Mendelian Inheritance in Humans Classroom Activity

Name:

Human have several single-gene traits that are easy to observe and which are controlled by genes that have only two alleles. Based on the specific combination of alleles (AA, Aa = dominant phenotype; aa = recessive phenotype) the trait is either present or absent. There is no "in between". Many other human traits, such as eye color and height are due to several pairs of genes, and the phenotypes show a continuous range of variation.

For the traits described in the following list, determine your phenotype and possible genotype.

Trait	Symbol	Dominant	Your	Your
		Phenotype	phenotype	genotype
1. Facial dimples: Best seen when smiling. With dominant	D, d	dimples		
phenotype, you may have a dimple only on one side, or on				
both.				
2. Bent little finger: A dominant allele causes the last joint	B, b	bent		
of the little finger to dramatically bend inward toward the 4 th				
finger. Lay both hands flat on a table relax your muscles, and				
note whether your have a bent or straight little finger.				
3. Eye Color: Eye color, as well as hair and skin color, is a	E, e	brown		
complex trait. The main pigment is melanin, and the more				
melanin, the darker the color. While the genetics of eye color				
is complex, alleles for the production of melanin dominate				
those for lack of melanin. Evaluate your eyes as either brown				
(M) or non-brown (m).				
4. Free ear lobe: Dominant trait is for lobes to hang free.	F. f	free		
With recessive phenotype, the lobes are attached directly to				
the head.				
5. Mid-digital hair: Some people have hair on the middle	H, h	hair		
segment of one or more of their fingers, while others don't.				
Any hair at all is the dominant phenotype. Complete absence				
of hair is recessive.				
6. Hand clasping: When the hands are clasped (without	L,l	left on top		
thinking about it!), most people place their left thumb on top				
of their right.				
7. Early Onset Myopia (childhood): Nearsightedness is a	M, m	myopic		
complex trait with at least 4 gene loci involved, however the				
heritability of myopia is very high and shows a dominant				
pattern.				
8. Achondroplasia: The most frequent or of short-limb	A, a	achondropla		
dwarfism. Affected individuals (Aa) exhibit short stature		sia		
cause by shortening of the limbs. (AA individuals die before				
birth).				
9. Chin cleft: A prominent cleft in the chin is inherited as the	C,c	cleft		
dominant phenotype. The cleft is due to the bond structure				
which underlies the Y-shaped fissure of the chin. Females				
appear to be less conspicuously affected than males.				
10. Achoo syndrome: Autosomal dominant trait also called	A, a	sneezing		
photo sneeze reflex, usually $2 - 3$ successive sneezes, when a				
dark-adapted person suddenly is exposed to bright light.				

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