## Human Pedigrees

## Exercises

Questions 1, use the pedigree chart shown below, And answer the questions:-
$\qquad$ 1. A male
$\qquad$ 2. A female
$\qquad$ 3. A marriage
$\qquad$ 4. A person who expresses the trait
$\qquad$ 5. A person who does not express the trait

$\qquad$ 6. A connection between parents and offspring
$\qquad$ 7. How many generations are shown on this chart?

Assuming the chart above is tracing the dominant trait of "White Forelock $(F)^{\prime \prime}$ through the family. $F$ is a tuft of white hair on the forehead.
$\qquad$ 8. What is the most likely genotype of individual "A"? (FF, Ff or ff?)
9. What is the most likely genotype of individual "C"? (FF, Ff or ff?) M.Sc. Aamal Muhsen \& M.Sc. Mazin E. Hadi

Question 2, A typical pedigree for a family that carries Falconi anemia.
Note that carriers are not indicated with half-colored shapes in this chart.


Analysis Questions.
To answer questions \#1-4, use the letter "f" to indicate the recessive Falconi anemia allele, and the letter " F " forthe normal allele.

1. What is Arlene's genotype? $\qquad$
2. What is George's genotype? $\qquad$
3. What are Ann \& Michael's genotypes? $\qquad$
4. Most likely, Sandra's genotype is $\qquad$ .

## Question 3 :

## Draw your own Pedigree

Condition of Interest: Albinism
Albinism is a condition in which there is a mutation in one of several possible genes, each of whichhelps to code for the protein melanin.. This gene is normally active in cells called melanocytes which are found in the skin and eyes. Albinism involves a significant reduction or absence of the production ofmelanin, giving affected individuals a lack of normal coloration to their skin/eyes.

Inheritance Pattern: normal melanin protein is produced by an autosomal dominant allele; albinism results from a lack of melanin and is caused by an autosomal recessive allele.

Use the letter $\mathbf{A}$ or a to represent dominant/recessive forms of albinism.
Two normally-pigmented parents have 3 children. The first child (a girl) and their second child (a boy) have normal pigmentation. Their third child (a girl) has albinism. That girl marries a normally pigmented male and they have four children. The first three (two girls and a boy) have normal pigmentation. Their fourth child (a girl) has albinism like her mother.


