Acronym	Name
ADR	Adverse Drug Reaction
AE	Adverse Event
APEX	Epic based electronic medical records at UCSF
CA	Coverage Analysis
CAP	Corrective Action Plan
CCRP	Certified Clinical Research Professional
CFR	Code of Federal Regulations
CHR	Committee on Human Research (former name for UCSF IRB)
CITI	Collaborative IRB Training Initiative
COI	Conflict of Interest
CRA	Clinical Research Associate
CRC	Clinical Research Coordinator
CRF	Case Report Form
CRO	Contract Research Organization
CRRM	Clinical Research Risk Manager
CRS	Clinical Research Services
CTA	Clinical Trials Agreement
CTCAE	Common Terminology Criteria for Adverse Events
CTSI	Clinical and Translational Science Institute
DA	Division Administrator
DM	Department Manager
DSMB	Data Safety Monitoring Board
DSMC	Data and Safety Monitoring Committee
DSMP	Data and Safety Monitoring Plan
DUA	Data Use Agreement
ECRF	Electronic Case Report Form
EDC	Electronic Data Capture
EMR	Electronic Medical Record
FDA	Food and Drug Administration
FWA	Federal wide Assurance
GCP	Good Clinical Practice
HIPAA	Health Insurance Portability and Accountability Act
HRPP	Human Research Protection Program
IB	Investigator's Brochure
ICF	Informed Consent Form
ICD	Industry Contracts Division
ICH	International Conference on Harmonization
IDE	Investigational Device Exemption
IND	Investigational New Drug
IRB	Institutional Review Board
iRIS	Online IRB application system at UCSF
ITA	Office of Innovation, Technology and Alliances
ITR	Investigational Trial Resource (Cancer Center)

MCA	Medicare Coverage Analysis
MOP	Manual of Procedures
MRN	Medical Record Number
MTA	Material Transfer Agreement
NIH	National Institutes of Health
NTF	Note to File
OCR	Office of Clinical Research
OHRP	Office for Human Research Protection
OSR	Office of Sponsored Research
PHI	Protected Health Information
PI	Principal Investigator
PRN	As Needed
QA	Quality Assurance
QC	Quality Control
QIU	Quality Improvement Unit
RFA	Research Financial Analyst
RMIS	Risk Management and Insurance Services
RMS	Research Management Services
RN	Registered Nurse
RSC	Research Services Coordinator
SAE	Serious Adverse Event
SEV	Site Evaluation Visit
SFVAMC	Veterans Administration Medical Center
SIV	Site Initiation Visit
SOC	Standard of Care
SOE	Schedule of Events
SOP	Standard Operating Procedure
Sub-I	Sub-Investigator