Acronyms & Abbreviations
(*E-IRB Specific)

AAHRPP	Association for Accreditation of Human Research Protection Programs
AA	Administrative Associate
AC*	Authorized to Provide Consent
ADAccount*	UK Link Blue Identification
AE	Adverse Event / Adverse Experience
AR	Administrative Regulation (UK)
CCTS	Center for Clinical and Translational Science
CFR	Code of Federal Regulations
CITI	Collaborative Institutional Training Initiative (online learning program)
COI	Conflict of Interest
CR	Continuation Review
CRF	Case Report Form
CRO	Clinical Research Organization (UK MCC)
CRSO	Clinical Research Services Office
CV	Curriculum Vitae
DA*	Department Authorization
DEV/EXC*	Deviation/Exception
DHHS (or HHS)	Department of Health and Human Services
DoD	Department of Defense
DoE	Department of Energy
DoED	Department of Education
DoJ	Department of Justice
DP*	Delegate Personnel
DSMB	Data and Safety Monitoring Board
DSMP	Data and Safety Monitoring Plan
EHS	Environmental Health & Safety
EMR	Electronic Medical Record
EPA	Environmental Protection Agency
FDA	Food and Drug Administration
FL	Full Review
FR	Final Review
FWA	Federal Wide Assurance
GCP	Good Clinical Practice
GIS	General Information Sheet
HHS (or DHHS)	Department of Health and Human Services
HIPAA	Health Insurance Portability and Accountability Act (Privacy Rule)
HRPP	Human Research Protection Program
HSP	Human Subject Protection
IBC	Institutional Biosafety Committee (UK)
ICH	International Conference on Harmonization GCP Guidelines
IDE	Investigational Device Exemption (application to FDA)

IDS	Investigational Drug Service (UK)
IND	Investigational New Drug (application to FDA)
IR	Initial Review
IRB	Institutional Review Board
LAR	Legally Authorized Representative
MCC	Markey Cancer Center (UK)
NA (or N/A)	Not Applicable
NCI	National Cancer Institute (NIH)
NIH	National Institute of Health
OHRP	Office for Human Research Protections
ORI	Office of Research Integrity (UK)
OSPA	Office of Sponsored Projects Administration (UK)
MCC	Markey Cancer Center
MR	Modification Request
PA	Professional Associate
PDO	Proposal Development Office
PersonID*	UK Employee Identification Number
PHI	Protected Health Information
PI	Principal Investigator
PST	Protocol Specific Training
PV	Protocol Violation
QIP	Quality Improvement Program
RDRC	Radioactive Drug Research Committee
RSC	Radiation Safety Committee (UK)
SD	Source Document
SFI	Significant Financial Interest
SOP	Standard Operating Procedure
SP	Study Personnel
TJC	The Joint Commission (previously JCAHO - Joint Commission on Accreditation of Healthcare Organizations)
UK	University of Kentucky
UP	Unanticipated Problem Involving Risks to Subjects or Others
VA	Veterans Affairs
VC*	View Comments
VPR	Vice President for Research
X or XX	Exempt Review
XP	Expedited Review