

Meiosis and genetics section 2: Predicting outcomes

12 questions

14. Which of the following best explains why gametes containing **recombinant** genotypes are **uncommon**? All other factors being equal, gamete production is unaffected by which of the following?
- a. crossing over
 - b. independent assortment
 - c. linkage
 - d. meiosis
15. Which of the following traits is governed by **multiple** alleles? For all subsequent items:
a. comparing two individuals with genotypes AaBb and Aabb
b. comparing three gametes produced by one of the same parents, specifically one that does not contain a crossover
 - i. no more than one allele for each trait
 - ii. two different alleles for each trait
 - iii. different responses to the same stimulus
 - iv. identical phenotype
c. prediction of inheritance for individuals **heterozygous** for either recessive traits or dominant traits only
16. If all gametes containing the combination **AB** or **ab** cannot survive the process of **meiosis**, which of the following would NOT be a possible result from **meiosis** in a cell containing **AB** genes?
- a. four daughter cells with **AB** genotypes
 - b. four daughter cells with **ab** genotypes
 - c. four daughter cells with **Ab** genotypes
 - d. four daughter cells with **aB** genotypes
17. In a cross between two pea plants, some offspring show the traits entirely represented. From this information, it can be inferred that the offspring are most likely to be
- a. recombinant
 - b. dominant
 - c. homozygous
 - d. heterozygous
18. Which of the following best describes the relationship between the traits shown in the Punnett square below?
- | | Red | Yellow |
|--------|-----|--------|
| Green | Red | Yellow |
| Yellow | Red | Yellow |
- a. dominant
 - b. codominant
 - c. incomplete dominance
 - d. recessive

Study Guide Section 2 Mendelian Genetics

Noel Cameron, Lawrence Schell

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Chase,Peter J. Russell,2005-06 This student resource contains chapter outlines of text material solutions to all end of chapter problems key terms suggestions for analytical approaches problem solving strategies and a variety of additional questions for student practice Also featured are questions that relate to chapter specific animations and iActivities

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Essential Medical Genetics Edward S. Tobias,Michael Connor,Malcolm Ferguson-Smith,2011-11-15 Adopted at Cambridge University Essential Medical Genetics provides students clinicians counsellors and scientists with the up to date information they need regarding the basic principles underlying medical genetics It also provides guidance on how to apply current knowledge in clinical contexts covering a wide variety of topics from genome structure and function to mutations screening and risk assessment for inherited disorders This sixth edition has been substantially updated to include for instance the latest information on the Human Genome Project as well as several new molecular genetic and chromosome analysis techniques In full colour throughout it includes a number of brand new features including a large number of self assessment questions Essentials chapter summaries further reading suggestions and case study scenarios introducing clinical situations An invaluable new section gives illustrated practical advice regarding how to choose the best available online genetic databases and also importantly how to most easily and most efficiently use them for a wide range of purposes Essential Medical Genetics is the perfect resource for a course on medical genetics and is now accompanied by a regularly

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Human Genetics and Genomics Bahar Taneri,Esra

Asilmaz,Türem Delikurt,Pembe Savas,Seniye Targen,Yagmur Esemen,2020-02-17 Finally meeting the need for a laboratory manual on human genetics this practical guide is the perfect companion title to all major standard textbooks on the subject The authors all have a high level research background and are actively involved in teaching and counseling Based on a standard curriculum in human genetics each chapter equals one practical unit of the course and topics range from basics in human inheritance to genetics in major disease clusters and from bioinformatics and personalized medicine to genetic counseling

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