AF 01-001/01	List of IRB SOPs
AF 02-001/01	Standard Operating Procedures Template
AF 03-001/01	Document History
AF 04-001/01	Log of SOP Recipients
AF 05-001/01	Request for Revision of an SOP
AF/01-002/01	Cover page of a Guideline (2 pages)
AF/02-002/01	Lists of Signatures
AF/03-002/01	Log of Guideline Distribution
AF/01-004/01	Confidentiality / Conflict of Interest Agreement Form
AF/02-004/01	Confidentiality Agreement form for Guest / Observer Attendees to IRB Meetings
AF/03-004/01	Confidentiality Agreement Form for Non-members Requesting Copies of IRB's Documents
AF/01-005/01	Training Record Form
AF/01-007/01	Checklist to determine if a proposed protocol qualifies for exemption from ethics review
AF/02-007/01	Exemption Letter Templates
AF/01-008/01	Contents of a Submitted Package (Checklist)
AF/02-008/01	Document Receipt Form
AF/03-008/01	Application Form for Initial Review
AF/04-008/01	Protocol Summary Sheet/Memorandum
AF/05-008/01	Application Form for Case Study/ Case Series
AF/01-010/01	Presentation template for the Primary Reviewers during a protocol review
AF/01-011/01	Examples of Non-significant Risk Device Studies
AF/02-011/01	Examples of Significant Risk Device Studies
AF/01-012/01	Study Assessment Form (5 pages)
AF/02-012/01	Study Assessment Form for case studies and case series
AF/03-012/01	Informed Consent Assessment Form
AF/03-012/01	Assessment Report Form
AF/04-012/01	IRB's Decision
AF/05-012/01	Action letter templates
AF/06-012/01	Approval Letter Template
AF/07-012/01	Guidance for reviewing a study protocol
AF/08-012/01	Informed Consent Process
AF/09-012/01	Informed consent Form (ADAPTED FROM WHO-GUIDELINE)
AF/10-012/01	Guides to Placebo Justification
AF/11-012/01	Criteria for Research Protocol Approval
AF/01-013/01	Re-submitted Protocol Review Form
AF/02-013/01	Application Form for Resubmitted Protocol Review
AF/01-014/01	Protocol Amendment Application Form
AF/01-015/01	Continuing Review Application Form (2 pages)
AF/01-016/01	Study Report Form
AF/02-016/01	Format for Review of Research Report
AF/03-016/01	Report Review Letter Template
AF/01-017/01	Deviation/Non Compliance/Violation Record
AF/01-018/01	Request Record Form
AF/01-019/01	Termination Memorandum
AF/01-020/01	Serious Adverse Event Report
AF/02-020/01	Unexpected Adverse Drug Reaction Report
AF/03-020/01	Safety Report Review Form
AF/01-021/01	Site Monitoring Visit Report
AF/01-022/01	Agenda format
AF/02-022/01	Form of IRB Meeting Minutes
AF/03-022/01	IRB Meeting Minutes Template

AF/01-024/01	Communication Record Form
AF/01-026/01	Document Request Form
AF/02-026/01	Logs of Requested IRB Documents
AF/01-027/01	Log of Requests for Copies of IRB documents
AF/02-027/01	Logs of Copies of Original Documents
AF/01-028/01	Audit and Inspection Checklist