Adult Cohort Data Collection

Short name	Long name	Additional information	Translated versions
SL	Screening Log		
EC	Enrollment Criteria		
BC	Baseline Eval - Coordinator		
BP	Baseline Eval – Patient	Health Behavior incorporated into this form	Spanish, Chinese, Korean
BI	Baseline Eval – Investigator		
QL	Quality of Life Questionnaire	Health Behavior incorporated into this form after baseline timepoint. SF-36v2™ Health Survey © 1996, 2000 by QualityMetric Incorporated and Medical Outcomes Trust. All Rights Reserved. SF-36® is a registered trademark of Medical Outcomes Trust. (SF-36v2 Standard, US Version 2.0)	QL: Spanish, Chinese, Korean, Vietnamese Health Behavior: Spanish, Chinese, Korean
SA	Symptom Assessment		Spanish, Chinese, Korean
FQ	Fatigue Questionnaire	© 2009 PROMIS Health Organization and PROMIS Cooperative Group	Spanish
FW	Follow-up Eval 12-week		
FF	Follow-up Evaluation		
SV	Special Visit Form		
FR	ALT Flare Resolution		
LB	Liver Biopsy		
FE	Follow-Up Events		
HC	HCC Form		
PQ	Pregnancy Questionnaire		
PP	Pregnancy Pre-delivery		
PF	Pregnancy Follow-up		
FB	Fibroscan		

Dataset	Description
Screen_info	Includes information about site transfers and after enrollment participant ineligibility along with the screening log information. This information is needed to identify the participants eligible for analyses and reasons for ineligibility. The last screening record for each consecutive participant who was not a transfer from another clinical site is used to determine screening eligibility.

Central lab datasets

Name	Description
hbrn_bcp_pc_ac_results	HBV Basal core promoter (BCP) and precore (PC) mutations on available samples. Some sample dates may have more than one result due to multiple aliquots drawn on the same day having been tested. If variable EXCLUDE =1, these were cases re-reviewed by the lab due to inconsistent results between IUPAC and Majority results within a sample or between different vials tested with the same sample date and after review it was decided to exclude that result for HBRN analyses.
hbrn_drug_resistance_ac_results	HBV drug resistant variant results on available samples.
hbrn_cdc_ac_results	HBV genotype and subtype results
hbrn_central_ac_results	Results for the following tests: HBV DNA (quantitative and log10), HBeAg quantitative, HBeAg qualitative, HBsAg quantitative, HBsAg qualitative, HBsAg confirmative, anti-HBs, anti-HDV
hbrn_supp_ac_results	Results for anti-HBe and anti-HBs completed on available samples for cases in which anti-HBe or anti-HBs were not done at the clinical site.
hbrn_pathology_ac_results	Pathology central read results when liver biopsy slides were obtained.
hbrn_event_adj_ac	The following clinical events underwent a central review by the HBRN Adjudication Committee to verify event, etiology when applicable, and date of onset/diagnosis. Refer to Adjudication materials and process_v1.8 for details. - ALT flare - Acute HBV - HBeAg loss - HBsAg LOSS - Cirrhosis - Hepatocellular carcinoma - Liver transplantation - Hepatic decompensation - Death

<u>Participant ID variables</u>

ID

- ID assigned at current site. Needed due to participants transferring from one HBRN site to another.

ORIG_ID

- ID at first enrollment in HBRN – if participant moved to a new site during the Cohort Study, a new ID was assigned at that current site. Use ORIG_ID to link participants' data across HBRN datasets.