	STANDARD OPERATING PROCEDURES			
LSU Health New Orleans	LIST OF ABBREVIATIONS			
Health Sciences Center & Healthcare Network	NUMBER	APPROVED BY	EFFECTIVE DATE	PAGE
	Abbreviations	Executive Director, ORS	07.21.2022	Page 1 of 2

ADR	Adverse Drug Reaction (same as Adverse Drug Experience)		
AE	Adverse Event (same as Adverse Experience, Adverse Reaction)		
CDA	Confidential Disclosure Agreement (same as Non-Disclosure Agreement)		
CFR	Code of Federal Regulations		
СОІ	Conflicts of Interest		
CRC	Clinical Research Coordinator		
CRNC	Clinical Research Nurse Coordinator		
CRF	Case Report Form (same as Electronic Case Report Form)		
CRO	Contract Research Organization		
СТА	Clinical Trial Agreement		
СТО	Clinical Trials Office		
DHHS	Department of Health and Human Services		
DSMB	Data & Safety Monitoring Board (same as Data & Safety Monitoring Committee, Data Monitoring Committee)		
DUA/DTUA	Data (Transfer and) Use Agreement		
eCRF	Electronic Case Report Form (same as Case Report Form)		
EHR	Electronic Health Record		
EMR	Electronic Medical Record		
FDA	Food and Drug Administration		
GCP	Good Clinical Practice		
HIPAA	Health Insurance Portability and Accountability Act of 1996		
HN or LSUHN	LSU Healthcare Network		
HRPP	Human Research Protection Program		
HSC or LSUHSC	LSU Health Sciences Center		
IB	Investigator's Brochure		
IBC	Institutional Biosafety Committee		
ICF	Informed Consent Form		
ICH	International Conference on Harmonization		
ICMJE	International Committee of Medical Journal Editors		
IDE	Investigational Device Exemption		
IND	Investigational New Drug		
IP	Investigational Product		
IRB	Institutional Review Board		
LAR	Legally Authorized Representative		

MTA	Material Transfer Agreement
NDA	Non-Disclosure Agreement (same as Confidential Disclosure Agreement)
NIH	National Institutes of Health
NPP	Notice of Privacy Practices
NSR/SR	Non-Significant Risk/Significant Risk Determination (for devices)
ОСМ	Office of Contract Management
OHRP	Office for Human Research Protections
OIP	Office of Innovation and Partnership
РНІ	Protected Health Information
PI	Principal Investigator
QA	Quality Assurance
QC	Quality Control
SAE	Serious Adverse Event
SIV	Site Initiation Visit
SOP	Standard Operating Procedure
SPA	Sponsored Projects Administration
Sub-I	Sub-Investigator (same as Co-Investigator)

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