FAQs

FAQ Categories:

- Biorepositories
- Consent
- Continuing Review and Expiration of IRB Approval
- iRIS
- Exemption from IRB Review
- External (Non-Duke) Sites
- FDA-Regulated Research
- General IRB Review Process
- Helpful Tools
- HIPAA
- Key Personnel
- Recruitment/Enrollment and Payment
- Reporting to the IRB
- Special Considerations
- Specialty Committees
- Vulnerable Populations