Participants identified via NHS

All studies will require a CRICBristol Risk Assessment

Who/what are your study subjects?

All Other Chief Investigators
Study type

All research except CTIMP or non-CE marked device research

CTIMP** and/or non-CE marked device*

CTIMP** and/or non-CE marked device*

Human Tissue only

All other research (i.e. not CTIMP, human tissue only or non-CE marked device)

What approvals do CRIC need to see?

Pathway F

- NHS REC approval
- NHS permission (R&D approval)
- HRA approval letter
- Sponsorship
- Terms and Conditions of User Agreement

Pathway G

- MHRA CTA (IMP) or device letter of no objection
- NHS REC approval
- Sponsorship
- Insurance/indemnity arrangements
- ARSAC (if applicable)
- Terms and Conditions of User Agreement

Pathway H

- MHRA CTA (IMP) or device letter of no objection
- NHS REC approval
- Sponsorship
- Insurance/indemnity arrangements
- ARSAC (if applicable)
- Terms and Conditions of User Agreement

Pathway J

- NHS REC approval
- Sponsorship
- Insurance/indemnity arrangements
- Terms and Conditions of User Agreement

Pathway K

- A University or equivalent ethics approval
- Terms and Conditions of User Agreement
- Sponsorship
- Insurance/indemnity arrangements

Key:
ARSAC: Administration of Radioactive Substances Advisory Committee
REC: Research Ethics Committee
MHRA: Medicines and Healthcare products Regulatory Agency
*Safety/efficacy of a medicine, foodstuff or placebo
CTA: Clinical Trial Authorisation

*For advice on non-CE marked device trials see http://www.mhra.gov.uk/Howweregulate/Devices/Clinical_trials/index.htm
**CTIMP: Clinical Trial of an Investigational Medicinal Product, see http://www.mhra.gov.uk/home/groups/lunit1/documents/websitesources/con009394.pdf