

Institution	Procedures and Policies for: (specimen acquisition, CTSA, tumor banking, IRB, etc.)	Brief Title	Current Version Date
Duke	Human Research Protection	Advertisement of Research Studies	3/4/2016
Duke	Human Research Protection	Amendments to IRB-Approved Studies	3/1/2016
Duke	Human Research Protection	Appropriate Study Personnel to Conduct the Consent Process	3/11/2019
Duke	Human Research Protection	Autopsy Authorizations	11/14/2007
Duke	Human Research Protection	Blood Drawing	12/13/2012
Duke	Human Research Protection	Case Reports	2/16/2016
Duke	Human Research Protection	Certificates of Confidentiality	12/22/2017
Duke	Human Research Protection	Children Who Are Wards of the State	2/19/2016
Duke	Human Research Protection	Research Involving Children	3/1/2016
Duke	Human Research Protection	Compensation for Study-Related Injury	1/3/2008
Duke	Human Research Protection	Compensation for Research Participation	2/15/2016
Duke	Human Research Protection	Conduct of an IRB Meeting	3/3/2016
Duke	Human Research Protection	Conflict of Interest for IRB Members & Consultants	6/5/2019
Duke	Human Research Protection	Consent and Its Documentation	2/1/2016
Duke	Human Research Protection	Consent Monitoring	3/1/2016
Duke	Human Research Protection	IRB Use of Consultants	3/1/2016
Duke	Human Research Protection	Continuing Review	3/7/2016
Duke	Human Research Protection	Determining Frequency of Continuing Review	3/2/2016
Duke	Human Research Protection	Contraceptive Use	7/2/2019
Duke	Human Research Protection	Corrective Actions for COI	6/5/2019
Duke	Human Research Protection	Data and Safety Monitoring	3/6/2016
Duke	Human Research Protection	Decision Making Capacity in Adults	3/3/2016
Duke	Human Research Protection	Department of Defense (DoD) Research	2/19/2016
Duke	Human Research Protection	Department of Education Research	3/1/2016
Duke	Human Research Protection	Emergency Use	3/2/2016
Duke	Human Research Protection	Engagement and Recruitment of Patients to a Research Protocol	3/1/2019
Duke	Human Research Protection	Expedited Review	2/19/2016
Duke	Human Research Protection	Expiration of IRB Approval and Closing Reports	2/29/2016
Duke	Human Research Protection	Reliance on the IRB of Another Institution, Organization, or an Independent IRB	3/3/2016
Duke	Human Research Protection	External Research by Duke Faculty or Staff	5/29/2011
Duke	Human Research Protection	External Research by Duke Trainees	3/3/2016
Duke	Human Research Protection	External Use of Duke Data by Former Students & Employees	8/31/2016
Duke	Human Research Protection	Fees Charged by the IRB	11/1/2016
Duke	Human Research Protection	Gadolinium Contrast Agents	9/10/2019
Duke	Human Research Protection	HIPAA as it Relates to Research	2/26/2016
Duke	Human Research Protection	HIPAA Exemption	3/1/2016
Duke	Human Research Protection	Holds, Suspensions, and Terminations	3/7/2016
Duke	Human Research Protection	Humanitarian Use Devices	3/8/2016
Duke	Human Research Protection	In Vitro Diagnostic Devices	3/1/2016
Duke	Human Research Protection	Institutional Official (IO)	2/24/2016
Duke	Human Research Protection	Institutional Oversight of Human Research	2/24/2016
Duke	Human Research Protection	Interaction with Compliance Offices by the IRB	2/19/2016
Duke	Human Research Protection	International Research	1/28/2016
Duke	Human Research Protection	Investigator Who Is Also a Sponsor	2/29/2016
Duke	Human Research Protection	Investigator/ Study Team Concerns	3/1/2016
Duke	Human Research Protection	IRB Approval Stamp on Consent Forms	8/19/2015
Duke	Human Research Protection	Key Personnel Qualifications	8/31/2017
Duke	Human Research Protection	Legal Counsel Opinion	3/1/2016
Duke	Human Research Protection	Legally Authorized Representatives (LAR)	3/1/2016
Duke	Human Research Protection	Local Physician Participation	2/23/2016
Duke	Human Research Protection	Mandatory State Reporting Requirements	3/7/2016
Duke	Human Research Protection	Membership, Voting and Quorum for the IRB	5/20/2016
Duke	Human Research Protection	Minutes of the IRB Meetings	3/7/2016
Duke	Human Research Protection	Modifications Processing Procedure	3/1/2016
Duke	Human Research Protection	Multi-Site Research	1/6/2017
Duke	Human Research Protection	Nomination and Evaluation of IRB Members	3/2/2016
Duke	Human Research Protection	Non-compliance	2/19/2016
Duke	Human Research Protection	Non-English-Speaking Subjects	1/11/2017
Duke	Human Research Protection	Pathology Tissue/Specimen Exclusions	n/a
Duke	Human Research Protection	Pediatric Risk Level	10/13/2014
Duke	Human Research Protection	Permissible Interim Activities	8/3/2016
Duke	Human Research Protection	Placement of a Research Consent Form in the Participant's Medical Record	2/18/2014
Duke	Human Research Protection	Planned Research in an Emergency Setting	3/1/2016
Duke	Human Research Protection	Pregnancy Testing	7/2/2019
Duke	Human Research Protection	Pregnant Partners of Research Subjects	11/5/2014
Duke	Human Research Protection	Pregnant Women, Fetuses, and Neonates	3/1/2016
Duke	Human Research Protection	Principal Investigator Qualifications	3/8/2016
Duke	Human Research Protection	Prisoners	2/24/2016
Duke	Human Research Protection	Processing Payments to Participants	n/a
Duke	Human Research Protection	Quality Improvement (QI) vs. Research (Policy and Checklist)	3/3/2016
Duke	Human Research Protection	Questionnaires by Telephone, Mail, or E-Mail	8/9/2004
Duke	Human Research Protection	Re-opening a Closed Study	1/6/2016
Duke	Human Research Protection	Recruiting Students, Employees (including study team members), Friends, & Family Members	3/19/2018
Duke	Human Research Protection	Reliance by External Sites on the DUHS IRB	3/1/2016
Duke	Human Research Protection	Reporting to DUHS Officials and Federal Agencies	8/30/2017
Duke	Human Research Protection	Research Databases, Specimen Repositories, and Contact Lists	3/7/2016
Duke	Human Research Protection	Research for Which Review by the DUHS HRPP Is Required	3/8/2016

Duke	Human Research Protection	Research Involving Devices	2/24/2016
Duke	Human Research Protection	Research Involving Drugs or Biologics	2/24/2016
Duke	Human Research Protection	Research Using Coded Private Information or Biospecimens	3/1/2016
Duke	Human Research Protection	Retention of Records	3/1/2016
Duke	Human Research Protection	Retrospective Research	10/14/2004
Duke	Human Research Protection	Scientific Validity of a Research Protocol	3/1/2016
Duke	Human Research Protection	Short Form Consent	3/2/2016
Duke	Human Research Protection	Significant Risk vs Non-Significant Risk Device Studies	3/4/2016
Duke	Human Research Protection	Specialty Committee Review	2/29/2016
Duke	Human Research Protection	State Law as It Relates to Research	3/7/2016
Duke	Human Research Protection	Subject Problems, Complaints, Concerns and Questions	3/1/2016
Duke	Human Research Protection	Thalidomide, Lenalidomide, or Their Analogs	10/11/2006
Duke	Human Research Protection	Unanticipated Problems (Prompt Reporting to the IRB)	8/9/2019
Duke	Human Research Protection	Undergraduates Conducting the Consent Process	11/29/2017
Duke	Human Research Protection	Undue Influence Upon the IRB	3/3/2016
Duke	Human Research Protection	Determining Which Studies Need Verification From Sources Other Than the Investigator	3/2/2016
Duke	Human Research Protection	Waiver for a Single Person (Single Person Exception)	2/26/2016
Duke	Human Research Protection	Waiver of Documentation of Consent	3/3/2016