

The National Coordinating Center for the Regional Genetics Networks

Glossaries of Genetics Terms

English – Add Language of Translation

2023 Edition

Compiled and edited by: Cynthia E. Roat, MPH

In collaboration with:
The Cross Cultural Health Care
Program (CCHCP)

Glossaries of Genetics Terms - Language

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Acknowledgments

Mission of the NCC

The National Coordinating Center for the Regional Genetics Networks (NCC) has been funded by the Health Resources and Services Administration (HRSA) since 2004. Our mission is to improve access to genetics for underserved populations. This mission is accomplished through different programs, many of which are educational tools and resources for non-genetics professionals. Learn more at https://nccrcg.org.

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Cindy Roat is an international consultant in language access in health care and patient navigation. She provides training and consulting for health care interpreters, providers, administrators, and patient navigators. Cindy compiled and edited this glossary for the NCC.

The Cross Cultural Health Care Program

The mission of The Cross Cultural Health Care Program (CCHCP) is to serve as a bridge between communities and health care institutions to advance access to quality health care that is culturally and linguistically appropriate. CCHCP provides resources, such as bilingual medical glossaries, as well as medical interpreter, patient navigator, and cultural competence training for individuals and institutions with the goal of systems change. Learn more at: https://xculture.org

CCHCP is supporting and coordinating the translation process of this glossary and is providing the glossary free of charge in its online store: https://xculture.org/store/

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The National Coordinating Center for the Regional Genetics Networks

Glossary of Autism Spectrum Disorder (ASD) Genetics Terms

English – Add Language of Translation

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Program (CCHCP)



Preface

The seven Regional Genetics Networks (RGNs), the National Genetics Education and Family Support Center (Family Center), and the National Coordinating Center for the Regional Genetics Networks (NCC) mission is to improve access to quality genetic services for medically underserved populations. This bilingual glossary of terms related to Autism Spectrum Disorders (ASD) is provided to help enable access to ASD medical services to limited English proficiency populations.

How to Use the Glossary

This glossary is organized alphabetically in English. The first column shows the term in English. The second column has the definition of the term in English. The third column has the translation of the term.

At the end of the translations is a list of references used for the definitions and places where you can find more information.

Glossary of Terms

English – Language of Translation

(n) – noun (v) – verb (adj) – adjective (adv) – adverb

Term	Definition	Translation
A-B-C analysis (n)	An approach to understanding behavior by examining the Antecedent (the cause), the Behavior, and the Consequence (the result). ¹	
Abstract language (n)	Any vocabulary or phrases with meanings that are not clearly stated. Examples include idioms like, "it's raining cats and dogs" or "break a leg" where the intended meaning does not match the literal meaning. There are also many examples of this in books and poetry where the author may describe something using abstract language like "it felt like a weight on my chest."	
Adaptive behavior or adaptive skills (n)	Includes communication, self-care, home living, social skills, community use, self-direction, health and safety, functional academics, leisure, and work. These are skills that help the person be successful in their environment and are learned skills rather than innate abilities. ²	
Age of majority (n)	The age established under state law when an individual is no longer a minor and has the right to make certain legal decisions without consent. ¹	
Anxiety disorder (n)	A pattern of constant worry or tension under many different circumstances. ¹	
Applied Behavior Analysis (ABA) (n)	ABA is the name of a professional field that uses principles of learning to increase performance of socially desirable behaviors. ABA practitioners carefully observe and measure behaviors and the context in which they occur in order to individualize teaching plans to improve specific behaviors. ABA is commonly used as one component in interventions for Autism Spectrum Disorders. ²	
Apraxia (n)	See "Dyspraxia." ²	

Term	Definition	Translation
Attention Deficit Hyperactivity Disorder (ADHD) (n)	A disorder that shows up in the areas of inattention, hyperactivity, and impulsiveness. It is evidenced by frequent shifting from one activity or focus to another, having difficulty organizing and completing tasks correctly, impulsive response, or failure to follow rules. It may occur with or without hyperactivity which includes behaviors such as excessive running, talking, fidenting, and/or rectlessness. ²	
Audiologist (n)	fidgeting, and/or restlessness. ² A specialist who evaluates for hearing loss as a potential cause or contributor to developmental delay, and designs/supports interventions to minimize the impact of hearing loss when it is found. ²	
Augmentative and Alternative Communication (AAC) (n)	Any method of communicating without speech, such as by signs, gestures, picture boards, or electronic or non-electronic devices. These methods can help individuals who are unable to use speech or who need to supplement their speech to communicate effectively. ²	
Autism Spectrum Disorder (ASD) (n)	A neurodevelopmental disorder. ASD symptoms are typically evident before a child is 3 years of age. The symptoms range from mild to severe – and individuals often have varied skills levels in different domains of functioning. Autism affects the person's overall development in 2 primary areas: 1. social communication, or the way a person uses gestures, body language, and language to communicate and interact socially and relate with others. 2. the presence of restricted, repetitive patterns of behavior, interests, or activities (e.g., repetitive motor movements, echolalia/repeating speech, idiosyncratic phrases, extreme distress at small changes/difficulty with transitions, strong attachment to unusual objects/topics, adverse or intense responses to sensory input). ²	
Aversive (adj)	An unwanted stimulus designed to change an individual's behavior through punishment. These should rarely be used in the treatment of autism. 1	
Behavior Intervention Plan (BIP) (n)	A plan to improve an individual's behavior, created based on the results of a Functional Behavior Assessment. ¹	
Behavioral disorder (n)	A condition in which behavior significantly deviates from acceptable norms. ¹	
Biomarker (n)	An indicator of a certain biological state. ¹	
Bipolar disorder (n)	A brain disorder that causes unusual shifts in mood, energy, activity levels, and the ability to carry out day-to-day tasks; also known as manic-depressive illness. ¹	
Board Certified Behavior Analyst (BCBA) (n)	A professional certified to provide ABA therapy by the Behavior Analyst Certification Board (BACB). ¹	

Term	Definition	Translation
Body language (n)	A form of non-verbal communication that is an important part of social communication. It includes things like gestures (head nodding or shaking), personal space, facial expressions, and eye contact. ⁵	
Case manager (n)	A professional from a school or service agency such as the Department of Developmental Disabilities who serves as a direct contact for families and helps gather resources, team members and ideas. ¹	
Catatonia (n)	A state in which a person does not move and does not respond to others. ¹	
Challenging behaviors (n)	Behaviors that are destructive and harmful to the individual or others (e.g., hitting, biting, running away), that prevent learning and cause others to label or isolate the individual for being odd or different. ¹	
Civil commitment (n)	A legal process in which an individual experiencing a mental health crisis is ordered into treatment against his or her will, including to a hospital. 1	
Cognitive behavioral therapy (n)	Type of therapy designed to help improve an individual's inappropriate or challenging behaviors by replacing the negative thoughts that cause these behaviors with positive thoughts. ¹	
Cognitive development (n)	How children learn to think, interpret information, make decisions, and solve problems. Areas of cognitive development include verbal reasoning (using language), non-verbal reasoning (using visual/spatial information), as well as executive control/functioning. ²	
Communication (n)	The developmental area that involves skills which enable people to understand (receptive communication) and share (expressive communication) thoughts and feelings. Waving goodbye, smiling, nodding, making eye-contact, using spoken language, following directions, and reading and writing are examples of communication. ²	
Communication disorder (n)	Difficulty with understanding and/or expressing messages. ² Communication disorders include problems with hearing, with making speech sounds (articulation), with having a clear voice (voice disorders), stuttering (fluency disorders), difficulty learning, knowing, and using grammar (language disorders), difficulty using language to get things done (social communication or pragmatic language disorders), and using language to learn (language-based learning disabilities such as dyslexia). ²	
Comorbid (adj)	Pertaining to a disease or disorder that occurs simultaneously with another. ¹	

Term	Definition	Translation
Compulsion (n)	The drive to do something in particular or in a particular way, such as the need to straighten all the forks at the dinner table. ¹	
Conservatorship (n)	The legal right given to a person to be responsible for the assets and finances of a person deemed fully or partially incapable of providing these necessities for himself or herself. ¹	
Crisis plan (n)	A document that outlines in specific detail the necessary strategies and steps that must be taken when a crisis occurs. ¹	
Data analysis (n)	The process of thoroughly inspecting information related to challenging behaviors in order to draw out useful information and conclusions that may result in strategies to improve behavior. ¹	
De-escalation (n)	The process of stopping a challenging behavior or crisis from intensifying and calming the situation. ¹	
Depression (n)	A mood disorder in which feelings of sadness, anger, or frustration interfere with everyday life for an extended period of time. ¹	
Developmental assessment (n)	A structured evaluation of a child's development in one or more areas including cognitive, language, motor, social/emotional, and adaptive. Professionals that conduct developmental assessments include developmental behavioral pediatricians, psychologists, speech language pathologists, occupational therapists, physical therapists, educators, and audiologists. Types of tests that are used during developmental assessments include interview, observation, questionnaires, standardized tests, and non-standardized tests. ²	
Developmental delay (n)	The term used to describe the condition of an infant or young child who is not achieving new skills in the typical time frame and/or is exhibiting behaviors that are not appropriate for his or her age. Some children who are developmentally delayed eventually have a specific diagnosis of a particular developmental disability. Other children with delays catch up with their typically developing peers. ²	
Developmental Disability (DD) (n)	A severe chronic disability that is attributed to a physical or mental impairment, other than the sole diagnosis of mental illness, or to a combination of mental and physical impairments, is manifested before the individual attains the age of 22, is likely to continue indefinitely, results in the inability to live independently without external support or continuing and regular assistance, reflects the need for a combination and sequence of special, interdisciplinary, or generic care, treatment, or other services that are planned and coordinated for that individual. ²	

Term	Definition	Translation
Developmental	Also known as a developmental-behavioral	
pediatrician (n)	pediatrician, these professionals have training and	
	experience to assess and treat a wide range of medical	
	and psychosocial aspects of a child's developmental	
	and behavioral difficulties. Their expertise may make	
	them a good choice for children with complicated	
	medical or developmental problems. ²	
Differential diagnosis (n)	Distinguishing between two or more diseases with	
	similar symptoms to identify which is causing	
Diamentian (n)	distress or challenging behavior. ¹	
Disruption (n)	An event that causes an unplanned deviation from a situation. ¹	
Down syndromo		
Down syndrome, also known as trisomy 21	The most common and readily identifiable chromosomal condition associated with intellectual	
(n)	disability. Children with Down syndrome typically have	
(")	developmental delays, but this can range from mild to	
	severe. Common physical signs of Down syndrome	
	include decreased muscle tone; short neck; flattened	
	facial profile and nose; small head, ears, and mouth;	
	upward slanting eyes; white spots on the colored part	
	of the eye (called Brushfield spots); wide, short hands	
	with short fingers; a single, deep, crease across the	
	palm of the hand; a deep groove between the first and	
	second toes. ²	
Dual diagnosis (n)	The identification of an additional mental health	
	disorder individuals with developmental disabilities. ¹	
Dysarthria (n)	A term used to describe the impact on speech	
	production of muscle weakness and/or reduced	
	muscle control due to neural damage. ²	
Dyspraxia (n)	A condition characterized by a difficulty with planning	
	and performing coordinated movements although	
	there is no apparent damage to muscles. Dyspraxia	
	can impact any motor system and will be described	
	based on the motor system affected (e.g., upper limb dyspraxia, speech/verbal dyspraxia). The term	
	"apraxia" is often used as a synonym for dyspraxia. ²	
Early intervention (n)	Programs or services designed to meet the	<u> </u>
	developmental needs of infants and toddlers (birth to	
	three years old) and their families.	
Echolalia or Echophrasia	The immediate and involuntary repetition of words or	
(n)	phrases just spoken by others, often a symptom of	
	autism or some types of schizophrenia. Also called	
	echophrasia. ³	
Elopement (n)	A situation in which an individual leaves a safe	
	place, a caretaker, or supervised situation, either	
	by "bolting," wandering or sneaking away.1	
Epilepsy (n)	A brain disorder in which a person has repeated	
	seizures (episodes of disturbed brain activity or	
	convulsions) over time. ¹	
Escalating (v)	Increasing or worsening rapidly.1	<u>I</u>

Term	Definition	Translation
Evidence-Based Practice	A teaching strategy or practice that has been proven	
(EBP) (n)	through research studies to improve skills or behaviors	
	for a certain population. ⁵	
Executive control /	A group of skills that helps people plan, organize,	
executive functioning (n)	control behavior, focus on multiple streams of	
	information at the same time, self-monitor, and revise	
	action plans as necessary. Acquiring these skills is one	
	of the most important tasks of the early childhood	
5 ()	years. ²	
Extinction (n)	A response used to eliminate a behavior that	
	involves ignoring a mild behavior when it is used for	
Fusion stiens brough (m)	attention. ¹	
Extinction burst (n)	Short term response to extinction in which there is a	
	sudden and temporary increase in the response's frequency, followed by an eventual decline. ¹	
Face blindness (n)	An impairment in the recognition of faces. ¹	
Fading (v)	Gradually reducing the number of prompts or types of	
rading (v)	prompts to encourage more independence for the	
	learner. ⁵	
Fecal digging (v)	The process in which an individual puts his fingers into	
	his rectum. ¹	
Fecal smearing (v)	The process in which feces are spread on property	
	or the individual himself.1	
Fetal Alcohol Spectrum	Children whose mothers drank during pregnancy, and	
Disorder (FASD) (n)	who have developmental impairment may be	
	diagnosed with one of several FASDs. The developing	
	brain is the organ most vulnerable to prenatal alcohol	
	exposure, but the range of impact of prenatal alcohol	
	exposure includes both physical and	
	neurodevelopmental impairment. These can include	
	growth difficulty, minor facial anomalies, general	
	developmental delays, seizures, learning disabilities,	
	ADHD, executive functioning and processing	
	challenges, as well as mental health or behavioral	
	challenges. The most severe FASD is Fetal Alcohol Syndrome (FAS – see FAS), but other FASDs are far	
	more common. ²	
Fetal Alcohol Syndrome	A permanent birth defect syndrome caused by	
(FAS) (n)	maternal drinking during pregnancy. FAS is	
(. 73) (11)	characterized by growth deficiency; a cluster of 3	
	minor facial abnormalities including a thin upper lip, a	
	smooth philtrum (i.e., the groove between nose and	
	upper lip); small eyes; and significant abnormalities in	
	brain development. FAS is the leading preventable	
	cause of intellectual disability and other	
	neurodevelopmental conditions. ²	
Function of behavior (n)	The purpose or reason behind a specific behavior for	
	an individual. ¹	

Term	Definition	Translation
Functional Behavior	The process of systematically determining the function	
Analysis (FBA) (n)	of behaviors, usually inappropriate, that are displayed	
	by people. Behaviors are defined, measured, and	
	analyzed in terms of what happened before and after	
	their occurrence. Over time the events before and	
	after the behavior occurs are systematically changed	
	in order to determine the function of the behavior for	
	the person displaying it. ²	
Functional	Effective and appropriate communication that an	
communication (n)	individual uses across his daily activities to meet his or	
	her needs.¹Can be verbal or non-verbal.	
Generalization (n)	The ability to use a target skill or behavior across the	
	intervention timespan or timeframe, setting, and	
	individuals (e.g., teachers, peers, parents). For	
	example, when teaching a child to greet others, a	
	teacher may initially implement the intervention when	
	the student first enters classroom. Although the child	
	may begin to independently greet others in the	
	context of entering the classroom, she may not be	
	able to do so when coming home from or entering a	
	different room in the school. If she can greet others	
	across settings and individuals, then she has engaged	
0 1 ()	in generalization of the skill. ⁵	
Gestures (n)	Body and hand movements used to communicate.	
	Examples include pointing, waving, opening arms to	
	demonstrate something is "big," banging a fist on a	
Cuardianchin (n)	table to emphasize a point or show anger, etc. ⁵	
Guardianship (n)	The legal right given to a person to be responsible for the food, health care, housing, and other	
	necessities of a person deemed fully or partially	
	incapable of providing these necessities for himself	
	or herself. ¹	
Hypersensitivity (n)	Overly intense or exaggerated response to sensation.	
Trypersensitivity (II)	It may include defensive responses (like covering ears	
	in anticipation of a sound, pulling away quickly when	
	touched lightly) or disliking or avoiding of certain	
	situations that involve an undesirable sensory	
	experience (like avoiding public restrooms because the	
	toilet flushing is aversive). ⁵	
Hyposensitivity (n)	Under exaggerated response to sensation. May	
,,,,	include lack of response entirely or a reduced or	
	slower response than expected. A child with	
	hyposensitivity may not stop working and turn around	
	when their name is called, may not seem to notice	
	that another child ran into them, may smile, or laugh	
	several seconds after you tickle them, may cry several	
	seconds after getting hurt or not at all, etc. ⁵	

Term	Definition	Translation
Idiosyncratic language/	Using language in unusual ways that may not be	
vocabulary (n)	obvious to the communication partner. This includes	
	using scripts from previous conversations or media	
	applied to different contexts like saying "C is for	
	cookie" to express that they like something. Or "Dora	
	loves waterfall!" to request to take a bath/shower.5	
Incontinence (n)	The (usually) involuntary passing of feces or urine,	
	generally not into a toilet or diaper. ¹	
Individualized Education	A written statement of a child's current level of	
Program (IEP) (n)	development (abilities and impairments) and an	
	individualized plan of instruction, including the goals,	
	the specific services to be received, the people who	
	will carry out the services, the standards, and	
	timelines for evaluating progress, and the amount and	
	degree to which the child will participate with non-	
	handicapped peers at school. The IEP is developed by	
	the child's parents and the professionals who	
	evaluated the child. It is required by the Individuals	
	with Disabilities Education Act (IDEA) for all children in	
	special education, ages three years and up. ²	
Individualized Family	A written plan describing the infant's or toddler's	
Service Plan (IFSP) (n)	current level of development, the family's strengths	
	and needs related to enhancement of the infant's or	
	toddler's development, goals for the child and the	
	other family members (as applicable), including the	
	criteria, procedures and time lines used to evaluate	
	progress (the IFSP should be evaluated and adjusted at	
	least once a year and reviewed at least every six months), and the specific early intervention services	
	needed to meet the goals (including the frequency and intensity and method of delivering services, the	
	projected date of initiating services and the	
	anticipated duration of services). The IFSP is	
	developed and implemented by the child's parents	
	and a multidisciplinary early intervention team (IFSP	
	Team). The Individualized Family Service Plan is	
	required by the Individuals with Disabilities Education	
	Act (IDEA) for all infants and toddlers receiving early	
	intervention services. ²	
Individuals with	The federal law reauthorized in 2004 that amends the	
Disabilities Education Act	Education for All Handicapped Children Act (Public	
(IDEA) (n)	Law 94-142). Part C of the law focuses on services to	
7	infants and toddlers who are at-risk or have	
	developmental disabilities. ²	

Term	Definition	Translation
Intellectual disability (n)	Characterized by significantly impaired intellectual	
	functioning, existing concurrently with related	
	limitations in two or more of the following applicable	
	adaptive skill areas: communication, self-care, home	
	living, social skills, community use, self-direction,	
	health and safety, functional academics, leisure, and	
	work. ²	
Interdisciplinary	A group of health care professionals from diverse	
evaluation team (n)	disciplines who form a team to work collaboratively in	
	conducting a cohesive patient evaluation. ²	
Intervention (n)	A strategy or process put in place in order to improve	
	or modify an individual's behavior e.g., medication,	
In the land of the	Applied Behavior Analysis).1	
Involuntary commitment (n)	A legal process in which an individual experiencing a mental health crisis is ordered into treatment	
(11)	against his or her will, including to a hospital. ¹	
Joint attention (n)	A set of early social communication skills used to show	
Joint attention (ii)	and/or share interest about an outside object or event	
	with a communication partner. They include initiating	
	joint attention where the child shifts their gaze and/or	
	gestures toward an object with the goal of getting	
	another person to notice it. They also include	
	responding to gaze shifts and gestures from other	
	people so that they can learn what others find	
	interesting. ⁵	
Language delay (n)	A disorder in which a learner's ability to understand	
	and/or use language is behind what would be	
	expected based on their age. ⁵	
Language disorder (n)	In children, this could mean trouble getting their	
	meaning or message across to others (expressive	
	language disorder), or understanding messages	
	coming from others (receptive language disorder), as	
	well as difficulty using language to get things done	
	(pragmatic language/social communication disorder). ²	
Learning disability (n)	A disorder that affects how a person learns and	
	understands primary skills such as reading, writing and math. ²	
Least Restrictive	The educational setting that permits a child with	
Environment (LRE) (n)	disabilities to derive the most educational benefit	
Environment (ERE) (II)	while participating in a regular educational	
	environment to the maximum extent appropriate. It is	
	presumed that a child with a disability will be	
	educated in the general education classroom, with	
	appropriate supports, unless the IEP Team deems	
	another setting as more appropriate. LRE is a	
	requirement under the IDEA. ²	
Maladaptive behavior (n)	A type of behavior that is often used to reduce	
, , ,	anxiety, but whose result does not provide	
	adequate or appropriate adjustment to the	
	environment or situation. ¹	

Term	Definition	Translation
Medicaid (n)	A government program that provides healthcare	
	coverage for low-income families and individuals	
	with disabilities in the United States. ¹	
Medical home (n)	A team-based healthcare delivery model led by a	
	physician that provides comprehensive and	
	continuous medical care to patients. ¹	
Mental health hold (n)	Involuntary hospitalization due to a mental health crisis. ¹	
Motor skill (n)	The learned ability to perform movements, such as	
WIOLOI SKIII (II)	holding the body upright to sit, using the hands to	
	manipulate small items, scooping food onto a spoon	
	and bringing it to the mouth, and moving the lips and	
	tongue to articulate different sounds. Fine motor skills	
	involve use of the small muscles of the body while	
	gross motor skills are associated with large muscle	
	movements. ²	
Neurodevelopmental	Problems affecting the development of the central	
conditions (n)	nervous system or brain, resulting in delayed or	
	unusual motor, speech, social or learning deficits.	
Nonverbal	Any form of or attempt at unspoken or "physical"	
communication (n)	communication. Examples are temper tantrums,	
	gestures, pointing and leading another person to a	
	desired object. ²	
Nutritionist (n)	Nutrition services are provided by Registered Dietitian	
	Nutritionists (RDNs). The nutritionist evaluates the	
	child's growth, energy intake, and nutritional status to	
	make sure they are getting appropriate nutrients from	
	their diet. A nutritionist often will work with a feeding	
	therapist (e.g., speech or occupational therapist), to	
	develop a plan when feeding problems are suspected. ²	
Obsession (n)	A repetitive thought or feeling dominated by a	
	particular idea, image, or desire, such as a person who	
	only wants to talk about elevators.1	
Obsessive Compulsive	An anxiety disorder in which people have unwanted	
Disorder (OCD) (n)	and repeated thoughts, feelings, ideas, or	
	sensations (obsessions) that make them feel driven	
<u> </u>	to do something (compulsions).1	
Occupational Therapist	A skilled healthcare provider that assists people across	
(OT) (n)	the lifespan participate in the things they want and	
	need to do through the therapeutic use of everyday	
	activities (occupations). Occupational therapy benefits	
	individuals who have physical, developmental, or cognitive limitations. With children, treatment is	
	geared toward development of fine motor skills (e.g.,	
	writing, cutting), gross motor skills (e.g., climbing	
	stairs, jumping), self-care (e.g., dressing, eating), and	
	play. OT is the primary profession that addresses	
	sensory processing differences. ²	
Ototoxic (n)	Damaging to the ears, causing sound sensitivities,	
Clotonic (II)	dizziness, or balance issues. ¹	
	dizziness, of balance issues.	

Term	Definition	Translation
Overcorrection (n)	A punishment mechanism for a challenging behavior that involves requiring an individual to engage in repetitive behavior to an excessive extent in an attempt to prevent the behavior from reoccurring. ¹	
Pediatric Autoimmune	Asubset of children and adolescents who have	
Neuropsychiatric	Obsessive Compulsive Disorder (OCD) and/or tic	
Disorders Associated with	disorders, and in whom symptoms worsen following	
Streptococcal infections	infections such as strep throat and scarlet fever.1	
(PANDAS) (n)		
Phenylketonuria (PKU)	An inherited disorder that increases the levels of a	
(n)	substance called phenylalanine in the blood. Phenylalanine is found in food proteins and in some artificial sweeteners. If PKU is not treated,	
	phenylalanine can build up to harmful levels in the	
	body, causing intellectual disability and other serious	
	health problems such as seizures, delayed	
	development, behavioral problems, and psychiatric	
	disorders. ²	
Physical Therapist (PT) (n)	A healthcare team member who treats conditions that	
	limit the ability to move and perform functional	
	activities of daily life. Physical therapists provide	
	services to improve body structures (e.g., muscles),	
	enhance functional activities (e.g., walking), and	
	promote improved participation in daily life. Goals for	
	children may include developing improved strength,	
	range of motion, coordination, balance, and	
Piec (a)	acquisition of new motor skills. ²	
Pica (n)	An eating disorder that involves eating things that	
Picture Exchange	are not food (e.g., dirt, plastic). A unique augmentative/alternative communication	
Communication Systems	intervention package that involves teaching an	
(PECS) (n)	individual to give a picture of a desired item to a	
(-20), (-1)	"communicative partner," and goes on to teach	
	discrimination of pictures and how to put them	
	together in sentences. ¹	
Polypharmacy (n)	The use of multiple medications by a patient. ¹	
Positive Behavior	An approach to helping people improve their difficult	
Supports (PBS) (n)	behavior by understanding what is causing it, and then	
	developing strategies to increase positive behaviors. 1	
Post-Traumatic Stress	An anxiety disorder that can occur after witnessing or	
Disorder (PTSD) (n)	experiencing a traumatic event. ¹	
Prematurity (n)	A premature birth is a birth that takes place more than	
	three weeks before a baby is due. Normally, a	
	pregnancy lasts about 40 weeks, so a premature birth	
	is one occurring before the start of the 37 th week of	
	pregnancy. ²	

Term	Definition	Translation
Prompting (v)	Any help given to a learner to assist in using a specific skill or behavior. Prompts can come in many different forms including visual, verbal, gestural, models, and partial or full physical. ⁵	
Psychologist (n)	Someone with a PhD or PsyD, trained in the evaluation, study and/or treatment of psychiatric and	
	cognitive disorders. A school psychologist is a professional trained in psychology and education who collaborates with children, educators, parents, and	
	other professionals to create healthy and supportive learning environments for students based on careful evaluation of cognitive, executive, and adaptive skills. ²	
Psychosis (n)	A loss of contact with reality that usually includes delusions and hallucinations. ¹	
Psychotropic (adj)	A medication or intervention that affects brain activity, behavior, or perception. ¹	
Puberty (n)	The process of physical changes that occur when a child's body matures into an adult. ¹	
Red flags (n)	Behaviors that cause caregivers concern in an area of a child's development. They warn you to stop, look, and think, and then observe and document. ²	
Reinforce (v)	To strengthen with additional material or support.1	
Reinforcement strategies	Methods used to promote or increase positive	
(n)	behavior by providing motivating reinforcers, such as praise, a favorite toy, a cookie, or a preferred activity. Also called a "reward" or an "incentive."	
Resilience (n)	An ability to recover from or adjust easily to change or a difficult situation. ¹	
Respite care (n)	A service that provides short-term breaks that can relieve stress, restore energy, and promote balance for caregivers. 1	
Restraints (n)	Physical restrictions immobilizing or reducing the ability of an individual to move their arms, legs, body, or head freely. This can be someone holding a	
	person so they cannot move, or it can be an item that restrains their body such as a device that holds their hands or bodies down. ¹	
Restricted or repetitive	Topics or tangible items that individuals with autism	
behaviors or interests (n)	pursue with great intensity and focus for long durations of time. ⁴	
Reward (n)	A prize, token, or preferred activity given to an individual for good behavior, designed to promote the same behavior in the future. ¹	
Risk factor (n)	Conditions that increase the likelihood of aggression. ¹	
Ritual (n)	A repetitive behavior that a person appears to use in a systematic way in order to promote calm or prevent anxiety, such as arranging all the pillows in a certain way before being able to settle in to sleep. ¹	

Term	Definition	Translation
Rumination (n)	The practice of (voluntarily or involuntarily) spitting	
	up partially digested food and re-chewing it, then	
	swallowing again or spitting it out. Rumination	
	often seems to be triggered by reflux or other	
	gastrointestinal concerns. ¹	
Schizophrenia (n)	A chronic, severe, and disabling brain disorder that	
	makes it hard for individuals to think clearly and tell	
Companies to the standard (s)	the difference between what is real and not real. ¹	
Screening test or tool (n)	An evaluation tool to identify children who are at-risk	
	for having or developing a developmental disability.	
	This is different from a diagnostic tool that is used to determine if a person has, or does not have, a	
	neurodevelopmental disability. ²	
Seclusion (n)	A situation in which an individual is isolated in a room	
	in response to a behavior they have exhibited. ¹	
Sedating (v)	Calming, sleep-inducing, or numbing an individual	
	experiencing challenging behaviors or struggling	
	during difficult situations. ¹	
Seizure disorder (n)	A seizure disorder, which can also be called epilepsy, is	
	a disorder in which brain activity is disturbed and	
	causes seizures. There are many types of seizures.	
	Some involve uncontrollable jerking movements or	
	losing awareness, but some can just look like a staring	
	spell. ⁵	
Self-advocacy (n)	The ability of an individual to communicate his or her	
	wants and concerns and make his or her own	
	decisions. ¹	
Sensory avoidance (n)	Blocking or staying away from something that is	
Canaami dafamahaanaa (n)	painful or bothersome. ¹	
Sensory defensiveness (n)	A tendency to react negatively or with alarm to	
	sensory input which is generally considered harmless or non-irritating. ¹	
Sensory input (n)	Any source that creates sensation and activates one	
Sensory input (ii)	or more of the senses -vision, smell, sound, taste,	
	and touch. ¹	
Sensory processing	Refers to difficulty detecting, organizing, or	
disorder (n)	responding to sensory information received and	
	interpreted in the brain via all seven senses and that	
	interferes with participation in daily life, development,	
	behavior, and social interactions. ²	
Sensory-seeking behavior	Behaviors caused by a need for additional	
(n)	stimulation of certain senses as a way of maintaining	
	attention or achieving a calmer state.1	
Sleep apnea (n)	A usually chronic, common disorder in which an	
	individual has one or more pauses in breathing	
	or shallow breaths up to 30 or more times per	
	hour during sleep, and results in daytime	
	sleepiness. ¹	

Term	Definition	Translation
Sleep disturbances (n)	Sleep disturbances can include a variety of issues	
	including difficulties falling asleep or staying asleep,	
	being on different sleep rhythms (e.g., awake at night),	
	or needing increased amounts of sleep. ⁵	
Social communication	Applies to children who have deficits in the social use	
disorder (n)	of language, but do not have the restricted interests or	
	repetitive behavior commonly found in autism	
	spectrum disorders. ²	
Social reciprocity (n)	The primary component of healthy social development	
	– acts of kindness – interactions. It is about "joint	
	attention" where there is purposeful language	
	including body language and pretend play. Lack of	
	social reciprocity is a key red flag, whether the child is	
	not engaged in the world around him. This concern	
Social work or social	should not be ignored. ² Social work practice is aimed at assisting individuals,	
worker (n)	groups, or communities to enhance or restore their	
worker (II)	capacity for social functioning and creating societal	
	conditions favorable to reach their goals. The primary	
	mission of the social work profession is to enhance	
	human well-being and help meet the basic human	
	needs of all people, with particular attention to the	
	needs and empowerment of people who are	
	vulnerable, oppressed, and living in poverty. ²	
Special Needs Parent	An advocate for parents of children with special needs	
Advocate (n)	who helps ensures that the child's rights and needs are	
. ,	met in school and in the community.1	
Specific learning disability	A disorder that manifests itself with a deficit in areas	
(n)	such as attention, reasoning, processing, memory,	
	communication, reading, writing, spelling, calculation,	
	coordination, social competence, and emotional	
	maturity. Often identified by a discrepancy between	
	school performance and the performance expected	
	based on overall intelligence. ²	
Speech generating device	A technological device that helps people who are	
or Voice output	unable to use speech to express their needs and	
technology (n)	exchange information with other people. ¹	
Speech sound disorder	Speech disorders in which some speech sounds in a	
(n)	child's native language are not produced, not	
	produced correctly, or are not used correctly. ²	
Speech-language	A clinician who assesses, diagnoses, treats, and helps	
pathologist (n)	prevent speech, language, cognitive, communication,	
	voice, swallowing, fluency, and other related	
	disorders. ²	

Term	Definition	Translation
Standardized test (n)	A test administered and scored in a consistent or	
	standard manner. It is administered in controlled	
	conditions that specify where, when, how, and for	
	how long children respond to the questions. In	
	standardized tests, the questions, conditions for administering, scoring procedures, and interpretations	
	are consistent. A well designed standardized test	
	provides an assessment of an individual's mastery of a	
	domain of knowledge or skill. ²	
Staring spells (n)	Occasions when an individual is in a trance staring into	
	space, which can often signal seizure activity.1	
Stereotypy (n)	Repetitive or ritualistic movements such as body	
	rocking or crossing and uncrossing of legs. ¹	
Stimulation (n)	Excitement or activity triggered by a stimulus either	
Complemental Consults	internally or externally.1	
Supplemental Security Income (SSI) (n)	A Federal income supplement program designed to help aged, blind, and disabled people who have little	
income (551) (II)	or no income, and provides cash to meet basic needs	
	for food, clothing, and shelter. ¹	
Tangibles (n)	Items or rewards that can be touched, such as a toy or	
	piece of candy.1	
Time delay (n)	A prompting procedure that fades prompts during	
	activities by having longer delays before a prompt is	
	provided.	
Tourette syndrome or	A neurological disorder characterized by tics, or	
Tourette's syndrome (n)	repetitive, stereotyped, involuntary movements and vocalizations. ¹	
Tracking scales (n)	A document or other tool used to track information	
Tracking scales (II)	such as changes in an individual's behaviors, side	
	effects of medications, school performance, etc. ¹	
Transition plan /	Part of IDEA, transition services means a coordinated	
transition services (n)	set of activities for a child with a disability that is	
	results-oriented and focused on improving the	
	academic and functional achievement of the child with	
	a disability in order to facilitate the child's movement	
	from school to post-school activities, including postsecondary education, vocational education,	
	integrated employment (including supported	
	employment), continuing and adult education, adult	
	services, independent living, or community	
	participation. Transition services will be based on the	
	individual child's needs, taking into account the child's	
	strengths, preferences, and interests. ²	
TRICARE (n)	The health care program for Uniformed Service	
	members, retirees, and their families worldwide. ¹	

Term	Definition	Translation
Visual cues (n)	An image or symbol presented to the learner that either prompts him to complete the target behavior/skill or assists in supporting the understanding of a particular activity, instruction, or direction. Visual cues can be used as prompts or supports. ⁵	
Visual schedules (n)	A display that shows the activities of a day or steps of an activity to support the learner in moving between activities or through activities. The displays can be made from objects, pictures, photos, or written text, and may be created for a full day, part of a day, or just the next two activities. ⁵	
Vocational rehabilitation (n)	A state and federally funded program designed to help people with disabilities become employed and to help those already employed perform more successfully through training, counseling, and other support methods.	
Voice output technology or Voice output communication aid or Speech generating device (n)	A technological device that helps people who are unable to use speech to express their needs and exchange information with other people. ¹	
Wraparound (n)	An integrated, multi-agency, community-based planning process designed to build teams of providers, family members and natural supports to help keep complex youth in their homes and communities. ¹	

References

- ¹ The definitions for these terms were taken from "Challenging Behaviors Glossary" published in 2012 by Autism Speaks. This glossary is part of a downloadable toolkit available at: https://www.autismspeaks.org/tool-kit/challenging-behaviors-tool-kit
- ² The definitions for these terms were taken from "Glossary of Terms Related to Neurodevelopmental Disabilities" by John Thorne, PhD, CCC-SLP. The "Glossary of Terms Related to Neurodevelopmental Disabilities" was adapted by John Thorne, PhD, CCC-SLP, from a number of in-depth resources (12/16) including:

http://www.inclusivechildcare.org/inclusion_glossary.cfm

http://www.ddrcco.com/resources-and-training/glossary-of-developmental-disability-terms.php http://autismnow.org/at-home/learn-and-understand-autism/autism-and-developmental-disabilities-glossary/

http://www.gennextmsp.org/wp-content/uploads/2015/05/ECS-Glossary-May2015.pdf http://www.ncca.biz/Aistear/pdfs/PrinciplesThemes_ENG/Glossary_ENG.pdf http://www.johnson-center.org/downloads/pdfs/What is a Standardized Test.pdf

- ³ The definition was taken from The American Heritage® Stedman's Medical Dictionary. Copyright ©2002,2001, 1995 by Houghton Mifflin Company. Published by Houghton Mifflin Company.
- ⁴ DSM-IV, 1994.
- Excerpted from Steinbrenner, J., Sam, A., Chin, J., Morgan, W., & AFIRM for Paras Team. (2019). *Introduction to ASD*. FPG Child Development Institute, University of North Carolina. Retrieved from https://afirm.fpg.unc.edu/introduction-asd

More information is available at:

Autism Society

Founded in 1965 by Dr. Bernard Rimland, Dr. Ruth Sullivan and many other parents of children with autism, the Autism Society is the leading source of trusted and reliable information about autism. Through its national network, the Autism Society has spearheaded numerous pieces of federal, state and local legislation, including the 2006 Combating Autism Act, the first federal autism-specific law. The Autism Society's website is one of the most visited websites on autism in the world and our on-line resource database, <u>Autism Source™</u>, and National Contact Center (800-3-AUTISM) provide information and service referrals to thousands of people each year. Look online at https://www.autism-society.org/.

Autism Speaks

Autism Speaks was founded in February 2005 by Bob and Suzanne Wright, grandparents of a child with autism. Recognizing the need for a powerful voice Bernie Marcus donated \$25 million to help financially launch the organization.

Building upon the legacy of three leading autism organizations, Autism Coalition for Research and Education (ACRE), the National Alliance for Autism Research (NAAR) and Cure Autism Now (CAN), who merged with the organization, Autism Speaks has made extraordinary advancements in the autism community.

Chief among these are increased global awareness of autism, better understanding of the breadth of the autism spectrum, and advocacy to increase research and access to care and support.

Today, Autism Speaks is dedicated to advancing research into causes and better treatments for autism spectrum disorders and related conditions both through direct funding and collaboration.

Look online at https://www.autismspeaks.org/.

<u>Centers for Disease Control and Prevention</u> (CDC)
 <u>CDC works 24/7</u> to protect America from health, safety, and security threats, both foreign and in the U.S. Whether diseases start at home or abroad, are chronic or acute, curable or preventable, human error or deliberate attack, CDC fights disease and supports communities and citizens to do the same.

CDC increases the health security of our nation. As the nation's health protection agency, CDC saves lives and protects people from health threats. To accomplish our mission, CDC conducts critical science and provides health information that protects our nation against expensive and dangerous health threats, and responds when these arise.

Look online at:

https://search.cdc.gov/search/index.html?query=autism%20spectrum%20disorder



The National Coordinating Center for the Regional Genetics Networks

Glossary of

Cancer Genetics Terms

English – Language of Translation

2023 Edition

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In collaboration with:
The Cross Cultural Health Care
Program (CCHCP)



Preface

The seven Regional Genetics Networks (RGNs), the National Genetics Education and Family Support Center (Family Center), and the National Coordinating Center for the Regional Genetics Networks (NCC) mission is to improve access to quality genetic services for medically underserved populations. This bilingual glossary of terms related to cancer genetics is provided to help enable access to cancer genetics medical services to limited English proficiency populations.

How to Use the Glossary

This glossary is organized alphabetically in English. The first column shows the term in English. The second column has the definition of the term in English. The third column has the translation of the term.

At the end of the translations of the cancer genetics terms is a page with Examples of Genes That are Often Checked in Genetic Testing, and a glossary of translations for Family Relationships Terms. It is very important for genetic counselors to know if a relative is a blood relative or a relative by marriage. There is also a page on Common Interpreting Errors, and a page on Analogies. Genetic counselors often use analogies to explain complex ideas in genetics.

Glossary of Terms

English – Spanish

(n) – noun (v) – verb (adj) – adjective (adv) – adverb

Term	Definition	Translation
Abnormal (adj)	Different from what is considered normal, average or expected; e.g., a gene sequence that is different than that found in most people.	
Benign (adj)	Something mild that does not threaten health or life. In cancer genetics, "benign" means "not cancerous."	
Blood test (n)	A test in which blood is drawn (usually from the patient's arm) and sent to a laboratory for analysis. In cancer genetics, blood tests provide cells for genetic sequencing.	
Breast cancer (n)	Cancer of the breast tissue. This cancer is more common in women, but it can affect men as well.	
Buccal swab (n)	A way to collect DNA from the cells on the inside of a person's cheek.	
CA-125 blood test (n)	A blood test used to look for early signs of ovarian cancer in women with a high cancer risk.	
Carrier (n)	A person who carries a genetic mutation in one of the two copies of a particular gene, regardless of whether they get cancer or not.	
Cell (n)	A small (microscopic) structure that forms the basic building block of every known living organism.	
Chromosome (n)	Thread-like structures located inside the nucleus of cells. In humans, there are 23 pairs of chromosomes, for a total of 46 chromosomes. Each chromosome is made of proteins and a single molecule of DNA, which carries genetic information.	
Colonoscopy (n)	A diagnostic test in which a flexible tube with a tiny camera on the end is inserted into the colon (large intestine) through the rectum in order to see the condition of the inside of the colon, or to take a small sample of tissue, or to cut out polyps (small growths).	
Condition (e.g., "genetic condition") (n)	A chronic (long-term) health issue.	
Consanguinity (n)	When parents are blood relatives to each other.	

Term	Definition	Translation
DCIS (ductal carcinoma in situ) (n)	A pre-cancerous growth in the breast, which is still within the milk ducts but has the potential of growing and spreading to other parts of the breast tissue. Sometimes called Stage 0 breast cancer.	
DTC (direct-to-consumer genetic testing) (n)	Genetic testing that is marketed directly to people at home through the internet or the TV etc. instead of through a medical provider.	
Deleterious mutation/ Disease-causing mutation (n)	A change in a person's DNA that may cause a medical condition. In cancer genetics, having a disease-causing mutation may increase the chance of getting cancer but does not mean that a person will definitely get cancer.	
DNA (deoxyribonucleic acid) (n)	The molecule that carries the genetic information of a cell. It provides the instructions used in the development, functioning and reproduction of the organism of which it is a part. The DNA is organized into chromosomes.	
DNA banking (n)	The secure, long-term storage of a person's genetic material. DNA banking allows families to have access to a deceased family member's DNA. When new genetic testing techniques are developed, doctors can test the banked DNA and use genetic information from the deceased person to tailor medical treatment for living descendants.	
DNA marker (also called a "genetic marker") (n)	A readily recognizable genetic trait, gene, or DNA segment.	
DNA mutation (n)	A change in the typical order of the chemicals that make up the DNA. Mutations or variants are often compared to misspelled words because chemicals that make up the DNA sequence are not in the expected order.	
DNA sequence (n)	The exact order of the chemicals that make up a DNA molecule.	
DNA sequencing (n)	The laboratory technique used to determine the exact order of the chemicals that make up a DNA molecule. This is one type of genetic testing.	

Term	Definition	Translation
Dominant (adj)	A genetic trait in which one copy of the gene is sufficient for a trait to be expressed; dominant traits can be inherited from a single parent.	
	In cancer genetics, dominant inheritance refers to conditions in which having only one altered copy of a particular gene pair is sufficient to cause an increased risk for cancer.	
Duplication (n)	A duplication occurs when part of a chromosome is copied abnormally, resulting in extra genetic material.	
Early age of onset (n)	Cancer diagnosed at an early age, usually before the age of 50.	
Egg (n)	The reproductive cells of a woman. When fertilized by sperm, the egg will grow into a baby.	
Environmental causes or factors (n)	Causes of illness that come from a person's surroundings, not from their genetic make-up. Environmental causes of illness include infection, trauma, diet, exposure to toxins, etc. Some environmental causes of illness can be avoided or changed, while genetic causes cannot.	
Familial (adj)	Belonging to a family; e.g., a familial trait is a trait that is shared among family members and may be due to genetic or environmental factors or both.	
Family history (n)	The medical history of the members of a family.	
Flip a coin (v)	A random decision-making tool used in the U.S. While a coin is flipped into the air and caught, a person predicts whether it will fall with the "heads" side up or the "tails" side up. If the coin falls as predicted, the person "wins." This expression is often used as a metaphor for any outcome that is random and has two possible outcomes, and to describe a situation in which each outcome is as likely as the other.	
Fragment (n)	A small piece; an incomplete part of a whole.	
Gene (n)	A specific sequence of DNA that determines specific traits in an individual.	

Term	Definition	Translation
Gene copy (n)	In human cells, DNA is arranged in 23 pairs of chromosomes, for a total of 46 chromosomes. These chromosomes contain all the genes that make up the DNA. One chromosome of each pair is inherited from each parent. As such, individuals have two copies of every gene, one inherited from the mother and one inherited from the father.	
Normal gene copy (n)	A normal copy of a gene is one that is found most frequently in the population. Also called the "working copy."	
Altered gene copy (n)	An altered copy of a gene is one that differs from the working copy and is considered to be "abnormal." Also called the "non-working copy."	
Gene deletion (n)	Having a piece of genetic information missing from a gene.	
General population (n)	"Most people."	
	For example, if you have the same risk of getting cancer as the general population, that means that you have the same chance of getting cancer as everyone else. This is in contrast to a "high risk population" who has a greater chance of getting cancer than everyone else.	
Generation (n)	The people who constitute a single step in a line of descent from an ancestor; a group of people born and living more or less at the same time.	
	Example: You, your brothers and sisters, all your spouses and your cousins are in the same generation. Your parents, your aunts and uncles and all their spouses form a previous generation. Your grandparents, their siblings and spouses form an even earlier generation. Your children and nieces and nephews form a later generation.	
Genetic (adj)	Having to do with inherited traits.	
Genetic counseling (n)	A discussion with a genetic counselor about the basic concepts of genetics, genetic conditions, the chances of being affected by a genetic condition or having a child with a genetic condition, and genetic testing and treatment.	

Term	Definition	Translation
Genetic counseling intern (n)	A genetic counseling student who has not yet completed his/her academic studies and is now practicing under the supervision of a more experienced counselor in preparation for providing genetic counseling services independently after obtaining his/her graduate degree.	
Genetic counselor (n)	A healthcare professional with a specialized graduate degree who works with people undergoing genetic testing. Genetic counselors provide information about genetic conditions, help patients understand their chances of being affected by a genetic condition or having a child with a genetic condition, and help them make informed decisions about testing and treatment. Genetic counselors also provide emotional support to patients and families.	
Genetic discrimination (n)	The act of refusing to provide, or charging more for, insurance or any service based on an individual's genetic make-up.	
Genetic factors (n)	Specific aspects of a person's genetic make-up that influence that person's health and development.	
Genetic information (n)	The information encoded in genes, which tells every cell in a body how to grow, what to do and how to reproduce.	
Genetic material (n)	All the parts of a cell that carry genetic information. Genetic material could include genes, parts of genes, a group of genes, a DNA molecule, a fragment of DNA, a group of DNA molecules, or the entire set of genetic instructions.	
Genetic predisposition (n)	An increased chance of a person developing a certain trait or disease based on that person's particular genetic makeup.	

Term	Definition	Translation
Genetic test (n)	A laboratory test designed to determine if a person has a gene mutation or a typical DNA sequence.	
Single gene test (n)	Single gene test: analysis of one particular gene.	
Gene panel test (n)	Gene panel test: analysis of several genes that have been shown to be associated with a particular condition; Example: breast cancer gene panel.	
Genetic trait (n)	A characteristic within a family that is passed down from parent to child genetically.	
Geneticist (n)	A doctor or scientist who studies genetics.	
Germline testing (n)	Germline testing refers to the analysis of a person's DNA, which he or she inherited from his or her parents. In biology and genetics, the germline is the group of cells that will pass on the genetic material to children, in other words, the cells from which the eggs and sperm come. After the egg and sperm come together to form a baby, these cells will then divide and multiply and will form the entire body. Germline testing is different from testing the DNA of someone's tumor, which may have a mutation that happened during the person's lifetime and was not inherited from his or her parents.	
Hereditary (adj)	Passed down from parent to child.	
Hereditary material (n)	Genetic material that is passed down from parent to child.	

Term	Definition	Translation
Informed consent (n)	The process of agreeing to a procedure or course of treatment after understanding what the procedure/treatment entails, the potential risks and benefits associated with it, and the other options available.	
Inheritance pattern (n)	The manner in which a particular genetic trait or disorder is passed from a parent to a child, e.g., autosomal dominant or recessive, X-linked dominant or recessive, or multifactorial.	
Inherited (adj)	Passed down from parent to child.	
Lynch syndrome (n)	An inherited genetic disorder that increases a person's risk of getting cancer of the colon, rectum, uterus, ovaries, and other cancers.	
Malignant (adj)	In cancer, this term means that the cancer cells or tumor are harmful and have the potential of spreading to other tissues or parts of the body.	
Metastasis (n)	The spreading of cancer from one organ to another.	
Molecule (n)	The smallest unit of a chemical compound that still has the properties of that compound. For example, a molecule of water is made up of two hydrogen atoms and one oxygen atom. Separately, they are just atoms, but when bonded together, they make a water molecule.	
Multifactorial (adj)	Due to a combination of genetic and non-genetic (environmental, hormonal, etc.) risk factors that act together to determine risk.	
Mutation (n)	A change in a gene, which can be deleterious (disease causing) or benign (non-disease-causing). In Spanish, the word is sometimes confused with "mutilation."	
Oncologist (n)	A physician who specializes in diagnosing and treating cancer.	
Oophorectomy	The surgical removal of one or both ovaries.	
Ovarian cancer (n)	Cancer of the ovaries, the organs in a woman that release eggs.	
Packets of genetic information (n)	A phrase genetic counselors use to describe genes or chromosomes.	

Term	Definition	Translation
Pattern (n)	A repeating arrangement or sequence; for example, the pattern of cancers in a family.	
Pedigree (n)	A family tree that can be used to trace the inheritance of specific genetic traits.	
Penetrance (n)	The probability that a specific genetic trait will be expressed if a person carries a mutation. "Complete penetrance" means that everyone who carries a particular gene mutation will show the trait related to that altered gene. "Incomplete penetrance" means that only some of the people who have the altered gene will actually show the related trait.	
Predisposition to cancer	Having a change in one of number of specific genes (a genetic mutation) that creates a higher than normal risk of a person developing cancer.	
Prognosis (n)	The most likely outcome of a disease process.	
Proliferation (cell proliferation) (n)	The controlled process by which a cell multiplies. Cancer arises when the process of cell division becomes uncontrolled.	
Prophylactic mastectomy (n)	A mastectomy is the surgical removal of the breast. "Prophylactic" means something done to prevent disease. A prophylactic mastectomy is the surgical removal of a healthy breast to prevent breast cancer. Prophylactic mastectomy is an option for individuals with a very high breast cancer risk, for example, women with a BRCA mutation.	
Protein (n)	A molecule made up of chains of amino acids. Proteins do most of the work in cells and are required for the structure, function, and regulation of the body's tissues and organs. Genes determine how specific amino acids are put together to form a specific protein.	
Random (adj)	Happening in an unpredictable way.	
Recessive (adj)	A genetic trait in which both copies of a gene are needed for the trait to be expressed; recessive traits are inherited from both parents, each one contributing one copy of the gene in question.	
	In cancer genetics, recessive inheritance refers to conditions in which both copies of a particular gene pair need to be altered to develop the disease.	

Term	Definition	Translation
Recurrence (n)	When something happens again.	
	In cancer, a return of the disease after treatment and after a period of time during which the cancer cannot be detected.	
Red flag (n)	A warning sign or a clue.	
Risk (n)	The chance that something will happen; in the case of cancer genetics, the chance of getting cancer.	
At risk (adj)	Has some possibility of getting cancer.	
At average risk (adj)	Has the same possibility of getting cancer as the general population.	
At high risk (adj)	Has a greater possibility of having a genetic condition than the general population.	
At higher risk than "X" (adj)	Has a greater possibility of getting cancer than the general population. Has a greater possibility of getting cancer than "X."	
Risk factor (n)	A circumstance that increases the risk of getting cancer.	
Runs in the family (e.g., cancer runs in the family) (v)	Is passed down from parents to children to grandchildren.	
Saliva (spit) test (n)	A genetic test done by collecting saliva (spit) instead of blood.	
Screen (v)	To test a person for a particular common health problem, even though the person has no symptoms to suggest that they have the problem.	
Screening test (n)	A test that looks for a common health problem even though the patient has no symptoms to suggest they have the particular disease.	
Sperm (n)	The reproductive cells of the man. When sperm fertilize a woman's egg, a baby develops.	
Sporadic (adj)	Random, once in a while.	
	In genetics, sporadic cancers are those caused by random chance or unknown factors in the environment.	

Term	Definition	Translation
Statistically significant (adj)	Not caused by chance.	
Syndrome (n)	A group of symptoms or features that consistently occur together or a condition characterized by a set of associated features. For example, a hereditary cancer syndrome refers to risk for a group of cancers all caused by a particular gene mutation. BRCA1 mutations increase risk for breast and ovarian cancers. This is called Hereditary Breast and Ovarian Cancer Syndrome. See also, Lynch Syndrome.	

Term	Definition	Translation
Test result (n)		
Negative (adj)	A negative result on a genetic test means that the laboratory did not find the specific genetic change that the test was designed to identify.	
True negative (n)	When a patient is being tested to see if he or she has a specific genetic change that is present in other family members, a negative result can be considered a "true negative." This means that instead of having an increased risk for cancer like other family members, they have the same risk as everyone else in the general population.	
Uninformative negative (n)	Sometimes a family has a strong history of a particular cancer, but no genetic change related to that cancer has been found in that family. An "uninformative negative" result means that a specific test was negative but is not conclusive because the "family" mutation has not been identified, and/or there may be changes that the test could not detect in the gene being analyzed, or in other genes that were not tested.	
Positive (adj)	A positive result on a genetic test means that the laboratory did find a specific genetic change that is associated with a hereditary cancer syndrome (a deleterious mutation). In cancer genetics, that means that the patient has an increased risk of getting the cancer(s) associated with the gene.	
Variant of uncertain significance (n)	If the laboratory finds a genetic change for which currently there is not enough information to know if this change is problematic or not, it reports a "variant of uncertain significance." Basically, this means that the laboratory found a genetic change, but they don't know what it means. All of these variants will eventually be reclassified as either a "positive" or "negative" result. Most become "negative" and are thought to represent natural variation between individuals.	
Polymorphism (n)	Everyone has some degree of commonly occurring genetic changes that are not associated with cancer. If the test finds this sort of change, it reports a "polymorphism."	
Trait (n)	A characteristic.	
Tumor (n)	An abnormal growth of cells; a tumor can be benign (not harmful) or malignant (harmful, cancer).	

Term	Definition	Translation
Tumor testing (n)	An analysis of the DNA of someone's tumor. This analysis is meant to identify changes that a person might have acquired in his/her tumor cells that are not present in the rest of his/her body cells and were not inherited from his/her parents. This testing is often done to help make treatment decisions.	
Typo (n)	A mistyped word; shortened slang for "typographical error." In genetic counseling "a typo" is commonly used as an analogy to refer to mutations or variants in a gene.	
Ultrasound (n)	An imaging method that uses high frequency sound waves to create a picture of something inside the body.	
Variant (n)	A version of something that differs from the norm. So, a genetic variant is a change to the usual genetic sequence.	

Examples of Genes That are Often Checked in Genetic Testing

Term	Definition	Translation
BRCA 1	"The Breast Cancer Genes."	
BRCA 2		
	The tumor suppressor genes that in mutated form tend	
	to be associated with an increased risk of certain	
	cancers, especially breast and ovarian cancers.	
MLH1 MSH2 MSH6 PMS2	"The Colon Cancer Genes."	
	These genes are involved in fixing mistakes that occur	
	when DNA replicates. An abnormality in one of these	
	genes can lead to Lynch Syndrome, a condition linked to	
	increased risk of cancer in the colon, rectum, uterus,	
	ovaries, as well as other organs.	
ATM	The ATM gene helps create a protein located in the cell	
	nucleus that controls the rate at which cells grow and	
	divide. The ATM protein also helps cells recognize	
	damaged DNA strands.	
	A mutation in the ATM gene can lead to increased risks	
	for cancer or a condition called ataxia-telangiectasia a	
	degenerative neurological condition that causes severe	
	disability.	
PALB2	Another gene in which mutations can lead to breast	
	cancer and possibly other cancers.	

Family Relationships

For genetic counselors, it is very important to know if a relative is a blood relative or a relative by marriage.

Term	Definition	Translation
Parent	Your mother or father.	
Mother	For genetic purposes, the woman whose egg was fertilized and grew to be you.	
Mother-in-law	Your husband or wife's mother.	
Stepmother	Your father's wife who is not your biological mother.	
Adoptive mother	A woman who is not your biological mother but who accepted legal responsibility for and raised you.	
Godmother	A woman chosen by your parents to be your spiritual guide in life, named as such through a ceremony in the Catholic church.	
Father	For genetic purposes, the man whose sperm fertilized the egg that grew to be you.	
Father-in-law	Your husband or wife's father.	
Stepfather	Your mother's husband who is not your biological father.	
Adoptive father	A man who is not your biological father but who accepted legal responsibility for and raised you.	
Godfather	A man chosen by your parents to be your spiritual guide in life, named as such through a ceremony in the Catholic church.	
Aunt	Your mother or father's sister or sister-in-law.	
Maternal aunt	Your mother's sister or sister-in-law.	
Paternal aunt	Your father's sister or sister-in-law.	

Term	Definition	Translation
Uncle	Your mother or father's brother or brother-in-law.	
Maternal uncle	Your mother's brother or brother-in-law.	
Paternal uncle	Your father's brother or brother-in-law.	
Cousin	Usually understood to be a first cousin.	
First cousin	Your aunt or uncle's child.	
First cousin once removed	Your aunt or uncle's grandchild or your first cousin's child	
Second cousin	The children of first cousins are second cousins to each other.	
Grandparent	Your parent's father or mother.	
Grandmother	Your mother or father's mother.	
Maternal grandmother	Your mother's mother.	
Paternal grandmother	Your father's mother.	
Grandfather	Your mother or father's father.	
Maternal grandfather	Your mother's father.	
Paternal grandfather	Your father's father.	
Great aunt	Your mother or father's aunt.	
Maternal great aunt	Your mother's aunt.	
Paternal great aunt	Your father's aunt.	
Great uncle	Your father or mother's uncle.	
Maternal great uncle	Your mother's uncle.	
Paternal great uncle	Your father's uncle.	

Term	Definition	Translation
Great grandparents	The parents of any of your grandparents.	
Great grandmother	The mother of any of your grandparents.	
Great grandfather	The father of any of your grandparents.	
Siblings	The children of your father and mother.	
Half siblings	Siblings (brothers and sisters) who have either the same mother and different fathers, or the same father but different mothers. Half siblings share some genetic similarity with you.	
Stepsiblings	The children of your stepmother but not your father; or the children of your stepfather but not your mother. Stepsiblings do not share any genetic similarity with you.	
Twins	Two siblings born at the same time.	
Identical twins	Twins who developed from the same egg and sperm, meaning that they are genetically identical. Also called monozygotic twins.	
Fraternal twins	Twins who developed from different eggs and sperm, meaning that they are genetically different and have the same number of shared genes as any other sibling. Also called dizygotic twins.	
Triplets	Three siblings born at the same time.	
Sister	A sibling who is a girl.	
Sister-in-law	Your brother's wife or your husband's sister.	
Half sister	A girl who is either the child of your father with a different mother, or the child of your mother with a different father.	
Stepsister	The daughter of your stepmother or stepfather and therefore not biologically related to you.	

Term	Definition	Translation
Brother	A sibling who is a boy.	
Brother-in-law	Your sister's husband or your wife's brother.	
Half brother	A boy who is either the child of your father with a different mother, or the child of your mother with a different father.	
Stepbrother	The son of your stepmother or stepfather and therefore not biologically related to you.	
Niece	Your sibling's daughter.	
Nephew	Your sibling's son.	
Spouse	Your husband or wife.	
Wife	The woman to whom you are married.	
Husband	The man to whom you are married.	
Children	Genetically speaking, the people who are produced from your egg or sperm.	
Daughter	Your child who is a girl.	
Daughter-in-law	Your son's wife.	
Stepdaughter	Your spouse's daughter who is not your biological child.	
Adoptive daughter	A girl for whom you have accepted legal responsibility and raised even though she is not your biological child.	
Goddaughter	A girl for whom you have accepted spiritual responsibility within a ceremony of the Catholic church; she may or may not be biologically related to you.	

Term	Definition	Translation
Son	Your child who is a boy.	
Son-in-law	Your daughter's husband.	
Stepson	Your spouse's son who is not your biological child.	
Adoptive son	A boy for whom you have accepted legal responsibility and raised even though he is not your biological child.	
Godson	A boy for whom you have accepted spiritual responsibility within a ceremony of the Catholic church. He may or may not be biologically related to you.	
Ancestor	A person from whom you are descended, usually more remote than a grandparent.	
Fiancé(e)	The person whom you have promised to marry.	
	Man: fiancé	
	Woman: fiancée	
Domestic partner	The person with whom you are living and have an intimate relationship, but to whom you are not married.	
Divorced	Having ended a marriage.	
Engaged	Having promised to marry someone.	
Widow	A woman whose husband has died.	
Widower	A man whose wife has died.	
Relative	A person in your family.	
Blood relation	A person who is related to you by blood not marriage, e.g., your sister, but not your sister-in-law; your mother but not your stepmother; your daughter but not your adopted daughter.	
Next of kin	Your closest living blood relation.	

Common Interpreting Errors

Source speech	Interpreted as	More accurate rendition
"Cancer just happens by chance."	"Cancer happens suddenly." "Cancer happens because you are unlucky. "	"Sometimes we don't know what causes cancer."
Cancer genes	Cancer cells	Genes that may cause cancer when altered (mutated).
"You are at higher risk for developing cancer."	Getting cancer	There is a greater possibility that you might get cancer.
"These genes protect us against cancer."	"They protect us, like the immune system They kill cancer cells "	"These genes help us to not get cancer."
Colonoscopy	Scan	Colonoscopy

Analogies

Genetic counselors often use analogies to explain complex ideas in genetics. An analogy explains a new concept by comparing it to something more common. A clue that an analogy is being used are the words "It's as if" and "It's like "

Examples:

"Genes are <u>like</u> an instruction book or an instruction manual. And in a book, there is a certain sequence to a story."

"It's as if you were reading through a long book and looking for one typo, for a letter that is mistaken."

"In the DNA there are molecule strings – like in this picture, each letter is a different molecule string."

"So a mutation is <u>like</u> a broken gene."

"So a mutation is like an unexpected change in the order of the letters."

"A gene is like a recipe, and mutations are like changes in that recipe."

"It's like chapters in a book, and like misspellings."

"Our cells are like libraries and our chromosomes are like books."

"Like a flip of a coin." (to explain that there is a 50% chance that a mutation will be passed on every time the patient has a child).

"Inside every cell are genes, which are <u>like</u> instruction books for our bodies; they tell our bodies how to grow and develop."

"It is kind of <u>like</u> if you a reading a book = and you notice that a word is spelled wrong; that is what this genetic test looks for."

Genetic counselors may mix the analogy in with more literal descriptions:

Example:

"We know that there are two in particular that when they don't work right, when there's a typo in the instruction manual, they can cause a high risk for breast and ovarian cancer."

Some analogies may not work with certain patients, due to linguistic or experiential differences.

Analogies that focus on genes as "letters in a book" may make no sense to speakers of character-based written languages such as Chinese, Japanese, and Korean.

"It's as if you were reading through a long book and looking for one typo, for a letter that is mistaken."

"So a mutation is <u>like</u> an unexpected change in the order of the letters."

"It's like chapters in a book, and like misspellings."

"It is kind of <u>like</u> if you a reading a book and you notice that a word is spelled wrong; that is what this genetic test looks for."

Analogies that focus on genes as a recipe may not work for patients who do not cook with written recipes.

"A gene is <u>like</u> a recipe, and mutations are like changes in that recipe."

Analogies that discuss random chance in terms of a coin toss may not work for patients from cultures where people don't flip coins to make decisions.

"Like a flip of a coin" (to explain that there is a 50% chance that a mutation will be passed on every time the patient has a child).



The National Coordinating Center for the Regional Genetics Networks

Glossary of Pediatric Genetics Terms

English – Language of Translation

2022 Edition

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The Cross Cultural Health Care
Program (CCHCP)



Preface

The seven Regional Genetics Networks (RGNs), the National Genetics Education and Family Support Center (Family Center), and the National Coordinating Center for the Regional Genetics Networks (NCC) mission is to improve access to quality genetic services for medically underserved populations. This bilingual glossary of terms related to Pediatric Genetics is provided to help enable access to pediatric medical services to limited English proficiency populations.

How to Use the Glossary

This glossary is organized alphabetically in English. The first column shows the term in English. The second column has the definition of the term in English. The third column has the translation of the term.

At the end of the translations of the pediatric genetics terms is a glossary of translations for family relationships terms. It is very important for genetic counselors to know if a relative is a blood relative or a relative by marriage.

Glossary of Terms

English - Spanish

(n) – noun (v) – verb (adj) – adjective (adv) – adverb

Term	Definition	Translation
Abnormal (adj)	Different from what is considered normal, average, or expected, e.g., a gene sequence that is different than that found in most people.	
Abortion (n)	The deliberate ending of a pregnancy by the removal of an embryo or fetus from the womb.	
	Technically, this is called an "induced abortion" in order to distinguish it from a "spontaneous abortion" which is also called a miscarriage or stillbirth. Also called a "termination of pregnancy" or just "termination."	
ACMG 59 (n)	Fifty-nine genes that the American College of Medical Genetics and Genomics has identified in which variances have a high risk of causing disease – but disease that can be mitigated through early detection or treatment. The ACMG recommends that variances in these genes should be reported to patients even if finding them was not the reason for genetic testing	
Adaptive skills (n)	Skills/abilities needed to perform daily activities, particularly those that allow an individual to live independently and/or work. Examples: brushing teeth, bathing, dressing, food shopping, and taking public transportation.	
Adoption (n)	The process of accepting legal responsibility for someone else's child to raise as your own.	
American College of Medical Genetics and Genomics (ACMG) (n)	A professional membership organization based in the United States that represents the interests of clinical geneticists, clinical laboratory geneticists, and genetic counselors. The ACMG focuses on improving patient care, establishing standards of care and laboratory policy, and educating members about advances in the field.	
Amniocentesis (n)	A procedure by which a sample of amniotic fluid is withdrawn from the amniotic sac. This is usually done by inserting a long needle through the abdominal and uterine walls with the guidance of ultrasound.	

Term	Definition	Translation
Amniotic fluid (n)	The liquid that surrounds a fetus as it develops in the uterus. This liquid contains skin cells that have sloughed off the fetus as well as other fetal cells.	
Anencephaly (n)	A neural tube defect that results in insufficient brain growth in a fetus. Babies with anencephaly usually die soon after birth or are stillborn.	
Benign (adj)	Something that does not threaten health or life. May refer to a type of change to the DNA that does not create health consequences.	
Biochemical testing (n)	Blood tests to identify elevated levels of certain naturally occurring substances/chemicals in the body that are linked to genetic conditions.	
Birth canal (n)	The passageway from the uterus through the cervix, the vagina, and the vulva through which a baby passes during the birth process.	
Birth defect (n)	A problem or physical difference with how the body works that is present at birth. Birth defects can be caused by genetic abnormalities, environmental influences, random chance, or by circumstances related to the birth process. Also called "congenital malformation" or "congenital anomaly."	
Blood test (n)	A test in which blood is drawn (usually from the patient's arm) and sent to a laboratory for analysis. In genetics, blood tests may provide cells for genetic sequencing, or blood samples may be used to assess things like protein or hormone levels.	
Brace (n)	A medical appliance that provides support to joints or bones, to weak muscles, or to strained ligaments.	
Buccal swab (n)	A way to collect DNA from the cells on the inside of a person's cheek.	
Carrier (n)	A person who carries a genetic mutation in one of the two copies of a particular gene, regardless of whether they get cancer or not.	
Cell (n)	The smallest (microscopic) functional unit of living organisms. All living things are composed of one or more cells. Within each cell are the structures needed for the cell to process energy, dispose of waste, reproduce, and perform specialized functions.	

Term	Definition	Translation
Cervix (n)	The narrow lower part of the uterus that connects to the vagina.	
Chorionic Villus Sampling (CVS) (n)	A procedure during pregnancy in which a sample of cells from the placenta is removed to check for possible genetic abnormalities.	
Chromosome (n)	Thread-like structures located inside the nucleus of cells. In humans, there are 23 pairs of chromosomes, for a total of 46 chromosomes. Each chromosome is made of a long strand of DNA, which carries genetic information.	
Chromosome test / Karyotype (n)	A test that looks at the number and arrangement of chromosomes in a cell.	
Cleft lip (n)	A congenital malformation that occurs when the upper lip does not form properly, causing an opening in the upper lip that can extend to the nose. This can occur together with a cleft palate or on its own.	
Cleft palate (n)	A congenital malformation where the roof of the mouth (palate) does not form properly, resulting in an opening into the nasal cavity. This can occur together with a cleft lip or on its own.	
Clinically significant (adj)	If a test result is "clinically significant," the result indicates a medical problem that can impact a person's life.	
Cognition (n)	The processes involved in thinking, learning, understanding, and remembering.	
Condition (e.g., "genetic condition") (n)	A long-term medical health issue.	
Consanguinity (n)	When parents are blood relatives to each other.	
Consultation model (n)	The model of care for patients with genetic conditions in which the genetic team acts as consultants, supporting the care being provided principally by the patient's Primary Care Provider.	
Continuing care model (n)	The model of care for patients with genetic conditions in which the genetic team takes primary responsibility for supervising and coordinating all care related to the patient's genetic condition.	

Term	Definition	Translation
Cystic fibrosis (n)	A progressive genetic condition that affects the exocrine glands (the glands that make sweat and digestive juices) and causes the production of thick, sticky mucus. This mucus blocks the pancreatic duct, the intestines, and the lungs, leading to persistent respiratory infections.	
De novo (adj)	Genetic changes that are not inherited but that occur during cell division.	
Deleterious mutation/ Disease-causing mutation (n)	A change in a person's DNA that may cause a medical condition.	
Deletion (n)	Having a section of genetic information (DNA) missing.	
Developmental assessment (n)	An evaluation of how an individual patient's intellectual, emotional, and social development compare with others of the same age.	
Diagnostic test (n)	A medical test that determines whether a patient has a particular medical problem. Diagnostic tests are often used when providers have a specific reason to believe that the medical problem may be present. Compare this to screening tests, which may be given routinely even if the provider has no reason to believe the patient has a certain problem. Screening tests often only report whether a patient is at an increased risk for the medical problem in question, whereas diagnostic tests report whether the problem is actually present.	
DNA (Deoxyribonucleic Acid) (n)	The molecule that carries the genetic information of a cell. It provides the instructions used in the development, functioning and reproduction of the organism of which it is a part. The DNA is organized into chromosomes.	
DNA marker (also called a "genetic marker") (n)	A readily recognizable genetic trait, gene, or DNA segment.	
DNA mutation (n)	A change in the typical sequence of the chemicals that make up the DNA, like the change in the order of letters in a word.	
	Mutations or variants are often compared to misspelled words because chemicals that make up the DNA sequence are not in the expected order.	

Term	Definition	Translation
DNA sequence (n)	The exact arrangement of the chemicals that make up a section of DNA.	
DNA sequencing (n)	The laboratory technique used to determine the exact arrangement of the chemicals that make up a section of DNA. This is one type of genetic testing.	
Dominant (adj)	A genetic trait in which one copy of the gene is sufficient for a trait to be expressed. In a dominant genetic condition, if one copy of the gene has a mutation, the person will be affected with the condition.	
Donor egg or sperm (n)	An egg (singular) or sperm (plural) donated by one person to be joined under laboratory conditions and implanted in a woman's uterus. The donor egg or sperm may come from the woman or man who will raise any resulting child, or they may come from a third party.	
Down syndrome (Trisomy 21) (n)	A genetic condition in which there are three copies of chromosome number 21 instead of two.	
	Down syndrome causes a distinct facial appearance, intellectual disability, developmental delays, and may be associated with thyroid or heart disease.	
Duplication (n)	Having an extra section of genetic information (DNA). A duplication occurs when part of a chromosome is copied abnormally, resulting in extra genetic material.	
Edwards syndrome (Trisomy 18) (n)	A genetic condition in which there are three copies of chromosome number 18 instead of two. Trisomy 18 is a very severe condition that causes problems with the brain, the heart, the kidneys, and the digestive tract. Most children affected by trisomy 18 die before or soon after birth.	
Egg (n)	The reproductive cells of a woman. When fertilized by sperm, the egg will grow into an embryo. Also called an "ovum," plural "ova."	
Embryo (n)	An unborn mammal, between conception and 8 weeks of gestation.	
Environmental causes or factors (n)	Causes of illness that come from a person's surroundings, not from their genetic make-up. Environmental causes of illness include infection, trauma, diet, exposure to toxins, etc. Some environmental causes of illness can be avoided or changed, while genetic causes cannot.	

Term	Definition	Translation
Exome sequencing (n)	A genetic test in which a patient's exons are sequenced and studied.	
Exon (n)	The part of the genome that codes for proteins. The majority of deleterious mutations occur in exons.	
Fallopian tube (n)	The tube that connects an ovary to the uterus.	
False negative (n)	A test result that finds no evidence of a condition when the condition actually does exist. For example, a false negative on a pregnancy test finds that the woman is not pregnant when, in fact, she is pregnant.	
False positive (n)	A test result that finds evidence of a condition when the condition does NOT actually exist. For example, a false positive on a pregnancy test finds that the woman is pregnant when, in fact, she is not.	
Familial (adj)	Belonging to a family, e.g., a familial trait is a trait that is shared by family members and may be due to genetic or environmental factors or both.	
Family history (n)	The medical history of the members of a biological family.	
Fertilization (n)	The joining of an egg and sperm to create the first cell that will develop into an embryo, then fetus, then baby.	
Fetal surgery (n)	Surgery conducted on a fetus while it is still in the uterus.	
Fetus(n)	An unborn mammal, between 8 weeks of gestation to birth.	
First trimester screening (n)	A blood test and ultrasound conducted at 10-13 weeks of pregnancy to screen for Down syndrome (trisomy 21), Edwards syndrome (trisomy 18), and certain other fetal problems.	
Flip a coin (v)	A random decision-making tool used in the U.S. While a coin is flipped into the air and caught, a person predicts whether it will fall with the "heads" side up or the "tails" side up. If the coin falls as predicted, the person "wins." This expression is often used as a metaphor for any outcome that is random and has two possible outcomes, and to describe a situation in which each outcome is as likely as the other.	

Term	Definition	Translation
Fragile X syndrome (n)	A genetic condition that affects the FMR1 gene so that it does not function properly. This syndrome causes intellectual disability, behavioral and learning challenges and various physical characteristics that are not life threatening.	
Fragment (n)	A small piece; an incomplete part of a whole.	
Gene (n)	A specific sequence of DNA that codes for one or many functions within the cell and body.	
Gene copy (n)	In human cells, DNA is arranged in 23 pairs of chromosomes, for a total of 46 chromosomes. These chromosomes contain all the genes that make up the DNA. One chromosome of each pair is inherited from each parent. As such, individuals have two copies of every gene, one inherited from the mother and one inherited from the father.	
Normal gene copy (n)	A normal copy of a gene is one that is found most frequently in the population. Also called the "working copy."	
Altered gene copy (n)	An altered copy of a gene is one that differs from the working copy and is considered to be "abnormal." Also called the "non-working copy."	
Gene deletion (n)	Having a piece of genetic information missing from a gene.	
Gene panel testing (n)	Genetic testing that looks at 2 or more specific genes known to be associated with particular conditions.	
Gene variant (n)	A change from the typical DNA sequences. A genetic variant can be benign, deleterious or of uncertain significance. Also called "mutation", although "genetic variant" is becoming the more common usage.	
General population (n)	"Most people."	
	For example, if you have the same risk of getting cancer as the general population, that means that you have the same chance of getting cancer as everyone else. This is in contrast to a "high risk population" who has a greater chance of getting cancer than everyone else.	

Term	Definition	Translation
Generation (n)	The people who constitute a single step in a line of descent from an ancestor; a group of people born and living more or less at the same time. Example: You, your brothers, and sisters, all your	
	spouses and your cousins are in the same generation. Your parents, your aunts, and uncles and all their spouses form a previous generation. Your grandparents, their siblings, and spouses form an even earlier generation. Your children and nieces and nephews form a later generation.	
Genetic alphabet (n)	A group name for four chemical bases, or "nucleotides," that form the basis of all genetic material. They are adenine (A), thymine (T), cytosine (C), and guanine (G).	
Genetic counseling (n)	A discussion with a medical professional with expertise in genetics about the basic concepts of genetics, genetic conditions, the chances of being affected by a genetic condition or having a child with a genetic condition, and genetic testing and treatment.	
Genetic counselor (n)	A healthcare professional with a specialized graduate degree who works with people who have concerns about genetic conditions in their family. Genetic counselors provide information about genetic conditions, help patients understand their chances of being affected by a genetic condition or having a child with a genetic condition, and help them make informed decisions about testing and treatment. Genetic counselors also provide emotional support to patients and families.	
Genetic discrimination (n)	Occurs when people are treated differently by their employer or insurance company due to their genetic makeup. There are federal and state laws that help protect against genetic discrimination.	
Genetic factors (n)	Specific aspects of a person's genetic make-up that influence that person's health and development.	
Genetic information (n)	The instructions encoded in DNA, which tells every cell in a body how to grow, what to do and how to reproduce.	
Genetic material (n)	All the parts of a cell that carry genetic information. Genetic material could include genes, parts of genes, a group of genes, a DNA molecule, a fragment of DNA, a group of DNA molecules, or the entire set of genetic instructions.	

Term	Definition	Translation
Genetic predisposition (n)	An increased chance of a person developing a certain trait or disease based on that person's particular genetic makeup.	
Genetic test (n)	A laboratory test designed to determine if a person has a gene mutation or a typical DNA sequence.	
Single site gene testing (n)	Single site gene testing: testing at a specific point in the genome. This is usually done to look for a mutation found in a family member or when one specific mutation is known to cause a genetic condition.	
Individual gene testing (n)	Individual gene testing: genetic analysis of the entirety of one specific gene.	
Gene panel test (n)	Gene panel test: analysis of several genes that have been shown to be associated with a particular condition, e.g., breast cancer gene panel.	
Microarray (n)	Microarray: genetic analysis that looks for unexpected numbers of specific genes (two copies of every gene code is usually expected).	
Exome sequencing (n)	Exome testing: a genetic test in which the exons of a patient's genetic code are sequenced and studied.	
Genome sequencing (n)	Genome sequencing: a genetic test in which the entire genetic code (the genome) of a patient is mapped out and studied.	
Genetic trait (n)	A characteristic within a family that is passed down from parent to child through their DNA.	
Geneticist (n)	A doctor or scientist who studies genetics.	
Genetics (adj)	The science of how an organism's genes interact with the environment to produce certain traits.	
Genetics (n)	The field of science and medicine that studies the biologic basis of heredity (how traits are passed from one generation to another) and how the instructions for life are used by all living organisms.	
Genome (n)	The entire genetic code of an individual.	
Genome sequencing (n)	A genetic test in which the entire genetic code of a patient is mapped out and studied.	

Term	Definition	Translation
Germline testing (n)	Germline testing refers to the analysis of a person's DNA, which he or she inherited from his or her parents.	
	In biology and genetics, the germline is the group of cells that will pass on the genetic material to children, in other words, the cells from which the eggs and sperm come.	
Guardianship (n)	A legal arrangement, supervised by a court, in which one person takes responsibility for a minor or an adult with a disability.	
Helix (n)	Something that is spiral in form. DNA is often described as being a "double helix."	
Hereditary (adj)	Passed down from parent to child.	
Hereditary material (n)	Genetic material that is passed down from parent to child.	
Huntington's disease (n)	A neurological disorder that causes physical and cognitive decline. Cognitive symptoms include problems with reasoning and controlling emotions. Physical symptoms include abnormal movements that cannot be controlled by the individual. This disorder is linked to a dominant gene variant, which means that a person inheriting only one copy of the affected gene will develop the disorder.	
Individualized Education Plan (IEP) (n)	A customized plan written and updated yearly for every child in public school special education classes. The IEP addresses current levels of educational performance, goals, and the special education and services that the child will need to meet those goals.	
In Vitro Fertilization (IVF) (n)	The fertilization of an egg by a sperm outside of a woman's body.	
	The process involves extracting eggs from a woman's ovaries, collecting sperm from a man, and combining a sperm and egg in a laboratory dish. The resulting fertilized egg is usually then implanted in a woman's uterus so that it can develop into a baby.	
Individual gene testing (n)	Genetic analysis of the entirety of one specific gene.	
Infertility (n)	The inability to have children.	

Term	Definition	Translation
Informed consent (n)	The process of agreeing to a procedure or course of treatment after understanding what the procedure/treatment entails, the potential risks and benefits associated with it, and the other options available.	
Inheritance pattern (n)	The manner in which a particular genetic trait or disorder is passed from a parent to a child, e.g., autosomal dominant or recessive, X-linked dominant or recessive, or multifactorial.	
Inherited (adj)	Passed down from parent to child.	
Insertion (n)	An extra segment of DNA added in at a place where it is not usually found.	
Intellectual disability (n)	A condition, varying in severity, in which a person has significant impairments in mental abilities, social skills, and core functions of daily living compared to others their age.	
Klinefelter's syndrome (47, XXY) (n)	A genetic condition in which a male has two copies of the X chromosome and one copy of the Y chromosome; compared to the typical chromosome makeup where a male has one X chromosome and one Y chromosome.	
Lynch syndrome (n)	An inherited genetic disorder that increases a person's risk of getting cancer of the colon, rectum, uterus, ovaries, and other cancers.	
Marker chromosome (n)	A small extra fragment of a chromosome detected when doing a chromosome test like a karyotype. Marker chromosomes can sometimes cause health or development problems, depending on how much and what genetic material is contained within.	
Maternal Serum Screening (MSS or Maternal serum alpha- fetoprotein test) (n)	A blood test available to pregnant women that identifies elevated risks for Down syndrome, trisomy 18 and neural tube defects. Usually conducted in the second trimester, between 14-20 weeks.	
Metastasis (n)	The spreading of cancer from one organ to another.	

Term	Definition	Translation
Microarray (n)	A high-resolution chromosome test that provides more information than a basic karyotype. A microarray measures the amount of chromosome material in a sample and can detect both large changes (e.g., if there is an extra or missing chromosome) as well as small changes (e.g., if there are very small pieces of chromosomes that are extra or missing). It does not look at the visual appearance or arrangement of chromosomes but measures the amount of genetic material.	
Miscarriage (n)	The spontaneous (not intentional) loss of a pregnancy. See "abortion." Also called pregnancy loss.	
Molecule (n)	The smallest unit of a chemical compound that still has the properties of that compound. For example, a molecule of water is made up of two hydrogen atoms and one oxygen atom. Separately, they are just atoms, but when bonded together, they make a water molecule.	
Mosaicism (n)	A condition in which some, but not all, cells in a sample show a genetic difference.	
Motor abilities (n)	The ability to move and use one's muscles.	
Multifactorial (adj)	Due to a combination of genetic and non-genetic (environmental, hormonal, etc.) risk factors that act together to determine risk.	
Mutation (n)	A change in a gene, usually deleterious. See "genetic variant."	
Non-Invasive Prenatal Testing (NIPT) (n)	A blood test available to pregnant women that identifies elevated risk for certain genetic conditions in the fetus. This test focuses on fragments of DNA from placental cells – which carry the fetus' genetic make-up – that are found in the mother's bloodstream.	
Nuchal thickening (n)	There is a pocket of fluid at the back of the neck of a fetus which can be measured in an ultrasound between 10-14 weeks gestation (called the nuchal translucency). If there is a large amount of fluid at this point, or if later in pregnancy the neck skin itself appears to be thicker, this is associated with a higher risk of chromosome problems and other rare genetic conditions.	

Term	Definition	Translation
Open Neural Tube Defect (ONTD) (n)	A birth defect of the spine, spinal cord, or brain that results from a hole in the spinal column not closing up when appropriate during early fetal development. The two most common types of ONTDs are spina bifida and anencephaly.	
Ova (n)	See "egg."	
Ovary (n)	The organ in a woman that stores and releases eggs. There are normally two.	
Packets of genetic information (n)	A phrase genetic counselors use to describe genes or chromosomes.	
Pathogenic (adj)	Disease causing.	
Pedigree (n)	A family tree that can be used to trace the inheritance of specific genetic traits.	
Penetrance (n)	The probability that a specific genetic trait will be expressed if a person carries a mutation. "Complete penetrance" means that everyone who carries a particular gene mutation will show the trait related to that altered gene. "Incomplete penetrance" means that only some of the people who have the altered gene will actually show the related trait.	
Placenta (n)	The organ that develops together with an embryo in a pregnant woman's uterus to nourish the embryo through the umbilical cord.	
Presymptomatic (adj)	Before symptoms appear.	
Prognosis (n)	The most likely outcome of a disease process.	
Protein (n)	A molecule made up of chains of <i>amino acids</i> . Proteins do most of the work in cells and are required for the structure, function, and regulation of the body's tissues and organs. Genes determine how specific amino acids are put together to form a specific protein.	
Random (adj)	Happening in an unpredictable way.	

Term	Definition	Translation
Recessive (adj)	A genetic trait in which both copies of a gene need to be altered for the trait to be expressed; recessive traits are typically inherited from both parents, each one contributing one copy of the gene in question. In prenatal genetics, recessive inheritance refers to	
	conditions in which both copies of a particular gene pair need to be altered to develop a specific disease or condition.	
Red flag (n)	A warning sign or a clue.	
Replicate (v)	То сору.	
Reproductive history (n)	The experiences a woman has had related to pregnancy and childbirth. A reproductive history usually includes age at the onset of menses (have a monthly period), age at the beginning of sexual intercourse, age at first conception, number of pregnancies, number of live births, number of miscarriages, number of abortions.	
Ring chromosome (n)	An abnormal formation of a chromosome in which the ends of two of the chromosome's arms have linked together to form a ring.	
Risk (n)	The chance that something will happen; in the case of prenatal genetics, the chance that the child will have a genetic condition.	
At risk (adj)	Has some possibility of having a genetic condition.	
At average risk (adj)	Has the same possibility of having a genetic condition as the general population.	
At high risk (adj)	Has a greater possibility of having a genetic condition than the general population.	
At higher risk than "X" (adj)	Has a greater possibility of having a genetic condition than "X."	
Risk factor (n)	A circumstance that increases the risk of having a genetic condition.	
Runs in the family (v)	Passed down from parents to children to grandchildren.	
Saliva (spit) sample (n)	A way to collect DNA from the cells in saliva (spit).	

Term	Definition	Translation
Screen (v)	To assess a person's risk for a particular health problem, even if the person has no symptoms to suggest that they have the problem.	
Screening test (n)	A test that looks to detect a particular health problem, even if a patient has no symptoms of that particular condition.	
	Prenatal screening tests aren't looking for genetic changes. They are designed to look at hormones, chemicals, other things that indicate a fetus might have a higher risk of a certain condition. They are not DNA results, do not give a definite answer about whether someone is affected, and do not provide a firm diagnosis. They are usually reported as a risk number (e.g., a 1 in chance), and if this number is above a certain cut-off point, the result will be categorized as "screen positive" or "screen negative."	
Semen (n)	A liquid produced by the testes, the prostate gland, the seminal vesicle, and the bulbourethral gland that carries, nourishes, and protects sperm cells on their way to fertilizing an egg. Also called seminal fluid.	
Seminal fluid (n)	See "semen."	
Sensory abilities (n)	The ability to see, hear, touch, taste, and smell.	
Sickle cell disease (n)	A genetic disorder caused by a mutation in a gene that helps to make hemoglobin, an important part of red blood cells. In sickle cell disease, red blood cells can become sickle-shaped (instead of round as they should be). These affected blood cells do not function properly, and die early, leaving a shortage of red blood cells, and can block blood flow causing pain.	
Single site analysis (n)	A genetic test that looks at only the part of a gene that is known to be associated with a certain disease.	
Somatic genetic changes (n)	Genetic changes in an individual that occur after conception. For example, exposure to the ultraviolet rays in sunlight can cause problems with replication of genetic material in cells, leading to changes in the gene that controls cell growth and then to skin cancer. These changes are somatic genetic changes.	
Special education (n)	Education that is tailored to individuals with developmental delays or other learning disabilities.	
Sperm (n)	The reproductive cells of the man. When sperm fertilize a woman's egg, a baby develops.	

Term	Definition	Translation
Sperm donation (n)	The process through which a man allows his sperm to be collected and used to fertilize the eggs of a woman who is not his sexual partner.	
Spina bifida (n)	A congenital defect of the spine in which part of the spinal cord is exposed through an opening in the bone structure. This may result in nerve damage and some degree of paralysis in the legs.	
Statistically significant (adj)	A measurement of whether the findings of research are meaningful. Refers to the likelihood that a relationship between two factors is linked by something other than chance.	
Supportive seating (n)	Seating, such as a wheelchair, that helps an individual function in their environment and that prevents worsening of their condition.	
Teratogen (n)	Something that has the potential to disturb the normal development of an embryo or fetus.	
Termination of pregnancy (n)	See "abortion."	

Term	Definition	Translation
Test result (n)		
Negative (adj)	A negative result on a genetic test means that the laboratory did not find the specific genetic change that the test was designed to identify.	
True negative (n)	When a patient is being tested to see if he or she has a specific genetic change that is present in other family members, a negative result can be considered a "true negative." This means that instead of having an increased risk for cancer like other family members, they have the same risk as everyone else in the general population.	
Positive (adj)	A positive result on a genetic test means that the laboratory did find a specific genetic change that is associated with a hereditary cancer syndrome (a deleterious mutation). In cancer genetics, that means that the patient has an increased risk of getting the cancer(s) associated with the gene.	
Variant of uncertain significance (n)	If the laboratory finds a genetic change for which currently there is not enough information to know if this change is problematic or not, it reports a "variant of uncertain significance." Basically, this means that the laboratory found a genetic change, but they don't know what it means. All of these variants will eventually be reclassified as either a "positive" or "negative" result. Most become "negative" and are thought to represent natural variation between individuals.	
Polymorphism (n)	Everyone has some degree of commonly occurring genetic changes that are not associated with cancer. If the test finds this sort of change, it reports a "polymorphism."	
Testes (n)	The organs in a man that create and release sperm. Also called "testicles." Singular testis.	
Toxic exposure (n)	Contact with something that is harmful or poisonous.	
Trait (n)	A characteristic of a person. In genetics, traits are aspects of a person defined or influenced by their genetic code, e.g., eye color, blood type, risk for certain diseases.	

Term	Definition	Translation
Translocation (n)	Translocations are a type of chromosome rearrangement. They can be "balanced," meaning that two chromosomes have pieces that have "swapped" with each other, but there is still the correct amount of chromosome material. Balanced translocations do not usually cause medical or development problems. They can also be "unbalanced" where pieces of chromosomes have traded places, but there is missing or extra chromosome material. Unbalanced translocations will often cause medical and development problems.	
Typo (n)	A mistyped word; shortened slang for "typographical error." In genetic counseling "a typo" is commonly used as an analogy to refer to mutations or variants in a gene.	
Ultrasound (n)	An imaging method that uses high frequency sound waves to create a picture of something inside the body, such as a fetus or baby.	
Uterus (n)	The organ in a woman's lower abdomen in which a fertilized egg develops into an embryo, then a fetus, then a baby. Also called "womb."	
Vagina (n)	The tube-like part of the female reproductive system that extends from the cervix to the outside of the body.	
Variant (n)	A version of something that differs from the norm. So, a genetic variant is a change to the usual genetic sequence.	
Vas deferens (n)	The tubes that lead from the testes to the urethra, through which sperm is ejaculated.	
X-linked (adj)	A trait that is influenced or determined by a gene on the "X" chromosome.	

Family Relationships

For genetic counselors, it is very important to know if a relative is a blood relative or a relative by marriage.

Term	Definition	Translation
Parent	Your mother or father.	
Mother	For genetic purposes, the woman whose egg was fertilized and grew to be you.	
Mother-in-law	Your husband or wife's mother.	
Stepmother	Your father's wife who is not your biological mother.	
Adoptive mother	A woman who is not your biological mother but who accepted legal responsibility for and raised you.	
Godmother	A woman chosen by your parents to be your spiritual guide in life, named as such through a ceremony in the Catholic church.	
Father	For genetic purposes, the man whose sperm fertilized the egg that grew to be you.	
Father-in-law	Your husband or wife's father.	
Stepfather	Your mother's husband who is not your biological father.	
Adoptive father	A man who is not your biological father but who accepted legal responsibility for and raised you.	
Godfather	A man chosen by your parents to be your spiritual guide in life, named as such through a ceremony in the Catholic church.	
Aunt	Your mother or father's sister or sister-in-law.	
Maternal aunt	Your mother's sister or sister-in-law.	
Paternal aunt	Your father's sister or sister-in-law.	
Uncle	Your mother or father's brother or brother-in-law.	
Maternal uncle	Your mother's brother or brother-in-law.	
Paternal uncle	Your father's brother or brother-in-law.	

Term	Definition	Translation
Cousin	Usually understood to be a first cousin.	
First cousin	Your aunt or uncle's child.	
First cousin once removed	Your aunt or uncle's grandchild or your first cousin's child	
Second cousin	The children of first cousins are second cousins to each other.	
Grandparent	Your parent's father or mother.	
Grandmother	Your mother or father's mother.	
Maternal grandmother	Your mother's mother.	
Paternal grandmother	Your father's mother.	
Grandfather	Your mother or father's father.	
Maternal grandfather	Your mother's father.	
Paternal grandfather	Your father's father.	
Great aunt	Your mother or father's aunt.	
Maternal great aunt	Your mother's aunt.	
Paternal great aunt	Your father's aunt.	
Great uncle	Your father or mother's uncle.	
Maternal great uncle	Your mother's uncle.	
Paternal great uncle	Your father's uncle.	
Great grandparents	The parents of any of your grandparents.	
Great grandmother	The mother of any of your grandparents.	
Great grandfather	The father of any of your grandparents.	

Term	Definition	Translation
Siblings	The children of your father and mother.	
Half siblings	Siblings (brothers and sisters) who have either the same mother and different fathers, or the same father but different mothers. Half siblings share some genetic similarity with you.	
Stepsiblings	The children of your stepmother but not your father; or the children of your stepfather but not your mother. Stepsiblings do not share any genetic similarity with you.	
Twins	Two siblings born at the same time.	
Identical twins	Twins who developed from the same egg and sperm, meaning that they are genetically identical. Also called monozygotic twins.	
Fraternal twins	Twins who developed from different eggs and sperm, meaning that they are genetically different and have the same number of shared genes as any other sibling. Also called dizygotic twins.	
Triplets	Three siblings born at the same time.	
Sister	A sibling who is a girl.	
Sister-in-law	Your brother's wife or your husband's sister.	
Half sister	A girl who is the either the child of your father with a different mother, or the child of your mother with a different father.	
Stepsister	The daughter of your stepmother or stepfather and therefore not biologically related to you.	
Brother	A sibling who is a boy.	
Brother-in-law	Your sister's husband or your wife's brother.	
Half brother	A boy who is the either the child of your father with a different mother, or the child of your mother with a different father.	
Stepbrother	The son of your stepmother or stepfather and therefore not biologically related to you.	
Niece	Your sibling's daughter.	
Nephew	Your sibling's son.	

Term	Definition	Translation
Spouse	Your husband or wife.	
Wife	The woman to whom you are married.	
Husband	The man to whom you are married.	
Children	Genetically speaking, the people who are produced from your egg or sperm.	
Daughter	Your child who is a girl.	
Daughter-in-law	Your son's wife.	
Stepdaughter	Your spouse's daughter who is not your biological child.	
Adoptive daughter	A girl for whom you have accepted legal responsibility and raised even though she is not your biological child.	
Goddaughter	A girl for whom you have accepted spiritual responsibility within a ceremony of the Catholic church; she may or may not be biologically related to you.	
Son	Your child who is a boy.	
Son-in-law	Your daughter's husband.	
Stepson	Your spouse's son who is not your biological child.	
Adoptive son	A boy for whom you have accepted legal responsibility and raised even though he is not your biological child.	
Godson	A boy for whom you have accepted spiritual responsibility within a ceremony of the Catholic church. He may or may not be biologically related to you.	
Ancestor	A person from whom you are descended, usually more remote than a grandparent.	
Fiancé(e)	The person whom you have promised to marry.	
	Man: fiancé	
	Woman: fiancée	

Term	Definition	Translation
Domestic partner	The person with whom you are living and have an intimate relationship, but to whom you are not married.	
Divorced	Having ended a marriage.	
Engaged	Having promised to marry someone.	
Widow	A woman whose husband has died.	
Widower	A man whose wife has died.	
Relative	A person in your family.	
Blood relation	A person who is related to you by blood not marriage, e.g., your sister, but not your sister-in-law; your mother but not your stepmother; your daughter but not your adopted daughter.	
Next of kin	Your closest living blood relation.	

Glossary of Pediatric Genetics Terms - Language

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The National Coordinating Center for the Regional Genetics Networks

Glossary of Prenatal Genetics Terms

English – Language of Translation

2022 Edition

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The Cross Cultural Health Care
Program (CCHCP)



Preface

The seven Regional Genetics Networks (RGNs), the National Genetics Education and Family Support Center (Family Center), and the National Coordinating Center for the Regional Genetics Networks (NCC) mission is to improve access to quality genetic services for medically underserved populations. This bilingual glossary of terms related to Prenatal Genetics is provided to help enable access to prenatal medical services to limited English proficiency populations.

How to Use the Glossary

This glossary is organized alphabetically in English. The first column shows the term in English. The second column has the definition of the term in English. The third column has the translation of the term.

At the end of the translations of the prenatal genetics terms are:

- A glossary of translations for family relationships terms. It is very important for genetic counselors to know if a relative is a blood relative or a relative by marriage.
- Examples of analogies used in genetics. Genetic counselors often use analogies to explain complex ideas in genetics. An analogy explains a new concept by comparing it to something more common.

Glossary of Terms

English – Spanish

(n) – noun (v) – verb (adj) – adjective (adv) – adverb

Term	Definition	Translation
Abnormal (adj)	Different from what is considered normal, average, or expected, e.g., a gene sequence that is different than that found in most people.	
Abortion (n)	The deliberate ending of a pregnancy by the removal of an embryo or fetus from the womb.	
	Technically, this is called an "induced abortion" in order to distinguish it from a "spontaneous abortion" which is also called a miscarriage or stillbirth. Also called a "termination of pregnancy" or just "termination."	
Adoption (n)	The process of legally taking someone else's child into your family to raise as your own child.	
Amniocentesis (n)	A procedure by which a sample of amniotic fluid is withdrawn from the amniotic sac. This is usually done by inserting a long needle through the abdominal and uterine walls with the guidance of ultrasound.	
Amniotic fluid (n)	The liquid that surrounds a fetus as it develops in the uterus. This liquid contains skin cells that have sloughed off the fetus as well as other fetal cells.	
Anencephaly (n)	A neural tube defect that results in insufficient brain growth in a fetus. Babies with anencephaly usually die soon after birth or are stillborn.	
Benign (adj)	Something that does not threaten health or life. May refer to a type of change to the DNA that does not create health consequences.	
	E.g., when discussing cancer, "benign" means "not cancerous."	
Birth canal (n)	The passageway from the uterus through the cervix, the vagina, and the vulva through which a baby passes during the birth process.	

Term	Definition	Translation
Birth defect (n)	A problem or physical difference with how the body works that is present at birth. Birth defects can be caused by genetic abnormalities, environmental influences, random chance, or by circumstances related to the birth process. Also called "congenital malformation" or "congenital anomaly."	
Blood test (n)	A test in which blood is drawn (usually from the patient's arm) and sent to a laboratory for analysis. In genetics, blood tests may provide cells for genetic sequencing, or blood samples may be used to assess things like protein or hormone levels.	
Buccal swab (n)	A way to collect DNA from the cells on the inside of a person's cheek.	
Carrier (n)	A person who has a genetic mutation in one of their two copies of a particular gene that is associated with a genetic condition. The other copy of the gene does not have a mutation, so that person usually does not have any medical issues related to the gene change.	
Cell (n)	The smallest (microscopic) functional unit of living organisms. All living things are composed of one or more cells. Within each cell are the structures needed for the cell to process energy, dispose of waste, reproduce, and perform specialized functions.	
Cervix (n)	The narrow lower part of the uterus that connects to the vagina.	
Chorionic Villus Sampling (CVS) (n)	A procedure during pregnancy in which a sample of cells from the placenta is removed to check for possible genetic abnormalities.	
Chromosome (n)	Thread-like structures located inside the nucleus of cells. In humans, there are 23 pairs of chromosomes, for a total of 46 chromosomes. Each chromosome is made of a long strand of DNA, which carries genetic information.	
Chromosome test / Karyotype (n)	A test that looks at the number and types of chromosomes in a cell.	
Cleft lip (n)	A congenital malformation that occurs when the upper lip does not form properly, causing an opening in the upper lip that can extend to the nose. This can occur together with a cleft palate or on its own.	

Term	Definition	Translation
Cleft palate (n)	A congenital malformation where the roof of the mouth (palate) does not form properly, resulting in an opening into the nasal cavity. This can occur together with a cleft lip or on its own.	
Clinically significant (adj)	If a test result is "clinically significant," the result indicates a medical problem that can impact a person's life.	
Condition (e.g., "genetic condition") (n)	A long-term medical health issue.	
Consanguinity (n)	When parents are blood relatives to each other.	
Cystic fibrosis (n)	A progressive genetic condition that affects the exocrine glands (the glands that make sweat and digestive juices) and causes the production of thick, sticky mucus. This mucus blocks the pancreatic duct, the intestines, and the lungs, leading to persistent respiratory infections.	
Deleterious mutation/ Disease-causing mutation (n)	A change in a person's DNA that may cause a medical condition.	
Deletion (n)	Having a section of genetic information (DNA) missing.	
Diagnostic test (n)	A medical test that determines whether a patient has a particular medical problem. Diagnostic tests are often used when providers have a specific reason to believe that the medical problem may be present. Compare this to screening tests, which may be given routinely, even if the provider has no reason to believe the patient has a specific problem. Screening tests often only report whether a patient is at an increased risk for the medical problem in question, whereas diagnostic tests report whether the problem is actually present.	

Term	Definition	Translation
DNA (Deoxyribonucleic Acid) (n)	The material that carries the genetic information of a cell. It provides the instructions used in the development, functioning and reproduction of the organism of which it is a part.	
	If you could stretch out the DNA of a chromosome and look at it through a microscope, it would look like a long ladder that is twisted into a spiral. The 'sides' of the ladder are made up of alternating phosphate and sugar groups. The 'rungs' are various combinations of two nitrogen bases: Adenine-Thymine and Cytosine-Guanine.	
	Individual sections of DNA that code for specific traits/functions are called genes.	
DNA marker (also called a "Genetic marker") (n)	A readily recognizable genetic trait, gene, or DNA segment.	
DNA mutation (n)	A change in the typical sequence of the chemicals that make up the DNA, like the change in the order of letters in a word.	
	Mutations or variants are often compared to misspelled words because chemicals that make up the DNA sequence are not in the expected order.	
DNA sequence (n)	The exact arrangement of the chemicals that make up a section of DNA.	
DNA sequencing (n)	The laboratory technique used to determine the exact arrangement of the chemicals that make up a section of DNA. This is one type of genetic testing.	
Dominant (adj)	A genetic trait in which one copy of the gene is sufficient for a trait to be expressed. In a dominant genetic condition, if one copy of the gene has a mutation, the person will be affected with the condition.	
Donor egg or sperm (n)	An egg (singular) or sperm (plural) donated by one person to be joined under laboratory conditions and implanted in a woman's uterus. The donor egg or sperm may come from the woman or man who will raise any resulting child, or they may come from a third party.	

Term	Definition	Translation
Down syndrome (Trisomy 21) (n)	A genetic condition in which there are three copies of chromosome number 21 instead of two.	
	Down syndrome causes a distinct facial appearance, intellectual disability, developmental delays, and may be associated with thyroid or heart disease.	
Duplication (n)	Having an extra section of genetic information (DNA). A <i>duplication</i> occurs when part of a chromosome is copied abnormally, resulting in extra <i>genetic</i> material.	
Edwards syndrome (Trisomy 18) (n)	A genetic condition in which there are three copies of chromosome number 18 instead of two.	
	Trisomy 18 is a very severe condition that causes problems with the brain, the heart, the kidneys, and the digestive tract. Most children affected by trisomy 18 die before or soon after birth.	
Egg (n)	The reproductive cells of a woman. When fertilized by sperm, the egg will grow into an embryo. Also called an "ovum," plural "ova."	
Embryo (n)	An unborn mammal, between conception and 8 weeks of gestation.	
Fallopian tube (n)	The tube that connects an ovary to the uterus.	
False negative (n)	A test result that finds no evidence of a condition when the condition actually does exist. For example, a false negative on a pregnancy test finds that the woman is not pregnant when, in fact, she is pregnant.	
False positive (n)	A test result that finds evidence of a condition when the condition does NOT actually exist. For example, a false positive on a pregnancy test finds that the woman is pregnant when, in fact, she is not.	
Familial (adj)	Belonging to a family, e.g., a familial trait is a trait that is shared among family members and may be due to genetic or environmental factors or both.	
Family history (n)	The medical history of the members of a biological family.	
Fertilization (n)	The joining of an egg and sperm to create the first cell that will develop into an embryo, then fetus, then baby.	
Fetal surgery (n)	Surgery conducted on a fetus while it is still in the uterus.	

Term	Definition	Translation
Fetus (n)	An unborn mammal, between 8 weeks of gestation to birth.	
First trimester screening (n)	A blood test and ultrasound conducted at 10-13 weeks of pregnancy to screen for Down syndrome (trisomy 21), Edwards syndrome (trisomy 18), and certain other fetal problems.	
Flip a coin (v)	A random decision-making tool used in the U.S. While a coin is flipped into the air and caught, a person predicts whether it will fall with the "heads" side up or the "tails" side up. If the coin falls as predicted, the person "wins." This expression is often used as a metaphor for any outcome that is random and has two possible outcomes, and to describe a situation in which each outcome is as likely as the other.	
Fragile X syndrome (n)	A genetic condition that affects the FMR1 gene so that it does not function properly. This syndrome causes intellectual disability, behavioral and learning challenges and various physical characteristics that are not life threatening.	
Fragment (n)	A small piece; an incomplete part of a whole.	
Gene (n)	A specific sequence of DNA that codes for one or many functions within the cell and body.	
General population (n)	"Most people."	
	For example, if you have the same risk of getting cancer as the general population, that means that you have the same chance of getting cancer as everyone else. This is in contrast to a "high risk population" who has a greater chance of getting cancer than everyone else.	
Generation (n)	The people who constitute a single step in a line of descent from an ancestor; a group of people born and living more or less at the same time.	
	Example: You, your brothers, and sisters, all your spouses and your cousins are in the same generation. Your parents, your aunts, and uncles and all their spouses form a previous generation. Your grandparents, their siblings, and spouses form an even earlier generation. Your children and nieces and nephews form a later generation.	

Term	Definition	Translation
Genetics (adj)	The science of how an organism's genes interact with the environment to produce certain traits.	
Genetic counseling (n)	A discussion with a medical professional with expertise in genetics about the basic concepts of genetics, genetic conditions, the chances of being affected by a genetic condition or having a child with a genetic condition, and genetic testing and treatment.	
Genetic counseling intern (n)	A genetic counseling student who has not yet completed his/her academic studies and is now practicing under the supervision of a more experienced counselor in preparation for providing genetic counseling services independently after obtaining his/her graduate degree.	
Genetic counselor (n)	A healthcare professional with a specialized graduate degree who works with people who have concerns about genetic conditions in their family. Genetic counselors provide information about genetic conditions, help patients understand their chances of being affected by a genetic condition or having a child with a genetic condition, and help them make informed decisions about testing and treatment. Genetic counselors also provide emotional support to patients and families.	
Genetic discrimination (n)	Occurs when people are treated differently by their employer or insurance company due to their genetic makeup. There are federal and state laws that help protect against genetic discrimination.	
Genetic factors (n)	Specific aspects of a person's genetic make-up that influence that person's health and development.	
Genetic information (n)	The instructions encoded in DNA, which tells every cell in a body how to grow, what to do and how to reproduce.	
Genetic material (n)	All the parts of a cell that carry genetic information. Genetic material could include genes, parts of genes, a group of genes, a DNA molecule, a fragment of DNA, a group of DNA molecules, or the entire set of genetic instructions.	
Genetic predisposition (n)	An increased chance of a person developing a certain trait or disease based on that person's particular genetic makeup.	
Genetic test (n)	A laboratory test designed to determine if a person has a change to their DNA.	

Term	Definition	Translation
Genetic trait (n)	A characteristic within a family that is passed down from parent to child through their DNA.	
Genetic variant (n)	A change from the typical DNA sequences. A genetic variant can be benign, deleterious or of uncertain significance. Also called "mutation," although "genetic variant" is becoming the more common usage.	
Geneticist (n)	A doctor or scientist who studies genetics.	
Hereditary (adj)	Passed down from parent to child.	
Hereditary material (n)	Genetic material that is passed down from parent to child.	
In Vitro Fertilization (IVF) (n)	The fertilization of an egg by a sperm outside of a woman's body.	
	The process involves extracting eggs from a woman's ovaries, collecting sperm from a man, and combining a sperm and egg in a laboratory dish. The resulting fertilized egg is usually then implanted in a woman's uterus so that it can develop into a baby.	
Infertility (n)	The inability to have children.	
Informed consent (n)	The process of agreeing to a procedure or course of treatment after understanding what the procedure/treatment entails, the potential risks and benefits associated with it, and the other options available.	
Inheritance pattern (n)	The manner in which a particular genetic trait or disorder is passed from a parent to a child, e.g., autosomal dominant or recessive, X-linked dominant or recessive, or multifactorial.	
Inherited (adj)	Passed down from parent to child.	
Insertion (n)	Having an extra segment of DNA added in at a place where it is not usually found.	
Intellectual disability (n)	A condition, varying in severity, in which a person has significant impairments in mental abilities, social skills, and core functions of daily living compared to others their age.	

Term	Definition	Translation
Klinefelter's syndrome (47, XXY) (n)	A genetic condition in which a male has two copies of the X chromosome and one copy of the Y chromosome; compared to the typical chromosome makeup where a male has one X chromosome and one Y chromosome.	
Marker chromosome (n)	A small extra fragment of a chromosome detected when doing a chromosome test like a karyotype. Marker chromosomes can sometimes cause health or development problems, depending on how much and what genetic material is contained within.	
Maternal Serum Screening (MSS or Maternal serum alpha- fetoprotein test) (n)	A blood test available to pregnant women that identifies elevated risks for Down syndrome, trisomy 18 and neural tube defects. Usually conducted in the second trimester, between 14-20 weeks.	
Microarray (n)	A high-resolution chromosome test that provides more information than a basic karyotype. A microarray measures the amount of chromosome material in a sample and can detect both large changes (e.g., if there is an extra or missing chromosome) as well as small changes (e.g., if there are very small pieces of chromosomes that are extra or missing). It does not look at the visual appearance or arrangement of chromosomes but measures the amount of genetic material.	
Miscarriage (n)	The spontaneous (not intentional) loss of a pregnancy. See "abortion." Also called pregnancy loss.	
Molecule (n)	The smallest unit of a chemical compound that still has the properties of that compound. For example, a molecule of water is made up of two hydrogen atoms and one oxygen atom. Separately, they are just atoms, but when bonded together, they make a water molecule.	
Mosaicism (n)	A condition in which some, but not all, cells in a sample show a genetic difference.	
Multifactorial (adj)	Due to a combination of genetic and non-genetic (environmental, hormonal, etc.) risk factors that act together to determine risk.	
Mutation (n)	A change in a gene, usually deleterious. See "genetic variant."	
	In Spanish, the word is sometimes confused with "mutilation."	

Term	Definition	Translation
Non-Invasive Prenatal Testing (NIPT) (n)	A blood test available to pregnant women that identifies elevated risk for certain genetic conditions in the fetus. This test focuses on fragments of DNA from placental cells – which carry the fetus' genetic make-up – that are found in the mother's bloodstream.	
Nuchal thickening (n)	There is a pocket of fluid at the back of the neck of a fetus which can be measured in an ultrasound between 10-14 weeks gestation (called the nuchal translucency). If there is a large amount of fluid at this point, or if later in pregnancy the neck skin itself appears to be thicker, this is associated with a higher risk of chromosome problems and other rare genetic conditions.	
Open Neural Tube Defect (ONTD) (n)	A birth defect of the spine, spinal cord, or brain that results from a hole in the spinal column not closing up when appropriate during early fetal development. The two most common types of ONTDs are spina bifida and anencephaly.	
Ova (n)	See "egg."	
Ovary (n)	The organ in a woman that stores and releases eggs. There are normally two.	
Penetrance (n)	The probability that a specific genetic trait will be expressed if a person carries a mutation. "Complete penetrance" means that everyone who carries a particular gene mutation will show the trait related to that altered gene. "Incomplete penetrance" means that only some of the people who have the altered gene will actually show the related trait.	
Placenta (n)	The organ that develops together with an embryo in a pregnant woman's uterus to nourish the embryo through the umbilical cord.	
Presymptomatic (adj)	Before symptoms appear.	
Prognosis (n)	The most likely outcome of a disease process.	
Protein (n)	A molecule made up of chains of <i>amino acids</i> . Proteins do most of the work in cells and are required for the structure, function, and regulation of the body's tissues and organs. Genes determine how specific amino acids are put together to form a specific protein.	
Random (adj)	Happening in an unpredictable way.	

Term	Definition	Translation
Recessive (adj)	A genetic trait in which both copies of a gene need to be altered for the trait to be expressed; recessive traits are typically inherited from both parents, each one contributing one copy of the gene in question. In prenatal genetics, recessive inheritance refers to	
	conditions in which both copies of a particular gene pair need to be altered to develop a specific disease or condition.	
Red flag (n)	A warning sign or a clue.	
Replicate (v)	То сору.	
Reproductive history (n)	The experiences a woman has had related to pregnancy and childbirth. A reproductive history usually includes age at the onset of menses (have a monthly period), age at the beginning of sexual intercourse, age at first conception, number of pregnancies, number of live births, number of miscarriages, number of abortions.	
Ring chromosome (n)	An abnormal formation of a chromosome in which the ends of two of the chromosome's arms have linked together to form a ring.	
Risk (n)	The chance that something will happen; in the case of prenatal genetics, the chance that the child will have a genetic condition.	
At risk (adj)	Has some possibility of having a genetic condition.	
At average risk (adj)	Has the same possibility of having a genetic condition as the general population.	
At high risk (adj)	Has a greater possibility of having a genetic condition than the general population.	
At higher risk than "X" (adj)	Has a greater possibility of having a genetic condition than "X."	
Risk factor (n)	A circumstance that increases the risk of having a genetic condition.	
Runs in the family (e.g., sickle cell disease runs in the family) (v)	When a certain condition or trait seems to appear in many members of a single family	

Term	Definition	Translation
Screen (v)	To assess a person's risk for a particular health problem, even if the person has no symptoms to suggest that they have the problem.	
Screening test (n)	A test that looks to detect a particular health problem, even if a patient has no symptoms of that particular condition.	
	Prenatal screening tests aren't looking for genetic changes. They are designed to look at hormones, chemicals, and other things that indicate a fetus might have a higher risk of a certain condition. They are not DNA results, do not give a definite answer about whether someone is affected, and do not provide a firm diagnosis. They are usually reported as a risk number (e.g., 1 in chance), and if this number is above a certain cut-off point, the result will be categorized as 'screen positive' or 'screen negative'.	
Semen (n)	A liquid produced by the testes, the prostate gland, the seminal vesicle, and the bulbourethral gland that carries, nourishes and protects sperm cells on their way to fertilizing an egg. Also called seminal fluid.	
Seminal fluid (n)	See "semen."	
Sickle cell disease (n)	A genetic disorder caused by a mutation in a gene that helps to make hemoglobin, an important part of red blood cells. In sickle cell disease, red blood cells can become sickle-shaped (instead of round as they should be). These affected blood cells do not function properly, and die early, leaving a shortage of red blood cells, and can block blood flow causing pain.	
Sperm (n)	The reproductive cells of the man. When sperm fertilize a woman's egg, a baby develops.	
Sperm donation (n)	The process through which a man allows his sperm to be collected and used to fertilize the eggs of a woman who is not his sexual partner.	
Spina bifida (n)	A congenital defect of the spine in which part of the spinal cord is exposed through an opening in the bone structure. This may result in nerve damage and some degree of paralysis in the legs.	
Statistically significant (adj)	A measurement of whether the findings of research are meaningful. Refers to the likelihood that a relationship between two factors is linked by something other than chance.	

Term	Definition	Translation
Teratogen (n)	Something that has the potential to disturb the normal development of an embryo or fetus.	
Termination of pregnancy (n)	See "abortion."	
Test results (n)		
Negative (adj)	A negative result on a diagnostic genetic test means that the laboratory did not find the specific genetic change that the test was designed to identify. A "screen negative" result on a screening test means that the person's risk of having whatever the test was designed to find is lower than the risk for most people.	
Positive (adj)	A positive result on a diagnostic genetic test means that the laboratory did find a specific genetic change that is associated with a hereditary condition (a deleterious mutation).	
	A "screen positive" result on a screening test means that the person's risk of having whatever the test screened for is significantly higher than expected. That still does not mean that the person definitely has that condition.	
Variant of uncertain significance (n)	If the laboratory finds a genetic change for which currently there is not enough information to know if this change is problematic or not, it reports a "variant of uncertain significance." Basically, this means that the laboratory found a genetic change, but they don't know what it means. Many of these variants will eventually be reclassified as either a "positive" or "negative" result. Most become "negative" and are thought to represent natural variation between	
Polymorphism (n)	individuals. Everyone has some degree of commonly occurring genetic changes that are not associated with medical problems. If the test finds this sort of change, it reports a "polymorphism."	
Testes (n)	The organs in a man that create and release sperm. Also called "testicles." Singular testis.	
Toxic exposure (n)	Contact with something that is harmful or poisonous.	

Term	Definition	Translation
Trait (n)	A characteristic of a person. In genetics, traits are aspects of a person defined or influenced by their genetic code. E.g., Eye color, blood type, risk for certain diseases.	
Translocation (n)	Translocations are a type of chromosome rearrangement. They can be 'balanced', meaning that two chromosomes have pieces that have 'swapped' with each other, but there is still the correct amount of chromosome material. Balanced translocations do not usually cause medical or development problems. They can also be 'unbalanced', where pieces of chromosomes have traded places, but there is missing or extra chromosome material. Unbalanced translocations will often cause medical and development problems.	
Typo (n)	A mistyped word; shortened slang for "typographical error." In genetic counseling "a typo" is commonly used as an analogy to refer to mutations or variants in a gene.	
Ultrasound (n)	An imaging method that uses high frequency sound waves to create a picture of something inside the body, such as a fetus or baby	
Uterus (n)	The organ in a woman's lower abdomen in which a fertilized egg develops into an embryo, then a fetus, then a baby. Also called "womb."	
Vagina (n)	The tube-like part of the female reproductive system that extends from the cervix to the outside of the body.	
Variant (n)	A version of something that differs from the norm. So, a genetic variant is a change to the usual genetic sequence.	
Vas deferens (n)	The tubes that lead from the testes to the urethra, through which sperm is ejaculated.	
X-linked (adj)	A trait that is influenced or determined by a gene on the "X" chromosome.	

Family Relationships

For genetic counselors, it is very important to know if a relative is a blood relative or a relative by marriage.

Term	Definition	Translation
Parent	Your mother or father.	
Mother	For genetic purposes, the woman whose egg was fertilized and grew to be you.	
Mother-in-law	Your husband or wife's mother.	
Stepmother	Your father's wife who is not your biological mother.	
Adoptive mother	A woman who is not your biological mother but who accepted legal responsibility for and raised you.	
Godmother	A woman chosen by your parents to be your spiritual guide in life, named as such through a ceremony in the Catholic church.	
Father	For genetic purposes, the man whose sperm fertilized the egg that grew to be you.	
Father-in-law	Your husband or wife's father.	
Stepfather	Your mother's husband who is not your biological father.	
Adoptive father	A man who is not your biological father but who accepted legal responsibility for and raised you.	
Godfather	A man chosen by your parents to be your spiritual guide in life, named as such through a ceremony in the Catholic church.	
Aunt	Your mother or father's sister or sister-in-law.	
Maternal aunt	Your mother's sister or sister-in-law.	
Paternal aunt	Your father's sister or sister-in-law.	
Uncle	Your mother or father's brother or brother-in-law.	
Maternal uncle	Your mother's brother or brother-in-law.	
Paternal uncle	Your father's brother or brother-in-law.	

Term	Definition	Translation
Cousin	Usually understood to be a first cousin.	
First cousin	Your aunt or uncle's child.	
First cousin once removed	Your aunt or uncle's grandchild or your first cousin's child	
Second cousin	The children of first cousins are second cousins to each other.	
Grandparent	Your parent's father or mother.	
Grandmother	Your mother or father's mother.	
Maternal grandmother	Your mother's mother.	
Paternal grandmother	Your father's mother.	
Grandfather	Your mother or father's father.	
Maternal grandfather	Your mother's father.	
Paternal grandfather	Your father's father.	
Great aunt	Your mother or father's aunt.	
Maternal great aunt	Your mother's aunt.	
Paternal great aunt	Your father's aunt.	
Great uncle	Your father or mother's uncle.	
Maternal great uncle	Your mother's uncle.	
Paternal great uncle	Your father's uncle.	
Great grandparents	The parents of any of your grandparents.	-
Great grandmother	The mother of any of your grandparents.	
Great grandfather	The father of any of your grandparents.	-

Term	Definition	Translation
Siblings	The children of your father and mother.	
Half siblings	Siblings (brothers and sisters) who have either the same mother and different fathers, or the same father but different mothers. Half-siblings share some genetic similarity with you.	
Stepsiblings	The children of your stepmother but not your father; or the children of your stepfather but not your mother. Stepsiblings do not share any genetic similarity with you.	
Twins	Two siblings born at the same time	
Identical twins	A twin that developed from the same egg and sperm, meaning that they are mostly genetically identical. Also called monozygotic twins.	
Fraternal twins	A twin who developed from a different egg and sperm than, meaning that he or she is genetically different and has the same number of shared genes as any other sibling. Also called dizygotic twins.	
Triplets	Three siblings born at the same time	
Sister	A sibling who is a girl.	
Sister-in-law	Your brother's wife.	
Half sister	A girl who is either the child of your father with a different mother, or the child of your mother with a different father.	
Stepsister	The daughter of your stepmother or stepfather and therefore not biologically related to you.	
Brother	A sibling who is a boy.	
Brother-in-law	Your sister's husband.	
Half brother	A boy who is either the child of your father with a different mother, or the child of your mother with a different father.	
Stepbrother	The son of your stepmother or stepfather and therefore not biologically related to you.	
Niece	Your sibling's daughter.	
Nephew	Your sibling's son.	

Term	Definition	Translation
Spouse	Your husband or wife.	
Wife	The woman to whom you are married.	
Husband	The man to whom you are married.	
Children	Genetically speaking, the people who are produced from your egg or sperm.	
Daughter	Your child who is a girl.	
Daughter-in-law	Your son's wife.	
Stepdaughter	Your spouse's daughter who is not your biological child.	
Adoptive daughter	A girl for whom you have accepted legal responsibility and raised even though she is not your biological child.	
Goddaughter	A girl for whom you have accepted spiritual responsibility within a ceremony of the Catholic church; she may or may not be biologically related to you.	
Son	Your child who is a boy.	
Son-in-law	Your daughter's husband.	
Stepson	Your spouse's son who is not your biological child.	
Adoptive son	A boy for whom you have accepted legal responsibility and raised even though he is not your biological child.	
Godson	A boy for whom you have accepted spiritual responsibility within a ceremony of the Catholic church. He may or may not be biologically related to you.	
Ancestor	A person from whom you are descended, usually more remote than a grandparent.	
Fiancé(e)	The person whom you have promised to marry.	
	Man: fiancé	
	Woman: fiancée	
Domestic partner	The person with whom you are living and have an intimate relationship, but to whom you are not married.	

Term	Definition	Translation
Divorced	Having ended a marriage.	
Engaged	Having promised to marry someone.	
Widow	A woman whose husband has died.	
Widower	A man whose wife has died.	
Relative	A person in your family.	
Blood relation	A person who is related to you by blood not marriage, e.g., your sister, but not your sister-in-law; your mother but not your stepmother; your daughter but not your adopted daughter.	
Next of kin	Your closest living blood relation.	

Glossary of Prenatal Genetics Terms - Language

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Analogies

Genetic counselors often use analogies to explain complex ideas in genetics. An analogy explains a new concept by comparing it to something more common. A clue that an analogy is being used are the words "It's as if" and "It's like "

Examples:

"Genes are <u>like</u> an instruction book or an instruction manual. And in a book, there is a certain sequence to a story."

"<u>It's as if</u> you were reading through a long book and looking for one typo, for a letter that is mistaken."

"So, a mutation is **like** an unexpected change in the order of the letters."

"A gene is **like** a recipe, and mutations are like changes in that recipe."

"It's like chapters in a book, and like misspellings."

"Our cells are like libraries and our chromosomes are like books."

"<u>Like</u> a flip of a coin." (to explain that there is a 50% chance that a mutation will be passed on every time the patient has a child).

"Inside every cell are genes, which are <u>like</u> instruction books for our bodies; they tell our bodies how to grow and develop."

"It is kind of <u>like</u> you a reading a book and you notice that a word is spelled wrong; that is what this genetic test looks for."

Some analogies may not work with certain patients, due to linguistic or experiential differences.

Analogies that focus on genes as "letters in a book" may make no sense to speakers of character-based written languages such as Chinese, Japanese, and Korean.

"It's as if you were reading through a long book and looking for one typo, for a letter that is mistaken."

"So a mutation is like an unexpected change in the order of the letters."

"It's like chapters in a book, and like misspellings."

"It is kind of <u>like</u> if you a reading a book and you notice that a word is spelled wrong; that is what this genetic test looks for."

Analogies that focus on genes as a recipe may not work for patients who do not cook with written recipes.

"A gene is <u>like</u> a recipe, and mutations are like changes in that recipe."

Analogies that discuss random chance in terms of a coin toss may not work for patients from cultures where people don't flip coins to make decisions.

"<u>Like</u> a flip of a coin" (to explain that there is a 50% chance that a mutation will be passed on every time the patient has a child).

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