

GENETICS PRACTICE 1: BASIC MENDELIAN GENETICS

Solve these genetics problems. Be sure to complete the Punnett square to show how you derived your solution.

1. In humans the allele for albinism is recessive to the allele for normal skin pigmentation. If two heterozygotes have children, what is the chance that a child will have normal skin pigment? What is the chance that a child will be albino?

	A	a	normal pigment:
A	AA	Aa	75% chance of AA & Aa
a	Aa	aa	albino: 25% chance of aa

- a. If the child is normal, what is the chance that it is a carrier (heterozygous) for the albino allele? (**CAREFUL**) 2/3 or 67%
2. In purple people eaters, one-horn is dominant and no horns is recessive. Show the cross of a purple people eater that is heterozygous for horns with a purple people eater that does not have horns. Summarize the genotypes & phenotypes of the possible offspring?

	H	h	
h	Hh	hh	50% chance of Hh (horned)
h	Hh	hh	50% chance of hh (no horn)

3. In humans, the brown-eye (B) allele is dominant to the blue-eye allele (b). If two heterozygotes mate, what will be the likely genotype and phenotype ratios of the offspring?

	B	b	75% chance of brown-eyed
B	BB	Bb	25% chance of blue-eyed
b	Bb	bb	25% chance of BB
			50% chance of Bb
			25% chance of bb

Ap Biology Genetics Practice 1 Basic Mendelian Genetics Answers

**Vinay Kumar, Abul K. Abbas, Jon C.
Aster**



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Instructor's Guide for Campbell's Biology Nina Caris, Harold T. Underwood, 1996 Vogel and Motulsky's Human Genetics Friedrich Vogel, Arno G. Motulsky, 2013-06-29 The first edition of this book published in 1979 was found useful by many students and was well received by the scientific community. Since the book was first written, human genetics has undergone dramatic developments, mainly due to the introduction of new concepts and techniques from molecular biology. Concurrently, basic scientists have become increasingly interested in problems of human genetics. More than 700 human genes have been mapped; genes of previously unsuspected complexity, such as the gene for factor VIII, have become known, and the structure of noncoding DNA sequences is being analyzed with the aim of understanding gene regulation. DNA diagnosis is being rapidly introduced into medical genetics. All this, as well as the extensive progress in most

other fields of human and medical genetics had to be considered in the preparation of this second edition. The book has been extensively revised and rewritten. A substantial new section dealing with gene and chromosomal structure at the molecular level has been added. The newer knowledge of molecular genetics has been incorporated and the conceptual and practical contribution of DNA methods for example in the hemoglobinopathies and in some other diseases is discussed. Many new figures and tables have been added and some illustrative material has been replaced. We have read carefully the many friendly and sometimes flattering reviews of the first edition.

Proceedings of Conference on Computers in the Undergraduate Curricula, 1977 **Biology** Michael R. Cummings, 1996 Robbins & Cotran Pathologic Basis of Disease - General Pathology, Vol 1: First Bangladesh Edition-E-Book Vinay Kumar, Abul K. Abbas, Jon C. Aster, 2017-05-12

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