

Non-Mendelian Genetics - Practice

1. In plants known as "four o'clocks", the allele for the dominant red-flower color is **incompletely dominant** over the allele for white-flowers. A gardener allows several heterozygous pink-flowered four o'clocks to self pollinate and collects 200 seeds.

Draw a Punnett square for the cross. Identify the flower color phenotypes and theoretical percentage.

	C^R	C^W
C^R	$C^R C^R$	$C^R C^W$
C^W	$C^R C^W$	$C^W C^W$

- | | |
|------------------|--------------------|
| <u>Phenotype</u> | <u>Percentages</u> |
| • <u>red</u> | • <u>25%</u> |
| • <u>pink</u> | • <u>50%</u> |
| • <u>white</u> | • <u>25%</u> |

2. Skin color in humans is determined by a polygenic inheritance system, possibly involving as many as 9 genes. For simplicity let's consider the influence of 2 genes: A and B, where the dominant allele darkens skin color. Suppose a woman who is $AABb$ mates with a man who is $Aabb$.

- A. List all of the possible genotypes of the gametes that could be produced by each the parents. Identify the percent ratios.

- Genotype: $AABb$ Ratio: $4/14$
 Genotype: $AABb$ Ratio: $4/16$
 Genotype: $AaBb$ Ratio: $4/16$
 Genotype: $Aabb$ Ratio: $4/16$

		Ab	Ab	ab	ab
$AABb$ $Aabb$	AB	$AABb$	$AABb$	$AaBb$	$AaBb$
	Ab	$AAbb$	$AAbb$	$Aabb$	$Aabb$
	AB	$AABb$	$AABb$	$AaBb$	$AaBb$
	Ab	$AAbb$	$AAbb$	$Aabb$	$Aabb$

2 genes → dihybrid cross

3. In rabbits, white coat color (C^W) and black coat color (C^B) are **codominant**, and both of these alleles are dominant over albino (c); heterozygotes ($C^W C^B$) are spotted.

Draw a Punnett Square that shows the genotypes and phenotypes of the offspring from a heterozygous black-coated rabbit and a homozygous white-coated rabbit?

	C^B	c
C^W	$C^B C^W$	$C^W c$
C^W	$C^B C^W$	$C^W c$

- 50% spotted $C^B C^W$ (codominance)
 50% white $C^W c$ (hybrid)

Study Mendelian Genetics Answer Key

**Anthea Maton, Jill D. Wright, Jean
Hopkins, Susan Johnson, David
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Mendelian Inheritance in Man Victor Almon McKusick,Clair A. Francomano,Stylianos E. Antonarakis,1990 A classic bibliographic guide to human genetics first published in book form in 1966 Each entry consists of six parts a preferred designation followed in parentheses by frequently used synonyms a brief description of the phenotype s the nature of the basic defect a resume of genetic information including mapping and molecular genetic details allelic variants and key references This edition contains 36 987 references mainly to the periodical literature and cites 54 623 authors Based on a continuously updated online version only four months separate closure of the file for the edition and the finished book Annotation copyrighted by Book News Inc Portland OR

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