety profiles for the available nucleoside analogues for chronic hepatitis B. Therefore, adverse events associated with oral anti-HBV therapy were very unlikely to substantially influence clinical outcomes in the present study. Sixth, we did not treat nucleoside analogue use as a time-dependent variable in the Cox model. Instead, we used the incident user design in this study, in which exposure time begins with the start of a new antiviral agent. This minimizes the potential for immortal person-time bias prior to treatment exposure, which may result in a downward trend toward understimation of the risk rate ratios. In conclusion, nucleoside analogue use was associated with a lower risk of HCC recurrence among patients with HBV-related HCC after liver resection.

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Author Contributions: Dr C.-Y. Wu had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis. Drs C.-Y. Wu and Chen contributed equally to this article as first authors; Drs C.-Y. Wu and Lin are both corresponding authors. Study concept and design: Chun-Ying Wu, Chen, Kuo, Lin. Acquisition of data: Chun-Ying Wu, Ho, Lin. Analysis and interpretation of data: Chun-Ying Wu, Chen, Ho, Hsu, Ming-Shiang Wu, Lin. Drafting of the manuscript: Chun-Ying Wu, Chen. Critical revision of the manuscript for important intellectual content: Chun-Ying Wu, Chen, Ho, Hsu, Kuo, Ming-Shiang Wu, Lin. Statistical analysis: Chun-Ying Wu, Chen, Ho, Hsu. Obtained funding: Chun-Ying Wu, Kuo. Administrative, technical, or material support: Chun-Ying Wu, Chen, Kuo, Ming-Shiang Wu, Lin. Study supervision: Chun-Ying Wu, Kuo, Lin.

Conflict of Interest Disclosures: All authors have completed and submitted the ICMJE Form for Disclosure of Potential Conflicts of Interest and none were reported.

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Online-Only Material: eTables 1 to 3 and an efigure are available at http://www.jama.com.

REFERENCES


