

Pedigree Analysis

Pedigree: A family tree describing the occurrence of heritable characteristics (genetic) and offspring across six generations as possible.

There is two a pedigree:

- Offspring are represented by squares

- Married are represented by circles

- An individual that has the trait (recessive) in the pedigree will have the circle or square shaded (shaded) in

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

- Married and wives are connected by horizontal lines

- Offspring come off the parents by vertical lines

- Consanguinity are indicated with Roman Numerals

- Individuals with in the generations are individually numbered

Knowing her and's offspring across one allele from each parent, a pedigree can be used to predict the genotype and phenotype of individuals.

Some Genetic Disorders

Albinism (Autosomal Recessive)

About one in every 17,000 people have albinism. These individuals fail to produce melanin, a photoprotective pigment. While melanin's role is protecting us from ultraviolet light (sunburns), it also has other important functions in the development of the nervous system and then color vision and of which can lower one's life.

Albinism is an autosomal recessive trait

Cystic Fibrosis (Autosomal Recessive)

The most common fatal genetic disease in the United States is cystic fibrosis, which strikes one out of every 2,500 children of European descent but is much rarer in other groups. One out of 25 whites (4%) are carriers. The normal allele for the gene codes for a membrane protein. The protein does not have the protein has the normal version that maintains between chloride and sodium ions normal. The membrane built up in the pancreas, lungs, digestive tract and other organs, conditions that form structural problems. Ultimately, individuals with cystic fibrosis die before their 40th birthday. Cystic fibrosis is the the it is clear more than 100,000 people, daily lives of patients is painful, and other pancreatic symptoms can prolong life. In the United States, more than half of the people with cystic fibrosis were born and their life spans are beyond.

Tay Sachs Disease (Autosomal Recessive)

Tay Sachs Disease is another fatal disorder 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 suffer seizures, blindness and deterioration of motor and mental performance. Inevitably, the child dies within a few years.