Scientific vs lay language

| abdomen | belly, stomach |
|----------------------|---|
| abdominal distention | bloating |
| absorb | take up fluids, take in |
| acid taste | sour taste |
| acidosis | condition when blood contains more acid than normal |
| | clearness, keenness, esp. of vision and airways |
| acuity | |
| acute | new, recent, sudden, urgent |
| | gall stones, which may cause upper abdominal pain and require |
| acute cholecystitis | hospitalization and surgery |
| adenopathy | swollen lymph nodes (glands) |
| adjuvant | helpful, assisting, aiding, supportive |
| agent | drug, medication |
| albumin | protein found in blood |
| allergic reaction | rash, hives, swelling, trouble breathing |
| alopecia | loss of hair |
| ambulate/ambulation/ | |
| ambulatory | walk, able to walk |
| analgesic | pain-relieving drug |
| anaphylaxis | serious, potentially life-threatening allergic reaction |
| | build up of fluid throughout the whole body, which occurs in |
| anasarca | severely ill people |
| | low number of red blood cells, can causes tiredness and |
| anemia | shortness of breath. May require a blood transfusion |
| | a drug or agent used to decrease the feeling of pain, or eliminate |
| anesthetic | the feeling of pain by putting you to sleep |
| angina | chest pain due to decreased oxygen getting to the heart. |
| anorexia | disorder in which person will not eat; lack of appetite |
| antecubital | related to the inner side of the forearm |
| antibody | protein made in the body in response to foreign substance |
| anticonvulsant | drug used to prevent seizures |
| antiemetic | medication to prevent nausea/vomiting |
| antilipemic | a drug that lowers fat levels in the blood |
| antimicrobial | drug that kills bacteria and other germs |
| antiretroviral | drug that works against the growth of certain viruses |
| antitussive | a drug used to relieve coughing |
| aplastic anemia | a disorder caused by decreased production of red blood cells |
| arrhythmia | irregular heart beat |
| | blood clot in an artery that blocks the artery. This could be serious |
| arterial thrombosis | and life threatening |
| arterial catheter | small tube placed in an artery |
| arthralgia | joint pain |
| | build up of fluid in the abdomen, which causes bloating and |
| | discomfort. This could require that the fluid be removed by a |
| accitoc | |
| ascites | procedure called paracentesis |
| aspiration | fluid entering the lungs, such as after vomiting |
| assay | lab test |
| asthenia | feeling weak and having no energy |

| | lung disease associated with tightening of air passages, making |
|----------------------|--|
| acthma | |
| asthma | breathing difficult |
| | this is when your immune system attacks normal cells in your |
| | body, including the cells that line your digestive tract. This may |
| | result in bleeding and inflammation of the esophagus, bowel |
| | (intestines), and lower gi tract (colon), which can cause bleeding, |
| | diarrhea and perforations (holes). This could be serious or life |
| | threatening. Hospitalization and treatment with medications |
| | (steroids) may be necessary. This can become severe and may |
| | require surgical removal of parts of the intestines or colon. These |
| | surgical procedures might result in your having a stoma (hole) |
| autoimmune enteritis | |
| axilla | armpit |
| | 1. Information gathered at the beginning of a study from which |
| | variations found in the study are measured. 2. A known value or |
| | quantity with which an unknown is compared when measured or |
| | assessed. 3. The initial time point in a clinical trial, just before a |
| | participant starts to receive the experimental treatment which is |
| | being tested. At this reference point, measurable values such as |
| | cd4 count are recorded. Safety and efficacy of a drug are often |
| baseline | determined by monitoring changes from the baseline values. |
| benefit | a valued or desired outcome; an advantage |
| benign | not malignant, without serious consequences |
| bilirubinemia | high levels of bilirubin in the blood |
| | the extent to which a drug or other substance becomes available |
| bioavailability | to the body |
| , | any therapeutic serum, toxin, anti-toxin, or analogous microbial |
| | product applicable to the prevention, treatment, or cure of |
| biologic | diseases or injuries |
| biopsy | removal and examination of tissue |
| | a randomized trial is "blind" if the participant is not told which arm |
| blind | of the trial he is on |
| bolus | a large amount given all at once |
| | the amount of calcium and other minerals in a given amount of |
| bone mass | bone |
| bowel perforation | perforation of the digestive system |
| bradycardia | slow heartbeat |
| | accumulation of fluid around the brain stem, this can be life |
| brain stem edema | threatening |
| bronchospasm | breathing distress caused by narrowing of the airways |
| carcinogenic | cancer-causing |
| cardiac | related to the heart |
| cardiac arrest | sudden, unexpected stopping of the heart. |
| cardiac effusion | collection of fluid around the heart |
| cardiac toxicity | damage to the heart |
| | heart muscle becomes damaged and the heart doesn't pump |
| cardiomyopathy | properly |
| cardiovascular | heart and blood vessels |

| cardioversion | return to normal heartbeat by electric shock |
|----------------------|---|
| | a research strategy that focuses on one case (an individual, a |
| | group, an organization, etc.) within its social context during one |
| case study | time period |
| catheter | a tube for withdrawing or giving fluids |
| central nervous | |
| system (CNS) | brain and spinal cord |
| cerebral trauma | damage to the brain |
| cessation | stopping |
| chemotherapy | treatment of disease, usually cancer, by chemical agents |
| chronic | continuing for a long time, ongoing |
| clinical | pertaining to medical care |
| | a probability sample that is determined by randomly selecting |
| | clusters of people from a population and subsequently selecting |
| cluster sample | every person in each cluster for inclusion in the sample |
| | having either a psychiatric disorder (e.g., psychosis, neurosis, |
| | personality or behavior disorders, or dementia) or a |
| | developmental disorder (e.g., mental retardation) that affects |
| | cognitive or emotional functions to the extent that capacity for |
| cognitively impaired | judgment and reasoning is significantly diminished |
| | a group of subjects initially identified as having one or more |
| cohort | characteristics in common who are followed over time |
| coma | unconscious state |
| | a method of providing experimental therapeutics prior to final FDA |
| compassionate use | approval for use in humans |
| | payment or medical care provided to subjects injured in research; |
| | does not refer to payment (remuneration) for participation in |
| compensation | research |
| | a legal term to indicate a person's capacity to act on one's own |
| | behalf; a person's ability to understand information presented, to |
| | realize the consequences of acting (or not acting) on that |
| competence | information, and to make a choice. |
| complete response | total disappearance of disease |
| | any factor that might serve as an alternative explanation for a |
| | study's result; confounding factors include non-randomized |
| confounding factor | samples, selection bias, and any arbitrary differences between |
| confounding factor | people that are being compared. |
| congenital | present before birth |
| conjunctivitis | redness and irritation of the thin membrane that covers the eye treatment phase intended to make a remission permanent (follows |
| consolidation phase | induction phase) |
| constipation | difficulty passing stools |
| contract | an agreement |
| | disadvantageous, perhaps dangerous; a treatment that should not |
| | be used in certain individuals or conditions due to risks. For |
| | instance, a drug may be contraindicated for pregnant women and |
| contraindicated | people with high blood pressure |
| contraindicated | Iheohie with high piona hiessare |

| | in many clinical trials, one group of patients will be given an |
|--|--|
| | experimental drug or treatment, while the control group is given |
| control group | either a standard treatment for the illness or a placebo |
| | |
| | research study in which the experimental treatment or procedure |
| controlled trial | is compared to a standard (control) treatment or procedure |
| | a non-probability sample that is determined by selecting |
| | participants that are readily accessible (convenient) to the |
| | researcher, (examples in studies of Stanford students might |
| | include going to an organizational meeting or hanging out outside |
| | of rastall and asking students exiting the lunchroom to take a |
| convenience comple | |
| convenience sample | survey) |
| cooperative group | association of multiple institutions to perform clinical trials |
| | related to the blood vessels that supply the heart, or to the heart |
| coronary | itself |
| | a relationship where two variables are associated (this can be |
| correlational | measured in terms of strength and direction using statistical tests) |
| relationship | but not causally related |
| | a type of clinical trial in which each subject experiences, at |
| cross-over design | different times, both the experimental and control therapy |
| culture | test for infection, or for organisms that could cause infection |
| cumulative | added together from the beginning |
| cutaneous | relating to the skin |
| debilitation | weakened condition |
| | |
| | |
| | giving participants previously undisclosed information about the |
| | giving participants previously undisclosed information about the research project following completion of their participation in |
| | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not |
| | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS |
| | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following |
| debrief | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study |
| | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or |
| | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is |
| debrief | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's |
| debrief deception | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals |
| debrief deception dehydrate | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids |
| debrief deception dehydrate dermatitis | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash |
| debrief deception dehydrate dermatitis dermatologic | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin |
| debrief deception dehydrate dermatitis | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin condition to grow worse |
| debrief deception dehydrate dermatitis dermatologic deteriorate | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin condition to grow worse refers to trials that are are conducted to find better tests or |
| debrief deception dehydrate dermatitis dermatologic | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin condition to grow worse |
| debrief deception dehydrate dermatitis dermatologic deteriorate | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin condition to grow worse refers to trials that are are conducted to find better tests or procedures for diagnosing a particular disease or condition |
| debrief deception dehydrate dermatitis dermatologic deteriorate | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin condition to grow worse refers to trials that are are conducted to find better tests or procedures for diagnosing a particular disease or condition frequent, loose watery stools, which can cause dehydration and |
| debrief deception dehydrate dermatitis dermatologic deteriorate | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin condition to grow worse refers to trials that are are conducted to find better tests or procedures for diagnosing a particular disease or condition |
| debrief deception dehydrate dermatitis dermatologic deteriorate diagnostic trials | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin condition to grow worse refers to trials that are are conducted to find better tests or procedures for diagnosing a particular disease or condition frequent, loose watery stools, which can cause dehydration and |
| debrief deception dehydrate dermatitis dermatologic deteriorate diagnostic trials diarrhea | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin condition to grow worse refers to trials that are are conducted to find better tests or procedures for diagnosing a particular disease or condition frequent, loose watery stools, which can cause dehydration and may require hospitalization and treatment with intravenous fluids |
| debrief deception dehydrate dermatitis dermatologic deteriorate diagnostic trials diagnostic trials diastolic distal | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin condition to grow worse refers to trials that are are conducted to find better tests or procedures for diagnosing a particular disease or condition frequent, loose watery stools, which can cause dehydration and may require hospitalization and treatment with intravenous fluids lower number in a blood pressure reading toward the end, away from the center of the body |
| debrief deception dehydrate dermatitis dermatologic deteriorate diagnostic trials diarrhea diastolic distal distal parathesias | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin condition to grow worse refers to trials that are are conducted to find better tests or procedures for diagnosing a particular disease or condition and may require hospitalization and treatment with intravenous fluids lower number in a blood pressure reading toward the end, away from the center of the body numbness and tingling in the hands and feet. |
| debrief deception dehydrate dermatitis dermatologic deteriorate diagnostic trials diagnostic trials diastolic distal | giving participants previously undisclosed information about the research project following completion of their participation in research. In studies involving deception, if the participants are not informed of the deception in the informed consent, the IRB-SBS requires a signed debrief form for each participant following completion of his/her participation in the study the intentional withholding of information from participants, or deception about the study's purpose and exact nature, that is deemed necessary by the researcher in order to meet the study's goals lose water or body fluids skin irritation, rash pertaining to the skin condition to grow worse refers to trials that are are conducted to find better tests or procedures for diagnosing a particular disease or condition frequent, loose watery stools, which can cause dehydration and may require hospitalization and treatment with intravenous fluids lower number in a blood pressure reading toward the end, away from the center of the body |

| | a clinical trial in which two or more doses of an agent (such as a |
|----------------------|--|
| | drug) are tested against each other to determine which dose |
| dose-ranging study | works best and is least harmful |
| | an experiment in which neither the participants nor the research |
| | staff who interact with them knows the memberships of the |
| | experimental or control groups. Also known as double-masked |
| double-blind design | design (see single-blind design and open design) |
| duration | length of time involved |
| dysplasia | abnormal cells |
| echocardiogram | using soundwaves for examination of the heart |
| edema | build up of fluid in the body causing swelling. |
| efficacy | effectiveness |
| electrocardiogram | electrical tracing of the heartbeat (ECG or EKG) |
| ÿ | changes in electrolytes (body salts), which usually do not cause |
| | any symptoms but that can sometimes cause fatigue, muscle |
| electrolyte changes | weakness, cramping, rigidity, irregular heart beat, or seizures |
| | may indicate inflammation of the pancreas, which could result in |
| elevated lipase, | abdominal pain and discomfort and could require hospitalization |
| amylase | and intravenous treatment |
| elevated uric acid | may worsen kidney function; cause joint pain (gout) and kidney |
| levels | |
| leveis | stones |
| aligibility aritaria | summary criteria for participant selection; includes inclusion and |
| eligibility criteria | exclusion criteria |
| | a legal status given to those individuals who have not yet attained |
| | the age of legal competency as defined by state law, but who are |
| | entitled to adult treatment because of assuming adult |
| | responsibilities such as being self-supporting and not living at |
| emancipated minor | home, marriage, or procreation |
| emesis | vomiting, throwing up |
| empirical | based on experimental data, not on a theory. |
| encephalopathy | disease of the brain that severely alters thinking. |
| endoscopic | |
| examination | examination of an internal part of the body with a lighted tube |
| endpoint | overall outcome that the protocol is designed to evaluate |
| enteral | by way of the intestines |
| enzyme | a chemical in the blood that causes chemical changes |
| | |
| | The branch of medical science that deals with the study of |
| epidemiology | incidence and distribution and control of a disease in a population. |
| epidural | outside the spinal cord |
| epistaxis | bloody nose |
| | fair or just; used in the context of selection of participants to |
| | indicate that the benefits and burdens of research are fairly |
| equitable | distributed |
| erythema | redness of the skin |
| ethnographic | |
| research | ethnography is the study of people and their cultures |
| evaluated, assessed | examined for a medical condition |
| excrete | discharge, pass |
| | |

| | refers to any of the FDA procedures, such as compassionate use, |
|-------------------------------|--|
| | parallel track, and treatment IND that distribute experimental |
| | drugs to participants who are failing on currently available |
| | ••••••••••••••••••••••••••••••••••••••• |
| ovponded ecopo | treatments for their condition and also are unable to participate in |
| expanded access | ongoing clinical trials |
| a sur a sina a su ta balanca. | a drug that is not FDA licensed for use in humans, or as a |
| experimental drug | treatment for a particular condition |
| | the events is an experimental design study that reactives treatment |
| | the group in an experimental design study that receives treatment |
| experimental group | in the form, or in various forms, of the independent variable |
| external | outside the body |
| extravasate | to leak outside of a planned area, such as out of a blood vessel |
| fatigue | feeling tired |
| fetus | unborn baby |
| fever | abnormally high body temperature |
| fibrillation | irregular beat of the heart or other muscle |
| fibrosis | scars |
| fibrous | having many fibers, such as scar tissue |
| | behavioral, social, or anthropological research involving the study |
| | of people or groups in their own environment and without |
| field research | manipulation for research purposes |
| fluoroscope | x-ray machine |
| fungus | form of infection |
| gait | walk |
| gastrointestinal | stomach and intestines |
| | pain prevention by giving drugs to cause loss of consciousness, |
| general anesthesia | as during surgery |
| | the ability to apply the results of a specific study to groups or |
| generalizability | situations beyond those actually studied |
| genetics* | the study of heredity |
| genomics* | the sudy of genes and their functions, and related techniques |
| genetics v. genomics | The main difference between genomics and genetics is that |
| · · | genetics scurtinizes the functioning and composition of the single |
| *(Based on W.H.O. | gene, whereas genomics addresses all genes and their inter- |
| definitions) | relationships in order to identify their combined influence on the |
| | growth and development of the organism |
| | (Based on W.H.O. definitions) |
| | tests to identify persons who have an inherited predisposition to a |
| | certain phenotype or who are at risk of producing offspring with |
| genetic screening | inherited diseases or disorders |
| genotype | the genetic constitution of an individual |
| gestational | pertaining to pregnancy |
| - | |
| | an individual who is authorized under applicable state or local law |
| guardian | to give permission on behalf of a child to general medical care |
| headache | pain in the head |
| heart palpitations | heart beats that are fast and hard |
| hematocrit | amount of red blood cells in the blood |
| hematoma | blood clot |
| hematuria | blood in urine |
| | |

| hemodynamic | T |
|--|--|
| measuring | measuring of blood flow |
| hemolysis | measuring of blood flow breakdown in red blood cells |
| | |
| homolytic uromic | red blood cells begin to dissolve, which leave wastes in the blood |
| hemolytic uremic | • |
| syndrome | and the kidneys are unable to get rid of excess fluid and wastes |
| hemoptysis | vomiting blood |
| hemorrhage | loss of blood (heavy bleeding) |
| hemorrhagic cystitis | inflammation of the bladder with severe bleeding |
| | needle placed in the arm with blood thinner to keep the blood from |
| heparin lock | clotting |
| hepatoma | cancer or tumor of the liver |
| | disease that can be transmitted to one's offspring, resulting in |
| heritable disease | damage to future children |
| | may cause fatigue, weight loss, rapid heartbeat, sweating, trouble |
| high thyroid function | with heat, nervousness |
| histopathologic | pertaining to the disease status of body tissues or cells |
| holter monitor | a portable machine for recording heart beats |
| hormone | a chemical in the body |
| | Individuals whose physiologic or behavioral characteristics and |
| | responses are the object of study in a research project. Under the |
| | federal regulations, human subjects are defined as: living |
| | individual(s) about whom an investigator conducting research |
| | obtains: (1) data through intervention or interaction with the |
| human subjects | individual; or (2) identifiable private information. |
| hypercalcemia | high levels of calcium in the blood |
| | high levels of potassium in the blood, which can cause the heart |
| hyperkalemia | to stop beating |
| hyperkeratosis | thickening of the skin, nails. |
| hypernatremia | high blood sodium level |
| hyperpigmentation | darkening of the skin |
| hyperpyrexia | high body temperature, a fever. |
| hypertension | high blood pressure |
| | excess amount of uric acid in the blood, gout, which can cause |
| hyperuricemia | |
| пурстансстна | |
| | pain in the joints |
| hypokalomia | decreased levels of potassium in the blood, which can cause |
| hypokalemia | decreased levels of potassium in the blood, which can cause irregular heart beat |
| | decreased levels of potassium in the blood, which can cause irregular heart beat low magnesium, which may result in muscle cramps, weakness, |
| hypokalemia hypomagnesemia | decreased levels of potassium in the blood, which can cause irregular heart beat low magnesium, which may result in muscle cramps, weakness, tremors or irregular heartbeat |
| hypomagnesemia | decreased levels of potassium in the blood, which can cause irregular heart beat low magnesium, which may result in muscle cramps, weakness, tremors or irregular heartbeat decreased levels of sodium in the blood, which can cause |
| | decreased levels of potassium in the blood, which can cause irregular heart beat low magnesium, which may result in muscle cramps, weakness, tremors or irregular heartbeat decreased levels of sodium in the blood, which can cause confusion, seizures, fatigue and low levels of consciousness |
| hypomagnesemia hyponatremia | decreased levels of potassium in the blood, which can cause irregular heart beat low magnesium, which may result in muscle cramps, weakness, tremors or irregular heartbeat decreased levels of sodium in the blood, which can cause confusion, seizures, fatigue and low levels of consciousness low phosphate, which may result in muscle weakness, bone pain, |
| hypomagnesemia hyponatremia hypophosphatemia | decreased levels of potassium in the blood, which can cause irregular heart beat low magnesium, which may result in muscle cramps, weakness, tremors or irregular heartbeat decreased levels of sodium in the blood, which can cause confusion, seizures, fatigue and low levels of consciousness |
| hypomagnesemia hyponatremia hypophosphatemia hypopigmentation / | decreased levels of potassium in the blood, which can cause irregular heart beat low magnesium, which may result in muscle cramps, weakness, tremors or irregular heartbeat decreased levels of sodium in the blood, which can cause confusion, seizures, fatigue and low levels of consciousness low phosphate, which may result in muscle weakness, bone pain, confusion and muscle breakdown |
| hypomagnesemia hyponatremia hypophosphatemia hypopigmentation / vitiligo | decreased levels of potassium in the blood, which can cause irregular heart beat low magnesium, which may result in muscle cramps, weakness, tremors or irregular heartbeat decreased levels of sodium in the blood, which can cause confusion, seizures, fatigue and low levels of consciousness low phosphate, which may result in muscle weakness, bone pain, confusion and muscle breakdown patches of the skin turn lighter than the surrounding skin |
| hypomagnesemia hyponatremia hypophosphatemia hypopigmentation / | decreased levels of potassium in the blood, which can cause irregular heart beat low magnesium, which may result in muscle cramps, weakness, tremors or irregular heartbeat decreased levels of sodium in the blood, which can cause confusion, seizures, fatigue and low levels of consciousness low phosphate, which may result in muscle weakness, bone pain, confusion and muscle breakdown patches of the skin turn lighter than the surrounding skin low blood pressure |
| hypomagnesemia hyponatremia hypophosphatemia hypopigmentation / vitiligo | decreased levels of potassium in the blood, which can cause irregular heart beat low magnesium, which may result in muscle cramps, weakness, tremors or irregular heartbeat decreased levels of sodium in the blood, which can cause confusion, seizures, fatigue and low levels of consciousness low phosphate, which may result in muscle weakness, bone pain, confusion and muscle breakdown patches of the skin turn lighter than the surrounding skin |

| | a testable statement of how two or more variables are expected to |
|---------------------|--|
| hypothesis | be related to one another |
| hypoxemia | a decrease of oxygen in the blood |
| hypoxia | a decrease of oxygen reaching body tissues |
| Пуроліа | surgical removal of the uterus, ovaries (female sex glands), or |
| hysterectomy | both uterus and ovaries |
| iatrogenic | caused by a physician or by treatment |
| idiopathic | of unknown cause |
| immunity | defense against, protection from |
| immunization | administration of a substance to prevent disease |
| immunoglobin | a protein that makes antibodies |
| immunological | |
| effects | effect on the immune system |
| | drug which works against the body's immune (protective) |
| | response, often used in transplantation and diseases caused by |
| immunosuppressive | immune system malfunction |
| | giving of drugs to help the body's immune (protective) system; |
| immunotherapy | usually used to destroy cancer cells |
| ininianounorapy | refers to a person's mental status and means inability to |
| | understand information presented, to appreciate the |
| | consequences of acting (or not acting) on that information, and to |
| incapacity | make a choice |
| inclusion/exclusion | the medical or social standards determining whether a person |
| criteria | may or may not be allowed to enter a clinical trial |
| | used as a legal term to indicate the inability to manage one's own |
| incompetence | affairs |
| induction | start |
| induration | hardening |
| indwelling | remaining in a given location, such as a catheter |
| infarct | death of tissue due to lack of blood supply |
| infectious disease | disease that is transmitted from one person to the next |
| inflammation | swollen, red, and painful |
| | the process of learning the key facts about a clinical trial before |
| informed consent | deciding whether or not to participate |
| | slow injection of a substance into the body, usually into the blood |
| infusion | by means of a catheter |
| ingestion | eating; taking by mouth |
| insomnia | inability to sleep |
| | confined, either voluntarily or involuntarily (e.g., a hospital, prison, |
| institutionalized | or nursing home) |
| interferon | drug which acts against viruses; antiviral agent |
| | occurring (regularly or irregularly) between two time points; |
| intermittent | repeatedly stopping, then starting again |
| interstitial | |
| pneumonitis, | inflammation of the lungs, which can cause shortness of breath |
| pneumonitis | and difficulty breathing |
| | includes both physical procedures by which data are gathered and |
| | manipulations of the participant or the participant's environment |
| intervention | that are performed for research purposes |
| | |

| intracatheter | small tube in a vein |
|----------------------|---|
| intramuscular | into the muscle; within the muscle |
| intraperitoneal | into the abdominal cavity |
| intrathecal | into the spinal fluid |
| intravenous (IV) | through the vein |
| | |
| intravesical | in the bladder |
| intubate | the placement of a tube into the airway |
| invasive procedure | puncturing, opening, or cutting the skin |
| investigational | a treatment method which has not been proven to be beneficial or |
| method | has not been accepted as standard care |
| | the individual(s) designated to have the appropriate level of |
| | authority and responsibility to direct the research project and/or |
| investigator | activity |
| irradiation | x-ray |
| | decreased oxygen in a tissue (usually because of decreased |
| ischemia | blood flow) |
| jaundice | yellowing of the skin |
| | Surgical procedure in which an incision is made in the abdominal |
| laparotomy | wall to enable a doctor to look at the organs inside. |
| Legally Authorized | |
| Representative | a person authorized either by statute or by court appointment to |
| (LAR) | make decisions on behalf of another person |
| lesion | wound or injury; a diseased patch of skin |
| lethargy | sleepiness, tiredness |
| leukopenia | low white blood cell count |
| lipid | fat |
| | creation of insensitivity to pain in a small, local area of the body, |
| local anesthesia | usually by injection of numbing drugs |
| localized | restricted to one area, limited to one area |
| | a study in which data are collected from the same sample at least |
| longitudinal atudu | two different times |
| longitudinal study | |
| | abnormal decrease in sugar in the blood, which can cause |
| low blood sugar / | weakness, fatigue, and if severe, can cause loss of |
| hypoglycemia | consciousness |
| | may cause fatigue, weight gain, fluid retention, feeling cold, |
| low thyroid function | decreased cognitive function |
| low white cell count | increased risk of infection |
| lumen | the cavity of an organ or tube (e.g., blood vessel) |
| | an x-ray of the lymph nodes or tissues after injecting dye into |
| lymphangiography | lymph vessels (e.g., in feet) |
| | a type of white blood cell important in immunity (protection) |
| lymphocyte | against infection |
| lymphoma | a cancer of the lymph nodes (or tissues) |
| malaise | a vague feeling of bodily discomfort, feeling badly |
| malfunction | condition in which something is not functioning properly |
| | cancer or other progressively enlarging and spreading tumor, |
| malignancy | usually fatal if not successfully treated |
| malignancy | |

| | and the second reached adulthood (as defined by state |
|-----------------------|---|
| | someone who has not reached adulthood (as defined by state |
| | law) but who may be treated as an adult for certain purposes (e.g. |
| mature minor | consenting to medical care) |
| medullablastoma | a type of brain tumor |
| megaloblastosis | change in red blood cells |
| metabolic acidosis | the body becomes more acid |
| metabolism | chemical changes which provide energy |
| | |
| metabolize | process of breaking down substances in the cells to obtain energy |
| metastasis | spread of cancer cells from one part of the body to another |
| | drug used to treat infections caused by parasites (invading |
| | organisms that take up living in the body) or other causes of |
| | anaerobic infection (not requiring oxygen to survive) mi |
| metronidazole | myocardial infarction, heart attack |
| minimal | slight |
| | a risk is minimal where the probability and magnitude of harm or |
| | discomfort anticipated in the proposed research are not greater, in |
| | and of themselves, than those ordinarily encountered in daily life |
| | or during the performance of routine physical or psychological |
| minimal risk | examinations or tests |
| | a risk is moderate when it includes non-public behavior or data |
| | and/or allows for connection of the response to the individual's |
| moderate risk | identity |
| monitor | check on; keep track of; watch carefully |
| | the collection and analysis of data as the project progresses to |
| | assure the appropriateness of the research, its design and |
| monitoring | participant protections |
| morbidity | undesired result or complication |
| mortality | death |
| motility | the ability to move |
| mucosa, mucous | moist lining of digestive, respiratory, reproductive, and urinary |
| membrane | tracts |
| | sores in the mouth and esophagus, which may be painful and |
| mucositis/stomatitis | cause difficulty swallowing |
| myalgia | muscle aches |
| myocardial | pertaining to the heart muscle |
| myocardial infarction | heart attack |
| | tube placed in the nose, reaching to the stomach |
| nasogastric tube | nci the national cancer institute |
| nausea | feeling sick to the stomach |
| necrosis | death of tissue |
| neoplasia/neoplasm | tumor, may be benign or malignant |
| neuroblastoma | a cancer of nerve tissue |
| | a neurologic deficit is a decrease in the function of the brain, |
| neurologic deficits | spinal cord, muscles, and/or nerves |
| neurological | pertaining to the nervous system |
| | damage to the nerves which can cause numbness, pain, and |
| neuropathy | weakness |
| nouropatity | Wouldood |

| | condition in which the number of white bloods cells called |
|---------------------|---|
| neutropenia | neutrophils is abnormally low |
| noninvasive | not breaking, cutting, or entering the skin |
| nosocomial | acquired in the hospital |
| | the proposition, to be tested statistically, that the experimental |
| | intervention has "no effect," meaning that the treatment and |
| null hypothesis | control groups will not differ as a result of the intervention |
| occlusion | closing; blockage; obstruction |
| | a drug prescribed for conditions other than those approved by the |
| off-label use | fda. |
| | the study of tumors or cancer |
| oncology | an experimental design in which both the investigator(s) and the |
| | participants know the treatment group(s) to which participants are |
| onon dooign | |
| open design | assigned |
| open-ended | survey questions that allow respondents to answer in their own |
| questions | words |
| anan lahal trial | a clinical trial in which doctors and participants know which drug or |
| open-label trial | vaccine is being administered |
| ophthalmic | pertaining to the eye |
| onnortuniotio | an infection caused by an organism that usually does not cause |
| opportunistic | illness, but causes disease when a person's immune response |
| infections | (resistance) to infection is impaired |
| oral administration | by mouth |
| | an FDA category that refers to medications used to treat diseases |
| orphan drugs | and conditions that occur rarely |
| orthopedic | pertaining to the bones |
| ostealgia | bone pain |
| osteopetrosis | rare bone disorder characterized by dense bone |
| osteoporosis | softening of the bones |
| ovaries | female sex glands |
| palpitation | rapid heart beat parameter measure |
| pancreatitis / | |
| inflammation of the | |
| pancreas | inflammation of the pancreas causing pain in the upper abdomen |
| pancytopenia | abnormal decrease in the levels of all type of blood cells |
| parenteral | given by injection |
| | individuals whose physiological or behavioral characteristics and |
| participant | responses are the object of study in a research project |
| patency | condition of being open |
| | making decisions for others against or apart from their wishes with |
| paternalism | the intent of doing them good |
| pathogenesis | development of a disease or unhealthy condition |
| peer review | review of a clinical trial by experts chosen by the study sponsor |
| percutaneous | through the skin |
| peripheral | not central |
| peripheral blood | vein blood |
| | the study of the way the body absorbs, distributes, and gets rid of |
| pharmacokinetics | a drug |
| phenotype | the physical manifestation of a gene function |

| phlebitis | irritation or inflammation of the vein |
|---------------------|---|
| placebo | an inactive substance; a pill/liquid that contains no medicine |
| P.00000 | a physical or emotional change, occurring after a substance is |
| | taken or administered, that is not the result of any special property |
| placebo effect | of the substance |
| placenta | afterbirth |
| plasma | fluid found in the blood |
| platelet | part of blood that causes clots |
| pialelel | collection of fluid around the lungs in the chest cavity, which can |
| playing offusion | . |
| pleural effusion | cause shortness of breath and may require treatment |
| | the entire group (or get or type) of people from which a recorder |
| nonulation | the entire group (or set or type) of people from which a researcher |
| population | samples, and to which she or he would ideally like to generalize |
| potential | possible |
| | increase or multiply the effect of a drug or toxin (poison) by giving |
| | another drug or toxin at the same time (sometimes an |
| potentiate | unintentional result) |
| potentiator | an agent that helps another agent work better |
| | refers to the testing of experimental drugs in the test tube or in |
| | animals - the testing that occurs before trials in humans may be |
| preclinical | carried out |
| prenatal | before birth |
| | |
| | refers to trials to find better ways to prevent disease in people who |
| prevention trials | have never had the disease or to prevent a disease from returning |
| | a person's capacity to control the extent, timing, and |
| | circumstances of shared oneself (physically, behaviorally, or |
| privacy | intellectually) with others |
| | includes information about behavior that occurs in a context in |
| | which an individual can reasonably expect that no observation or |
| | recording is taking place, and information which has been |
| | provided for specific purposes by an individual and which the |
| private information | individual can reasonably expect will not be made public |
| | a subset of the population chosen in such a way that every |
| | member of the population has a known (nonzero) chance of being |
| probability sample | selected into the sample |
| prognosis | outlook, probable outcomes |
| prone | lying on the stomach |
| prophylaxis | use of drugs to prevent disease |
| | |
| | studies designed to observe outcomes or events that occur after |
| | the group of participants has been identified. prospective studies |
| | do not have to involve manipulation or intervention but may be |
| prospective studies | purely observational or involve only the collection of data instead. |
| prosthesis | artificial part, most often limbs, such as arms or legs |

| [| la succession de la succe |
|-------------------------------|--|
| | name, address, elements of dates related to an individual (e.g., |
| | birthdate), email address, numbers; telephone, fax, social |
| | security, medical record, health beneficiary/health insurance, |
| | certificate or license numbers, vehicle, account numbers, |
| | characteristics, or codes (e.g., global positioning system (gps) |
| | readings), web urls, internet protocol (io) addresses, biometric |
| protected health | identifiers (e.g. voice, fingerprints), full face photographs or |
| information | comparable images |
| proteinuria | excess protein in the urine |
| protocol | a study plan on which all clinical trials are based |
| proximal | closer to the center of the body, away from the end |
| pruritis | itchy skin |
| psychosis | nervous breakdown |
| pulmonary | pertaining to the lungs |
| · · · | a blood clot that causes a sudden blockage in a lung artery, |
| pulmonary embolism | usually due to a blood clot that traveled to the lung from the leg |
| | tissue in the lungs becomes stiff making breathing difficult, |
| | resulting in shortness of breath, and if severe, can cause heart |
| pulmonary fibrosis | failure |
| pulmonary | abnormally high blood pressure in the blood vessels in the lungs, |
| hypertension | which makes it harder to pump blood into the lungs |
| | an experimental design that is missing one or more aspects of the |
| quasi-experiment | (classic) controlled experiment. |
| radiation therapy | x-ray or cobalt treatment |
| random | by chance (like the flip of a coin) |
| | a method based on chance by which study participants are |
| randomization | |
| Tanuomization | assigned to a treatment group |
| Devine evidie Ovine dine rece | an autoimmune disorder causing blood vessels to spasm when |
| Raynaud's Syndrome | |
| recombinant | formation of new combinations of genes |
| reconstitution | putting back together the original parts or elements |
| | the period during which a trial is attempting to identify and enroll |
| recruiting | participants |
| recruitment status | indicates the current stage of a trial |
| recur | happen again |
| refractory | not responding to treatment |
| regeneration | re-growth of a structure or of lost tissue |
| regimen | pattern of giving treatment |
| relapse | the return of a disease |
| reliability | the degree to which a measure yields consistent results |
| remission | disappearance of evidence of cancer or other disease |
| | payment for participation in research; this is different from |
| | compensation, which typically refers to payment for research- |
| remuneration | related injuries |
| renal | pertaining to the kidneys |
| replicable | possible to duplicate |
| | a sample in which the participants closely match the |
| representative | characteristics of the population, and thus, all segments of the |
| sample | population are represented in the sample |
| | |

| a systematic investigation (i.e., the gathering and analysis information) designed to develop or contribute to generalize the sect research knowledge resect remove or cut out surgically difficulty breathing with low levels of oxygen in the blood, we could be parisus and life threatening and require you to be | |
|---|-----------|
| research knowledge resect remove or cut out surgically difficulty breathing with low levels of oxygen in the blood, with low levels of oxygen in the blood. | Zable |
| resect remove or cut out surgically difficulty breathing with low levels of oxygen in the blood, | |
| difficulty breathing with low levels of oxygen in the blood, | |
| | |
| | |
| could be serious and life threatening and require you to ha | |
| tube inserted into your windpipe that is hooked up to a ma | achine to |
| respiratory failure help you breathe | |
| research participants, who fill out a survey, are interviewe | |
| participate in an experiment, are observed in a naturalistic | setting, |
| respondents or who are otherwise studied | |
| rhabdomyolysis rhabdomyolysis is a breakdown of muscle fibers. | |
| rigors chills and shivering | |
| saline salt water solution | |
| sample a subset of a given population used for research purposes | S |
| sarcoma a type of cancer | |
| screening examination, test | |
| secretion release | |
| sedative a drug to calm or make less anxious | |
| seizures convulsions | |
| seminoma a type of testicular cancer (found in the male sex glands) | |
| sequentially in a row, in order | |
| side effects any undesired actions or effects of a drug or treatment | |
| simultaneous at the same time | |
| typically, a study design in which the investigator, but not | the |
| participant, knows the identity of the treatment assignment | nt. |
| occasionally the participant, but not the investigator, know | /s the |
| single-blind design assignment. also known as single-masked design | |
| a non-probability sample that is created by using member | s of the |
| group of interest to identify other members of the group (f | or |
| example, asking a participant at the end of an interview for | or |
| snowball sample suggestions about who else to interview) | |
| social systematic manipulation of, or experimentation in, social of | or |
| experimentation economic systems; used in planning public policy | |
| somnolence sleepiness | |
| a sample, as of human tissue, blood or urine, used for dia | Ignostic |
| specimen or pathological analyses | |
| an instrument to measure the amount of air taken into and | b |
| spirometer exhaled from the lungs | |
| staging an evaluation of the extent of the disease | |
| a treatment plan that the majority of the medical communi | ity would |
| standard of care accept as appropriate | |
| a treatment currently in wide use and approved by the FD | A, |
| considered to be effective in the treatment of a specific dis | sease or |
| | |

| | the probability that an event or difference occurred by chance |
|---|--|
| | alone. In clinical trials, the level of statistical significance depends |
| statistical | on the number of participants studied and the observations made, |
| significance | as well as the magnitude of differences observed |
| stenosis | narrowing of a duct, tube, or one of the blood vessels in the heart |
| Stevens-Johnson | skin condition that causes painful blisters and sores of the skin |
| syndrome | and mucous membranes, especially in the mouth |
| stimuli | something which causes a change |
| stomatitis | mouth sores, inflammation of the mouth |
| | arrange in groups for analysis of results (e.g., stratify by age, sex, |
| stratify | etc.) |
| , | a data collection method in which an interviewer reads a |
| | standardized interview schedule to the respondent and records |
| structured interview | the answers |
| | a primary or secondary outcome used to judge the effectiveness |
| study endpoint | of a treatment |
| | stunned state in which it is difficult to get a response or the |
| stupor | attention of the subject |
| subclavian | under the collarbone |
| subcutaneous | under the skin |
| supine | lying on the back |
| supine position | lying on the back |
| supplement | add |
| | general medical care aimed at symptoms, not intended to improve |
| supportive care | or cure underlying disease |
| | a study in which the same data are collected from all members of |
| | the sample using a highly structured questionnaire and analyzed |
| survey | using statistical tests |
| syndrome | a condition characterized by a set of symptoms |
| | top number in blood pressure; pressure during active contraction |
| systolic | of the heart |
| T-lymphocytes | type of white blood cells |
| tachycardia | fast heart rate |
| ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,, | capable of causing malformations in a fetus (developing baby still |
| teratogenic | inside the mother's body) |
| testes/testicles | male sex glands |
| | a general explanation about a specific behavior or set of events |
| | that is based on known principles and serves to organize related |
| theory | events in a meaningful way |
| | |
| therapy | treatment intended and expected to alleviate a disease or disorder |
| | Low number of platelets, which may cause bleeding and bruising. |
| | May require a blood transfusion. Bleeding may be serious or life |
| thrombocytopenia | threatening. |
| thrombosis | clotting |
| thrombus | blood clot |
| tinnitus | ringing in the ears |
| | |

| | a method for deciding on the strength of a drug or solution; |
|-----------------------|--|
| titration | gradually increasing the dose |
| topical | on the surface |
| | applied to a certain area of the skin and reducing pain only in the |
| topical anesthetic | area to which applied |
| toxicity | side effects or undesirable effects of a drug or treatment |
| transdermal | through the skin |
| transiently | temporarily |
| trauma | injury; wound |
| | refers to trials which test new treatments, new combinations of |
| treatment trials | drugs, or new approaches to surgery or radiation therapy |
| | an unanticipated problem involving risk to human participants or |
| | others, is one that (1) was unforeseen at the time of its |
| Unanticipated | occurrence, and (2) indicates that participants or others are at an |
| Problem | increased risk of harm |
| uptake | absorbing and taking in of a substance by living tissue |
| urticaria | hives |
| | the degree to which a measure assesses what we think it is |
| validity | assessing |
| valvuloplasty | plastic repair of a valve, especially a heart valve |
| | any characteristic or trait that can vary from one person to another |
| | (race, sex, academic major) or for one person over time (age, |
| variable | political beliefs) |
| varices | enlarged veins |
| vasospasm | narrowing of the blood vessels |
| | a carrier that can transmit disease-causing microorganisms |
| vector | (germs and viruses) |
| venipuncture | needle stick, blood draw, entering the skin with a needle |
| vertical transmission | spread of disease |
| visual disturbances | inability to see properly. |
| | free of coercion, duress, or undue inducement. Used in the |
| | research context to refer to a subject's decision to participate (or |
| voluntary | to continue to participate) in a research activity |
| | persons who are wards of the state or any other agency, |
| ward | institution, or entity |