

Pedigree Worksheet

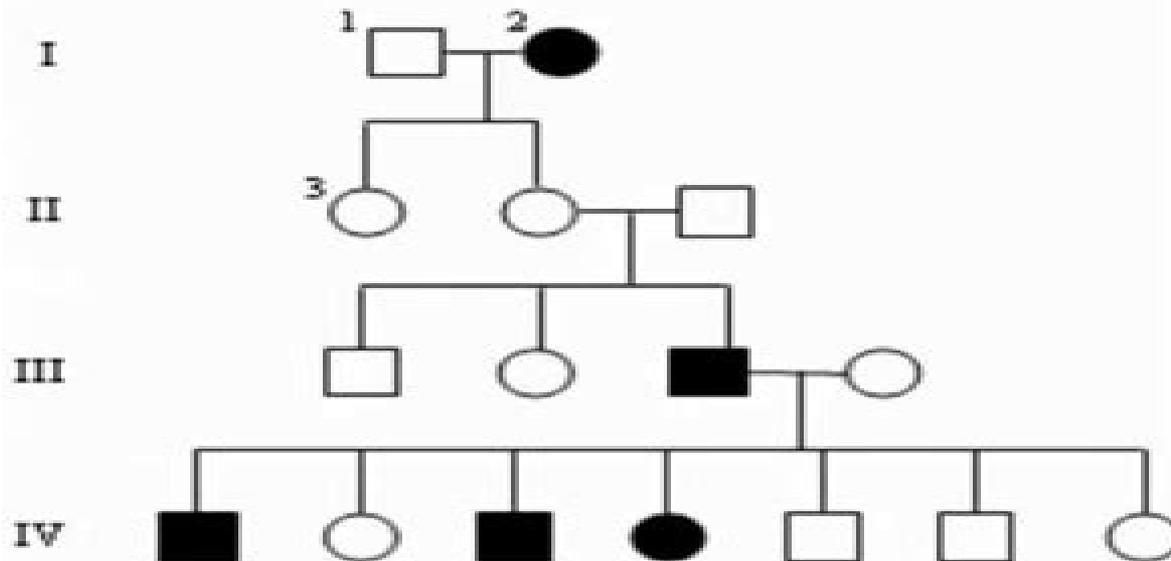
Name: _____ P _____

A sex-linked recessive characteristic is determined by an allele that is carried only on the X chromosomes. The shorter Y chromosome does not carry an allele for a sex-linked trait. Since there is only one X in his genotype, XY, a male who carries a particular recessive allele on the X chromosome will have the sex-linked condition. A female who carries a recessive allele in one X chromosome will not show the condition if there is a dominant allele on her other X chromosome. She will express the recessive condition only if she inherits two recessive alleles – one from each parent. Her chances of inheriting the condition are thus greatly reduced. One sex linked trait is hemophilia, a condition in which the blood does not clot properly. Most people who have hemophilia are men.

A. Interpreting a human pedigree. (In Class)

1. Number all individuals on the pedigree at the top of each shape.
2. How many males are there? _____
3. How many males have hemophilia? _____
4. How many female are there? _____
5. How many females have hemophilia? _____
6. How many marriages are there? _____
7. How many children did the first couple (couple in row I) have? _____
8. How many children did the third couple (couple in row III) have? _____
9. How many generations are there? _____
10. How many members are there in the fourth generation? _____

The genotypes of the males in a pedigree for sex-linked inheritance are easy to determine since normal blood clotting (N) is dominant and hemophilia is recessive (n). Since these alleles are on the X chromosome only, a male represented by a clear square will have the genotype $X^N Y$. A darkened square will be $X^n Y$. Label *under* each individual on the pedigree. Females with hemophilia have an easy genotype to identify. They are all $X^n X^n$. Both recessive alleles must be present for a female to have hemophilia. If one dominant allele is present (X^N), the individual would be normal for clotting. Females who do not show the trait for hemophilia may be homozygous dominant ($X^N X^N$) or heterozygous ($X^N X^n$). A heterozygous female is called a carrier. Examination of offspring can often determine which genotype the parents have. If any child (son or daughter) has hemophilia, then the female must be heterozygous. If her son has hemophilia, he has genotype $X^n Y$. He inherited the Y from the father, so the other allele is his genotype (X^n) had to come from the mother. If a daughter has hemophilia ($X^n X^n$), she inherited an X^n from each parent, thus making the genotype for the normal mother $X^N X^n$.



11. What is the genotype for Individual #1?

Phenotype?

12. What is the genotype for Individual #10?

Phenotype?

13. What is the genotype for the first born male in generation III?

Phenotype?

14. How many females have the genotype $X^N X^n$?

15. What would be the genotype for the female who marries into the family in generation III?

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