

Genetics Packet ~ Punnett Square Practice **KEY**

Basics

1. The following pairs of letters represent alleles of different genotypes. Indicate which pairs are Heterozygous and which are Homozygous. Also indicate whether the homozygous pairs are Dominant or Recessive (*note heterozygous pairs don't need either dominant nor recessive labels.)

A. BB = Homozygous dominant

D. gg = Homozygous recessive

B. Bb = Heterozygous

E. aa = Homozygous recessive

C. Cg = Heterozygous

F. Ee = Heterozygous

2. In humans, brown eye color (B), is dominant over blue eye color (b). What are the phenotypes of the following genotypes?

A. Bb = Brown eyes

B. BB = Brown eyes

C. bb = blue eyes

Monohybrid Crosses with Complete Dominance

3. A heterozygous smooth pea pod plant is crossed with a wrinkled pea pod plant. There are two alleles for pea pod, smooth and wrinkled. Use R for seed texture. Predict the offspring from this cross.

a. What is the genotype of the parents? Rr x rr

b. Set up a Punnett square with possible gametes.

	R	r
r	Rr	rr
r	Rr	rr

c. Fill in the Punnett square for the resultant offspring.

d. What is the predicted genotypic ratio for the offspring? 1 Rr : 1 rr

e. What is the predicted phenotypic ratio for the offspring? 1 smooth : 1 wrinkled

f. If this cross produced 50 seeds how many would you predict to have a wrinkled pod? 25

4. In humans, achondroplasia "dwarfism" (D) is dominant over normal (d).

A homozygous dominant (DD) person dies before the age of one.

A heterozygous (Dd) person is dwarfed. A homozygous recessive individual is normal.

A heterozygous dwarf man marries a heterozygous dwarf woman...

	D	d
D	DD	Dd
d	Dd	dd

a. What is the probability of having a normal child? 1/3... 33.3%

b. What is the probability that the next child will also be normal? 1/3... 33.3%

each child is a new shot at the same punnett square!

c. What is the probability of having a child that is a dwarf? 2/3... 66.6%

d. What is the probability of having a child that dies at one from this disorder? 25%

Genetics Lab 53 Answers

M Woodhall



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25 Years of p53 Research Pierre Hainaut, Klas G. Wiman, 2007-10-05 1 Communication awareness and access to information Given the complexity of the field and the fact that data pertaining to each particular aspects of p53 biology or deregulation are scattered in many different publications it is extremely difficult to access the full scale of relevant information of any specific p53 related topic Review articles despite their fundamental role in disseminating knowledge usually focus only on general mechanisms and do not discuss in detail the many variations that can occur with respect to cell type particular mutation type as well as biological activation context Books such as this one may help in this task by putting into perspective both general considerations on the p53 pathway and more specific information on various aspects of p53 In the longer term however open access to p53 complexity will require the development of knowledge bases accessible through the web and using simple navigation tools to guide users towards the specific information they need Several efforts are currently being developed in that direction They need to be strengthened and better integrated within the rapidly growing galaxy of web based information sources on molecular and individual variations in cancer 2 Reference functional assays and structural analysis Given the huge diversity of cellular and animal models for wild type or mutant p53 functions it will be important to set up standard universally accepted assays to measure critical p53 protein functions *Review* Oak Ridge National Laboratory, 1992 **A Fatal Inheritance** Lawrence Ingrassia, 2024-05-14 Nominated for the National Book Critics Circle Award in Nonfiction Named a best book of the year by Amazon NPR and Kirkus Weaving his own moving family story with a sweeping history of cancer research Lawrence Ingrassia delivers an intimate gripping tale that sits at the intersection of memoir and medical thriller Ingrassia lost his mother two sisters brother and nephew to cancer different cancers developing at different points throughout their lives And while highly unusual his family is not the only one to wonder whether their heartbreak is the result of unbelievable bad luck or if there might be another explanation Through meticulous research and riveting storytelling Ingrassia takes us from the 1960s when Dr Frederick Pei Li and Dr Joseph Fraumeni Jr first met not yet knowing that they would help make a groundbreaking discovery that would affect cancer patients for decades to come to present day as Ingrassia and countless others continue to unpack and build upon Li and Fraumeni's initial discoveries and to understand what this means for their families In the face of seemingly unbearable loss Ingrassia holds onto hope He urges us to fight like Charlie his nephew who battled cancer his entire life starting with a rare tumor in his cheek at the age of two and to look toward the future as gene sequencing screening protocols CRISPR gene editing and other developing technologies may continue to extend lifespans and perhaps one day even offer cures **Environmental Health Perspectives**, 1991 *Counseling About Cancer* Katherine A. Schneider, 2011-12-08 Important scientific discoveries and ever changing guidelines for how to identify and manage patients with hereditary cancer syndromes are constantly evolving This Third Edition of *Counseling About Cancer* is completely updated and expanded to feature five entirely new chapters on

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