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ABBREVIATIONS

Below is a list of common abbreviations that will be used throughout this Investigator's Manual.

ABPI	Association of the British Pharmaceutical Industry
BAC	Bioethics Advisory Committee
CAPA	Corrective Action and Preventive Action
CFR	US Code of Federal Regulations
CIOMS	Council for International Organisations of Medical Sciences
CIRB	SingHealth Centralised Institutional Review Board
CITI	Collaborative Institutional Training Initiative
CRC	Clinical Research Coordinator
CRF	Case Report Form
CRM	Clinical Research Materials
CRU	Clinical Research Unit
СТА	Clinical Trial Authorisation
CTC	Clinical Trial Certificate
CTN	Clinical Trial Notification
DCF	Data Collection Form
DHHS	US Department of Health and Human Services
DNA	Deoxyribonucleic acid
DR	Department Representative
DSMB	Data Safety Monitoring Board
DSRB	Domain Specific Review Board
FCOI	Financial Conflict Of Interest
FDA	US Food and Drug Administration
GCP	Good Clinical Practice
HRPP	Human Research Protection Programme
HBR	Human Biomedical Research
HBRA	Human Biomedical Research Act
HIV	Human Immunodeficiency Virus
HSA	Health Sciences Authority

ICFInformed Consent FormICHInternational Council for HarmonisationICOIInstitutional Conflict Of InterestIDEInvestigational Device ExemptionINDInvestigational New Drug ApplicationIQInstitution OfficerIRInstitution RepresentativeIRBInstitutional Review BoardMDMedical DeviceMOHMinistry of Health SingaporeMPMedicinal ProductNHGNational Medical Ethics CommitteeNUSNational Medical Ethics CommitteeNUSNational Ouriersity of SingaporeOHRPPNHG Office of Human Research Protection ProgrammePCRProper Conduct of ResearchPDPAPersonal Data Protection ActPDPAPersonal Data Protection CommissionPIPrincipal InvestigatorQAQuality ImprovementRECNHG Research Ethics CommitteeRIAResearch InstitutionROAMNHG Research Online Administration & Management SystemSAESerious Adverse EventSBESocial, behavioural and educational (modules from CITI)SDCSingapore Dental CouncilSOPStandard Operating ProceduresSTDSexually Transmitted DiseasesTPTherapeutic Product		
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IDEInvestigational Device ExemptionIDEInvestigational New Drug ApplicationINDInvestigational New Drug ApplicationIOInstitution OfficerIRInstitution RepresentativeIRBInstitutional Review BoardMDMedical DeviceMOHMinistry of Health SingaporeMPMedicial ProductNHGNational Healthcare GroupNMECNational Medical Ethics CommitteeNUSNational University of SingaporeOHRPPNHG Office of Human Research Protection ProgrammePCRProper Conduct of ResearchPDPAPersonal Data Protection ActPDPCPersonal Data Protection CommissionPIPrincipal InvestigatorQAQuality ImprovementRECNHG Research Ethics CommitteeRIResearch InstitutionROAMNHG Research Online Administration & Management SystemSAESerious Adverse EventSBESocial, behavioural and educational (modules from CITI)SDCSingapore Medical CouncilSOPStandard Operating ProceduresSTDSexually Transmitted Diseases	ICH	International Council for Harmonisation
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RIResearch InstitutionROAMNHG Research Online Administration & Management SystemSAESerious Adverse EventSBESocial, behavioural and educational (modules from CITI)SDCSingapore Dental CouncilSMCSingapore Medical CouncilSOPStandard Operating ProceduresSTDSexually Transmitted Diseases	QI	Quality Improvement
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SBESocial, behavioural and educational (modules from CITI)SDCSingapore Dental CouncilSMCSingapore Medical CouncilSOPStandard Operating ProceduresSTDSexually Transmitted Diseases	ROAM	NHG Research Online Administration & Management System
SDC Singapore Dental Council SMC Singapore Medical Council SOP Standard Operating Procedures STD Sexually Transmitted Diseases	SAE	Serious Adverse Event
SMC Singapore Medical Council SOP Standard Operating Procedures STD Sexually Transmitted Diseases	SBE	Social, behavioural and educational (modules from CITI)
SOP Standard Operating Procedures STD Sexually Transmitted Diseases	SDC	Singapore Dental Council
STD Sexually Transmitted Diseases	SMC	Singapore Medical Council
	SOP	Standard Operating Procedures
TP Therapeutic Product	STD	Sexually Transmitted Diseases
	TP	Therapeutic Product

UPIRTSO	Unanticipated Problems Involving Risks To Subjects or Others
USADR	Unexpected Serious Adverse Drug Reactions