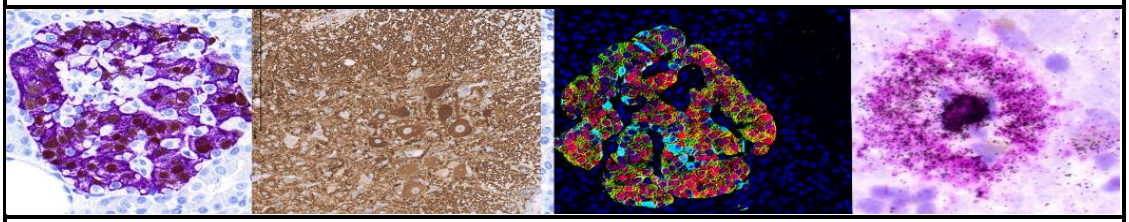
Biosamples Procurement Form

Biosamples Collection Questions	Project Collection Specifications
Company name:	
Company reference #:	
Talkbio project reference #:	
Disease or healthy diagnosis:	
Collected before or after surgery:	
Clinical inclusion criteria:	
Clinical exclusion criteria:	
Any treated /treatment-naïve donors acceptable:	
Sample format:	
Sample volume and aliquots:	
How many samples/sets/pools are needed:	
Shipment address:	
Desirable timeframe for delivery:	
Sample storage @ -20° C / -80° C / -150° C:	
Shipment conditions (ambient, +4° C, dry ice, LN2):	
Prospective collection with informed consent:	
Retrospective collection with consent waiver acceptable:	
Any additional testing, specifications or comments:	

CLINICAL DATA PACKAGES & LABORATORY TESTING

Please provide as much information as possible, clinical inclusion and exclusion criteria can be specified, treated and treatment-naive donors available, clinical data packages are provided as standard, and additional clinical and laboratory testing is available on demand.



PATIENT CONSENT & ETHICS POLICY

All prospective biosamples collections are approved by an Institutional Review Board or Independent Ethics Committee, who operate in accordance with local regulations (including ICH, HIPAA, and GCP), ensuring strict ethical standards consistent with the guidelines of the World Health Organization and ISBER, and patient confidentiality and safety, are protected. Biosamples are de-identified and consented for use in a wide range of research including the development of commercial products and services.