

Introduction to Genes and Alleles



Every living organism has **genes**, which are sections of DNA that carry instructions for specific traits, like eye color or blood type. Each gene exists in different **versions**, called **alleles**.

For example, a gene for flower color might have one allele for purple and another for white. You inherit one allele from each parent, so you can have two of the same (**homozygous**) or two different (**heterozygous**) alleles.

These combinations determine how traits are expressed, following patterns first discovered by **Gregor Mendel**, the father of genetics.

Biologists refer to **dominant** and **recessive** alleles to describe how traits are inherited and expressed. A **dominant allele** (denoted with a capital letter) is one that shows its effect even if only one copy is present in the pair of alleles (**heterozygous**). In contrast, a **recessive allele** (denoted with a lowercase letter) only shows its effect when an individual has two copies of it (**homozygous**). For example, if the allele for brown eyes is dominant and the allele for blue eyes is recessive, a person with one of each will have brown eyes. Dominance does not mean that the allele is better or more common – just that it masks the effect of the recessive allele when both are present.

In biology, the **genotype** refers to the genetic makeup of an organism – the specific combination of alleles it carries for a particular trait. The **phenotype** is the physical or observable expression of that trait, such as eye color, blood type, or the presence of dimples. While the genotype provides the instructions, the phenotype is the result of how those instructions are carried out. For example, a person with one dominant allele for brown eyes and one recessive allele for blue eyes (genotype: Bb) will have the phenotype of brown eyes. The environment can also influence phenotype, but it is primarily determined by genotype.

Punnett squares are a simple but powerful tool used by biologists to predict the possible genetic outcomes of a cross between two individuals. By arranging the alleles from each parent along the top and side of a grid, a Punnett square shows all the potential combinations of alleles in their offspring. This helps determine the **probability** of inheriting particular traits. For example, if two heterozygous parents (Bb) are crossed, a Punnett square can show that their children have a 25% chance of being homozygous dominant (BB), 50% chance of being heterozygous (Bb), and 25% chance of being homozygous recessive (bb). This makes Punnett squares especially useful when studying inheritance patterns and understanding Mendelian genetics.

	B	b
B	BB 	Bb 
b	Bb 	bb 

Introduction To Genes Vocabulary Review

Wolfgang Guggemos



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