9. Answer the questions for the following pedigree:





	For person 2, the genotype is ^{b.} ¹⁷ , and the phenotype is ^{c.} <u>no disease (tay sachs)</u> .
	For person 1, the genotype is ^{d.} .
	How did you determine this? e. Since they don't have the disease they must have one "T". This person must have given a "t" allele to their oldest son who has the disease "tt"
1.	What are the chances that person 3's children will be normal? f. <u>[Vic there of rooms]</u> Consider a model in which there are three gene pairs of alleles; a dominant allele in any pair adds pigment to the skin. Use the letters A , B , C to indicate pigment formation and a , b , c to indicate lack of pigment formation.
	What is the genotype for the darkest individual? a. ABBCC
	What is the genotype for the lightest individual? ^{b.}
	What is the genotype for the offspring from a cross of the individuals from a and b? c_{-} ABDCC
2.	How does the skin color of this person compare to either of the parents? ^d . The phenotype would be an even mixture with a skin color shade between light and dark. A man with blood type A reproduces with a woman who has blood type B. Their child has blood type O. Give the genotype of all persons involved: man ^{a.} $A = A = A = A = A = A = A = A = A = A $
3.	If a child has AB blood and the father has type B blood, what could the genotype of the mother be?

^{14.} If a child has BO blood and the father has type O blood, what could the genotype of the mother be?



15. Both a man and a woman have sickle-cell trait. List all phenotypes among the offspring, as well as the chance (percent) of each occurring. a. ss = normal = 25%, b. ss = trait = 50%, c. s, s, = disease = 25%.