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Reporting Summary

Nature Research wishes to improve the reproducibility of the work that we publish. This form provides structure for consistency and transparency in reporting. For further information on Nature Research policies, see Authors & Referees and the Editorial Policy Checklist.

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For	all statistical analys	es, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.
n/a	Confirmed	
	The exact sam	ple size (n) for each experimental group/condition, given as a discrete number and unit of measurement
	A statement of	on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
	The statistical Only common to	test(s) used AND whether they are one- or two-sided ests should be described solely by name; describe more complex techniques in the Methods section.
\boxtimes	A description	of all covariates tested
\boxtimes	A description	of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
	A full descript AND variation	ion of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
	For null hypot Give P values as	thesis testing, the test statistic (e.g. F , t , r) with confidence intervals, effect sizes, degrees of freedom and P value noted exact values whenever suitable.
\boxtimes	For Bayesian a	analysis, information on the choice of priors and Markov chain Monte Carlo settings
\boxtimes	For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes	
\boxtimes	\boxtimes Estimates of effect sizes (e.g. Cohen's d , Pearson's r), indicating how they were calculated	
	'	Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.
So	ftware and c	ode
Poli	cy information abo	ut <u>availability of computer code</u>
Da	ata collection	Provide a description of all commercial, open source and custom code used to collect the data in this study, specifying the version used OR

state that no software was used.

Data analysis

Provide a description of all commercial, open source and custom code used to analyse the data in this study, specifying the version used OR state that no software was used.

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors/reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Research guidelines for submitting code & software for further information.

Data

Policy information about availability of data

All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A list of figures that have associated raw data
- A description of any restrictions on data availability

The crystal structures presented in this work have been deposited in the Protein Data Bank (PDB) with accession codes 6J10 [https://www.rcsb.org/structure/6J10]. Data supporting the findings of this manuscript are available from the corresponding authors upon reasonable request. A reporting summary for this article is available as a Supplementary Information file. The source data underlying Figs 1-7 and Supplementary Figs 1, 2, 8, 11-15 can be provided as a Source Data file.

Field-spe	cific reporting		
Please select the or	ne below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.		
∑ Life sciences	Behavioural & social sciences Ecological, evolutionary & environmental sciences		
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Life scier	ices study design		
All studies must dis	sclose on these points even when the disclosure is negative.		
Sample size	We chose the sample size based on literatures in the field.		
Data exclusions	ot applicable		
Replication	Experiment data was well repeated.		
Randomization	Not applicable		
Blinding	Not applicable		
Reportin	g for specific materials, systems and methods		
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Materials & exp	perimental systems Methods		
n/a Involved in th	e study n/a Involved in the study		
Antibodies	ChIP-seq		
Eukaryotic Palaeontol			
	d other organisms		
Human res	earch participants		
Clinical dat			
Antibodies			
Antibodies used	Anti-HBV core antibody from DAKO company		
Validation	Reference paper (Park et al., 2003, Journal of Biological Chemistry)		
Eukaryotic c	ell lines		
Policy information			
Cell line source(s			
Authentication	Commercially available and freely accessible		
Mycoplasma con	ramination We have tested.		
Commonly misidentified lines (See ICLAC register) HepG2 cells, Hepatoblastoma, Genetic profile by array comparative genomic hybridization (CGH) analysis was used for the HepG2 cell line identification in the previous report (Lopez-Terrada et al., 2009, Human Pathology).			

Animals and other organisms

Policy information about studies involving animals; ARRIVE guidelines recommended for reporting animal research

Laboratory animals

Human liver-chimeric uPA/SCID (PXB) mice and C57BL/6 mice

Wild animals	Provide details on animals observed in or captured in the field; report species, sex and age where possible. Describe how animals were caught and transported and what happened to captive animals after the study (if killed, explain why and describe method; if released, say where and when) OR state that the study did not involve wild animals.
Field-collected samples	Mice were kept in laboratory animal research center of CHA University. Animal protocols were performed in accordance with the guidelines of the committee for animal experiments in CHA University. Infection, extraction of serum sample and sacrifice were performed under ether anesthesia.

Ethics oversight	All animal protocols were approved by the Institutional Animal Care and Use Committee of CHA University (IACUC-180090).				
Note that full information on the approval of the study protocol must also be provided in the manuscript.					
Flow Cytometry					
Plots					
Confirm that:					
The axis labels state the marker and fluorochrome used (e.g. CD4-FITC).					
The axis scales are clearly visible. Include numbers along axes only for bottom left plot of group (a 'group' is an analysis of identical markers).					
All plots are contour plots with outliers or pseudocolor plots.					
A numerical value for nur	mber of cells or percentage (with statistics) is provided.				
Methodology					
Sample preparation	Standard single cell preparation method				
Instrument	FACS Canto II				
Software	FlowJo				
Cell population abundance	Almost all				
Gating strategy	Not applicable				
Tick this box to confirm the	hat a figure exemplifying the gating strategy is provided in the Supplementary Information.				