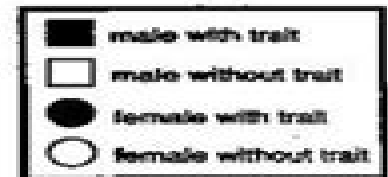


By studying a human pedigree, you can determine whether a trait is dominant or recessive. To interpret the three pedigrees below, use the same key shown at the right. Of course, the individual with the trait could be homozygous dominant or heterozygous dominant.



_____ How do you know? _____

P₁

P₂

F₁

F₂

The pedigree chart illustrates the inheritance of a trait across four generations:

- P generation:** A male (square) and a female (circle) are shown. The male is affected (filled square), and the female is unaffected (open circle).
- F₁ generation:** They have four children: two males and two females. The first male is affected (filled square), and the other three are unaffected (open shapes).
- F₂ generation:** The first male from the F₁ generation (affected) is mated with an unaffected female. They have four children: one affected male, one unaffected male, and two unaffected females.
- F₃ generation:** The first male from the F₂ generation (affected) is mated with an unaffected female. They have four children: one affected male, one unaffected male, and two affected females (filled circles).

Explain. _____

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The Practical Guide to the Genetic Family History Robin L. Bennett, 2011-09-20 HELPS YOU DEVELOP AND ASSESS PEDIGREES TO MAKE DIAGNOSES EVALUATE RISK AND COUNSEL PATIENTS The Second Edition of The Practical Guide to the Genetic Family History not only shows how to take a medical family history and record a pedigree but also explains why each bit of information gathered is important It provides essential support in diagnosing conditions with a genetic component Moreover it aids in recommending genetic testing referring patients for genetic counseling determining patterns of inheritance calculating risk of disease making decisions for medical management and surveillance and informing and educating patients Based on the author's twenty five years as a genetic counselor the book also helps readers deal with the psychological social cultural and ethical problems that arise in gathering a medical family history and sharing findings with patients Featuring a new Foreword by Arno Motulsky widely recognized as the founder of medical genetics and completely updated to reflect the most recent findings in genetic medicine this Second Edition presents the latest information and methods for preparing and assessing a pedigree including Value and utility of a thorough medical family history Directed questions to ask when developing a medical family history for specific disease conditions Use of pedigrees to identify individuals with an increased susceptibility to cancer Verification of family medical information Special considerations when adoptions or gamete donors are involved Ethical issues that may arise in recording a pedigree Throughout the book clinical examples based on hypothetical families illustrate key concepts helping readers understand how real issues present themselves and how they can be resolved This book will enable all healthcare providers including physicians nurses medical social workers and physician assistants as well as genetic counselors to take full advantage of the pedigree as a primary tool for making a genetic risk assessment and providing counseling for patients and their families

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