



**Reminders**

**+NETT/SIREN Study Coordinator Call**  
November 3rd at 1pm ET

**+Virtual IM**  
November 9th @10am ET  
November 12th @3pm ET

**+NETT/SIREN Steering Committee Call**  
November 25th at 12pm ET

**OSU & Hennepin—Participant Enrolled**

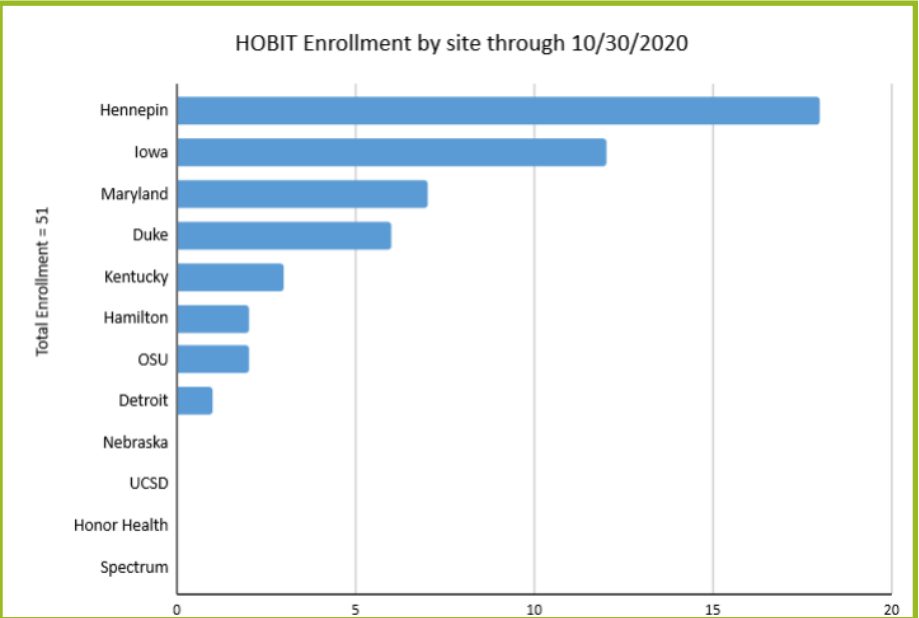
Great job to both OSU & Hennepin County Medical Center on their recent enrollments!

**Kentucky—Complete with EFIC**

Kudos to Kentucky on completing EFIC activities!

**St. Mary's—Chamber Modifications & Contract Complete, IRB Application Submitted, EFIC activities and Simulation Scheduled**

Thank you to St. Mary's on making great progress towards opening to enrollment!



## HOBIT Updates

### HOBIT / BioHOBIT Virtual Investigator Meeting

We will be holding a Virtual HOBIT / BioHOBIT Investigator Meeting Monday November 9, 10:00am – 12:00pm (Eastern) and Thursday November 12, 3:00pm – 5:00pm (Eastern). The meeting will be offered at two different times to accommodate schedules.

The PI and Primary Study Coordinator from each site will be required to attend one of the offered times. Other study team member are welcome to attend as well.

Appointment requests for both dates have been sent to everyone. Please respond to the appointment request for the time that you are available to attend.

### HOBIT Informed Consent Form (Version 12 Oct 2020)

A new HOBIT Informed Consent Form (*Advarra IRB Approved Version 12 OCT 2020*) was recently released for use. This new form contains provisions for collecting blood and CSF samples from HOBIT subjects. Edits to the consent form include:

- “Your loved one” has been replaced with "Your family member (or a person you represent)".
- Inclusion of blood and cerebrospinal fluid sample collection information, risks, and sample storage.
- Inclusion of information about FITBIR.
- Inclusion of statement for alternatives of participation.
- Grammatical updates

Please use (*Advarra IRB Approved Version 12 OCT 2020*) for enrolling all new HOBIT subjects. Your site specific eConsent link is currently updated to reflect the new change. Study teams will be able to find the paper copy as well as the ICF Approval Notice in WebDCU.

### BioHOBIT Shipping Address for Kits

In anticipation of the ancillary study BioHOBIT starting soon, we would like to ship blood draw kits to sites. If you have not done so, please send Natalie Fisher (brownnat@med.umich.edu) the address of where we should send kits to your site.

