

Rising alcohol-related liver cancer prompts new prediction tool

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A. ROC comparison



	Internal cohort	External cohort
Cut-off value = 90		
Sensitivity (%)	99.8 (99.5 - 99.9)	99.8 (99.1 - 100.0)
Specificity (%)	4.1 (3.6 - 4.7)	4.2 (3.0 - 5.6)
PPV (%)	27.9 (27.8 - 28.0)	38.4 (38.1 - 38.7)
NPV (%)	98.2 (95.3 - 99.3)	97.7 (85.6 - 99.7)
Cut-off value = 160		
Sensitivity (%)	51.7 (49.4 - 53.9)	51.3 (47.3 - 55.3)
Specificity (%)	84.0 (83.0 - 85.0)	82.7 (80.3 - 85.0)
PPV (%)	54.6 (52.7 - 56.4)	64.0 (60.3 - 67.4)
NPV (%)	82.4 (81.7 - 83.0)	85.9 (83.5 - 87.9)

B. HCC prediction by nomogram

	0		Leve viele energy	Nomogram-estimated patients distribution		Nomogram- estimated	Actually-observed HCC incidence	
90)	Ŭ	Ĭ	Low-risk group	Internal	External	incidence	Internal	External
Low (0-9			T	3.0%	2.6%	<5%	1.8%	2.3%
Medium risk (91-160)	90		Medium-risk group ÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅÅ	71.3%	67.2%	5%-50%	18.3%	27.0%
High risk (161-240)	240		High-risk group	25.7%	30.2%	>50%	54.6%	63.6%

(A) Validation of the nomogram scores. The nomogram score of each patient in both internal and external cohort was calculated and plotted by ROC. (B) HCC risk stratification by the nomogram. Patients with ALD could be categorized into low-risk group (0–90 points) with less than 5% probability of HCC, intermediate-



risk group (91–160 points) with about 20.0–50% probability of HCC and highrisk group (161–240 points) with over 50% probability of HCC. ALD, alcoholassociated liver disease; AUC, area under the curve; HCC, hepatocellular carcinoma; NPV, negative predictive value; PPV, positive predictive value; ROC, receiver operating characteristic curve. Credit: Chang B, Tian H, Huang A, et al.

Liver cancer is the sixth most common cancer and the third most frequent cause of cancer-related death globally. However, its distribution and causes vary greatly across different regions. While areas like Eastern Asia and sub-Saharan Africa see the most cases, the reasons behind them differ significantly.

In <u>high-income countries</u>, liver cancer has been on the decline thanks to widespread newborn hepatitis B vaccination and antiviral drugs. Meanwhile, <u>low-income countries</u> witness a worrying rise, often linked to increased hepatitis B and C infections and injectable drug use.

While viral hepatitis remains a major concern, another factor is gaining attention: <u>alcohol consumption</u>. Studies show that chronic alcohol consumption can directly cause about 10% of cancer cases in men and 3%, respectively, in women. In fact, a study at the Mayo Clinic revealed that alcoholic cirrhosis was the main culprit in 29% of patients with hepatocellular carcinoma (HCC), the most common liver cancer type.

But the study doesn't just sound the alarm; it also offers a potential solution. Researchers identified key risk factors for HCC in people with <u>alcohol-related liver disease</u>, including <u>heavy drinking</u>, age, diabetes, male sex, and liver cirrhosis. Based on these factors, they developed a novel tool called a nomogram that can predict HCC risk with high accuracy and ease of use.





AILD, autoimmune-induced liver disease (including autoimmune hepatitis, primary biliary cirrhosis and primary sclerosing cholangitis); ALD, alcoholassociated liver disease; DILI, drug-induced liver injury; HB, hepatitis B; HC, hepatitis C; HCC, hepatocellular carcinoma; OLD, other liver diseases (including non-alcoholic fatty liver disease, Wilson's disease, Buddi-Chiari syndrome, liver injury with unknown reasons, etc). Credit: Chang B, Tian H, Huang A, et al.

This nomogram could be a game-changer for doctors, allowing them to personalize treatment plans and identify individuals at the highest risk for HCC. Early intervention could save lives and prevent unnecessary suffering.

This study underscores the growing public health concern of alcoholrelated liver cancer. The newly developed nomogram offers a valuable tool for doctors to identify high-risk individuals and personalize



treatment plans, potentially saving lives and preventing unnecessary suffering.

While the study provides valuable insights, it acknowledges limitations like its retrospective nature and the need for further validation in larger, prospective studies. Additionally, incorporating other factors like smoking, genetics, and dietary habits could further improve the prediction model.



(A) Change of aetiology in the entire enrolled patients with chronic liver diseases. (B) Change of aetiology in the patients with alcohol-associated liver



disease (ALD). (C) Change of aetiology in the patients with HCC. (D) Change of aetiology in the patients with ALD-related HCC. AILD, autoimmune-induced liver disease; DILI, drug-induced liver injury; HBV, hepatitis B virus; HCC, hepatocellular carcinoma; HCV, hepatitis C virus. Credit: Chang B, Tian H, Huang A, et al.

The researchers also highlight the need for future research on <u>non-alcoholic fatty liver disease</u> and its link to liver cancer, as this area remains under-investigated.

Overall, this study shines a light on the rising threat of alcohol-related <u>liver cancer</u> and offers a promising tool for early detection and intervention. Further research and public health efforts are crucial to combat this growing health challenge.

The findings are **<u>published</u>** in the journal *eGastroenterology*.

More information: Binxia Chang et al, Prevalence and prediction of hepatocellular carcinoma in alcohol-associated liver disease: a retrospective study of 136 571 patients with chronic liver diseases, *eGastroenterology* (2024). DOI: 10.1136/egastro-2023-100036

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