AgingResearchBiobank

Informed Consent Questionnaire for Specimens

Specimen Informed Consent

1. Has an Institutional Review Board reviewed and verified that submission of the collection to the Biorepository for subsequent sharing with non-Study investigators for research purposes is consistent with the informed consent of study participants?

Yes     [ ]
No      [ ]

1. Does the Study informed consent document include restrictions on the use of the biospecimens?

Yes     [ ]
No      [ ]

1. If Yes, provide a list of restrictions.

Click here to enter text.

1. Submit a document summarizing any Study and/or site-specific restrictions/changes to the Study informed consent template regarding the use and storage of biospecimens with this Questionnaire.

The summary should include information on:

* restrictions on biospecimen use by non-Study investigators, if applicable
* restrictions on biospecimen use by research topic (e.g., disease or organ specific), genetic use restrictions, commercial entities, etc., if applicable
* changes to the restrictions for biospecimen use over time, including the effective date (s)
* site specific changes to the Study informed consent related to biospecimens
* how data and samples already collected will be managed if a participant withdraws consent to continue participation in the study