CP-CTNet Acronym List

Acronym	Description
ACC	Awarded Contracting Center
ACRP	Association of Clinical Research Professionals
ADL	Activities of Daily Living
AE	Adverse Event
AIF	Action Item (Site Response) Form
AO	Affiliated Organization
AQuIP	Accrual Quality Improvement Program
BA/BE	Bioavailability/Bioequivalence
CA No.	NCI unique five or six digit number that identifies the specific protocol application
CAP	College of American Pathologists
caDSR	Cancer Data Standards Repository
CCRA	Certified Clinical Research Associate
CCSA	CCS Associates, Inc. (DCP Regulatory Contractor)
CDA	Confidential Disclosure Agreements
CDE	Common Data Element
CDISC	Clinical Data Interchange Standards Consortium
CDMS	Clinical Data Management System
CFID	Central Files Inventory Database
CFR	Code of Federal Regulations
CIRB	Central Institutional Review Board
CITI	The Collaborative Institutional Training Program
CLIA	Clinical Laboratory Improvement Amendments
CLP	Clinical Laboratory Permit
COA	Certificate of Analysis
CoC	Certificate of Confidentiality
COV	Closeout Visit
CPC	Cancer Prevention and Control
CP-CTNet	Cancer Prevention – Clinical Trials Network
CR	Change Request
CRA	Clinical Research Associate
CRADA	Cooperative Research and Development Agreement
CRF	Case Report Form
CRO	Contract Research Organization
CSA	Clinical Supply Agreement
CSR	Center for Scientific Research (at NIH)
CTA	Clinical Trial Agreement
CTAC	Clinical Trials and Translational Research Advisory Committee
CTCAE	Common Terminology Criteria for Adverse Events
CTRO	Clinical Trials Reporting Office
CTRP	Clinical Trials Reporting Program
CV	Coefficient of Variation
CV(s)	Curriculum Vita(e)
DBA	Database Administrator

CP-CTNet Acronym List

Acronym	Description
dbGaP	Database of Genotypes and Phenotypes
DCP	Division of Cancer Prevention at the National Cancer Institute
DCPCR	DCP Collaboration Repository
DCP PIO	DCP's Protocol Information Office
DE	Data Entry
DEA	Division of Extramural Activities
DHHS	Department of Health and Human Services
DMACC	CP-CTNet Data Management, Auditing and Coordinating Center
DMC	Data Monitoring Committee (also known as Data and Safety Monitoring Board)
DMP	Data Management Plan
DSA	Drug Shipment Authorization
DSMB/DSMC	Data Safety Monitoring Board/Committee
DSMP	Data and Safety Monitoring Plan
DTL	Delegation of Tasks Log
EDC	Electronic Data Capture
EHR	Electronic Health Record
E-Learning	Electronic Learning
EMR	Electronic Medical Record
FDA	Food and Drug Administration
FDF	Financial Disclosure Form
FOA	Funding Opportunity Announcement
FTP	File Transfer Protocol
FWA	Federal-wide Assurance (Number)
GCP	Good Clinical Practice
GDS	Genomic Data Sharing
GMS	Grants Management Specialist
GPA	Genomic Program Administrator
HIPAA	Health Insurance Portability and Accounting Act of 1966
HSP	Human Subjects Protection
IAM	Identity and Access Management
IATA	International Air Transport Association
IB	Investigator's Brochure
ICD	Informed Consent Document
ICF	Informed Consent Form
ICH	International Conference on Harmonization
IDE	Investigational Device Exception
IEC	Independent Ethics Committee
IMS	Information Management Services, Inc. (DCP Contractor)
IND	Investigational New Drug
IRB	Institutional Review Board
IT	Information Technology
JCAHO	Joint Commission on Accreditation of Healthcare Organizations
LAO	Lead Academic Organization

CP-CTNet Acronym List

Acronym	Description
LNV	Lab Normal Value
LOI	Letter of Intent
MDS	Minimum Dataset
MedRA	Medical Dictionary for Regulatory Activities
MIMP	Multi-Institutional Monitoring Plan
ML	Medical License
MM	Medical Monitor
MOP	Manual of Operations
MRIGlobal	DCP Agent Repository Contractor
MTA	Material Transfer Agreement
N/A	Not Applicable
NC	Nurse Consultant
NCAB	National Cancer Advisory Board
NCI	National Cancer Institute
NCICBIIT	NCI Center for Biomedical Informatics and Information Technology
NCTN	National Clinical Trials Network
NDA	New Drug Application
NIH	National Institutes of Health
NOS	Not Otherwise Specified
NTF	Note-to-file
OBA	Office of Biotechnology Activities
OD	Office of the Director at the NCI
OER	Office of Extramural Research, NIH
OGA	Office of Grants Administration
OHRP	Office of Human Research Protections
OMB	Office of Management and Budget
PA	Program Announcement
PD	Project Director
PD	Protocol Deviation
PDF	Portable Document Format
PHI	Personal Health Information or Protected Health Information
PHS	Public Health Service
PI	Principal Investigator
PID	Participant Identification (Number)
PII	Personally Identifiable Information
PK/PD	Pharmacokinetics/Pharmacodynamic
PSW	Protocol Submission Worksheet
QA	Quality Assurance
QC	Quality Control
RRA	Recruitment, Retention and Adherence
RDC	Remote Data Capture (System)
RCR	Registration Credential Repository
RDE	Remote Data Entry

CP-CTNet Acronym List

Acronym	Description
SAE	Serious Adverse Event
SAS	Analytics software used to manage, analyze and report on data
SC	Site Coordinator
SDR	Source Data Review
SDV	Source Data Verification
SEB	Surrogate Endpoint Marker
SFTP	Secure File Transfer Protocol
SI	Sub-investigator
SIV	Study Initiation Visit
SL	Scientific Lead
SME	Subject Matter Expert
SOC	System Organ Class
SoCRA	Society of Clinical Research Associates
SOP	Standard Operating Procedure
SOW	Statement of Work
SVAR	System Variable Attribute Report
TAC	Treatment Assignment Code
TAD	Treatment Assignment Description
TOC	Table of Contents
URL	Uniform Resource Locator (internet address of resource)
US, USA	United States, United States of America
VA	Veterans Affairs
WGS/WES	Whole Genome Sequencing/Whole Exome Sequencing