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POGIL - Meiosis

How does sexual reproduction lead to genetic variation?

Withy?

Cells reproduce through mitosis to make exact copies of the original cell. This is done for growth and repair. Sexually-reproducing organisms have a second form of cell division that produces reproductive cells with half the number of chromosomes. This process is called **meiosis**, and without it, humans, oak trees, beetles, and all other sexually reproducing organisms would be vastly different than they are today.

Model 1 - Meiosis I



- 1. According to Model 1, the cells that go through meiosis are found in which type of organs?
- Considering what you already know about mitosis in cells, what event must take place during interphase before a cell proceeds to division?
- 3. What two structures make up a single replicated chromosome?
- 4. In Model 1, how many replicated chromosomes does the cell contain during prophase?

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processes in nondividing cells **Meiosis and Gametogenesis**, 1997-11-24 In spite of the fact that the process of meiosis is fundamental to inheritance surprisingly little is understood about how it actually occurs There has recently been a flurry of research activity in this area and this volume summarizes the advances coming from this work All authors are recognized and respected research scientists at the forefront of research in meiosis Of particular interest is the emphasis in this volume on meiosis in the context of gametogenesis in higher eukaryotic organisms backed up by chapters on meiotic mechanisms in other model organisms. The focus is on modern molecular and cytological techniques and how these have elucidated fundamental mechanisms of meiosis Authors provide easy access to the literature for those who want to pursue topics in greater depth but reviews are comprehensive so that this book may become a standard reference Key Features Comprehensive reviews that taken together provide up to date coverage of a rapidly moving field Features new and unpublished information Integrates research in diverse organisms to present an overview of common threads in mechanisms of meiosis Includes thoughtful consideration of areas for future investigation Recombination and Meiosis Richard Egel, Dirk-Henner Lankenau, 2007-11-13 This fascinating volume addresses the processes and mechanisms taking place in the cell during meiosis and recombination It covers multicellular eukaryotes such as Drosophila Arabidopsis mice and humans Once per life cycle mitotic nuclear divisions are replaced by meiosis I and II reducing chromosome number from the diploid level to a haploid genome reshuffling the homologous chromosomes by their centromeres and recombining chromosome arms by crossing over Controlling Events in Meiosis Clive W. Evans, Hugh G. Dickinson, 1984 Keeney, 2010-07-19 Each generation in a sexually reproducing organism such as a fly or a mouse passes through the bottleneck of meiosis which is the specialized cell division that gives rise to haploid reproductive cells sperm eggs spores etc The principal function of meiosis is to reduce the genome complement by half which is accomplished through sequential execution of one round of DNA replication followed by two rounds of chromosome segregation Within the extended prophase between DNA replication and the first meiotic division in most organisms homologous maternal and paternal chromosomes pair with one another and undergo homologous recombination which establishes physical connections that link the homologous chromosomes until the time they are separated at anaphase I Recombination also serves to increase genetic diversity from one generation to the next by breaking up linkage groups The unique chromosome dynamics of meiosis have fascinated scientists for well over a century but in recent years there has been an explosion of new information about how meiotic chromosomes pair recombine and are segregated Progress has been driven by advances in three main areas 1 genetic identification of meiosis defective mutants and cloning of the genes involved 2 development of direct physical assays for DNA intermediates and products of recombination and 3 increasingly sophisticated cy logical methods that describe chromosome behaviors and the spatial and temporal patterns by which specific proteins associate with meiotic chromosomes Holt Biology: Meiosis and sexual reproduction, 2003 **Meiosis** Scott Keeney, 2016-08-23 Each generation in a sexually

reproducing organism such as a fly or a mouse passes through the bottleneck of meiosis which is the specialized cell division that gives rise to haploid reproductive cells sperm eggs spores etc The principal function of meiosis is to reduce the genome complement by half which is accomplished through sequential execution of one round of DNA replication followed by two rounds of chromosome segregation Within the extended prophase between DNA replication and the first meiotic division in most organisms homologous maternal and paternal chromosomes pair with one another and undergo homologous recombination which establishes physical connections that link the homologous chromosomes until the time they are separated at anaphase I Recombination also serves to increase genetic diversity from one generation to the next by breaking up linkage groups The unique chromosome dynamics of meiosis have fascinated scientists for well over a century but in recent years there has been an explosion of new information about how meiotic chromosomes pair recombine and are segregated Progress has been driven by advances in three main areas 1 genetic identification of meiosis defective mutants and cloning of the genes involved 2 development of direct physical assays for DNA intermediates and products of recombination and 3 increasingly sophisticated cy logical methods that describe chromosome behaviors and the spatial and temporal patterns by which specific proteins associate with meiotic chromosomes Chromosome Morphology and Meiosis in Some Hexapolid Wheats Ahmedbasha S., 1968 Mitosis and Meiosis Gary Parker, 1968 **Chapter Resource 7** Meiosis/Reproduction Biology Holt Rinehart & Winston, Holt, Rinehart and Winston Staff, 2004-01-01 Looking at Chromosomes John McLeish, Brian Snoad, 1959

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