University of California, San Diego
Human Research Protections Program
Institutional Review Board
Standard Operating Policies and Procedures

Section 8.1
Abbreviations

ADE	Adverse Drug Event/Experience
AE	Adverse Event
CFR	Code of Federal Regulations
COI	Conflict of Interest
CTA	Clinical Trial Agreement
CRA	Clinical Research Associate
CRF	Case Report Form
CRO	Contract Research Organizations
ESCRO	Embryonic Stem Cell Research Oversight
FDA	Food and Drug Administration
GCP	Good Clinical Practice
HDE	Humanitarian Device Exemption
HERC	Human Exposure Review Committee
HRPP	UCSD Human Research Protections Program
HUD	Humanitarian Use Device
IBC	Institutional Biosafety Committee
ICF	Informed Consent Form
ICH	International Conference on Harmonization
IDE	Investigational Device Exemption
IND	Investigational New Drug
IRB	Institutional Review Board
IRC	Independent Review Committee
MTA	Materials Transfer Agreement
NSR	Non-significant Risk
OCGA	Office of Contract and Grant Administration
OHRP	Office for Human Research Protections
PI	Principal Investigator
PRMC	Protocol Review and Monitoring Committee
SAE	Serious Adverse Event
SR	Significant Risk
SOM	School of Medicine
SOPP	Standard Operating Policies and Procedures
UCSD	University of California, San Diego
UPR	Unanticipated Problem Involving Risk to Subjects or Others