

Genetics Case Study Comparison

Case Study	Genetic Basis	Phenotype	Mode of Inheritance	Carrier Status	Genetic Testing	Prevention
Phenylketonuria (PKU)	Autosomal recessive mutation in the phenylalanine hydroxylase (PAH) gene.	Intellectual disability, growth retardation, and a musty body odor.	Autosomal recessive.	Carrier status is determined by genetic testing.	Genetic testing can identify carriers.	Early diagnosis and dietary restriction of phenylalanine.
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Genetics Case Study Comparison Answer Key

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