

Exercise 10: Linkage and Mapping

8. Four different female *Drosophila* were isolated, each homozygous for the autosomally-linked gene(s) *scd* (for scute) *sup* (for supernumbering). These genes are in the same cluster, with a distance of 50 cM. Genes in the *scd* subfamily are very similar, with 8.0% homology, respectively, below them. The percentage recombination for each female's genotype are shown below. Each female's genotype was homozygous *scd*¹, while *sup* was *sup*¹, *sup*², *sup*³, or *sup*⁴ (increasingly more different). These females are heterozygous with respect to the previously defined *scd* subfamily. What is the order of the genes from left to right?

Female id.		Female id.	
<u>scd</u> <u>sup</u>	Type of recombination	<u>scd</u> <u>sup</u>	Type of recombination
<u>scd</u> <u>sup</u>	relative frequency	<u>scd</u> <u>sup</u>	relative frequency
1. <u>scd</u> <u>sup</u>		1. <u>scd</u> <u>sup</u>	
2. <u>scd</u> <u>sup</u>		2. <u>scd</u> <u>sup</u>	
3. <u>scd</u> <u>sup</u>		3. <u>scd</u> <u>sup</u>	
4. <u>scd</u> <u>sup</u>		4. <u>scd</u> <u>sup</u>	
5. <u>scd</u> <u>sup</u>		5. <u>scd</u> <u>sup</u>	
6. <u>scd</u> <u>sup</u>		6. <u>scd</u> <u>sup</u>	
7. <u>scd</u> <u>sup</u>		7. <u>scd</u> <u>sup</u>	
8. <u>scd</u> <u>sup</u>		8. <u>scd</u> <u>sup</u>	

9. Assume that both the *scd* and *sup* genes are completely dominant to the alleles for *scd* and *sup* respectively. Consider the following crosses between the subfamilies (you have to fill out the first column, the frequencies for the subfamilies are in parentheses). Fill in the subfamily (1-4 frequency), if it is statistically different, not paired gene and reciprocal with a different subfamily combination. The *scd* subfamily has either been substituted for the reciprocal subfamily and remains the same. Fill in the rest of the table.

Female subfamily	Male
Female 1	Male 1
Female 2	Male 1
Female 3	Male 1
Female 4	Male 1
Female 1	Male 2
Female 2	Male 2
Female 3	Male 2
Female 4	Male 2

Section 3 Gene Linkage And Mapping Study Guide Answers

BM King



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Study Guide and Solutions Manual for Students, to Accompany General Genetics David Freifelder, Leon A. Snyder, 1985

Study Guide for Genetics, Third Edition, Daniel L. Hartl Rowland H. Davis, Stephen J. Weller, John P. Carulli, 1994

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the past 50 years have been dramatically changed the understanding and management of inherited disorders and are beginning to have a major impact on the practice of medicine overall The rapidity of these advances means that clinicians and scientists in the field are often unfamiliar with the key research that has led to many developments that now are accepted and familiar Few have time to search or the original papers which are scattered and often difficult to obtain This collection has been edited mainly for medical geneticists and genetics researchers who wish to learn more about how their field originated and developed Brief clearly written commentaries on each paper and section place the work in its current context and serve to unify the different parts of the book They also help make it a readable and authoritative source of information The papers chosen fall into several groups First are classic descriptions of important genetic disorders often from the pre mendelian era The following sections deal with the definition of human mendelian inheritance the origins of human cytogenetics the early development of the human gene map and the transition from biochemical genetics to human molecular genetics the relatively recent studies that have shown how mendelian principles are increasingly modifiable and finally advances in the treatment and management of genetic disorders which are placed in their social context

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