

Summary of Chapter 10, Sections E and F of the
Corporate Policy for Protection of Human Research Subjects

The research coordinator can only obtain consent from the subject if there are no clinical issues and/or medical treatment questions posed by the subject.

The researcher coordinator can begin screening procedures, if the ICF is signed by the subject and themselves and if the screening procedures are non-investigational (assuming the screening procedures are described in the informed consent form).

The physician investigator will obtain the subject's consent to the screening procedures if the subject has clinical issues and or medical treatment questions.

The physician investigator will obtain the informed consent of the patient to participation in the investigational study and will sign the consent (attesting to other activities as well – see informed consent form template) before any investigational procedures take place with the subject.