Article type : Original Paper

Title: Human genomics of acute liver failure due to hepatitis B virus infection: an exome sequencing study in liver transplant recipients

Short running title: Human genomics of fulminant hepatitis B

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This is the author manuscript accepted for publication and has undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the Version of Record. Please cite this article as [doi: 10.1111/jvh.13019](https://doi.org/10.1111/jvh.13019)

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Keywords

Acute Liver Failure (ALF); Fulminant hepatitis; Hepatitis B Virus (HBV); Exome sequencing

Abstract

Acute liver failure (ALF) or fulminant hepatitis is a rare, yet severe outcome of infection with hepatitis B virus (HBV) that carries a high mortality rate. The occurrence of a life-threatening condition upon infection with a prevalent virus in individuals without known risk factors is suggestive of pathogen-specific immune dysregulation. In the absence of established differences in HBV virulence, we hypothesized that ALF upon primary infection with HBV could be due to rare deleterious variants in the human genome. To search for such variants, we performed exome sequencing in 21 previously healthy adults who required liver transplantation upon fulminant HBV infection and 172 controls that were positive for anti-HBc and anti-HBs but had no clinical history of jaundice or liver disease. After a series of hypothesis-driven filtering steps, we searched for putatively pathogenic variants that were significantly associated with case-control status. We did not find any causal variant or gene, a result that does not support the hypothesis of a shared monogenic basis for human susceptibility to HBV-related ALF in adults. This study represents a first attempt at deciphering the human genetic contribution to the most severe clinical presentation of acute HBV infection in previously healthy individuals.

Background

Hepatitis B virus (HBV) is a common human pathogen that attacks the liver and can cause both acute and chronic disease. There is high inter-individual variability in the clinical presentation of HBV infection, which ranges from self-limited to fulminant

acute disease, and from mild chronic hepatitis to liver cirrhosis and hepatocellular carcinoma [1] . Differences in viral or environmental factors only explain a fraction of this variability [2–5] . Previous studies have identified some human genetic factors that play a modulating role in the clinical course of HBV infection [6,7] . However, our understanding of host genetic influences on the disease is still very limited.

Fulminant hepatitis or acute liver failure (ALF) is defined as the rapid development of liver injury leading to severe impairment of the synthetic capacity and to hepatic encephalopathy in patients without previous liver disease [8,9] . ALF due to HBV infection, or fulminant hepatitis B, is observed in less than 0.1% of infected individuals but carries a high mortality and is an indication for urgent liver transplantation [10–15] .

Such an unusual clinical presentation fits the definition of an extreme phenotype. Electing patients with extreme phenotype increases the power to detect causal gene variants as these patients are more likely to carry alleles with profound functional consequences that are otherwise very rare in the population, due to purifying selection [16–20] . In this study, we used exome sequencing and statistical analysis in a cohort of 21 cases and 172 controls to search for human genetic variants conferring extreme susceptibility to HBV. Cases were previously healthy adults who required liver transplantation for fulminant hepatitis B and controls were HBV-infected adults who did not develop fulminant hepatitis (Figure 1).

Methods

Study participants

Twenty-one liver transplant recipients who developed ALF due to fulminant HBV infection were recruited in the transplantation units of the University Hospitals of Lausanne, Zurich, Bern, Geneva, and Melbourne. Patients with fulminant hepatitis B due to reactivation after withdrawal of anti-HBV drugs and patients with pre-existing liver diseases, known immune deficiency or other chronic conditions were excluded. The following demographic and clinical information were collected: age at transplantation date, gender, and ethnicity. For each study participant, we obtained 3ml of blood in EDTA vacutainer tubes and 2.5ml blood in PAXgene blood RNA tubes. Samples were immediately frozen at -70°C, and then shipped and analyzed in

batch.

Control population

One hundred seventy-two controls were selected from our in-house database of exome-sequenced individuals. They were adults of European ancestry, who were positive for anti-HBc and anti-HBs, but had no clinical history of jaundice or liver disease. The controls were HBV-eliminated at the time of blood collection for exome-sequencing.

DNA sequencing and alignment

Genomic DNA was extracted from whole blood using QIAgen DNeasy Blood and Tissue kit. Cluster generation was performed using Illumina TruSeq PE Cluster Kit v5 reagents. Libraries were sequenced as 100-basepair long, paired-end reads on Illumina HiSeq 2500 using TruSeq SBS Kit v5 reagents. Sequencing reads were processed using CASAVA v1.82, and aligned to the human reference genome hg19 using BWA [21,22] version 0.6.2. PCR duplicates were removed using Picard 1.27-1 (<http://picard.sourceforge.net/>). We used Samtools [23] Visualization of aligned reads.

Variant calling

We used Genome Analysis Toolkit (GATK) [24,25] version 3.1-1 to call single nucleotide variants (SNVs) and small insertion and deletions (indels) from duplicate-marked bam files. We used HaplotypeCaller for multi-sample variant calling on all samples following GATK best practice.

Variant effect prediction, frequency estimation and filtering

We used SnpEff [26] version 4.3T to predict the functional impact of variants. As a single variant can have several predicted effects, we only considered the most severe effect for each variant according to SnpEff order of impact severity. We used genome aggregation database (gnomAD) to assign minor allele frequency (MAF) to variants (gnomAD, includes 123,136 exome sequences and 15,496 whole-genome sequences) [27]. For variants that were not present in gnomAD were assigned $MAF=1-e8$ to avoid having $-\log(0)$ in the following burden analysis. Only biallelic variants that were flagged as PASS by GATK, and were called in all cases and all controls were

included in the analysis. Known polymorphic genes and genes in noisy alignment regions were excluded from the analysis [28–30] . We restricted all the downstream analyses to protein modifying variants (missense, inframe indels, frame-shift indels, splice-site disrupting, nonsense). All analyses were done on both rare ($MAF \leq 0.01$) and low-frequency ($MAF \leq 0.05$) variants. We refer to variants that passed above filtering criteria as putatively pathogenic variants.

Single variant association tests

We used Fisher's exact test to look for association of single variants with case-control status. Each variant was given an allele count based on the number of alternate alleles $G_{ij} \in \{0,1,2\}$, where G_{ij} is the genotype of variant j in individual i . We summarized the reference and alternate allele counts for cases and controls, into 2x2 contingency tables. These tables were analyzed using one-tailed Fisher's exact test. We used Bonferroni correction to correct for multiple testing.

Gene burden association tests

Gene burden test was performed using GMMAT [31] version 0.7-1, a generalized linear mixed model framework as follows:

$$H = W\alpha + C\beta + M ; \quad M \sim N \left(0, \sum_{k=1}^K \tau_k V_k \right)$$

Where H is an n -vector of case-control status for n individuals, W is an n -vector of gender covariate, and C is an n -vector of the gene burden scores. M is an n -vector of random effects. τ_k is the variance component parameters and V_k are known $n \times n$ matrices. We ran this model using an $n \times n$ kinship coefficients matrix calculated using PC-Relate [32] . We used Bonferroni correction to correct for multiple testing. To calculate the gene burden scores, we used two different methods:

i) Binary collapsing method

Each gene was given a burden score of zero if no putatively pathogenic variant was present in the gene and a gene burden score of one otherwise:

$$C_i = \begin{cases} 1 & \text{if } \sum_{j=1}^m G_{ij} > 0 \\ 0 & \text{if } \sum_{j=1}^m G_{ij} = 0 \end{cases}$$

Where $G_{ij} \in \{0,1,2\}$ is the genotype of variant j in individual i , and C_i is the gene burden score for individual i . This approach is based on the Cohort Allelic Sum Test (CAST) method [33].

ii) Weighted sum collapsing method

First, each gene was given a burden score as follows:

$$C_i = \sum_{j=1}^m (G_{ij} * -\log_{10}(AF_i\text{-gnomAD}))$$

Where $j \in \{0,1,2, \dots, m\}$ is the m th variant per gene, and $G_{ij} \in \{0,1,2\}$ is the genotype of variant j in individual i , $AF_i\text{-gnomAD}$ is the minor allele frequency of j in gnomAD, and C_i is the gene burden score for individual i . This approach is based on the Madsen and Browning weighted sum method [34].

Results

Study participants

Of the 21 cases, 13 (62%) were female. The median age at transplantation was 36.5 years (range 22-58). Of 21 cases, 16 (76%) were European, four (19%) were Asian and one (5%) was African (Supplementary Figure 1).

Exome sequencing, variant calling and variant filtering

Exome sequencing data were generated from DNA extracted from whole blood for all study participants. On average per sample, 96% of reads passing filtering criteria were unique (not marked as duplicate). Ninety-seven percent of unique reads could be aligned to the human reference genome GRCh37. The mean on-bait coverage was 73x, with 99% of target bases reaching at least 2x coverage, 97% of target bases achieving at least 10x coverage and 84% achieving at least 30x coverage. 205,642 variants were detected after GATK quality control filtering including 520 novel variants. The average transition to transversion ratio (Ti/Tv) was 2.66, and the average heterozygous to homozygous ratio was 1.5. A total of 38,062 low-frequency

variants ($MAF \leq 0.05$) passed filtering criteria including 31,620 rare variants ($MAF \leq 0.01$, Table 1).

Single variant associate analysis

All putatively pathogenic variants were tested for association with case-control status using Fisher's exact test. We first restricted the analysis to rare variants ($MAF \leq 0.01$) and European cases only. One variant passed the Bonferroni correction threshold (p-value $< 1.6e-6$). The variant was a missense SNV in *IGSF3* (rs78806598, p-value=1.8e-18). Visualizing the aligned reads for this variant convinced us that this variant is called due to misalignment. This gene was excluded from our further analyses. Expanding the analysis to include the five non-European cases and the low-frequency variants did not lead to discovery of any significant associations (Supplementary Tables 1-3).

Gene-based association analysis

11,595 genes were included in the gene burden analysis. Two different burden scores were calculated for each gene using the approaches described in the methods section. Using the weighted sum method, *SLC29A1* had the lowest p-value (p-value=1.7e-5). CTSW had the lowest p-value in binary collapsing method (p-value=1.8e-5). However, none of these genes passed the Bonferroni correction threshold (p-value $< 2.5e-6$ for 20,000 protein coding genes, Supplementary Tables 4-5). Including the low-frequency variants (12,295 genes in total) did not change these results. The top associations including low-frequency variants were *ADAM32* (p-value=1.7e-5) and *PREX2* (p-value=8.7e-6) in for weighted sum method binary collapsing method respectively (Supplementary Tables 6-7). Overall, the results from the two collapsing methods and the results between rare variants ($MAF \leq 0.01$) and low-frequency variants ($MAF \leq 0.05$) analyses were highly concordant (Figure 2, Supplementary Figure 2). The highest correlation ($r^2=0.927$, CI:0.924-0.929) was observed between the results of weighted sum and binary collapsing methods for rare variants. The lowest correlation ($r^2=0.739$, CI:0.731-0.747) was observed between the results of binary collapsing method for rare and low-frequency variants (Figure 2).

Discussion and Conclusion

The role of human genetic factors in susceptibility to fulminant hepatitis B is poorly

understood. Monogenic defects in key immune genes and pathways have been shown to cause extreme susceptibility to other common pathogens in apparently healthy individuals [7,35,36]. A prime example is herpes simplex encephalitis (HSE), the most common form of sporadic viral encephalitis in the western world, which is only observed in an extremely low fraction of people infected with type 1 herpes simplex virus (HSV-1). Children who develop HSE upon primary HSV-1 infection are not particularly susceptible to other infections, and children with other primary immunodeficiencies are not more susceptible to HSE [36]. Since 2006, multiple genetic variants have been causally linked with HSE [37–43]. Similarly, ALF only occurs in $< 1/1000$ of individuals after primary infection with HBV. Because this is an extremely rare clinical event, we hypothesized that it could be the first manifestation of a rare monogenic defect, resulting in pathogen-specific immune dysregulation.

We used exome sequencing to systematically search for rare, putatively pathogenic variants that could explain extreme susceptibility to HBV infection. We analyzed the genetic variants present in the exomes of 21 liver transplant recipients and compared them to 172 controls who were exposed to HBV but did not develop fulminant hepatitis B. First, we performed a single variant association analysis using Fisher's exact test. Fisher's exact test is a conservative test of association but guarantees type I error control for small sample sizes [44]. We found one significant association (p -value $< 1.4e-6$) in one gene: *IGSF3*. However, the manual inspection of the mapped reads in the region demonstrated that this variant was wrongly called, due to a mapping error. False-positive incidental findings are a major problem in small-scale exome sequencing studies [28]. Previous studies have proposed guidelines to avoid misinterpretations and erroneous reports of potential causality due to false-positive findings [29,30]. Our results show that even after applying these guidelines, it is important to ensure the quality of final findings by visualizing the mapped regions and manually verifying the quality of each variant call.

Rare variant association studies are usually underpowered. To enrich association signals and reduce the penalty of multiple testing correction, it is common to aggregate information across multiple rare variants within a region (gene, exon, sliding window, etc.) and test for the association of all variants in the region with the

phenotype of interest [45] . We performed gene-based association analysis using two different aggregation methods: weighted sum collapsing and binary collapsing. Both methods assume that all the variants included in the test have the same direction of effect (increasing disease risk in our scenario) and thus are underpowered to detect disease-gene associations if variants exert their effects in opposite directions. Binary collapsing assumes that all putatively pathogenic variants have the same effect size. Weighted sum collapsing assumes that rarer variants have larger effect sizes and that the risk of disease is a function of the sum of the variant effect sizes. We did not find any genes to be significantly associated with case-control status. The p-values and the top ranked genes in both analyses were highly concordant (Figure 2, Supplementary Figure 2). The high correlation between the p-values of weighted sum and binary collapsing methods suggests that most individuals carry only one putatively pathogenic variant per gene. This implies that larger sample sizes or linkage studies in families with multiple affected individuals will be needed to increase statistical power for detecting potential associations between rare variants and HBV-related ALF.

We did not identify any genetic variant conferring monogenic susceptibility to fulminant hepatitis B in adults. Our results suggest that ALF upon primary infection with HBV is likely to be multifactorial. This conclusion is in line with a previous exome sequencing study of fulminant hepatitis A, which also failed to find any convincing casual gene or genetic variant [46] . Our failure to detect a Mendelian cause for fulminant hepatitis B, despite previous success for comparable phenotypes, could be due to a number of factors and limitations of our study: 1- The severe liver injury observed in patients with fulminant hepatitis B can be due to opposite pathogenic mechanisms: an inefficient innate immune response, which is unable to prevent viral replication, activate the adaptive immune system and clear the virus; and an over-activation of innate immune signaling pathways leading to cytokine storm and uncontrolled inflammation [47–49] . This implies that genetic variants with opposite effects (e.g. gain-of-function and loss-of-function variants in the same gene or pathway) could contribute synergistically to the disease. Such a genetic architecture would be extremely difficult to identify. 2- Our study was performed in adults, while most previous examples come from pediatric studies. A previous twin study has shown that the estimated heritability of many immune parameters decreases with age, suggesting that the cumulative influence of environmental exposures alters

the role of human genetics in susceptibility to infectious diseases in older patients [50]
. 3- Environmental factors have been implicated in the pathogenesis of HBV disease
and ALF. Our study design prevented an in-depth evaluation of the potential
contribution of environmental risk factors such as alcohol consumption [51] . 4- Due
to our recruitment criteria, we did not have access to information about the viral
genome. HBV genetic variation has previously been shown to be associated with
disease severity and infection outcome, but these results remain controversial. In
particular, mutations in the pre-core and basal core promoter regions of the HBV
genome were associated with ALF in some studies [52–54] , but not in others [54–
57] . The inclusion of viral genome information might allow for the stratification of
patients based on known HBV mutations, thus increasing the signal-to-noise ratio in
human genetic analyses.

This study represents the first attempt at identifying human genetic variants involved
in the pathogenesis of fulminant hepatitis B in previously healthy individuals. The
absence of any conclusive finding indicates that ALF due to primary HBV infection is
unlikely to be the result of a single monogenic disorder, and that a more complex
genetic architecture is probably involved, intermixed with viral and environmental
factors. Going forward, studies that aim at identifying the genetic causes of fulminant
hepatitis B will need to include more patients and to better characterize them at the
molecular level (e.g. to stratify them based on specific immune activation markers
measured during acute disease). Inclusion of matching controls with proven HBV
infection (HBsAg positive), who could clear the infection in the absence of antiviral
therapy, could allow for better control of confounding factors such as vaccination
history and increase the power to detect human genetic contributors to ALF. To
obtain a more complete description of human genetic variation, full genome
sequencing would be preferable, which will allow the exploration of non-coding
variants, large structural variants and exonic variants that are not well-covered by
current exome capture methods. Finally, a parallel evaluation of the viral genome and
of any potentially interfering factor will be necessary, as individual susceptibility to
HBV is the result of a complex interplay between host, pathogen and environment.

Declarations

Ethics approval and consent to participate

The study was approved by the responsible institutional Human Research Ethics Committees in Switzerland and Australia. Each study participant provided written informed consent for genetic testing.

Consent for publication

Not applicable.

Availability of data and materials

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Competing interests

The authors declare that they have no competing interests.

Funding

The study was funded by grants from the Fondation Lausannoise pour la Transplantation d'Organes and the Novartis Foundation. Additional support was provided by the Swiss National Science Foundation (SNF Professorship grant PP00P3_133703 to JF). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Authors contributions

J.F. designed research; S.A, N.C, P.S, C.H, N.S, E.G, B.M, P.A, A.J.T, D.M, and J.F performed research; N.S, E.G, B.M, P.A, A.J.T, and D.M. contributed new reagents/analytic tools; S.A, N.C, P.S, and C.H analyzed data; and S.A, N.C, and J.F. wrote the paper.

Acknowledgements

Sequencing was performed at the Lausanne Genomic Technologies Facility of the University of Lausanne. The computations were performed at the Vital-IT (<http://www.vital-it.ch>) Center for high-performance computing of the SIB Swiss Institute of Bioinformatics. We would like to thank the study participants, as well as the study nurses, physicians and laboratories who participated in the recruitment.

References

1. MacLachlan JH, Locarnini S, Cowie BC (2015) Estimating the global prevalence of hepatitis B. *Lancet* 386: 1515–7. doi:10.1016/S0140-6736(15)61116-3.
2. Karvellas C, Cardoso F, Gottfried M, Reddy, Hanje, et al. (2017) HBV-Associated Acute Liver Failure After Immunosuppression and Risk of Death. *Clinical gastroenterology and hepatology : the official clinical practice journal of the American Gastroenterological Association* 15: 113–122. doi:10.1016/j.cgh.2016.06.008.
3. Aragri M, Alteri C, Battisti A, Carlo D, Minichini C, et al. (2016) Multiple Hepatitis B Virus (HBV) Quasispecies and Immune-Escape Mutations Are Present in HBV Surface Antigen and Reverse Transcriptase of Patients With Acute Hepatitis B. *The Journal of infectious diseases* 213: 1897–905. doi:10.1093/infdis/jiw049.
4. Zhang A, Wan Z, You S, Liu H, Zhu B, et al. (2013) Association of Hepatitis B Virus Mutations of A1846T and C1913A/G With Acute-on-Chronic Liver Failure Development From Different Underlying Chronic Liver Diseases. *Hepatitis monthly* 13: e12445. doi:10.5812/hepatmon.12445.
5. Goldstein S, Alter M, Williams I, Moyer L, Judson F, et al. (2002) Incidence and risk factors for acute hepatitis B in the United States, 1982-1998: implications for vaccination programs. *The Journal of infectious diseases* 185: 713–9. doi:10.1086/339192.
6. Matsuura K, Isogawa M, Tanaka Y (2016) Host genetic variants influencing the clinical course of hepatitis B virus infection. *J Med Virol* 88: 371–9. doi:10.1002/jmv.24350.
7. Kenney A, Dowdle J, Bozzacco L, McMichael T, Gelais C, et al. (2017) Human Genetic Determinants of Viral Diseases. *Annual review of genetics* 51: 241–263. doi:10.1146/annurev-genet-120116-023425.
8. Trey C, Davidson CS (1970) The management of fulminant hepatic failure. *Prog Liver Dis* 3: 282–98.
9. Ritt DJ, Whelan G, Werner DJ, Eigenbrodt EH, Schenker S, et al. (1969) Acute hepatic necrosis with stupor or coma. An analysis of thirty-one patients. *Medicine (Baltimore)* 48: 151–72.
10. Papaevangelou G, Tassopoulos N, Roumeliotou-Karayannis A, Richardson C (1984) Etiology of fulminant viral hepatitis in Greece. *Hepatology* 4: 369–72.
11. Schiødt F, Davern T, Shakil O, McGuire B, Samuel G, et al. (2003) Viral

- Hepatitis-Related Acute Liver Failure. *The American Journal of Gastroenterology* 98: 448–453. doi:10.1111/j.1572-0241.2003.t01-1-07223.x.
12. Oketani M, Uto H, Ido A, Tsubouchi H (2014) Management of hepatitis B virus-related acute liver failure. *Clin J Gastroenterol* 7: 19–26. doi:10.1007/s12328-013-0447-1.
13. Fujiwara K, Mochida S, Matsui A, Nakayama N, Nagoshi S, et al. (2008) Fulminant hepatitis and late onset hepatic failure in Japan. *Hepatol Res* 38: 646–57. doi:10.1111/j.1872-034X.2008.00322.x.
14. Lee W, Squires R, Nyberg S, Doo E, Hoofnagle J (2008) Acute liver failure: Summary of a workshop. *Hepatology* 47: 1401–1415. doi:10.1002/hep.22177.
15. Ostapowicz G, Fontana RJ, Schiødt FV, Larson A, Davern TJ, et al. (2002) Results of a prospective study of acute liver failure at 17 tertiary care centers in the United States. *Ann Intern Med* 137: 947–54. doi:10.7326/0003-4819-137-12-200212170-00007.
16. Guey L, Kravic J, Melander O, Burt N, Laramie J, et al. (2011) Power in the phenotypic extremes: a simulation study of power in discovery and replication of rare variants. *Genetic epidemiology* 35: 236–46. doi:10.1002/gepi.20572.
17. Lander, Botstein (1989) Mapping mendelian factors underlying quantitative traits using RFLP linkage maps. *Genetics* 121: 185–99.
18. Barnett I, Lee S, Lin X (2013) Detecting rare variant effects using extreme phenotype sampling in sequencing association studies. *Genetic epidemiology* 37: 142–51. doi:10.1002/gepi.21699.
19. Li D, Lewinger J, Gauderman W, Murcray C, Conti D (2011) Using extreme phenotype sampling to identify the rare causal variants of quantitative traits in association studies. *Genetic epidemiology* 35: 790–9. doi:10.1002/gepi.20628.
20. Gibson G (2012) Rare and common variants: twenty arguments. *Nature reviews Genetics* 13: 135–45. doi:10.1038/nrg3118.
21. Li H, Durbin R (2009) Fast and accurate short read alignment with Burrows-Wheeler transform. *Bioinformatics* 25: 1754–60. doi:10.1093/bioinformatics/btp324.
22. Li H, Durbin R (2010) Fast and accurate long-read alignment with Burrows-Wheeler transform. *Bioinformatics* 26: 589–95. doi:10.1093/bioinformatics/btp698.
23. Li H, Handsaker B, Wysoker A, Fennell T, Ruan J, et al. (2009) The Sequence Alignment/Map format and SAMtools. *Bioinformatics* 25: 2078–9. doi:10.1093/bioinformatics/btp352.

24. McKenna A, Hanna M, Banks E, Sivachenko A, Cibulskis K, et al. (2010) The Genome Analysis Toolkit: a MapReduce framework for analyzing next-generation DNA sequencing data. *Genome research* 20: 1297–303. doi:10.1101/gr.107524.110.
25. Auwera GA Van der, Carneiro MO, Hartl C, Poplin R, Angel G Del, et al. (2013) From FastQ data to high confidence variant calls: the Genome Analysis Toolkit best practices pipeline. *Curr Protoc Bioinformatics* 43: 11.10.1–33. doi:10.1002/0471250953.bi1110s43.
26. Cingolani P, Platts A, Wang L e, Coon M, Nguyen T, et al. (2012) A program for annotating and predicting the effects of single nucleotide polymorphisms, SnpEff: SNPs in the genome of *Drosophila melanogaster* strain w1118; iso-2; iso-3. *Fly* 6: 80–92. doi:10.4161/fly.19695.
27. Exome Aggregation Consortium (ExAC) (n.d.) Cambridge, MA (URL: <http://exac.broadinstitute.org>) [August 2016].
28. Kohane I, Hsing M, Kong S (2012) Taxonomizing, sizing, and overcoming the incidentalome. *Genetics in medicine : official journal of the American College of Medical Genetics* 14: 399–404. doi:10.1038/gim.2011.68.
29. MacArthur, Manolio, Dimmock, Rehm, Shendure, et al. (2014) Guidelines for investigating causality of sequence variants in human disease. *Nature* 508: 469–76. doi:10.1038/nature13127.
30. Fajardo K, Adams D, Program N, Mason C, Sincan M, et al. (2012) Detecting false-positive signals in exome sequencing. *Human mutation* 33: 609–13. doi:10.1002/humu.22033.
31. Chen H, Wang C, Conomos M, Stilp A, Li Z, et al. (2016) Control for Population Structure and Relatedness for Binary Traits in Genetic Association Studies via Logistic Mixed Models. *American journal of human genetics* 98: 653–66. doi:10.1016/j.ajhg.2016.02.012.
32. Conomos M, Reiner A, Weir B, Thornton T (2016) Model-free Estimation of Recent Genetic Relatedness. *American journal of human genetics* 98: 127–48. doi:10.1016/j.ajhg.2015.11.022.
33. Morgenthaler S, Thilly W (2007) A strategy to discover genes that carry multi-allelic or mono-allelic risk for common diseases: a cohort allelic sums test (CAST). *Mutation research* 615: 28–56. doi:10.1016/j.mrfmmm.2006.09.003.
34. Madsen B, Browning S (2009) A groupwise association test for rare mutations using a weighted sum statistic. *PLoS genetics* 5: e1000384.

- doi:10.1371/journal.pgen.1000384.
35. Casanova J-LL (2015) Human genetic basis of interindividual variability in the course of infection. *Proc Natl Acad Sci USA* 112: E7118–27. doi:10.1073/pnas.1521644112.
36. Alcaïs A, Abel L, Casanova J-LL (2009) Human genetics of infectious diseases: between proof of principle and paradigm. *J Clin Invest* 119: 2506–14. doi:10.1172/JCI38111.
37. Zhang S-YY, Jouanguy E, Ugolini S, Smahi A, Elain G, et al. (2007) TLR3 deficiency in patients with herpes simplex encephalitis. *Science* 317: 1522–7. doi:10.1126/science.1139522.
38. Andersen LL, Mørk N, Reinert LS, Kofod-Olsen E, Narita R, et al. (2015) Functional IRF3 deficiency in a patient with herpes simplex encephalitis. *J Exp Med* 212: 1371–9. doi:10.1084/jem.20142274.
39. Herman M, Ciancanelli M, Ou Y-HH, Lorenzo L, Klaudel-Dreszler M, et al. (2012) Heterozygous TBK1 mutations impair TLR3 immunity and underlie herpes simplex encephalitis of childhood. *J Exp Med* 209: 1567–82. doi:10.1084/jem.20111316.
40. Guo Y, Audry M, Ciancanelli M, Alsina L, Azevedo J, et al. (2011) Herpes simplex virus encephalitis in a patient with complete TLR3 deficiency: TLR3 is otherwise redundant in protective immunity. *J Exp Med* 208: 2083–98. doi:10.1084/jem.20101568.
41. Pérez de Diego R, Sancho-Shimizu V, Lorenzo L, Puel A, Plancoulaine S, et al. (2010) Human TRAF3 adaptor molecule deficiency leads to impaired Toll-like receptor 3 response and susceptibility to herpes simplex encephalitis. *Immunity* 33: 400–11. doi:10.1016/j.immuni.2010.08.014.
42. Sancho-Shimizu V, Pérez de Diego R, Lorenzo L, Halwani R, Alangari A, et al. (2011) Herpes simplex encephalitis in children with autosomal recessive and dominant TRIF deficiency. *J Clin Invest* 121: 4889–902. doi:10.1172/JCI59259.
43. Casrouge A, Zhang S-YY, Eidenschenk C, Jouanguy E, Puel A, et al. (2006) Herpes simplex virus encephalitis in human UNC-93B deficiency. *Science* 314: 308–12. Available: <http://www.ncbi.nlm.nih.gov/pubmed/16973841>.
44. Li B, Leal S (2008) Methods for detecting associations with rare variants for common diseases: application to analysis of sequence data. *American journal of human genetics* 83: 311–21. doi:10.1016/j.ajhg.2008.06.024.

45. Lee S, Abecasis G, Boehnke M, Lin X (2014) Rare-variant association analysis: study designs and statistical tests. *American journal of human genetics* 95: 5–23. doi:10.1016/j.ajhg.2014.06.009.
46. Long D, Fix OK, Deng X, Seielstad M, Luring AS (2014) Whole genome sequencing to identify host genetic risk factors for severe outcomes of hepatitis A virus infection. *J Med Virol* 86: 1661–8. doi:10.1002/jmv.24007.
47. Han Q, Zhang C, Zhang J, Tian Z (2013) The role of innate immunity in HBV infection. *Semin Immunopathol* 35: 23–38. doi:10.1007/s00281-012-0331-y.
48. Maini MK, Gehring AJ (2016) The role of innate immunity in the immunopathology and treatment of HBV infection. *J Hepatol* 64: S60–70. doi:10.1016/j.jhep.2016.01.028.
49. Jung M-CC, Pape GR (2002) Immunology of hepatitis B infection. *Lancet Infect Dis* 2: 43–50.
50. Brodin P, Jojic V, Gao T, Bhattacharya S, Angel C (2015) Variation in the human immune system is largely driven by non-heritable influences. *Cell*.
51. Gitto S, Vitale G, Villa E, Andreone P (2014) Update on Alcohol and Viral Hepatitis. *J Clin Transl Hepatol* 2: 228–33. doi:10.14218/JCTH.2014.00030.
52. Hou J, Lin Y, Waters J, Wang Z, Min J, et al. (2002) Detection and significance of a G1862T variant of hepatitis B virus in Chinese patients with fulminant hepatitis. *J Gen Virol* 83: 2291–8. doi:10.1099/0022-1317-83-9-2291.
53. Inoue J, Ueno Y, Nagasaki F, Wakui Y, Kondo Y, et al. (2009) Enhanced intracellular retention of a hepatitis B virus strain associated with fulminant hepatitis. *Virology* 395: 202–9. doi:10.1016/j.virol.2009.09.028.
54. Cui X-JJ, Cho Y-KK, Song B-CC (2015) Influence of the basal core promoter and precore mutation on replication of hepatitis B virus and antiviral susceptibility of different genotypes. *J Med Virol* 87: 601–8. doi:10.1002/jmv.24117.
55. Sterneck M, Günther S, Santantonio T, Fischer L, Broelsch CE, et al. (1996) Hepatitis B virus genomes of patients with fulminant hepatitis do not share a specific mutation. *Hepatology* 24: 300–6. doi:10.1002/hep.510240203.
56. Chen Y, Michitaka K, Matsubara H, Yamamoto K, Horiike N, et al. (2003) Complete genome sequence of hepatitis B virus (HBV) from a patient with fulminant hepatitis without precore and core promoter mutations: comparison with HBV from a patient with acute hepatitis infected from the same infectious source. *J Hepatol* 38: 84–90.

57. Liu C-JJ, Kao J-HH, Lai M-YY, Chen P-JJ, Chen D-SS (2004) Precore/core promoter mutations and genotypes of hepatitis B virus in chronic hepatitis B patients with fulminant or subfulminant hepatitis. J Med Virol 72: 545–50. doi:10.1002/jmv.20024.

Table 1: Total number of rare ($MAF \leq 0.01$) and rare plus low-frequency ($MAF \leq 0.05$) variants that passes quality control and filtering criteria.

Effect	$MAF \leq 0.01$	$MAF \leq 0.05$
Inframe indel	321	368
Frameshift indel	880	954
Missense SNV	29,549	35,756
Splice site acceptor SNV	152	177
Splice site donor SNV	177	200
Nonsense	541	607
Total variants	31,620	38,062
Total genes	11,595	12,295

Figure Legends

Figure 1: **Overview of data production and data analysis pipeline.** MAF: minor allele frequency, GATK: Genome Analysis Tool Kit

Figure 2: **Comparison between different gene burden analysis methods and different MAF thresholds.** The circles below each plot show the top ten associated genes in the two compared analyses and the number of shared genes between the two sets: A) correlation between p-value for rare variants ($MAF \leq 0.01$) using weighted sum method (light green circle) and binary collapsing method (light red circle) B) correlation between p-value for binary collapsing method using $MAF \leq 0.01$ (light red circle) and $MAF \leq 0.05$ (dark red circle).

Supplementary Tables legends

Supplementary Tables 1-3: **Fisher's exact test results for single variant association analysis** for: 1- Rare variants ($MAF \leq 0.01$), 16 European cases and 172 controls, 2-

Rare variants ($MAF \leq 0.01$), all 21 cases and 172 controls, 3- Low-frequency variants ($MAF \leq 0.05$), all 21 cases and 172 controls. Column names: chromosome, position, reference allele, alternate allele, variant ID, gene, number of putatively pathogenic alleles in cases, number of putatively pathogenic alleles in controls, number of non-putatively pathogenic alleles in cases, number of non-putatively pathogenic alleles in controls

Supplementary Tables 4-7: **Gene burden association results** for: 4- Rare variants ($MAF \leq 0.01$) and weighted sum method, 5- Rare variants ($MAF \leq 0.01$) and binary collapsing method, 6- Rare variants ($MAF \leq 0.05$) and weighted sum method, 7- Rare variants ($MAF \leq 0.05$) and binary collapsing method . Column names: gene, score, variance, p-value

Supplementary Figures legends

Supplementary Figure 1: **Principal component analysis (PAC)** A-B) PCA analysis for 21 cases, 172 controls and continental populations from 1000 genomes project, C-D) PCA analysis for 21 cases and 172 controls.

Supplementary Figure 2: **Comparison between different gene burden analysis methods and different MAF thresholds**. The circles below each plot show the top ten associated genes in the two compared analyses and the number of shared genes between the two sets: A) correlation between p-value for low-frequency variants ($MAF \leq 0.05$) using weighted sum method (dark green circle) and binary collapsing method (dark red circle). B) correlation between p-value for weighted sum method using $MAF \leq 0.01$ (light green circle) and $MAF \leq 0.05$ (dark green circle).

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Title:

Human genomics of acute liver failure due to hepatitis B virus infection: An exome sequencing study in liver transplant recipients

Date:

2019-02-01

Citation:

Asgari, S., Chaturvedi, N., Scepanovic, P., Hammer, C., Semmo, N., Giostra, E., Muellhaupt, B., Angus, P., Thompson, A. J., Moradpour, D. & Fellay, J. (2019). Human genomics of acute liver failure due to hepatitis B virus infection: An exome sequencing study in liver transplant recipients. JOURNAL OF VIRAL HEPATITIS, 26 (2), pp.271-277. <https://doi.org/10.1111/jvh.13019>.

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