Glossary of Research Terms

**Aggregate Data** is data not limited to one participant, which is tracked across time, organizations, populations, or other variables. (Source: National Center for Biotechnology Information)

**Asset Mapping** is a visual representation of a location’s resourceful sites and services to its community. Asset mapping allows one to identify a community’s unmet needs for a certain resource, which is useful in determining how to build upon already existing assets or creation of new ones. Asset mapping promotes community empowerment, partnership, and involvement. (Source: UCLA Center for Health Policy Research)

**Cohort** is a set of people studied over a period of time. People within a cohort embody certain shared characteristics and are examined for investigation on a certain outcome like disease, death, etc. (Source: National Center for Biotechnology Information)

**Cohort Study/Panel Study** is a type of epidemiological study in which sets of a certain population are identified to have, at one point, been exposed or unexposed to a certain factor that is hypothesized to influence the occurrence of a certain outcome. (Source: UCLA School of Public Health)

**Datasets** are a collection of data that is organized for studying. Most commonly, a single database table with a single statistical data matrix is used. (Source: Wikipedia)

**Demographics** are the statistical characteristics or qualities (such as age, sex, income, etc.) of populations used as identifiers. (Source: Wikipedia)

**Environmental Scanning** is a tool for obtaining and organizing data of an organization that allows the study for certain events and trends; environmental scanning helps plan the initiation and future of a project, provide evidence of the directionality of an organization, or raise awareness of issues. (Source: National Center for Biotechnology Information)

**Federal Poverty** is a certain income level measured annually by the Department of Health and Human Services that is used to determine one’s eligibility for certain programs and benefits; a reference to the Census Bureau poverty thresholds that are characterized by family size, number of children, or whether or not elderly. Such thresholds are used for calculating the number of people in poverty. (Sources: U.S. Centers for Medicare & Medicaid Services, U.S. Department of Health & Human Services)

**Geocoding** is an assigned code that associates an individual to a geographic location. (Source: UCLA Clinical and Translational Science Institute)

**Informed Consent** is a process in which an individual gives consent to a treatment or procedure after being informed of the risk and benefits of a procedure or treatment, and its alternatives.
Informed consent could also involve the participant’s voluntary release of confidential information after being informed of the purpose of the information’s disclosure. (Source: UCLA Center for Mental Health in Schools)

**Institutional review board (IRB)** is a group that reviews and monitors biomedical research involving human subjects. IRB has the authority to approve or disapprove research. IRB ensures that the appropriate steps are taken to protect the rights and welfare of the research subjects. IRBs use a group process in its review of research protocols or related materials. (Source: U.S. Food and Drug Administration)

**Longitudinal Study** is a study of a certain outcome that is measured on a cohort repeatedly across a set time period. Time may be measured in minutes, days, or years. (Source: UCLA School of Public Health)

**Meta-analysis** is a process in which research across individual studies is synthesized using a formal scientific process; a combination of evidence across various studies done using an organized, formal method. (Source: UCLA Department of Statistics)

**Needs Assessment** is a systematic set of procedures that determine an entity’s needs, examine its nature and causes of need, and set priorities for future action to appease such needs. (Source: U.S. Department of Education)

**Poor birth outcome** is the risk of mortality and a variety of health and developmental problems for an infant. A birth outcome is considered poor if the infant’s health risk is at a higher percentage than expected or normal. Length of gestation and birth weight and are two common factors in considering whether an infant’s birth outcome is poor. An infant is considered preterm if born before 37 completed gestation weeks. Birth weight is determined by length of gestation as well as fetal growth. Infants may be born with low birth weight if born prematurely. (Source: Nurse Family Partnership)

**Qualitative Study** is the method of research expressed using words and concepts rather than numerical expression. Qualitative data is often used to acquire an understanding of a particular event, and answer the “how” or “why” of this event. (Source: UCLA Center for Health Policy Research)

**Quantitative Study** is the method of research expressed using numbers and percentages rather than words. Quantitative data is often used to answer the “who, what, when, and where” questions of an assessment. (Source: UCLA Center for Health Policy Research)

**Query** is request or question for information concerning a set of data. (Source: Merriam-Webster Dictionary)
**Institutional Repository** is an organizational service in which digital materials created by members of an institution are preserved for members of the organization to access or distribute. (Source: [Association of Research Libraries](https://www.arl.org))

**Risk-assessment** is the assessment of the probability of harm in a physical, psychological, social, or economic sense that occurs as a result of participating in a research study. Assessment involves determining the magnitude and significance of such harm. (Source: [UCLA Office of Human Research Protection Program](https://ohrpp.ucla.edu))

**Statistical significance** is a mathematical technique used in the field of statistics that measures whether certain results are likely to be true. Statistical significance is calculated as the probability that something is occurring due to chance, which is expressed as a P-value. A P-value less than 0.05 (p<0.05) means that a result is unlikely to be due to chance, and is thus likely to be true. (Source: [U.S. Department of Health and Human Services](https://www.hhs.gov))

**Surveillance** is the collection of data that is used for health planning and promotion, and disease control and prevention. Surveillance is used to monitor changes for health factors and detect the onset of outbreaks. (Source: [UCLA School of Public Health](https://www.ucla.edu))

**Survey Instrument** is a tool for implementing scientific methods and protocol consistently for obtaining data from respondents. (Source: [Office of Behavioral and Social Sciences Research](https://obssr.od.nih.gov))
Sources

Association of Research Libraries
Merriam-Webster Dictionary
National Center for Biotechnology Information
Nurse Family Partnership
Office of Behavioral and Social Sciences Research
UCLA Center for Health Policy Research
UCLA Center for Mental Health in Schools
UCLA Clinical and Translational Science Institute
UCLA Department of Statistics
UCLA Office of Human Research Protection Program
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