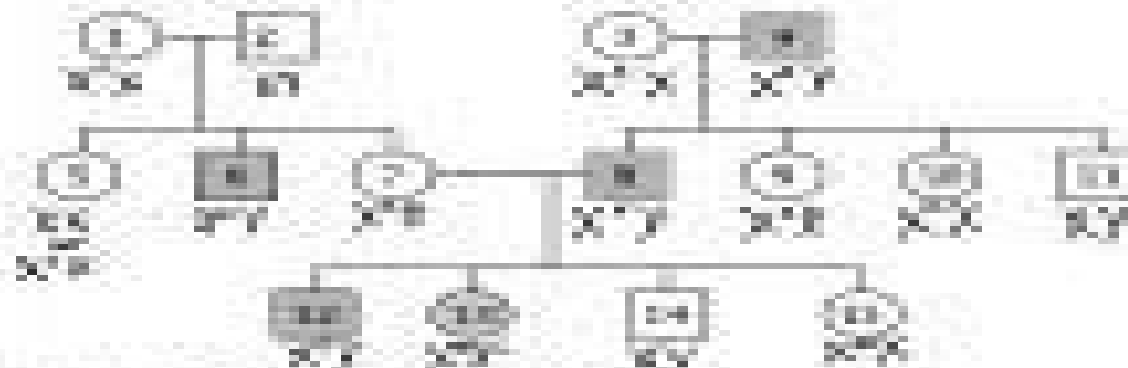


Genetics Practice Problems: Pedigree Tables

Remember the following when working pedigree tables:

- 1) Circles are females and squares are males.
- 2) A shaded circle or square indicates that a person has the trait.

The pedigree chart below is the inheritance of color blindness. Shaded individuals are colorblind. First, determine the probable genotype of person 1 = 11. Then answer the questions below the table.



What do you know about the genotype of the mother of 11?

These people (11) had a color blind son (12). His dad is normal, his color blind mom, from his mother. Since the mother is not shaded she must be a carrier.

Number 21 is colorblind just like her father. Where could she get her alleles for color blindness?

She inherited her alleles for color blindness from her mother. Must have been that 11 is heterozygous. From their father.

Number numbers 25 and 26 were colorblind. How did they have a colorblind son (27)?

Both mothers carry the disease for color blindness.

What must be the genotype of the parents of a colorblind daughter (34)?

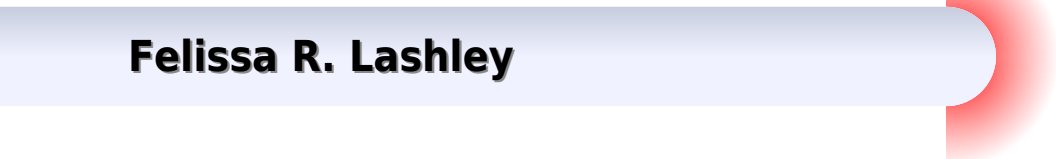
To have a color blind daughter, she must have the recessive alleles from each parent. Her father must be $X^c Y$ and mother must have $X^c X^c$ or $X^c X^N$.

If number 23 marries a normal man, what is the probability that their sons will be colorblind?

$X^N X^N \times X^N Y \rightarrow$ all of their sons will be normal with no color blind.

Genetics Practice Problems Pedigree Tables Key

Felissa R. Lashley



Genetics Practice Problems Pedigree Tables Key:

Genetics in Practice Jo Haydon, 2007-10-22 Nurses are increasingly being asked to provide information when a genetic diagnosis is made whether to the individual patient or to members of his or her family This book provides a comprehensive introduction to present day genetic services to help professionals feel more confident when dealing with queries or advising where to go for further information Comprehensive case studies are used to explain the various models of inheritance and explore the possibilities for families following genetic diagnosis Family history taking risk assessment basic biology of chromosomes and genes and laboratory techniques are all described Issues of ethnicity and ethics are addressed The book also discusses the development of the role of the genetic nurse counselor Molecular Pathology in Clinical Practice Debra G.B. Leonard, 2016-02-02 This authoritative textbook offers in depth coverage of all aspects of molecular pathology practice and embodies the current standard in molecular testing Since the successful first edition new sections have been added on pharmacogenetics and genomics while other sections have been revised and updated to reflect the rapid advances in the field The result is a superb reference that encompasses molecular biology basics genetics inherited cancers solid tumors neoplastic hematopathology infectious diseases identity testing HLA typing laboratory management genomics and proteomics Throughout the text emphasis is placed on the molecular variations being detected the clinical usefulness of the tests and important clinical and laboratory issues The second edition of Molecular Pathology in Clinical Practice will be an invaluable source of information for all practicing molecular pathologists and will also be of utility for other pathologists clinical colleagues and trainees Maternal-Newborn Davis Essential Nursing Content + Practice Questions Sheila Whitworth, Taralyn McMullan, 2017-03-08 Too much information Too little time Here s everything you need to succeed in your maternal newborn nursing course and prepare for course exams and the NCLEX Succinct content reviews in outline format focus on must know information while case studies and NCLEX style questions develop your ability to apply your knowledge in simulated clinical situations A 100 question final exam at the end of the book You ll also find proven techniques and tips to help you study more effectively learn how to approach different types of questions and improve your critical thinking skills Rosenberg's Molecular and Genetic Basis of Neurological and Psychiatric Disease, Seventh Edition Roger N. Rosenberg, Juan M. Pascual, 2024-10-04 Rosenberg s Molecular and Genetic Basis of Neurologic and Psychiatric Disease Seventh Edition provides a comprehensive introduction and reference to the foundations and key practical aspects relevant to neurologic and psychiatric disease This volume has been thoroughly revised and includes newly commissioned chapters on ethics genetic counselling and genet therapy for the central nervous system disorders A favorite of over four generations of students clinicians and scholars this new edition retains and expands the informative concise and critical tone of the previous edition This is an essential reference for general medical practitioners neurologists psychiatrists geneticists and related professionals and for the neuroscience and neurology research community at large Both volumes combined provide a

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