Appendix A: Children
This form is required when children/wards are involved in the research. Children are defined as individuals who have not attained the legal age for consent to treatments or procedures involved in the research, under the applicable law of the jurisdiction in which the research will be conducted. See Research Involving Children for guidance.

Appendix B: Pregnant Women and Neonates
This form is required when pregnant women/ neonates are research subjects. See the guidance on Pregnant Women and Neonates for more information.

Appendix C:
This form is required when prisoners are research subjects. See the guidance on Prisoners for more information.

Appendix D: Waivers of Consent and Signatures or Alterations
This form is required when a waiver of informed consent, alteration of consent, waiver of documentation of consent, or alteration or waiver of protected health information (PHI) is needed.

Appendix E: Multi-Site Research
A multi-site study is a study that involves multiple institutions engaged in the research project. This form is required when the Northern Arizona IRB will review research activities for an investigator or research staff who are not affiliated with the University. See Ceded IRB Review for more information. This form is not needed if each research site will obtain their own IRB approval.

Appendix F: Drugs and Devices
This form is required when the Human Research is a clinical investigation of a drug or device, which is "any experiment that involves a test article on one or more human subjects that either (1) requires FDA approval, or (2) is intended to be submitted to or inspected by the FDA for research or a marketing permit." (21 CFR 50.3(c))