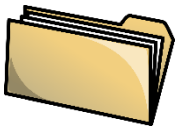
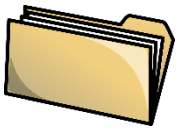
Policies, Procedures & Guidelines

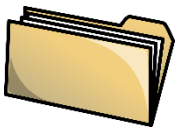
A – Z



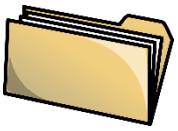
Amendment to Study Documentation



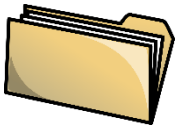
Breach of Participant Confidentiality



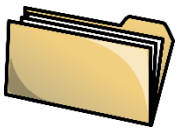
Clinical Trial Sample Analysis in University Laboratories



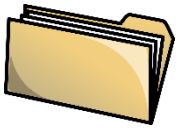
Complaints from Research Participants



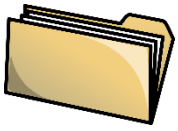
Contracting for CTIMPs



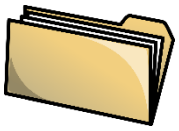
Convening of Trial Steering Committees and Data Monitoring Committees for Clinical Trials



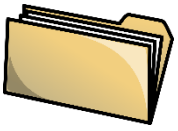
Creation, Control, Amendment and Storage of SOPs



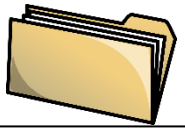
Data Management, Collection, Validation and Storage



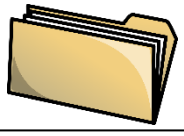
Delegation of Responsibilities



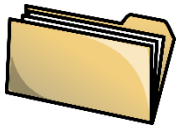
Development and Review of Research Plan/Study Protocol



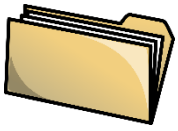
Education, Training and Experience



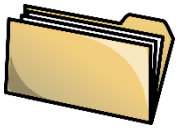
End of Study Declaration, Early Termination and Final Report



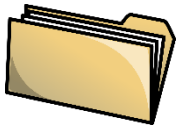
The Ethical Approval of Research



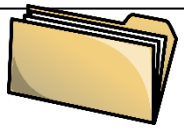
Indemnity and Sponsorship of Research Studies



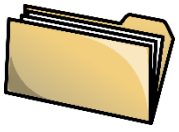
Informed Consent for Research



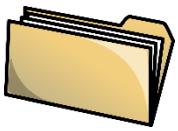
Maintaining Laboratory Books



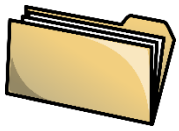
Matters of Non-Compliance with Study Protocol



Monitoring of Research Studies

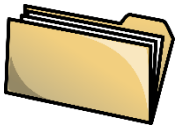


Preparation, Completion, Signing and Correcting Case Report Forms



Production of Progress Reports

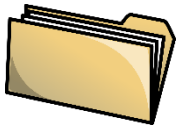
Registration of Clinical Trials



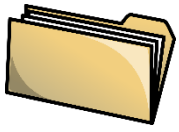
Reporting and Managing Research Related Adverse Events

Research Governance Audit

Risk Assessment of Research Studies



Setting Up, Maintaining and Archiving Research Files



Sponsor Green Light