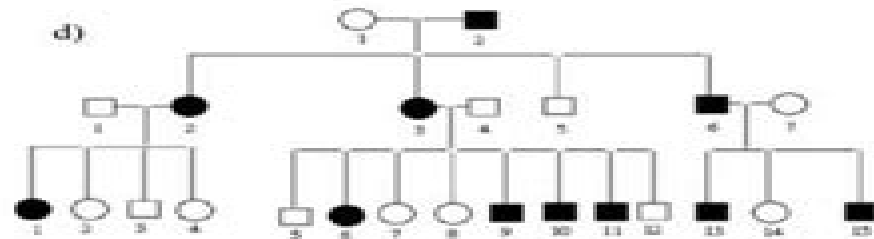
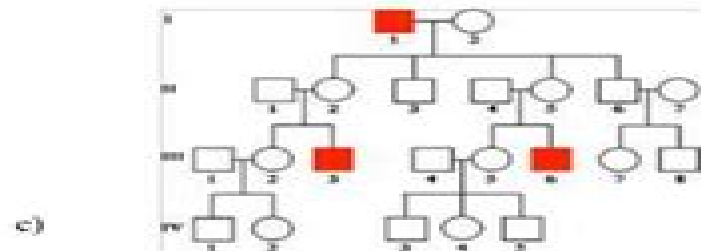
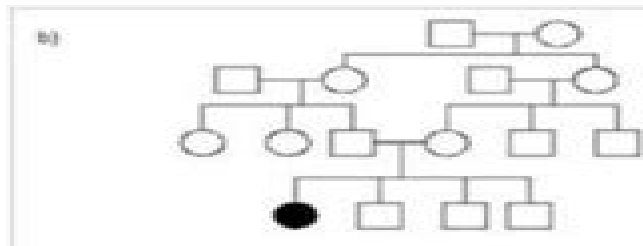
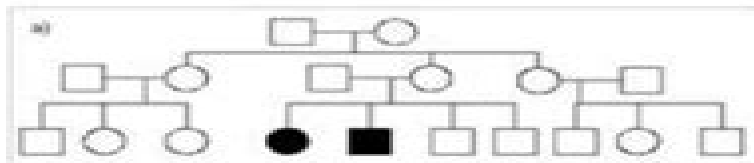


Human Pedigree Analysis Problem Sheet

Background: Autosomes are chromosomes 1-22 (for humans, non sex). Sex chromosomes represent the 23rd pair of chromosomes (in humans). There are a number of different types of human pedigrees that you may encounter, however, there are only a few different modes of inheritance that you will need to be familiar. The following pedigrees show you different examples of human traits that can be traced through generations. See if you can identify the modes of inheritance, and answer any questions directly related to each pedigree. While you are working on this, keep the following clues in mind:

| Clues for Autosomal Inheritance | |
|--|--|
| Recessive <ul style="list-style-type: none"> individual expressing trait has 2 normal parents two affected parents can not have an unaffected child | Dominant <ul style="list-style-type: none"> every affected person has at least one affected parent each generation will have affected individuals |
| Clues for Sex-linked Inheritance | |
| Recessive <ul style="list-style-type: none"> no father-to-son transmission predominantly males affected trait may skip generations | |

- For each of the pedigrees below, identify the mode of inheritance and provide at least 2 reasons for your choice.



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Human Genes and Genomes Leon E. Rosenberg, Diane Drobnis Rosenberg, 2012-05-21 In the nearly 60 years since Watson and Crick proposed the double helical structure of DNA the molecule of heredity waves of discoveries have made genetics the most thrilling field in the sciences The study of genes and genomics today explores all aspects of the life with relevance in the lab in the doctor's office in the courtroom and even in social relationships In this helpful guidebook one of the most respected and accomplished human geneticists of our time communicates the importance of genes and genomics studies in all aspects of life With the use of core concepts and the integration of extensive references this book provides students and professionals alike with the most in depth view of the current state of the science and its relevance across disciplines Bridges the gap between basic human genetic understanding and one of the most promising avenues for advances in the diagnosis prevention and treatment of human disease Includes the latest information on diagnostic testing population screening predicting disease susceptibility pharmacogenomics and more Explores ethical legal regulatory and economic aspects of genomics in medicine Integrates historical classical genetics approach with the latest discoveries in structural and functional genomics

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