

Pedigree Analysis

Pedigree: A family tree describing the occurrence of heritable characteristics (genotype and offspring) across an intergenerational population.

How to read a pedigree:

Offspring are represented by squares

Parents are represented by circles

An individual that has the trait is marked in the pedigree with the circle or square shaded (shaded) in

Individuals who are carriers for the trait (heterozygous) have their circle or square half shaded

Married and unions are connected by horizontal lines

Offspring come off the parents by vertical lines

Consanguinity are indicated with Roman Numerals

Individuals with no genealogy are individually numbered

Knowing how to read offspring across one allele from each parent, a pedigree can be used to predict the genotype and phenotype of individuals

New Genetic Disorders

Albinism (Autosomal Recessive)

Albinism is every 17,000 people have albinism. These individuals lack tyrosinase enzyme, a phenotypic pigment. While melanin's role is protecting us from ultraviolet light (sunburn), it also has other important functions in the development of the retina and bones and hair color (melanin of which our hair is made)

Albinism is an autosomal recessive trait

Cystic Fibrosis (Autosomal Recessive)

The most common fatal genetic disease in the United States is cystic fibrosis, which affects one out of every 2,500 children of European descent (but is much less in other groups). The rate of 25 alleles (CF) are carrier. The normal allele for the gene codes for a membrane protein. The protein does not have the protein for the normal version that connects between ducts and other organs. The membrane opens the pancreas, lungs, digestive tract and other organs, conditions that form (cystic fibrosis). However, individuals with cystic fibrosis die before they 40th birthday. Cystic fibrosis is the the at a slow pace than clogged airways, dehydration of respiratory to prevent infections, and other pancreatic treatment can prolong life. In the United States, more than half of the people with cystic fibrosis were carrier and their last disease is beyond

Tay Sachs Disease (Autosomal Recessive)

Tay Sachs Disease is another fatal disease inherited as a recessive allele. The disease is caused by a dysfunctional enzyme that fails to break down lipids (fats). The symptoms of Tay Sachs usually show up a few months after birth. The infant begins to exhibit mental, blindness and degeneration of motor and mental performance. Inevitably, the child dies within a few years