1. In <u>Drosophila</u> the genes *A B D* are linked. You cross a strain that is homozygous for *d* with a strain that is homozygous for *ab*. Below are the phenotypes and progeny of a test cross. Outline the cross and determine the map for the genes. Calculate the coefficient of coincidence and the interference.

<u>Phenotype</u>	<u>Number</u>
ABd	389
abD	413
ABD	60
abd	68
aBd	29
AbD	34
Abd	3
aBD	4

2. In <u>Drosophila</u> the genes *XYZ* are linked. You cross a strain that is homozygous for *x z* with a strain that is homozygous for *y*. Below are the phenotypes and progeny of a test cross. Outline the cross and determine the map. Calculate the coefficient of coincidence and the interference.

<u>Phenotype</u>	<u>Number</u>
Xyz	13
XYZ	220
XyZ	2480
xyZ	170
xYz	2429
XYz	178
xYZ	19
xyz	233

3. In Tasmanian devils, three recessive mutations are h (hairy nostrils), r (rat tail), and f (short fang). Outline a test cross between a devil with hairy nostrils and rat tail with a devil with short fangs. From the data below, calculate the map, the coefficient of coincidence, and the interference.

<u>Phenotype</u>	<u>Number</u>
HRF	73
hrf	63
Hrf	96
hRF	110
HrF	2
hRf	2
hrF	306
HRf	348

4. You are conducting genetic research on rare Tasmanian fuzz puppies. To your dismay, you find that your arch rival, an unscrupulous geneticist at the University of Rochester has just published a genetic map of the very same mutants that you are studying. Her genetic map is as follows:

Rd Bt Fz ■ 10.4 ■ 6.2 ■

Rd is a mutation that gives fuzz puppies blood-shot eyes. Bt is a mutation that produces black tongues. Fz is a mutation that causes the fuzzy hair on their ears to be straight. All mutations are recessive. You decide to repeat her experiment and cross a wild-type strain with a strain that is homozygous recessive for all traits. The results of your cross is below. Is her map consistent with yours?

Phenotype Number

rd bt fz	310
RD BT FZ	328
RD bt fz	40
rd BT FZ	44
rd bt FZ	35
RD BT fz	37
rd BT fz	4
RD bt Fz	4

5. Three <u>Drosophila genes map as follows</u>:



Using the map, calculate the expected F2 progeny for the cross

**ABD** x **abd** in which a test cross was performed with the F1 progeny. Assume a coefficient of 1.0. and a total of 3000 progeny. How would the numbers change if the coefficient of coincidence was 0.8? Determine whether the data below is consistent with your map.

<u>Number</u>	
	1121
	1139
	257
	266
	87
	112
	8
	10
	Number

6. You cross a dihybrid cross with a strain of flies that is a and a strain that that is b. The results of the F2 are given. Use X<sup>2</sup> to help you determine if a and b are linked.

<u>Phenotype</u>	<u>Number</u>	
AB		1757
Ab		583
aB		605
ab		225

7. You cross a dihybrid cross with a strain of flies that is a and a strain that it b. The results of the F2 are given. Use  $X^2$  to help you determine if a and b are linked.

<u>Phenotype</u>	Number	
AB		1827
Ab		424
aB		386
ab		298

## EUKARYOTIC MAPPING PRACTICE PROBLEMS - SOLUTIONS

F1 ABd/abD x abd/abd

Test Cross

Number	
389	highest number
413	= nonrecombinants
60	intermediate number
68	= single XO
29	intermediate number
34	= single XO
3	smallest number
4	= double XO
	<u>Number</u> 389 413 60 68 29 34 3 4

In the double XO's, B is the gene that has changed linkage. Therefore, the sequence is --A - - - B - - - D - - -.

A - B =  $[(29 + 34 + 3 + 4) / 1000] \times 100 = 7 \text{ cM}$ 

B - D = [ (60 + 68 + 3 + 4) / 1000 ] x 100 = 13.5 cM

Coef. Coinc = obsvd 2XO's / exp 2XO's

= (7/1000) / (.007 x .135)

interference = 1 - coef. coinc. = 1 - 0.74 = 0.26 = pos interf.

2. P1 xYz/xYz x XyZ/XyZ

F1	xYz/XyZ	Х	xyz/xyz
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Test Cross			
<u>Phenotypes</u>	Number		
xYz	2429	) highest number	
<u>XyZ</u>	2480	) = nonrecombinants	
xyZ	17(	) intermediate number	
<u>XYz</u>	178	3 = single XO	
XYZ	220	) intermediate number	
xyz	233	3 = single XO	
Xyz	13	smallest number	
<u>xYZ</u>	19	= double XO	

In the double XO's, Z is the gene that has changed linkage. Therefore, the sequence is -X - -Z - -Y - -.

X - Z = [ (170 + 178 + 13 + 19) / 5742 ] x 100 = 6.62 cM

Z - Y = [ (220+ 233 + 13 + 19) / 5742 ] x 100 = 8.45 cM

Coef. Coinc = obsvd 2XO's / exp 2XO's

 $= (32 / 5742) / (.066 \times .0845)$ 

interference = 1 - coef. coinc. = 1 - 1.0 = 0 = no interf.

## 3. P1 hrF/hrF x HRf/Hrf

## F1 hrF/HRf x hrf/hrf

Test Cross		
<u>Phenotypes</u>	Number	
HRF	73	intermediate number
<u>hrf</u>	63	= single XO
Hrf	96	intermediate number
hRF	110	= single XO
HrF	2	smallest number
hRf	2	= double XO
hrF	306	highest number
HRf	348	= nonrecombinants

In the double XO's, His the gene that has changed linkage. Therefore, the sequence is  $\,$  -- F - - - R - - .

H - F = [ (73 + 163 + 12 + 2) / 1000 ] x 100 = <u>14 cM</u>

F - R = [ (96 + 110 + 2 + 2) / 1000 ] x 100 = <u>21 cM</u>

Coef. Coinc = obsvd 2XO's / exp 2XO's

 $= (4 / 1000) / (0.14 \times 0.21)$ 

= <u>0.136</u>

interference = 1 - coef. coinc. = 1 - 0.136 = 0.864 = pos interf.

4. When comparing results of two experiments, you must compare data, not maps. Therefore, you must calculate the data you would have expected based on your competetors map. These numbers become your expected values, and your numbers become your observed values in a X<sup>2</sup> test.

Of 802 progeny, 10.4% should show a recombination event between Rd and Bt:

 $802 \times 0.104 = 83.4 = \text{single} + \text{double recombinants}$ 

6.2% should show a recombination event between Bt and Fz:

 $802 \times 0.062 = 49.7 = \text{single} + \text{double recombinants}$ 

The double recombinants will be

802 x 0.104 x 0.062 = 5.2

The Rd - Bt single recombinants will be

83.4 - 5.2 = 78.2

The <u>Bt - Fz single recombinants</u> will be

49.7 - 5.2 = 44.5

Finally, the nonrecombinants will be

802 - 78.2 - 44.5 - 5.2 = 674.1

Now we are ready to set up the X2 table. But don't forget that each of the four classes of phenotypes that you have calculated are actually 8 pairs of reciprocals. Therefore, you must take the numbers and divide them in half, one for one of the pair, and one for the other.

<u>PHENOTYPE</u>	<u>OBSERVED</u>	<u>EXPECTED</u>	<u>O - E</u>	(0 - E) <sup>2</sup>	<u>DIV EXPT</u>
rd bt fz	310	337.05	-27.05	731.70	2.17
RD BT FZ	328	337.05	-9.05	81.90	0.24
RD bt fz	40	39.10	0.90	0.81	0.02
rd BT FZ	44	39.10	4.90	24.01	0.61
rd bt FZ	35	22.25	12.75	162.56	7.31
RD BT fz	37	22.25	14.75	217.56	9.78
rd BT fz	4	2.60	1.40	1.96	0.75
RD bt Fz	4	2.60	1.40	1.96	0.75
				_	

χ<sup>2</sup> = 21.64

degrees of freedom = 7 probability = 0.05 Crit.  $X^2 = 14.067$ 

 $H_{\mbox{\scriptsize O}}$  = there is no significant difference between observed and expected values

$$X2 = 21.64 > 14.067$$

Therefore, reject the  $H_0$ . Your data does not agree with your competetor's map. But who is correct?????

5. Of 3000 progeny, 15.7% should show a recombination event between A and B:

 $3000 \times 0.157 = 471 = \text{single} + \text{double recombinants}$ 

3.9% should show a recombination event between B and D:

 $3000 \times 0.039 = 117 = single + double recombinants$ 

The double recombinants will be

3000 x 0.157 x 0.039 = 18

The <u>A - B single recombinants</u> will be

471 - 18 = 453

The <u>B - D single recombinants</u> will be

117 - 18 = 99

Finally, the nonrecombinants will be

3000 - 453 - 117 - 18 = 2430

Now we are ready to set up the X2 table. But don't forget that each of the four classes of phenotypes that you have calculated are actually 8 pairs of reciprocals. Therefore, you must take the numbers and divide them in half, one for one of the pair, and one for the other.

<u>PHENOTYP</u>	<u>EXPECTED</u>	<u>EXPECTED</u>	<u>OBS - EXP</u>	(O - E) <sup>2</sup>	<u>DIV BY</u>
<u>E</u>				<u></u>	<u>EXP</u>
ABD	1121	1215.0	-94.00	8836.00	7.27
abd	1139	1215.0	-76.00	5776.00	4.75
aBD	257	226.5	30.50	930.25	4.11
Abd	266	226.5	39.50	1560.25	6.89
ABd	87	49.5	37.50	1406.25	28.41
abD	112	49.5	62.50	3906.25	78.91
AbD	8	9.0	-1.00	1.00	0.11
aBd	10	9.0	1.00	1.00	0.11
				x <sup>2</sup> =	130.57
degrees of freedom = 7		probability =	0.05 Crit. X <sup>2</sup>	<sup>2</sup> =14.067	

 $H_{\text{O}}$  = there is no significant difference between observed and expected values

X2 = 130.57> 14.067

Therefore, reject the H<sub>0</sub>. Your data is inconsistent

If the coefficient of coincidence is 0.8, that means you are only seeing 80% of the double recombination that you expect. Therefore, the double recombinants will be

0.8 x [3000 x 0.157 x 0.039] = 0.8 x 18 = <u>14</u>

You then subtract 14 instead of 18 from the other classes of recombinants.

6. If the two genes are <u>unlinked</u> then they would segregate in a 9:3:3:1 ratio. If they were linked, then there would be a substantial departure from this ratio, but without knowing their distance, we could not predict the phenotype numbers. Therefore, we will test to see if they are unlinked.

<u>Phenotype</u>	<u>Observed</u>	<u>Expected</u>	<u>O - E</u>	<u>(0 - E )<sup>2</sup></u>	<u>/ Exp</u>
AB	1757	1782.9	-25.90	670.81	0.38
Ab	583	594.4	-11.40	129.96	0.22
aB	605	594.4	10.60	112.36	0.19
ab	225	198.1	26.90	723.61	3.65
				X <sup>2</sup> =	4.44
			~		

degrees of freedom = 3 probability = 0.05 Crit.  $X^2$  =7.815

 $H_{\mbox{\scriptsize O}}$  = there is no significant difference between observed and expected values

Therefore, accept the H<sub>o</sub>. The genes are unlinked

7. If the two genes are <u>unlinked</u> then they would segregate in a 9:3:3:1 ratio. If they were linked, then there would be a substantial departure from this ratio, but without knowing their distance, we could not predict the phenotype numbers. Therefore, we will test to see if they are unlinked.

<u>Phenotype</u>	<u>Observed</u>	<u>Expected</u>	<u>O - E</u>	<u>(0 - E)</u> 2	<u>/ Exp</u>
AB	1827	1651	176.00	30976.00	18.76
Ab	424	550	-126.00	15876.00	28.87
aB	386	550	-164.00	26896.00	48.90
ab	298	183	115.00	13225.00	72.27

χ<sup>2</sup> = 168.80

degrees of freedom = 3

probability = 0.05 Crit.  $X^2 = 7.815$ 

 $H_{\text{O}}$  = there is no significant difference between observed and expected values

Therefore, reject the  ${\rm H}_{\rm O}$ . The genes are probably linked with some degree of recombination between them.