Acronym List

ADR	Adverse Drug Reaction
AE	Adverse Event
BRI	Bloorview Research Institute
CAREB	Canadian Association of Research Ethics Boards
CEO	Chief Executive Officer
CFR	Code of Federal Regulations (US)
COI	Conflict of Interest
DSMB	Drug Safety Monitoring Board
FDA	US Food and Drug Administration
GCP	Good Clinical Practice
HHS	US Department of Health and Human Services
HIROC	Healthcare Insurance Reciprocal Corporation of Canada
ICH	International Conference on Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use
LPI	Local Principal Investigator
OHRP	Office for Human Research Protections (HHS)
PHI	Protected Health Information
RAC	Research Advisory Committee
REB	Research Ethics Board
SAE	Serious Adverse Event
SOP	Standard Operating Procedure
SUSAR	Suspected Unexpected Serious Adverse Reaction
TAHSN	Toronto Academic Health Sciences Network
TCPS2	Tri-Council Policy Statement: Ethical Conduct for Research Involving Humans
US	United States