IRBNet Document Type and Document Description

Instructions: When uploading documents IRBNet will require you to choose a document type classification. Below is a list of common documents we require and the IRBNet equivalent document type classification.

Document Description	IRBNet Document Type
Abstract/Summary of Recent literature related to the study	Continuing Review/Progress Report
Adverse Event Table	Adverse Event Report
Assent Form for Minors age 12-17	Child Assent
Cancer Center Scientific Review Committee Approval or Waiver	Other
Conflict of Interest Committee Review	Other
Consent Form	Consent Form
Data Safety Monitoring Board / Committee (DSMB/DSMC) Reports	Continuing Review/Progress Report
Debriefing script for deception studies	Consent Form
Drug/Package Insert	Other
Eligibility Screening Script	Consent Form
FDA Form for Investigator initiated investigational drug clinical trial	Protocol
Federal Grant Application / Proposal describing the study	Proposal
HRP-211 Application for Initial Review	Application Form
HRP-212 Continuing Review Progress Report for closing the study	Closure/Final Report
HRP-212 Continuing Review Progress Report for continual renewal of the study	Continuing Review/Progress Report
HRP-213 Modification	Amendment/Modification
HRP-214 Reportable New Information *For adverse event reporting	Adverse Event Report
HRP-214 Reportable New Information *For non-adverse event reporting	Reportable Event (Non-AE)

IRBNet Document Type and Document Description

Document Description	IRBNet Document Type
HRP-503 Protocol or Description of Study	Protocol
HUD Patient Information	Other
International IRB / Ethics Board Approval	Letter
Investigational Devices Record Form	Protocol
Investigational Drug Information Form with the PI signature	Protocol
Investigator's Brochure	Investigator's Brochure
IRB Fee Invoicing Form	Other
Letter from FDA or DHHS regarding the drug or device	Protocol
Letter of Action with reviewer comments/concerns	Letter
Letter of Information for Minors age 7-11	Child Assent
Miscellaneous	Other
Operating Room Resource	Other
Questionnaire, Survey, Assessment, Interview Script	Questionnaire/Survey
Radiation Use Committee Approval or Waiver	Other
Recruitment materials – Flyer, Brochure, Ads, Letter of Inquiry, Telephone/Oral Recruitment Script, etc.	Advertisement
Reportable New Information (RNI) Summary Table	Protocol Deviation / Violation Report
Research Personnel List	Application Form
Response memo from the PI addressing IRB comments/concerns	Letter
Site/School Approval Letter	Letter
Sponsor or Cooperative report relevant to the renewal	Continuing Review/Progress Report
Sponsor Protocol or Investigator Initiated Study Protocol	Protocol
Surrogate Self-Assessment Checklist	Consent Form