AAHRPP	Association for the Accreditation of Human Research Protection Programs
AE	Adverse Event
APB	Administrative Panel on Biosafety
CCTO	Cancer Clinical Trials Office
CFR	Code of Federal Regulations
CITI	Collaborative IRB Training Initiative
COI	Conflict of Interest
CQI	Continuous Quality Improvement
CTSU	Clinical and Translational Science Unit (formerly GCRC)
DHHS	Department of Health and Human Services (or HHS)
DSMB/C	Data Safety Monitoring Board/Committee
EH & S	Environmental Health and Safety
FDA	Food and Drug Administration
FWA	Federalwide Assurance
HIPAA	Health Insurance Portability and Accountability Act (1996)
HRPP	Human Research Protection Program
HSR	Human Subjects Research
IDE	Investigational Device Exemption
IND	Investigational New Drug
IRB	Institutional Review Board
LAR	Legally Authorized Representative
LPCH	Lucile Packard Children's Hospital at Stanford
MTA	Material Transfer Agreement
NIH	National Institutes of Health
OHRP	Office for Human Research Protections
PAIRE	Palo Alto Institute for Research and Education
PD/PI	Protocol Director
PI	Principal Investigator
PHI	Protected Health Information
PHS	Public Health System
PRIM&R	Public Responsibility in Medicine and Research
RCO	Research Compliance Office
SAE	Serious Adverse Event
SHC	Stanford Hospital and Clinics
Spectrum	Stanford Center for Clinical and Translational Education and Research
SIR	Sponsor Investigator Research
UP	Unanticipated Problem (Involving Risks to Participants or Others)
VA	Veterans Administration
VAPAHCS	Veterans Affairs Palo Alto Health Care System