Acronym List

| **Acronym** | **Description** |
| --- | --- |
| ACR | Account Change Request |
| ACRP | Association of Clinical Research Professionals |
| AE | Adverse Event |
| AIF | Action Item (Site Response) Form |
| AIS | Auditing and Informatics Support |
| AMV | Annual Monitoring Visit |
| AQuIP | Accrual Quality Improvement Program |
| AQuIP OARS | Accrual Quality Improvement Program Online Accrual Reporting System |
| CAP | College of American Pathologists |
| caDSR | Cancer Data Standards Repository |
| CCDS | Consortia Clinical Data Subsystem |
| CCRA | Certified Clinical Research Associate |
| CCSA | CCS Associates, Inc. (DCP Regulatory Contractor) |
| CDEs | Common Data Elements |
| CDT | Clinical Data Transfer (Website) |
| 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 |
| CLO | Consortium Lead Organization |
| COR | Contracting Officer’s Representative |
| Co-COR | Co-Contracting Officer’s Representative |
| COV | Closeout Visit |
| CPC | Cancer Prevention and Control |
| CR | Change Request |
| CRA | Clinical Research Associate |
| CRF | Case Report Form |
| CTA | Clinical Trial Agreement |
| CV(s) | Curriculum Vita(e) |
| DARF | Drug Accountability Record Form |
| DBA | Database Administrator |
| DCP | Division of Cancer Prevention at the National Cancer Institute |
| DCPCR | DCP Collaboration Repository |
| DCP PIO | DCP’s Protocol Information Office |
| DE | Data Entry |
| DESK | DCP Enterprise System Knowledgebase |
| DHHS | Department of Health and Human Services |
| 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 |
| FTP | File Transfer Protocol |
| FWA | Federal-wide Assurance (Number) |
| GCP | Good Clinical Practice |
| HIPAA | Health Insurance Portability and Accounting Act of 1966 |
| HSP | Human Subjects Protection |
| IB | Investigator’s Brochure |
| ICD | Informed Consent Document |
| ICF | Informed Consent Form |
| ICH | International Conference on Harmonization |
| 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 |
| LNV | Lab Normal Value |
| MDS | Minimum Dataset |
| MIMP | Multi-Institutional Monitoring Plan |
| ML | Medical License |
| MM | Medical Monitor |
| MRIGlobal | DCP Contractor |
| NC | Nurse Consultant |
| NCI | National Cancer Institute |
| NCICBIIT | NCI Center for Biomedical Informatics and Information Technology |
| NDA | New Drug Application |
| NIH | National Institutes of Health |
| NTF | Note-to-file |
| OBA | Office of Biotechnology Activities |
| OHRP | Office of Human Research Protections |
| OMB | Office of Management and Budget |
| PD | Protocol Deviation |
| PDF | Portable Document Format |
| PHI | Personal Health Information or Protected Health Information |
| PI | Principal Investigator |
| PID | Participant Identification (Number) |
| PIO | Protocol and Information Office |
| PK | Pharmacokinetics |
| PO | Participating Organization |
| PSP | Project Specific Procedure |
| QA | Quality Assurance |
| QC | Quality Control |
| RDC | Remote Data Capture (System) |
| RDE | Remote Data Entry |
| SAE | Serious Adverse Event |
| SAS | Statistical Analysis System |
| SDR | Source Data Review |
| SDV | Source Data Verification |
| SFTP | Secure File Transfer Protocol |
| SI | Sub-investigator |
| SIV | Study Initiation Visit |
| SM | Scientific Monitor |
| SME | Subject Matter Expert |
| SoCRA | Society of Clinical Research Associates |
| SOP | Standard Operating Procedure |
| SOW | Statement of Work |
| TRI | Technical Resources International, Inc. (DCP Monitoring Contractor) |
| US | United States |
| VA | Veterans Affairs |