

Monthly updates from the Human Research Protections Office

#### **New Educational Resources**

Below are new documents that have been uploaded to our website. Click on the blue document image to view on our website.



## **Comprehensive Informed Consent Guidance Document**

The informed consent process requires ongoing information exchange beyond just obtaining signatures. See our detailed guidance for essential information on consent documentation, signatures procedures and special population considerations.



### **Single IRB Review Reliance Agreement Overview**

A reliance agreement allows one IRB (IRB of Record) to review research for multiple sites, reducing duplicate oversight. Several factors determine what IRB will serve as the IRB of Record. Click on document for overview.



#### IRB Submission Guideline for Human Sample Genetic Testing

See detailed guidance for specific requirements on sample handing, analysis methods and participant protections.



THIS Friday,

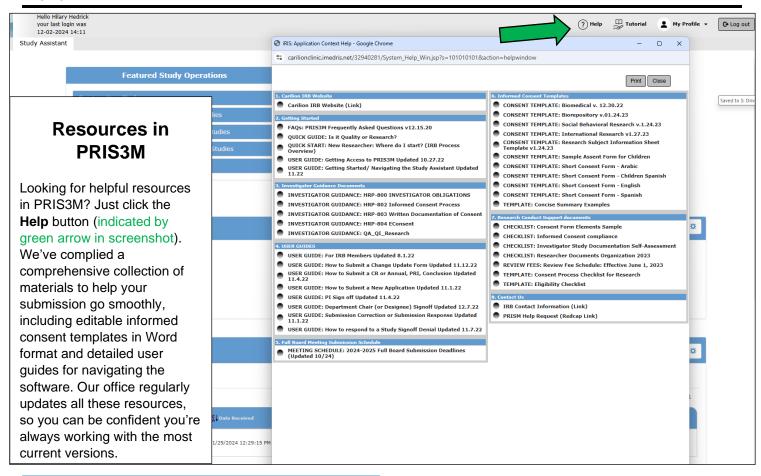
**January 17** 

12-1 PM

**LIVE** Webinar



# The Review





# Join Our Team: IRB Regulatory Affairs Administrator Position Now Open

As our office continues to grow, we're creating an additional IRB Regulatory Affairs Administrator position. This role will be instrumental in evaluating study submissions, ensuring regulatory compliance, and providing guidance to investigators and research teams. Interested candidates can click <a href="https://example.com/here">here</a> to apply!

Don't forget to visit and bookmark our Learning Channel. Click on the YouTube icon below.





Educational needs, inquires, and/or to schedule trainings, email hhhedrick@carilionclinic.org