

BioIVT Ethical Procurement of Human & Animal Biological Specimens for Research Use

Human Biological Specimens for Research Use

BioIVT conducts its human research activities in accordance with regulations surrounding human research subject safety and protection which include, ethical principles that have their origin in the Declaration of Helsinki and are consistent with Good Clinical Practice (GCP). Pursuant to these regulations, prospective biological sample collections are administered under Institutional Review Board (IRB) approved protocols and IRB-approved investigators and sites. Informed consent is central to enrollment of research subjects under applicable regulations and IRB standards. Under these requirements, BioIVT ensures that research subjects are adequately consented using the current IRB-approved consent document and provided information on the use of their samples and data, prior to study participation.

Animal Specimens for Research Use

BioIVT ensures that sites responsible for the housing and care of animals for research use at minimum, meet United States Department of Agriculture (USDA) Animal Welfare Act (AWA) requirements and where applicable, Institutional Animal Care and use Committee (IACUC) approvals and requirements.



Amanshe Slaney
Research Compliance & Regulatory Affairs



Jeffrey Gatz
CEO