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Culturing Techniques For Drosophila

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7516 Genetics: Scavardaand Hartl MATERIALSANDMETHODS Recoveryofarosy (ry) MutationinD. Simulans. Astrain Of D. Simulans Carrying Avermilion (v) Mutationwasprovided By Victoria Finnerty. Males Ofthis Strain Were Subjected To Mutagenesis With Ethyl Methanesulfonate (21) Prior To Mat- Ing With V Females. Approximately 300 F1 Offspring From Treated 3th, 2024

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Embryo10 (FIG.1a). This Work Was Revolutionary, because It Was The First Mutagenesis In Any Multicellular Organ-ism That Attempted To Find Most Or All Of The Mutations That Affect A Given Process, and Because It Was One Of The First Screens For Phenotypes In The Embryo Rather Tha 1th, 2024

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Drosophila Schneider 2 (S2) Cells - University Of Washington

S2 Cells Should Be Subcultured To A Final Density Of 2 To 4 X 106 Cells/ml. Do Not Split Cells Below A Density Of 0.5 X 106 Cells/ml. For Example, 2 Ml Of Cells From A 75 Cm2 Flask At A Density Of 2 X107 Cells/ml Sh 3th, 2024

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Under Normal Conditions Each Cyst Of Meiocytes In D. Melanogaster Males Contains 16 Cells That Are The Result Of Four Relatively Synchronous Spermatogo- Nial Divisions. Intercellular Cytoplasmic Bridges (ring Canals), The Products Of Incomplete Spermatogoni 1th, 2024

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Cross 3: P Generation = Wild Type Male * White Eye Female Parental Cross F1 Cross 2. Refer To The Punnett Squares Above. In The Boxes Below, Record The Expected Ratios For The Genotypes And Phenotypes Of The F1 And F2 Generations In The Experiment. Cross 1: P Generation = Sepia Male * Wild Type Female /P1 Genotypes: 1th, 2024

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See Cross #1 And Cross #2 Backcrossing The F1 Generation With The Homozygote Recessive Mutant Is Another Cross That Can Be Set Up To Demonstrate The Mode Of Inheritance. In This Case The Back Cross Progeny Would Be Expected To Segregate Into Two Classes (mutant And Wild Type) Of Equal Size. Monohybrid Cross For A Character 1th, 2024

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