

Human Tissue Standard Operating Procedures

The standard operating procedures for Human Tissue are:

- [Receipt, Labelling, Tracking and Storage of Human Tissue](#) (pdf)
- [Disposal of Human Tissue](#) (pdf)
- [Internal Audit of Human Tissue Authority Licenced Premises](#) (pdf)
- [Recording Human Tissue Act Related Adverse Events](#) (pdf)
- [Assessment of Risk to Personnel for the Handling of Human Tissue](#) (pdf)
- [Assessment of Risk to Human Tissue](#) (pdf)
- [External Transfer and Export of Relevant Material](#) (pdf)
- [Import of Relevant Material](#) (pdf)
- [Transportation of Human Tissue](#) (pdf)
- [Informed Consent for Research](#) (pdf)
- [Research Governance Audit](#) (pdf)
- [Complaints from Research Participants](#) (pdf)
- [Withdrawal of Consent](#) (pdf)

Relevant Research Standard Operating Procedures

Research standard operating procedures can be found [here](#).

Relevant Forms

- [HTA Adverse Events Form v 5.0](#) (docx)
- [Health and Safety Risk Assessment Guidance and Risk Assessment Form](#)
- [Request for External Transfer or Export of Human Tissue Form](#) (doc)
- [Authority to Import Form](#) (doc)
- [Consent Form Template](#) (Consent forms may vary according to the nature of the study. The template consent form is suitable for most studies but may require alteration. Text in red or within square brackets indicates wording that should only be included if appropriate for the study.)
- [Research File Content Checklist](#) (doc)
- [Disposal Form Template](#) (doc)

Relevant Guidelines

- [Guidelines for Research Files](#)