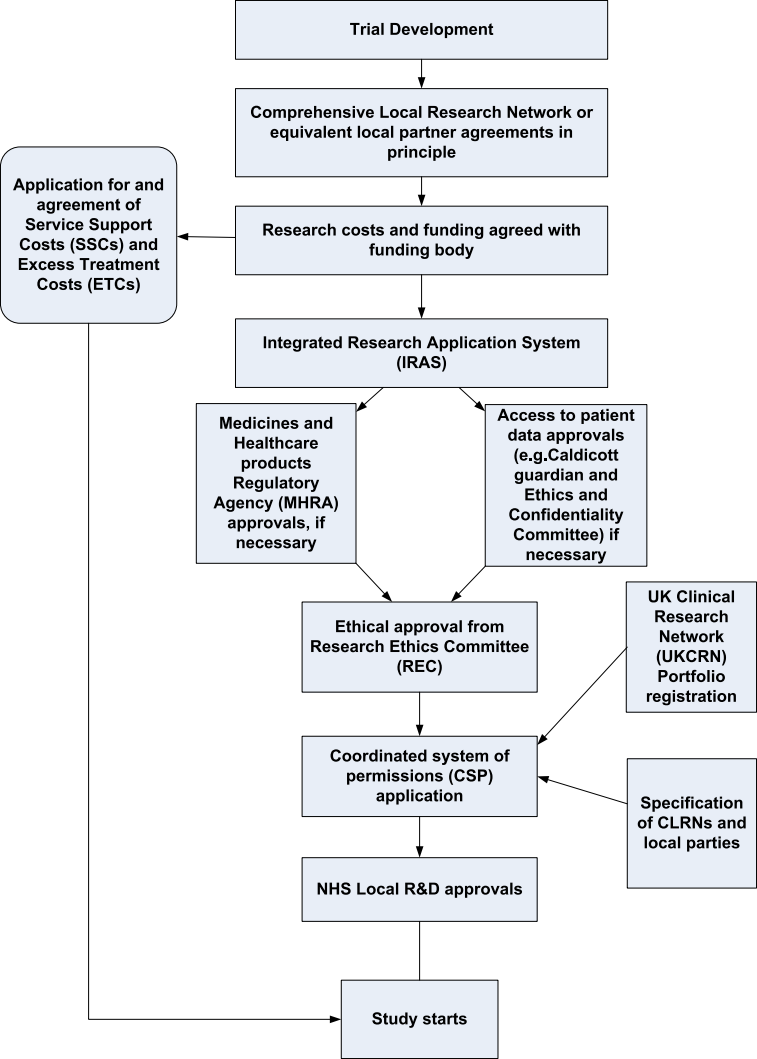
Trial Development
Comprehensive Local Research Network or equivalent local partner agreements in principle
Application for and agreement of Service Support Costs (SSCs) and Excess Treatment Costs (ETCs)
Research costs and funding agreed with funding body
Integrated Research Application System (IRAS)
Medicines and Healthcare products Regulatory Agency (MHRA) approvals, if necessary
Access to patient data approvals (e.g.Caldicott guardian and Ethics and Confidentiality Committee) if necessary
Ethical approval from Research Ethics Committee (REC)
UK Clinical Research Network (UKCRN) Portfolio registration
Coordinated system of permissions (CSP) application
Specification of CLRNs and local parties
NHS Local R&D approvals
Study starts