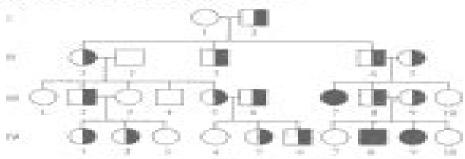
Pedigree Worksheet 1

Sickle Cell Anemia

Analyse the pedigree for sickle-cell anersia. Use the key below to the left to help. Remember, sickle-cell anersia is a recessive blood disorder. After analysing, answer the questions.

| State | Streets
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| Contro



- 1. How many deservations are represented in the projecte?
- In generation I, which parent is between your for the recessive abote?
- How many offspring from generation I inherited the sickle-cell allele?
- 4. Which individual in generation it marries a spouse who is homography dominant?
- 5. In which generation does the first case of sixtle-cell assersia appear?
- 6. Which appropriation contains the result male contains?
- T. Home any individuals had investuous relationships? If so, which ones?
- 8. Can two certiers produce an individual with sickle-cell assente?
- Can a normal homopagous individual produce offspring with sickle-call argents?
- 10. Which parents produce two children with siddle-cell aromia?
- If individual IV-1 mates with a male center for siddle-cell trait, what is the chance their first offspring, will have siddle-cell anomal?
- If individual IV-10 mates with another individual, is there any way an offspring could have sixide-cell ancesta? Explain.

Pedigree Worksheet Sickle Cell Anemia Answers

Alvin Silverstein, Virginia B.
Silverstein, Laura Silverstein Nunn

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Basic Genetics .1998-04-13 Sickle Cell Anemia and Your Child Roland B. Scott, Althea D. Kessler, 1960 Answers to Common Ouestions about Sickle Cell Disease ,1973 Sickle Cell Anemia and Sickle Cell Trait Job Corps (U.S.),1972 Adolescents with Sickle Cell Anemia and Sickle Cell Trait National Institutes of Health (U.S.), 1978 Sickle Cell **Anemia** Judy Monroe Peterson, 2008-08-15 Sickle cell anemia is a genetic disease of the blood It is caused by a defect in one gene of a person Genes are the elements in cells that carry the information that determines traits such as hair or eye color In sickle cell anemia a defect in the gene controls how hemoglobin is made This defect can be passed from parents to their children Students explore the history of sickle cell anemia the pioneering doctors who studied its cause and early treatments They also investigate hemoglobin S who gets sickle cell and how the gene mutation is inherited. They learn about the different types of sickle cell disease and treatments including blood transfusions and bone marrow transplants and some of the promising new research in medicines and gene therapy Questions & Answers Missouri. Department of Health, Missouri. Sickle Cell Anemia Program, 1987* Sickle cell anemia Rumi Michael Leigh, 2022-08-21 Sickle Cell Anemia Things You Should Know Questions and Answers explains this inherited blood disorder in a simple question and answer format covering causes symptoms complications and treatments Inside you will learn what sickle cell anemia is how it affects red blood cells and hemoglobin and why these cells become rigid sticky and sickle shaped The book explains how sickle cell anemia shortens the life of red blood cells and disrupts normal blood flow leading to serious health problems It covers symptoms such as extreme pain fatigue infections slow growth delayed puberty swelling in the hands and feet and vision problems Complications are also discussed including stroke pulmonary hypertension organ damage blood clots gallstones priapism ulcers blindness and pregnancy risks Readers will also find explanations of diagnostic methods from blood tests to prenatal testing and treatment options such as medications blood transfusions and bone marrow transplant This book is suitable for anyone who wants to understand sickle cell anemia its effects on the body and current approaches to treatment in a direct and easy to read way Sickle Cell Anemia and Sickle Cell Trait United States. Department of Labor.1972 Sickle Cell Trait New York Heart Association. Committee on Sickle Cell Anemia, Children's Blood Foundation, New York (State). Department of Health, 1972 Sickle Cell Anemia National Institutes of Health (U.S.), 1965 Planning for Sickle Cell Anemia in Pennsylvania Pennsylvania. Irvis Committee on Sickle Cell Anemia, 1973

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