CDISC MRCT Controlled Terminology, 2025-09-26

 $Source: NCI\ EVS\ Terminology\ Resources\ website: http://www.cancer.gov/cancertopics/cancerlibrary/terminologyresources/cdisc$

NCI Code	CDISC Submission Value	Codelist Name	CDISC Definition	Codelist Extensible
C203912	MRCT Center Clinical Research Glossary	MRCT Center Clinical Research Glossary	The terminology relevant to the plain language glossary of the Multi-Regional Clinical Trial (MRCT) Center of Brigham and Women's Hospital and Harvard.	

NCI Code: C203912, Codelist extensible:

	C203912 MRCT Center Cl			
	NCI Code CDISC Subm	•	CDISC Definition	NCI Preferred Term
C203914	additive effect		The combined effect when two or more things are used together. (https://mrctcenter.org/glossaryterm/additive-effect/)	Additive Effect
C25729 C41331	adherence adverse event		Following the study directions and requirements. (https://mrctcenter.org/glossaryterm/adherence/) Any health problem that happens during a study. (https://mrctcenter.org/glossaryterm/adverse-	Adherence Adverse Event
C41332	adverse reaction		event/) A health problem that happens during a study and is reported as possibly caused by the study	Adverse Reaction
			treatment. (https://mrctcenter.org/glossaryterm/adverse-reaction/)	
C142392	anonymize		Remove, change, or hide personal details to protect participant privacy. (https://mrctcenter.org/glossaryterm/anonymize/)	Anonymization
C16295	antibody		A protein made by the body to fight an illness or infection. (https://mrctcenter.org/glossaryterm/antibody/)	Antibody
C268	antigen		A substance that causes the body's immune system to react. (https://mrctcenter.org/glossaryterm/antigen/)	Antigen
C70800	approval (IRB/ethic	.s)	Official decision by an ethics committee to allow a research study involving human participants to begin. (https://mrctcenter.org/glossaryterm/approval-irb-ethics/)	Institutional Review Board Approva
C15538	arm		A group of participants in a research study who all receive the same study treatment. (https://mrctcenter.org/glossaryterm/arm/)	Protocol Treatment Arm
C203934	assent form		A document used to explain the details of a research study to children or people who are unable to	Assent Form
C161418	assent		give legal consent. (https://mrctcenter.org/glossaryterm/assent-form/) Willingness to take part in a research study by someone who is not able to give legal consent.	Informed Assent
C25217	assessment		(https://mrctcenter.org/glossaryterm/assent/) Information that is collected and analyzed from a study participant.	Assessment
C142400	baseline assessme	ent	(https://mrctcenter.org/glossaryterm/assessment/) Information that is collected and analyzed from a study participant at the start of a study.	Baseline Assessment
C209461	basket trial		(https://mrctcenter.org/glossaryterm/baseline-assessment/) A research study that tests one study treatment for different diseases and conditions.	Basket Trial
	benefits of a resea	and advade.	(https://mrctcenter.org/glossaryterm/basket-trial/)	
C203915		Cristudy	The ways a research study might help the participant and others. (https://mrctcenter.org/glossaryterm/benefits-of-a-research-study/)	Benefits of a Research Study
C28232	bias (research)		Flaws in the way a study is designed, done, or analyzed that lead to one conclusion being favored over another. (https://mrctcenter.org/glossaryterm/research-bias/)	Bias
C179748	biobanking (resear	ch)	Storing biological samples from participants for future research. (https://mrctcenter.org/glossaryterm/biobanking-research/)	Biospecimen Storage
C16342	biomarker		Something in the body that is measured as an indicator of personal health or disease. (https://mrctcenter.org/glossaryterm/biomarker/)	Biomarker
C37932	birth control		A way to prevent pregnancy. (https://mrctcenter.org/glossaryterm/birth-control/)	Contraception
C49068	blinding		A way to keep participants, and often study staff, from knowing which study treatment a participant is assigned. (https://mrctcenter.org/glossaryterm/blinding/)	Blinded
C28221 C15197	blood draw case-control study		Taking a sample of blood by using a needle. (https://mrctcenter.org/glossaryterm/blood-draw/) Research that uses existing data to compare a group of participants with a disease or trait to a	Phlebotomy Case-Control Study
C142422	clinical benefit		group without the same disease or trait. (https://mrctcenter.org/glossaryterm/case-control-study/) A health change that researchers measure to show that the study treatment helps the study	Clinical Benefit
C51811	Clinical Research (Coordinator	participants. (https://mrctcenter.org/glossaryterm/clinical-benefit/) A research staff member who helps manage studies. (https://mrctcenter.org/glossaryterm/clinical-	Clinical Coordinator
	(CRC)	2001 UII IULUI	research-coordinator-crc/)	
C142435	clinical research		A type of science that uses people's data to study health, illness, behaviors, or conditions in careful and defined ways. (https://mrctcenter.org/glossaryterm/clinical-research/)	
C71104	clinical trial		A research study that tests drugs, devices and treatments to see if they are safe and work in people. (https://mrctcenter.org/glossaryterm/clinical-trial/)	Clinical Trial
C85499 C61512	clinician cohort		A health care provider. (https://mrctcenter.org/glossaryterm/clinician/) A group of study participants that are similar in some way.	Clinician Cohort
C203916	Comparative Effect	tivonoss	(https://mrctcenter.org/glossaryterm/cohort/) A study comparing two or more treatments. (https://mrctcenter.org/glossaryterm/comparative-	Comparative Effectiveness
	Research (CER)	iveriess	effectiveness-research-cer/)	Research
C142458	comparator		Something that is compared to the study treatment. (https://mrctcenter.org/glossaryterm/comparator/)	Comparator
C209471	compensation (stud	dy)	Money and other forms of payment that may be given to participants for completing study activities. (https://mrctcenter.org/glossaryterm/compensation-study/)	Study Compensation
C39490 C17204	compliance Computerized Tom	ography (CT)	Following research requirements. (https://mrctcenter.org/glossaryterm/compliance/) A way to take pictures of the inside of a person's body using a type of radiation and a computer.	Patient Compliance Computed Tomography
	scan concomitant medic		(https://mrctcenter.org/glossaryterm/computerized-tomography-ct-scan/)	Concomitant Agent
C70902		auons	Non-study medicines that are allowed to be taken at the same time as the study treatment. (https://mrctcenter.org/glossaryterm/concomitant-medications/)	Ü
C153144 C53324	conduct confidence interval		To do a study or procedure. (https://mrctcenter.org/glossaryterm/conduct/) The defined range of numbers used to describe where the results are expected to fall.	Study Conduct Confidence Interval
C16466	confidentiality		(https://mrctcenter.org/glossaryterm/confidence-interval/) Protecting personal information from people who should not have access.	Confidentiality
C209462	confounding		(https://mrctcenter.org/glossaryterm/confidentiality/) When the study outcome is influenced by outside conditions that were not expected by the study	Confounding Variable
C16468	consent form		researchers. (https://mrctcenter.org/glossaryterm/confounding/) A document used to explain the planned research before a person decides whether or not to join a	Consent Form
		0	study. (https://mrctcenter.org/glossaryterm/consent-form/)	
C54148	Contract Research (CRO)	Organization	A group that is paid by the study sponsor to support research studies. (https://mrctcenter.org/glossaryterm/contract-research-organization-cro/)	Contract Research Organization
C28143	control group		The people in a study who do not receive the study treatment or do not have the condition being studied. (https://mrctcenter.org/glossaryterm/control-group/)	Control Group
C48834 C82637	correlation crossover trial		When two or more measures are linked. (https://mrctcenter.org/glossaryterm/correlation/) A type of study where each participant receives all the study treatments but in a randomly assigned	Correlation Crossover Study
		nmmittee/Data	order. (https://mrctcenter.org/glossaryterm/crossover-trial/)	-
C142489	Data Monitoring Co and Safety Monitor		An independent group of experts that reviews study data to make sure that patient safety is protected. (https://mrctcenter.org/glossaryterm/data-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-committee-data-and-safety-monitoring-com	Data Monitoring Committee
C25474	(DMC/DSMB) data		monitoring) Information collected from or about people taking part in a research study.	Data
C203935	database (research	1)	(https://mrctcenter.org/glossaryterm/data/) Information that is collected and organized to be used for research.	Research Database
C176257	decentralized clinic		(https://mrctcenter.org/glossaryterm/database-research/) A type of research study where some or all of the participant's study-related activities happen at	Decentralized Clinical Trial
- *:	222		places other than a single study site. (https://mrctcenter.org/glossaryterm/decentralized-clinical-trial/)	
C142444	discontinue (partici	pant)	To remove a study participant from a study. (https://mrctcenter.org/glossaryterm/discontinue- participant/)	Study Subject Discontinuation
C49502	discontinue (study	treatment)	To stop a study treatment in a participant. (https://mrctcenter.org/glossaryterm/discontinue-study-	Drug Withdrawn
C17747	disease progressio	n	treatment/) An illness getting worse over time. (https://mrctcenter.org/glossaryterm/disease-progression/)	Disease Progression
C17751	disease-free surviv	al	The length of time after treatment that a person lives without the illness coming back. (https://mrctcenter.org/glossaryterm/disease-free-survival/)	Disease-Free Survival
C90475	dose escalation stu	ıdy	A kind of study where increasing amounts of a study treatment are given to different groups to find the best dose. (https://mrctcenter.org/glossaryterm/dose-escalation-study/)	Titration Study
C15228	double-blind study		A study that is set up so that the study treatment that each participant receives is not known by the participants or the researchers. (https://mrctcenter.org/glossaryterm/double-blind-study/)	Double Blind Study
C203919	drug holiday		A time period decided between the participant and study team when a medication is stopped and	Drug Holiday
C15986	drug therapy		then re-started. (https://mrctcenter.org/glossaryterm/drug-holiday/) The use of medicine to treat a disease, condition, or symptom.	Pharmacotherapy
C203920	e-consent (form)		(https://mrctcenter.org/glossaryterm/drug-therapy/) An electronic version of an informed consent form. (https://mrctcenter.org/glossaryterm/e-consent-	Electronic Consent Form
C142522	effectiveness		form/) How well a treatment works. (https://mrctcenter.org/glossaryterm/effectiveness/)	Effectiveness
C88183	efficacy		How well a study treatment works in the study. (https://mrctcenter.org/glossaryterm/efficacy/)	Efficacy
C16112	eligibility criteria		The reasons a person can be included in, or excluded from, a study. (https://mrctcenter.org/glossaryterm/eligibility-criteria/)	Clinical Trial Eligibility Criteria
0.02	Emergency Use At (EUA)	ıthorization	A process to make a treatment or vaccine available during a public health emergency, before all research is complete, and before full approval is granted.	Emergency Use Authorization
C96966			(https://mrctcenter.org/glossaryterm/emergency-use-authorization-eua/) A measure of the expected effect of the study treatment.	End Point
C96966	endnoint			
C96966 C25212	endpoint		(https://mrctcenter.org/glossaryterm/endpoint/)	
C96966 C25212 C37948	enroll		(https://mrctcenter.org/glossaryterm/endpoint/) The action of a participant joining the study after providing informed consent. (https://mrctcenter.org/glossaryterm/enroll/)	Enrollment
C96966 C25212	·		(https://mrctcenter.org/glossaryterm/endpoint/) The action of a participant joining the study after providing informed consent.	

	C203912	MRCT Center Clinical Research			
	NCI Code	Glossary CDISC Submission Value	CDISC Synonym	CDISC Definition	NCI Preferred Term
C203921 C142533		equivalent (effect) eSignature		The same or almost the same result. (https://mrctcenter.org/glossaryterm/equivalent-effect/) A way for a person to sign their name using a computer or device, instead of signing on paper.	Equivalent Effect Electronic Signature
C25370		exclusion criteria		(https://mrctcenter.org/glossaryterm/esignature/) A list of reasons a person cannot be included in a study.	Exclusion Criteria
				(https://mrctcenter.org/glossaryterm/exclusion-criteria/)	
C98722		expanded access		A process for a doctor to request an unapproved treatment for a seriously ill patient. (https://mrctcenter.org/glossaryterm/expanded-access/)	Expanded Access Study
C147146		exploratory research		A process to find facts that can guide the design of future studies. (https://mrctcenter.org/glossaryterm/exploratory-research/)	Exploratory Research
C154589		focus group		A group interview to learn what people think about a topic. (https://mrctcenter.org/glossaryterm/focus-group/)	Focus Group
C25515		frequency		How often something happens over a period of time. (https://mrctcenter.org/glossaryterm/frequency/)	Temporal Frequency
C142429		generalizability		How research results can apply to people who were not part of the study. (https://mrctcenter.org/glossaryterm/generalizability/)	Clinical Generalizability
C15709		genetic testing		A medical test that could identify a health risk to them or their biological family members by looking at their genes (DNA). (https://mrctcenter.org/glossaryterm/genetic-testing/)	Genetic Testing
C93150		hazard ratio		A measure of risk that compares two treatments in the same study. (https://mrctcenter.org/glossaryterm/hazard-ratio/)	Hazard Ratio
C49651		healthy volunteer		A study participant who does not have a disease or condition, including the one being studied.	Healthy Subject
C27998		hereditary		(https://mrctcenter.org/glossaryterm/healthy-volunteer/) A parent's features and traits being passed to their biological children before birth.	Hereditary
C70665		human subject		(https://mrctcenter.org/glossaryterm/hereditary/) A person who participates in a research study. (https://mrctcenter.org/glossaryterm/human-subject/)	Human Study Subject
C142668 C17930		hypothesis immune response		An idea that is tested in a research study. (https://mrctcenter.org/glossaryterm/hypothesis/) The body's reaction to a substance, illness, or infection.	Research Hypothesis Immune Response Process
C16726		incidence		(https://mrctcenter.org/glossaryterm/immune-response/) Number of new cases or events during a period of time.	Incidence
C25532		inclusion criteria		(https://mrctcenter.org/glossaryterm/incidence/) A list of requirements a person must meet to take part in a study.	Inclusion Criteria
C16735		informed consent		(https://mrctcenter.org/glossaryterm/inclusion-criteria/) The process of learning and discussing the details of a research study before deciding whether to	Informed Consent
				take part. (https://mrctcenter.org/glossaryterm/informed-consent/)	
C15388		infusion		A way to give a fluid to the study participant, usually through a vein. (https://mrctcenter.org/glossaryterm/infusion/)	Infusion Procedure
C16741		Institutional Review Board (IRB)		A team of people who review studies to protect the rights and welfare of study participants. (https://mrctcenter.org/glossaryterm/institutional-review-board-irb/)	Institutional Review Board
C82667		Investigational Device Exemption (IDE)		An approval given by the United States Food & Drug Administration to use a medical tool, instrument, test, or method in a study that enrolls people.	Investigational Device Exemption
C72968		investigational device		(https://mrctcenter.org/glossaryterm/investigational-device-exemption-ide/) A medical tool, instrument, or test that has not yet been approved for the reason being studied in	Investigational Medical Device
C202579		investigational medicine		the research. (https://mrctcenter.org/glossaryterm/investigational-device/) A treatment or drug that is not yet approved for the condition being studied.	Investigational Medicinal Product
C96090		Investigational New Drug (IND)		(https://mrctcenter.org/glossaryterm/investigational-medicine/) An application to the United States Food & Drug Administration (FDA) to get permission to use a	Investigational New Drug
-		application		drug in a research study that enrolls people. (https://mrctcenter.org/glossaryterm/investigational-new-drug-ind-application/)	Application
C41161		investigational product		A drug, device, vaccine or other treatment being tested in a study. (https://mrctcenter.org/glossaryterm/investigational-product/)	Protocol Agent
C25936 C16032		investigator long-term follow-up (research)		A person who leads a research study. (https://mrctcenter.org/glossaryterm/investigator/) When researchers continue to collect data from study participants after initial research activities are	Investigator
		, , ,		completed. (https://mrctcenter.org/glossaryterm/long-term-follow-up-research/)	
C15273		longitudinal study		Research that collects data from the same participants over a long time. (https://mrctcenter.org/glossaryterm/longitudinal-study/)	Longitudinal Study
C16809		Magnetic Resonance Imaging (MRI)		A way to take pictures of the inside of a person's body with a machine that uses strong magnets and radio waves. (https://mrctcenter.org/glossaryterm/magnetic-resonance-imaging-mri/)	Magnetic Resonance Imaging
C191278		masking (study treatment)		Hiding or covering what the study treatments look like so participants cannot tell the difference. (https://mrctcenter.org/glossaryterm/masking-study-treatment/)	Masking
C165770		master protocol		An overall research plan that guides sub-studies that have their own research questions. (https://mrctcenter.org/glossaryterm/master-protocol/)	Master Protocol
C25564 C53319		maximum mean		The most or largest amount. (https://mrctcenter.org/glossaryterm/maximum/) The average. (https://mrctcenter.org/glossaryterm/mean/)	Maximum Arithmetic Mean
C28007		median		The middle number in a set of numbers when listed in order from lowest to highest. (https://mrctcenter.org/glossaryterm/median/)	Median
C16830		medical device		A test kit, implant, or other item that can be used, worn, or inserted to prevent, diagnose, monitor, or treat a disease, condition, or symptom. (https://mrctcenter.org/glossaryterm/medical-device/)	Medical Device
C25570 C156663		minimal minimur minor	m	Very small. (https://mrctcenter.org/glossaryterm/minimal/) Someone considered too young to give legal consent. (https://mrctcenter.org/glossaryterm/minor/)	Minimum Minor Person
C61256		monitor		To observe, check or evaluate something in a study over time. (https://mrctcenter.org/glossaryterm/monitor/)	Monitoring
C184382		morbidity (rate)		The number of people who develop a disease or illness in a group over time. (https://mrctcenter.org/glossaryterm/morbidity-rate/)	Morbidity Rate
C16880		mortality (rate)		The number of deaths in a group of people over time. (https://mrctcenter.org/glossaryterm/mortality-rate/)	Mortality Rate
C16104		multicenter trial		A study that takes place at more than one research center.	Multi-Institutional Clinical Trial
C35681		negative test result		(https://mrctcenter.org/glossaryterm/multicenter-trial/) A test result that shows a person does not have what was tested for.	Negative Test Result
C203922		negligible		(https://mrctcenter.org/glossaryterm/negative-test-result/) So small that it has little to no impact. (https://mrctcenter.org/glossaryterm/negligible/)	Negligible
C201294 C184386		non-compliance non-inferiority trial		Not following research requirements. (https://mrctcenter.org/glossaryterm/non-compliance/) A study to test if a study treatment works about as well as another treatment for the same condition.	Subject Noncompliance Non-Inferiority Trial
C142450		objective		(https://mrctcenter.org/glossaryterm/non-inferiority-trial/) A purpose or goal of a study. (https://mrctcenter.org/glossaryterm/objective/)	Clinical Trial Objective
C16084		observational study		A study that collects health information about study participants without giving a treatment. (https://mrctcenter.org/glossaryterm/observational-study/)	Observational Study
C16932		odds ratio		The chance of a health event happening in one group compared with the chance of the same event	Odds Ratio
C125600		off-label		happening in another group. (https://mrctcenter.org/glossaryterm/odds-ratio/) The use of a treatment in a different way or for a condition other than what it is approved for. (https://mrctcenter.org/glossaryterm/off.jabel/)	Off-Label Treatment
C49659		open-label		(https://mrctcenter.org/glossaryterm/off-label/) A type of study where participants and research staff know which treatment participants are being	Open Label Study
C221715		opt-in		given. (https://mrctcenter.org/glossaryterm/open-label/) To agree to optional study activities or optional data uses that are not required for the study.	Subject Opt-In for Optional Activities
C221714		opt-out		(https://mrctcenter.org/glossaryterm/opt-in/) To decline optional study activities or optional data uses that are not required for the study.	Subject Opt-Out for Optional
C204098		outcome (of study)		(https://mrctcenter.org/glossaryterm/opt-out/) A description of the overall results of the study. (https://mrctcenter.org/glossaryterm/outcome-of-	Activities Study Outcome Description
C93407		outcome measure		study/) The way that a study endpoint is measured. (https://mrctcenter.org/glossaryterm/outcome-	Study Outcome Measurement
C44185		p-value (probability value)		measure/) A number that researchers use to show that a result did not occur by chance.	P-Value
C83083		participant code		(https://mrctcenter.org/glossaryterm/p-value-probability-value/) A set of numbers, letters, or both that is used to identify a participant instead of their personal	Subject Identifier
C95401		Patient Reported Outcomes (PROs)		information. (https://mrctcenter.org/glossaryterm/participant-code/) The information that patients share about their own health or well-being to answer questions in a	Patient Reported Outcome
		. , ,		rine information that patients share about their own realin or well-being to answer questions in a study. (https://mrctcenter.org/glossaryterm/patient-reported-outcomes-pros/) Evaluation by independent experts. (https://mrctcenter.org/glossaryterm/peer-review/)	Peer Review
C16963 C90492		peer review personal data		Any information that is related to a person, including information that can identify them.	Personal Information
C49662		Pharmacodynamic (PD) study		(https://mrctcenter.org/glossaryterm/personal-data/) A study that measures the effects of a drug on the human body.	Pharmacodynamic Study
C15299		Pharmacokinetic (PK) study		(https://mrctcenter.org/glossaryterm/pharmacodynamic-pd-study/) A study that measures what happens to a drug in a person's body over time.	Pharmacokinetics
C142637		pharmacovigilance		(https://mrctcenter.org/glossaryterm/pharmacokinetic-pk-study/) A process to detect, review, and make decisions about drug safety to protect patients.	Pharmacovigilance
C48281		phase		(https://mrctcenter.org/glossaryterm/pharmacovigilance/) A step in the overall clinical research process to test a new drug or treatment.	Trial Phase
C15303		pilot study		(https://mrctcenter.org/glossaryterm/phase/) A small study that is done to test a process before starting a larger study.	Pilot Study
C753		placebo		(https://mrctcenter.org/glossaryterm/pilot-study/) Something that looks like the treatment being studied, but doesn't contain any medicine.	Placebo
C203925		placebo-controlled study		(https://mrctcenter.org/glossaryterm/placebo/) A study with two or more groups where one group is given a placebo.	Placebo-Controlled Trial
C203925		platform trial		A study with two of more groups where the group is given a placebo. (https://mrctcenter.org/glossaryterm/placebo-controlled-study/) A research study that tests and compares two or more study treatments for a disease or condition,	Platform Trial
J2U3400		pianomi mai		with study treatment groups being added or removed during the study period.	r Muonn mai

C203912 **MRCT Center Clinical Research** Glossary **CDISC Synonym** NCI Code **CDISC Submission Value CDISC Definition NCI Preferred Term** (https://mrctcenter.org/glossaryterm/platform-trial/) C38758 A test result that shows a person has what was tested for positive test result Positive Finding C142640 post-market surveillance Continuing to collect and analyze information about the risks and benefits of medicine and devices Postmarketing Surveillance after they have been approved for patient use. (https://mrctcenter.org/glossaryterm/post-marketsurveillance/) When participants can still receive a study treatment after their participation has ended C187706 post-trial access Continued Access Study (https://mrctcenter.org/glossaryterm/post-trial-access/) C142642 A study to test a treatment in the lab or in animals before testing it in people preclinical study Preclinical Study (https://mrctcenter.org/glossaryterm/preclinical-study/) C17010 prevalence Number of known cases or events in a group. (https://mrctcenter.org/glossaryterm/prevalence/) Prevalence A study measure that is used to answer the main research question. (https://mrctcenter.org/glossaryterm/primary-endpoint/) C94496 primary endpoint Primary Endpoint C19924 The main researcher who oversees a research study and makes sure it is done as planned. principal investigator Principal Investigator (https://mrctcenter.org/glossaryterm/principal-investigator/) C54154 probability The likelihood or chance that something might happen. Probability (https://mrctcenter.org/glossaryterm/probability/) C98769 procedures (for participants) The activities that participants will be asked to do during the research study Physical Medical Procedure (https://mrctcenter.org/glossaryterm/procedures-for-participants/) The length of time without a person's illness getting worse. C28234 progression-free survival Progression-free Survival (https://mrctcenter.org/glossaryterm/progression-free-survival/) C142646 prospective study Research that uses new data collected from participants. Prospective Study (https://mrctcenter.org/glossaryterm/prospective-study/) C142451 protocol A complete description of the research plan and procedures. Clinical Trial Protocol (https://mrctcenter.org/glossaryterm/protocol/) C119264 A person who is legally allowed to make research decisions for someone else. proxy Proxv (https://mrctcenter.org/glossaryterm/proxy/) C142654 pseudonymize Replace personal details with a code so that data are protected. Pseudonymization (https://mrctcenter.org/glossaryterm/pseudonymized/) C17047 Quality of Life (QOL) How someone feels and functions day to day. (https://mrctcenter.org/glossaryterm/quality-of-life-Quality of Life qol/) A list of questions for study participants to answer as part of the study C17048 questionnaire Questionnaire (https://mrctcenter.org/glossaryterm/questionnaire/) C25196 randomization A way to use chance to place study participants into different study treatment groups. Randomization (https://mrctcenter.org/glossaryterm/randomization/) C46079 randomized controlled trial Research that uses chance for participants to be assigned to the study treatment group or a Randomized Controlled Clinical comparison group. (https://mrctcenter.org/glossaryterm/randomized-controlled-trial/) Trial Real World Data (RWD) Information from many different sources used for health research purposes. Real-world Data C165830 (https://mrctcenter.org/glossaryterm/real-world-data-rwd/) C165831 Real World Evidence (RWE) Real-world Evidence Findings from analyzing real world data. (https://mrctcenter.org/glossaryterm/real-world-evidence-When a study is open to new participants joining the study. C221713 recruiting (status) Study is Recruiting (https://mrctcenter.org/glossaryterm/recruiting-status/) An organized list of research information. (https://mrctcenter.org/glossaryterm/registry-study/) Study Registry C93453 registry (study) The chance of a harmful event happening in one study group compared to another. (https://mrctcenter.org/glossaryterm/relative-risk/) C93152 relative risk Relative Risk C203936 A collection of participant data and samples stored for future research. Research Repository repository (research) (https://mrctcenter.org/glossaryterm/repository-research/) C203930 results (study) Study Results Findings from the study. (https://mrctcenter.org/glossaryterm/results-study/) C53312 retrospective study Research that uses already existing data. (https://mrctcenter.org/glossaryterm/retrospective-study/) Retrospective Study C203928 risk-benefit ratio A comparison of the possible bad and potential good things that could happen if a participant joins Risk-Benefit Ratio a research study. (https://mrctcenter.org/glossaryterm/risk-benefit-ratio/) C142718 risks of a research study The possible harms of being in a research study. (https://mrctcenter.org/glossaryterm/risks-of-a-Subject Risk Body fluids or tissues collected from a participant during a research study. C70699 sample (study) Biospecimen (https://mrctcenter.org/glossaryterm/sample-study/) Sample Size C53190 sample size The number of participants in a study or study group. (https://mrctcenter.org/glossaryterm/sample-C142678 schedule of assessments A chart that lists the study activities and when they will happen during a study. Schedule Of Assessments (https://mrctcenter.org/glossaryterm/schedule-of-assessments/) C49628 When a person cannot join a study because they do not fit its eligibility criteria. Trial Screen Failure screen failure (https://mrctcenter.org/glossaryterm/screen-failure/) C48262 Tests and questions to find out if a person can join a study screening Trial Screening (https://mrctcenter.org/glossaryterm/screening/) C139173 A measure used to answer other important questions in the study that are not the main research Secondary Endpoint secondary endpoint question. (https://mrctcenter.org/glossaryterm/secondary-endpoint/) How well a medical test can accurately identify people who have a disease or trait. (https://mrctcenter.org/glossaryterm/sensitivity-medical-test/) C41394 sensitivity (medical test) Diagnostic Sensitivity C142685 Serious Adverse Drug Reaction A health issue, possibly caused by a medication, that leads to hospital care, lasting medical Serious Adverse Drug Reaction problems, life-threatening conditions, or death. (https://mrctcenter.org/glossaryterm adverse-drug-reaction-sadr/) Serious Adverse Event (SAE) C41335 A health issue that happens during a study and leads to hospital care, lasting medical problems, Serious Adverse Event life-threatening conditions, or death. (https://mrctcenter.org/glossaryterm/serious-adverse-event-C116527 sham (procedure) A medical process that looks and feels like the study treatment, but is not expected to have a health Sham Intervention effect on the condition being studied. (https://mrctcenter.org/glossaryterm/sham-procedure/) side effect A health change that is not the intended effect of the treatment and usually considered a problem. Side Effect C2861 (https://mrctcenter.org/glossaryterm/side-effect/) C28233 single-blind study A study that is set up so that the study treatment each participant receives is not known by the Single Blind Study participants but is known by the researchers. (https://mrctcenter.org/glossaryterm/single-blind-C41395 How well a medical test can accurately identify people who do not have a disease or trait. (https://mrctcenter.org/glossaryterm/specificity-medical-test/) specificity (medical test) Diagnostic Specificity C70793 The group that is in charge of, or pays for, a research study. Clinical Study Sponsor sponsor (https://dev.mrctcenter.org/glossaryterm/sponsor-3/) standard of care The usual treatment given to patients for an illness. (https://mrctcenter.org/glossaryterm/standard-C94396 **Best Practice** of-care/) Statistical Significance Results that are very unlikely to have occurred by chance C61040 statistically significant (https://mrctcenter.org/glossaryterm/statistically-significant/) C15320 study design The way a study is set up to answer the study question. (https://mrctcenter.org/glossaryterm/study-Study Design C215672 study doctor A person with relevant health training who makes sure clinical study procedures are done correctly, Clinical Trial Physician and participants are safe and cared for. (https://mrctcenter.org/glossaryterm/study-doctor/) How likely it is that a study can be completed. (https://mrctcenter.org/glossaryterm/study-feasibility/) C209473 Study Feasibility study feasibility A set of numbers, letters, or both, that is used to identify a study. C83082 study identifier Study Identifier (https://mrctcenter.org/glossaryterm/study-identifier/) C25218 study intervention A treatment given to the participants in a study. (https://mrctcenter.org/glossaryterm/study. C203929 study life cycle The steps of a research study from beginning to end. (https://mrctcenter.org/glossaryterm/study-life- Study Life Cycle study monitor A qualified person who checks that a research study is being done safely and is following the study Study Monitor C41201 plan correctly. (https://mrctcenter.org/glossaryterm/study-monitor/) C142710 study participant A person who joins a research study. (https://mrctcenter.org/glossaryterm/study-participant/) Study Participant C70833 All the participants in a study. (https://mrctcenter.org/glossaryterm/study-population/) Study Population study population C80403 An approved location where research study activities take place. study site Study Site (https://mrctcenter.org/glossaryterm/study-site/) C142737 study statistician A person who uses math to help design a study and interpret the data. Trial Statistician (https://mrctcenter.org/glossaryterm/study-statistician/) C198230 substudy A study with a smaller group of participants already enrolled in the main study. Substudy (https://mrctcenter.org/glossaryterm/substudy/) A study to test if a study treatment works better than another treatment for the same condition. C142722 superiority trial Superiority Trial (https://mrctcenter.org/glossaryterm/superiority-trial/) C203931 synergistic effect When two or more things combined have a greater effect than when their individual effects are Synergistic Effect added together. (https://mrctcenter.org/glossaryterm/synergistic-effect/) When research is stopped earlier than planned. (https://mrctcenter.org/glossaryterm/termination-of-C142739 termination (of a research study) How much a participant or group of participants can accept a study treatment's unwanted effects so Study Intervention Tolerability they can keep taking it. (https://mrctcenter.org/glossaryterm/tolerability/) tolerability C203932 Treatment Effect C209469 treatment effect How much a study treatment changes a condition, symptom, or function. ((https://mrctcenter.org/glossaryterm/treatment-effect/) A research study that tests and compares two or more study treatments for one disease or Umbrella Trial C209470 umbrella trial condition. (https://mrctcenter.org/glossaryterm/umbrella-trial/) A medical substance that helps the body's immune system build protection against an infection or C923 vaccine Vaccine disease. (https://mrctcenter.org/glossaryterm/vaccine/) Choosing to participate in research without feeling pressured. C203933 voluntary participation Voluntary Participation

С	C203912	MRCT Center Clinical Research Glossary			
N	ICI Code	CDISC Submission Value	CDISC Synonym	CDISC Definition	NCI Preferred Term
				(https://mrctcenter.org/glossaryterm/voluntary-participation/)	
C42872		wash-out		A time before starting a study treatment when a person stops taking other medicines. (https://mrctcenter.org/glossaryterm/wash-out/)	Washout Period
C49634		withdraw		To stop being a participant in a study. (https://mrctcenter.org/glossaryterm/withdraw/)	Withdrawal by Subject
C38101		X-ray		A way of taking pictures of the inside of a person's body using X-ray radiation. (https://mrctcenter.org/glossaryterm/x-ray/)	X-Ray Imaging