10. Informed Consent - Secondary Use of Existing Data/Records/Specimens

Completion of this section is required based on the response provided to questions 1-1.1, 5.2 and 5.3.

10.3* What type of informed consent will be obtained from subjects for the use of their data, records and/or specimens?

Select all that apply:

- Request for waiver of informed consent/parental permission/legally authorized representative consent
- Pre-existing consent(s) covers this activity
- Re-consent/assent subjects for use of existing data/records/specimens for a new research purpose

10.3.1* Describe terms and conditions of original consent process.

[Blank field]

10.3.2 Upload any documents related to consent, assent, permission, and or debriefing documents, including oral scripts here.

[Blank field]

10.3.3* Describe the plan to assent, consent, or re-consent the subject or the subject's legally authorized representative.

[Blank field]

Full waiver is very common for secondary-use studies

Activates 10.3.1 and 10.3.2 below. This applies to data/biospecimens obtained from some repositories, such as UMMS Central Biorepository. This answer often pairs with "Yes, always" to HIPAA authorization question 25-1.3.

Activates 10.3.3 and section 10-1.