Informed Consent Process

Informed Consent is the knowing consent of an individual or his/her legally authorized representative which is obtained without undue inducement or element of force, fraud, deceit, duress, or other forms of constraint or coercion. Informed consent is not just a form or signature, but a process of information exchange that includes:

⇒ subject recruitment materials
⇒ verbal instructions
⇒ written materials
⇒ questions/answer session
⇒ agreement documented by signature

The investigator must obtain legally effective written informed consent prior to enrolling a subject in a research project unless a specific exemption or waiver has been approved by the IRB, thereby waiving the requirement for informed consent or the requirement to obtain signed informed consent. Unless the IRB waives the requirement, informed consent must be documented by the use of a written consent form approved by the IRB, and signed by the subject or the subject’s legally authorized representative. The IRB reviews all informed consent documents for adherence to Federal regulations regarding the required elements of informed consent and for assurance of the adequacy of the information contained in the informed consent. The IRB has the authority to observe or to have a third party observe the consent process and the research. (Refer to the quality improvement policy).

Whenever the IRB requires documentation of informed consent, before a subject can participate in the research, the consent document must be signed and personally dated by the subject or by the subject's legally authorized representative, and by the person who conducted the informed consent discussion. During this period of prospective enrollment, the investigator (or qualified designee) must ascertain, either during the preliminary telephone interview (interest query), face-to-face encounter or by review of the medical history, the subject’s ability to provide consent (HIPAA regulations regarding sharing/access to PHI apply). Before participation in the research, the subject or the subject's legally authorized representative must be given a copy of the signed and dated written informed consent form and any other written information provided to the subjects. During a subject’s participation in the research, the subject or the subject’s legally authorized representative must be given a copy of the signed and dated consent form updates and a copy of any authorizations to PHI, or amendments to the written information originally provided.
Level

The informed consent document must be written using language that can be understood by someone reading at the seventh grade level. Medical terminology should be avoided or defined. The consent form is a statement addressed to the participant and should read as such. Separate forms may be required for different participant groups (parents, children) as well as for release of particular types of information (photographs, audio recordings video recordings).

Version Date

All Informed Consent Forms must bear a version date in the footer on each page of the consent. The version date must be updated whenever a revision is made to the informed consent document. (Refer to modification policy)

Elements

The informed consent document must contain all the required elements of Informed Consent, as well as any pertinent additional elements

Required Elements in Informed Consent: seeking informed consent the following information shall be provided to each subject (§46.116, 21CFR50, 21CFR56):

⇒ A statement (introduction) that the study involves research, an explanation of the purposes of the research and the expected duration of the subject’s participation, a description of the procedures to be followed, and identification of any procedures which are experimental [§.116(a)(1)] [§.50.25(a)(1)].

⇒ Possible Risks/Discomforts - state any known risks, side effects [§.116(a)(2)] [§.50.25(a)(2)]. In double-blinded studies, risks or possible reactions should be listed separately for each agent in each arm of the study.

⇒ Possible Benefits - describe potential benefits which might be expected by the subject and society in general [§.116(a)(3)] [§.50.25(a)(3)]. If the individual will receive no benefit this must be stated.

⇒ Financial Costs - list possible financial costs to participant [§.50.25(b)(3)]

⇒ A disclosure of appropriate alternative procedures or courses of treatment, if any, that might be advantageous to the subject [§.116(a)(4)] [§.50.25(a)(4)]. If there are no alternatives, so state. (If there are alternatives, please describe them.)
⇒ Confidentiality - A statement describing the extent, if any, to which confidentiality of records identifying the subject will be maintained [§.116(a)(5)] [§.50.25(a)(5)].

⇒ Voluntary Participation - note that participation is voluntary and subject may refuse to participate with no penalties [§.50.25(a)(8)]. List point of contact by name and phone number to call to terminate participation [§.116(a)(7-8)] [§.50.25(a)(7)] (two names and two different telephone numbers are required).

⇒ For research involving more than minimal risk, an explanation as to whether any compensation [§.50.25(a)(3)] and an explanation as to whether any medical treatments are available if injury occurs [§.50.25(a)(6)], and if so, what they consist of, or where further information may be obtained [§.116(a)(6)]

⇒ Injury / Complications - An explanation of whom to contact for answers to pertinent questions about the research and research subjects’ rights, and whom to contact in the event of a research-related injury to the subject [§.116(a)(7)] [§.50.25(a)(7)]. Describe in detail how complications will be handled.

⇒ A statement that participation is voluntary, refusal to participate will involve no penalty or loss of benefits to which the subject is otherwise entitled, and the subject may discontinue participation at any time without penalty or loss of benefits to which the subject is otherwise entitled [§.116(a)(8)] [§.50.25(a)(8)].

⇒ Consent - by signing the consent form, the subject certifies that the document has been read to them, that they understand and that they have received a copy. The individual has been given the opportunity to ask questions and to discuss participation with the investigator.

⇒ Signatures / Dates - Each page of the informed consent must be date-stamped by the IRB Director or Coordinator and initialed by the subject.

Additional Elements of Informed Consent: Additionally, one or more of the following elements of information must also be provided to each subject if required as indicated below:

⇒ The consent process must include a statement that the particular treatment or procedure may involve risks to the participant which are currently unforeseeable UNLESS the risk profile of all research-related interventions is well known and the research involves no investigational drugs or devices.

⇒ The consent process must disclose that the particular treatment or procedure may involve risks to the embryo or fetus, if the participant is or may become pregnant, which are currently unforeseeable UNLESS the research excludes
women of child bearing potential and pregnant women or the risk profile of all research interventions or interactions on embryos and fetuses is foreseeable or there is no reasonable expectation that this research causes risks to fetuses or embryos.

⇒ The consent process must disclose anticipated circumstances under which the participant’s participation may be terminated by the investigator without regard to the participant’s consent UNLESS there are no anticipated circumstances under which the participant’s participation will be terminated by the investigator without regard to the participant’s consent.

⇒ The consent process must disclose any additional costs to the participant that may result from participation in the research UNLESS there are no costs to the participant that may result from participation in the research.

⇒ The consent process must disclose the consequences of a participant’s decision to withdraw from the research and procedures for orderly termination of participation by the subject UNLESS there are no adverse consequences (physical, social, economic, legal, or psychological) of a participant’s decision to withdraw from the research.

⇒ The consent process must disclose that significant new findings developed during the course of the research which may relate to the participant’s willingness to continue participation will be provided to the participant UNLESS significant new findings during the course of the research which may relate to the participant’s willingness to continue participation are unlikely.

⇒ The consent process must disclose the approximate number of participants involved in the study UNLESS the approximate number of participants involved in the study is not important to a decision to take part in the research.

For applicable clinical trials, the consent process must disclose “A description of this clinical trial will be available on http://www.ClinicalTrials.gov, as required by U.S. Law. This Web site will not include information that can identify you. At most, the Web site will include a summary of the results. You can search this Web site at any time.”

**VA Required Paragraphs**

The only informed consent document that the VA can recognize is the VA Form 10-1086. A VA Form 10-1086 must be used as the consent form for all VA research. The VA Form 10-1086 must incorporate all the elements required by regulations. In addition, the following statements are required to be inserted in VA 10-1086s.

⇒ VA 10-1086 must contain this statement if the research involves an investigational drug with an IND or a medical device with an IDE) A verbatim statement: “I have been told because this study involves articles regulated by...
the FDA, the FDA may inspect research identifying me as a subject of this investigation.”

⇒ VA 10-1086 consents must contain a statement that a veteran—subject will not be required to pay for care received as a subject in a VA research project except if they are in an eligibility category that requires they pay a co-pay for medical services that are not part of the study.

Example 1: “You will not be charged for any treatments or procedures that are part of this study.

However, if you are required to make co-payments for services provided by the VA or if you receive treatment that is part of your usual medical care, you or your third-party payer (e.g., insurance company) may be billed.”

1) In the confidentiality section, the 10-1086 must include a statement that the Government Accounting Office (GAO), the ETSU/VA IRB, R&D, FDA, OHRP, DHHS, ORO, and any other applicable institutions may have access to the records

2) VA 10-1086 must contain this Injury/Complications paragraph "According to VA Regulations [38CFR17.85(a)] the medical facility shall provide necessary medical care to a research subject injured as a result of participation in a research project. However, no additional compensation has been set aside. You have not waived any legal rights or released the VA or its agents from liability for negligence by signing this form.”

3) VA 10-1086 must contain this statement required for veteran subjects "If you are a veteran taking part in a research study at the James H. Quillen VAMC, a copy of your signed/dated consent form will be placed in your medical record.”

4) VA 10-1086 must contain an adequate description of any payment. (must include timing; method of payment; and if subject is being paid by VA check through Austin, the informed consent must note that the social security number will be required to process the check and that payments of any amount will be reported to the IRS and may be counted as income)

5) VA 10-1086 must contain this statement if the researcher believes that bodily fluids, substances or tissues of a research subject could lead to the development of a commercially valuable product "I authorize the use of my bodily fluids, substances or tissues for research purposes.”

6) VA 10-1086 must contain Signature and date lines for the following

   a) subject or the subject’s legally-authorized representative, and

   b) If required by the IRB, witness whose role is to witness the subject’s or the subject’s legally-authorized representative’s signature, and
c) person obtaining the informed consent

7) If the sponsor or IRB requires a witness to the consenting process in addition to the witness to the subject’s signature and if the same person needs to serve both capacities, a note to that effect is placed under the witness’s signature line.

8) VA consents must include information about where and how a veteran could verify the validity of a study and authorized contacts.

9) The name of the study, the name of the PI, and the sponsor of the study

10) Clearly defines for the subject the risks related to the research and, therefore, need to be discussed with the research team, versus risks of usual care provided by the subject’s health care provider.

11) Includes language advising subjects to review the risks of usual care with their health care providers.

12) The protocol and consent document are consistent with the HIPAA Authorization.

13) Required in both the ICD and HIPAA if the study is collaborative: The data resulting from this study are to be used in a collaborative study that combines VA data with non-VA data. The data are to be disclosed to the Coordinating Center site (insert name) where the data will be combined and analyzed for the study.

14) If the specimens are to be retained after the end of the study for future research, the consent must disclose where the specimens will be retained, who will have access to them, and how long they will be retained. (All applicable policies, including organizations, VA and other federal requirements must be met for handling, use and storage of biologic specimens and data.)

15) If any of the data are to be retained after the end of the study for future research, the consent must disclose where the data will be retained, and who will have access to the data. (All applicable policies, including organizations, VA and other federal requirements must be met regarding the use and storage of data.)

16) If the participant will be re-contacted for future research, whether within a VA facility or outside a VA facility, the consent must disclose this information.

17) If the participant will receive a report of aggregate results or any results specific to the participant, the consent must disclose this information.

18) If the research includes taking photographs or making video or phone recordings that will be used for research purposes, the consent document must include information describing any photographs, video, and/or audio recordings to be taken or obtained for research purposes, how the photographs, video, and/or audio recordings will be used for the research, and whether the photographs, video, and/or audio recordings will disclosed outside VA. The consent for research does not give legal authority to disclose the photographs, video, and/or
audio recordings outside VA. A HIPAA Authorization is needed to make such disclosures.

19) Any real or apparent conflict of interest by investigators where the research will be performed

For VA studies, in the event that someone other than the investigator will be conducting the consent interview or obtaining consent, the investigator must provide a formal and prospective delegation of the responsibility of obtaining informed consent (in the protocol or the IRB submission forms). The delegate must have received appropriate training (completed CITI IRB requirements as well as protocol-specific training by the PI). The person, who must be a member of the study team, must be knowledgeable about the research to be conducted and the consenting process, and must be able to answer questions about the study.

The informed consent form must be signed and dated by the subject or the subject’s legally authorized representative, and the person obtaining the informed consent.

Compensation

Compensation in the form of payments to subjects for their participation in a research study must be IRB approved. The amount must be commensurate with the expected contributions of the subject. The amount and terms of the payment (check or cash, etc., as well as timing of receipt of compensation) must be stated precisely. The Informed Consent form should reflect a fair and appropriate amount that does not place undue pressure (coercion) on the volunteer.

For Non-English Speaking Subjects

Regulations require that informed consent be obtained in a language that is understandable to the subject (or to the subject’s legally authorized representative). Validated translations of consent forms must be available for non-English speaking subjects. To address possible questions or concerns raised by the prospective subject, a qualified translator must be present and may act as a witness. Documentation of the qualifications of the translator must be added to the research records and available for administrative or QI auditing upon request.

When a full-length ICD embodying all the required elements is required by the IRB to document the consent process, that form must be written in a language understandable to the subject. The IRB requires that the appropriately translated ICD be submitted to the IRB for review and approval prior to their use in enrolling subjects. When informed consent is documented in accordance with HHS regulations at 45 CFR 46.117 (b) (1), the written ICD should embody, in language understandable to the subject, all elements necessary for legally effective informed consent. The IRB may use expedited review
procedures in approving such documents if the English language ICD has already been approved, and the investigator attests in writing to the accuracy of the translation.

**Telephone Consenting is Not Recommended**

For non-VA Studies, an investigator may, however, conduct a preliminary telephone interview to query a participant’s interest in possibly participating in the research. If initial contact with prospective study subjects is to be made by telephone, a script of the phone contact is to be reviewed and approved by the IRB **prior to use**. Similarly if initial contact is to be made by mail, the content of the mailing script and list must be reviewed and approved by the IRB **prior to initiation**.

Following the telephone interview, and under certain circumstances (to be determined by the IRB with evidence of need), the investigator can fax an IRB approved informed consent document to the participant for his or her review. Documentation of this process is critical. The signature page must be witnessed by an individual known to the IRB or notarized, and reflect both the date and time of each signature. Enrollment may begin once the Principal Investigator receives a copy of the signed and notarized consent document. The original document must be immediately (within 24 hours) forwarded to the study file or patient record as appropriate, to be placed in the patient’s medical record upon discharge. The date and time that the consent document (signed, notarized original copy) was entered into the patient chart must additionally be noted in the progress notes.

In VA studies, during the recruitment process, researchers must make initial contacts with veterans in person and/or by letter prior to any telephone contact, unless there is written documentation that the participant is willing to be contacted by phone about the study in question or a specific kind of research. (E.g. If the prospective participant has diabetes, the participant may indicate a desire to be notified of any diabetes-related research studies.) The initial contact must provide a telephone number or other means that veterans can use to verify the study constitutes VA research. (One source of information about clinical trials is [http://www.clinicaltrials.gov](http://www.clinicaltrials.gov)).

**Legally authorized representative**

A legally authorized representative shall be an individual or judicial or other body authorized under applicable law to consent on behalf of a prospective subject to the subject's participation in the procedure(s) involved in the research. Legally authorized representative is synonymous with legally acceptable representative. [21 CFR §50.3(l)] [45 CFR §46.102(c)]. A signature line for legally authorized representatives may only be included on the consent document if the IRB approves the enrollment of participants based on the permission of a legally authorized representative. For VA studies, consent by legally authorized representative is limited to situations where the prospective
participant is incompetent or has impaired decision-making capacity, as determined and documented in a patient’s medical record in a signed and dated progress note.

For cognitively impaired veterans subjects who would be enrolled in research studies at a Department of Veteran Affairs Facility, permission is obtained from a health care agent appointed by the person in a Durable Power of Attorney for Health Care (DPAHC) or similar document, court-appointed guardians of the person, or from next-of-kin in the following order: Spouse, Adult child (18 years or older), parent, adult sibling (18 years or older), grandparent, or adult grandchild (18 years or older) or close friend.

For non-VA studies, in the case of an incompetent individual or an individual who lacks decision-making capacity, the individuals’ health care decision maker (LAR) is designated in order of preference as one of the following:

A. Court-appointed Conservator or Guardian of the individual with authority to make health care decisions for the patient

B. Person named in the patient’s Durable Power of Attorney for Health Care (DPAHC)

C. If the patient does not have a court-appointed guardian or conservator, AND does not have a person authorized to act under a Durable Power of Attorney for Health Care, then both of the following must be true for the individual identified to serve as the surrogate decision-maker for this patient:

1. The person identified above is an adult who has exhibited special care and concern for the patient, who is familiar with the patient’s personal values AND who is reasonably available to serve as a surrogate.

2. It appears as though the person can make health care decisions for the patient in accordance with the patient’s individual health care instructions, if any, and other wishes, if known to the health care decision maker. IF the patient has not given individual health care instructions, and the patient’s specific wishes are not known, the health care decision maker can make a determination of the patient’s desires or best interests in light of the patient’s personal values and beliefs to the extent they are known.

This person may include, in order of descending preference, the patient’s spouse, the patient’s adult child, the patient’s parent, the patient’s adult sibling, any other adult relative of the patient or another adult who satisfies the requirements listed above.

The investigator must indicate in the application that he/she is requesting to utilize consent of a health care decision maker. The IRB must approve the use of an LAR. The IRB will review the rationale for this request, and ensure there are appropriate safeguards in place.

In addition, if research involving adults who are unable to consent is being conducted outside of the state of Tennessee, the investigator must consult Legal Counsel about which individuals are “legally authorized representatives” when the research is
conducted in that jurisdiction. Researchers must submit a copy of this legal opinion to the IRB.

Investigators must obtain a copy of the court order if a court appointed conservator or guardian gives consent. Investigators must obtain a copy of the DPAHC if Person named in the patient’s Durable Power of Attorney for Health Care (DPAHC) gives consent. In addition, if an individual identified to serve as the surrogate decision-maker for this patient gives consent, the investigator must document additional information evidencing the person’s qualifications to serve as a surrogate. That information must include how long this person has lived with the patients, how long this person has known the patient and how often this person sees the patient, and any other evidence of the appropriateness of the selected surrogate.

Exculpatory language

Exculpatory language is prohibited. Informed consent, whether oral or written documents, may not contain any exculpatory language through which the subject is made to waive or appear to waive any of the subject’s legal rights, or releases or appears to release the investigator, the sponsor, ETSU, VAMC or its agents from liability for negligence.

Requiring a Witness Signature on the Consent Form

The institution and the IRB reserve the right to require the signature of a witness on informed consent documents, as a matter of policy, under certain situations. Both the institution and the IRB have the authority to require protections for human subjects that exceed the minimum standards required under federal or state regulations. If the sponsor or the IRB require a witness to the consent process who also witnesses the signature, a note to that effect must be added to the consent document under the witness’s signature line.

For VA studies, the witness cannot be the person who obtained consent from the participant, but may be a member of the study team or may be a family member.

HIPAA Authorizations

When the HIPAA Authorization is embedded in the body of the ICD the IRB shall be responsible for reviewing both the content of the Authorization and its appropriateness to the research. When the HIPAA Authorization is attached to the ICD as an addendum (preferred), the IRB Chair, designee of the Chair, or the IRB Coordinator shall be responsible for the review.
Children

For participants < than 18 years of age, their parents or legal guardian are the legally authorized representative who may grant permission for their participation in research. When research is conducted in the state of Tennessee, children are all individuals under the age of 18 without exception.

In addition, if research involving children is being conducted outside of the state of Tennessee, the investigator must consult Legal Counsel to determine the definition of who is a “child” when the research is conducted in that jurisdiction. Researchers must submit a copy of this legal opinion to the IRB.

According to the Tennessee Department of Children Services (DCS), applicable policies by virtue of the court order granting DCS legal custody of certain children (e.g., foster children) that department is the agency that is authorized to grant permission for participation in research in their custody. The decision of whether to grant permission for research is made on a case-by-case basis by DCS. In such cases the PI must obtain a copy of the court order from DCS. The case manager (including the case manager’s supervisor(s) or Regional Administrators’ designee(s), the foster parent, and the contract agency caseworker are authorized by DCS to sign consent for routine medical care.

DCS is authorized under Tennessee State law to consent to health care for the child, and therefore can serve as a guardian as defined in Subpart D.

Applicable State Laws regarding Reporting Requirements

Mandatory Reporting of Abuse: Any person who has knowledge of or is called upon to render aid to any child who is suffering from or has sustained any wound, injury, disability, or physical or mental condition is required to report the harm immediately by telephone to the:

⇒ Judge having juvenile jurisdiction over the child;
⇒ County office of the department;
⇒ Sheriff of the county where the child resides; or
⇒ Chief Law enforcement official of the municipality where the child resides.
⇒ The report will include (to the extent known) the name, address, and age of the child, the name address of the person responsible for the care of the child, and the facts requiring the report.

Mandatory Reporting of Sexually Transmitted Disease: Every physician or other person who makes a diagnosis of, treats or prescribes for a case of sexually transmitted disease designated as reportable is required to report the case immediately to the
Department of Health. Reports include the name, address, age, sex, race, stage of disease, treatment, and control of the disease. Children 13 years of age or younger must be reported to the Department of Health. Reporting is required for children where sexual abuse is suspected regardless of injury to the Department of Health and the Department of Health will notify the Department of Children’s Services.

Mandatory Reporting of Cancer: All laboratories, hospitals, facilities and practitioners are required to report to the Department of Health within 6 months of diagnosis any cancer in an individual if making the initial diagnosis. The report includes the diagnosis, occupation, family history and personal habits of the person diagnosed with cancer.

Because of these laws, IRB members must evaluate studies to determine whether disclosure of the implications of the laws is required for legally effective informed consent. Because of these laws, investigators must ensure that the consent process provides participants with accurate information concerning required reporting. Investigators are also responsible for compliance with these regulations.

**Waiver of Informed Consent**

DHHS provides for waiving or altering elements of informed consent under certain conditions [§.116(c)-(f)]. FDA has no such provisions because the types of studies that would qualify for waiver or alteration are either not regulated by FDA or are covered by the Emergency Treatment provision of FDA Regulation 21 CFR 50.23.

Waiver of informed consent cannot be given when research is subject to FDA regulation.

The IRB may approve a consent procedure which does not include, or which alters, some or all of the elements of informed consent, or waive the requirement to obtain informed consent provided one of the following sets of conditions exists and is documented:

A. 45 CFR 46.116(c) (Must meet one of the following criteria from section 1 as well as criteria number 2)

1. The research or demonstration project is to be conducted by or subject to the approval of state or local government officials and is designed to study, evaluate, or otherwise examine at least one of the following:
   
a) public benefit or service programs; or
   
b) procedures for obtaining benefits or services under these programs; or
   
c) possible changes in or alternatives to those programs or procedures; or
   
d) possible changes in methods or levels of payment for benefits or services under those programs **AND**
2. The research could not practicably be carried out without the waiver or alteration.

B. 45 CFR 46.116(d) (Must meet all four criteria detailed below)

1. The research involves no more than minimal risk to the subjects; **and**

2. The waiver or alteration will not adversely affect the rights and welfare of the subjects **and**

3. The research could not practicably be carried out without the waiver or alteration **and**

4. Whenever appropriate, the subjects will be provided with additional pertinent information after participation.

If a waiver is granted, the minutes will document both the justification for and the approval of the waiver, along with any controverted issues raised by the IRB.

A waiver of parental permission (or student consent if the student is an adult) may not be granted if the study involves funding from the Department of Education and the study involves a survey, analysis, or evaluation that reveals information concerning the following categories:

A. political affiliations or beliefs of the student or the student’s parent;

B. mental or psychological problems of the student or the student’s family;

C. sex behavior or attitudes;

D. illegal, anti-social, self-incriminating, or demeaning behavior;

E. critical appraisals of other individuals with whom respondents have close family relationships;

F. legally recognized privileged or analogous relationships, such as those of lawyers, physicians, and ministers;

G. religious practices, affiliations, or beliefs of the student or student’s parent; or

H. income (other than that required by law to determine eligibility for participation in a program or for receiving financial assistance under such program)

In addition, all instructional materials, including teacher’s manuals, films, tapes, or other supplementary material which will be used in connection with any survey, analysis, or evaluation as part of any applicable program shall be available for inspection by the parents or guardians of the children.
Waiver of Documentation of Informed Consent

Under certain conditions, the IRB can waive the requirement that the subject sign the consent form. However, waiver of documentation of informed consent does not constitute waiver of informed consent. The IRB may waive the requirement for the investigator to obtain a signed consent form for some or all subjects if it finds either (46 CFR 117(c):

⇒ That the only record linking the subject and the research would be the consent document, and the principal risk would be potential harm resulting from a breach of confidentiality. Each subject will be asked whether the subject wants documentation linking the subject with the research, and the subject’s wishes will govern.

OR

⇒ That the research presents no more than minimal risk of harm to subjects and involves no procedures for which written consent is normally required outside of the research context.

In cases where the documentation requirement for informed consent is waived, the IRB may require the investigator to provide participants with a written statement regarding the research.

In addition, the IRB minutes will document required determinations regarding waiver of requirement for written documentation of informed consent. The minutes will also document the protocol specific findings justifying the determinations.

Observation of the Consent Process

Per 45 CFR 46, the IRB has the authority to observe or have a third party observe the consent process.

⇒ Situations where observation of informed consent may be requested include:

⇒ Studies where the capacity of the participant to provide informed consent may be questionable

⇒ Previous investigator serious or continuing non-compliance

⇒ High risk studies, such as Phase I trials

⇒ Complaint(s) received about the informed consent process

⇒ Any others as determined necessary by the IRB