



## UK CLL Biobank Governance Committee Terms of Reference

### Overview of the biobank

The UK CLL Biobank was established in 2008 as a national biorepository linked to multiple CLL trials and observational studies developed through the NCRI CLL Study Group. Located in the University of Liverpool GCP laboratory and funded by Blood Cancer UK, its primary purpose is to drive and facilitate laboratory research to elucidate CLL biology and heterogeneity including variable therapy outcomes. Sample are collected, transported, processed and stored to GCP standards, with sample access overseen by the NCRI CLL Study Group.

### Purpose of the governance committee

1. To provide a pathway of accountability to biobank stakeholders
2. To provide advice on biobank strategy and operations

### Governance committee membership

Member	Role	Organisation
Prof Russell Patmore	Chair of governance committee	Hull University Teaching Hospitals
Prof Andrew Pettitt	Biobank Director	University of Liverpool
Dr Melania Oates	Biobank Manager	University of Liverpool
Dr Lisa Flaherty	GCP Lab QA manager	University of Liverpool
Dr Hannah Davies	HTA lead and Sponsor representative	University of Liverpool
Ms Natasha Greatorex	Head of trial management	University of Leeds
Dr Piers Patten	Chair of NCRI CLL Study Group	Kings College Hospital
Prof Jon Strefford	Biobank user and UK CLL Forum rep	University of Southampton
Prof Chris Pepper	Biobank user and UK CLL Forum rep	University of Sussex
Dr Pauline Robbe	Biobank user	University of Oxford
Mr Frank Mercer	Patient and CLL Support rep	N/A
Dr Sarah McDonald	Observer	Blood Cancer UK
<i>funder representative</i>	Observer	

Membership will be reviewed every 3 years or earlier as requested by individual members

### Frequency of governance committee meetings

- Annually with additional ad-hoc meetings as required

### Quoracy for governance committee meetings

The committee will be considered quorate if at least 5 members are in attendance including

- The independent Chair (or deputy nominated by the Chair)
- The Biobank Director or Manager
- Sponsor representative
- At least one representative from the NCRI CLL Study Group or UK CLL Forum
- Representative from CLL Support



### Objectives

1. To receive assurance that the biobank is operating in accordance with all legal requirements and the principles of Good Clinical Practice for Laboratories
2. To be aware of the outcome of all internal audits and external inspections and any corrective and preventative actions (CAPAs)
3. To be aware of any quality incidents and resulting actions
4. To advise on all aspects of biobank operation including
  - a. Sample collection, transport, processing and storage
  - b. Visibility and accessibility of samples to researchers
  - c. Engagement with clinical trials units
  - d. Quality assurance and governance
5. To advise on all aspects of biobank strategy including
  - a. Meeting the future needs of researchers
  - b. Financial sustainability
  - c. Publicity and communication
6. To advise on specific issues raised by the Director, Manager, Sponsor, users or other stakeholders