WEEK 7 PROBLEMS

Problems From Chapter 7

- 7.1 What is the difference between a selected marker and a counterselected marker? Why are both necessary in an Hfr X F- mating?
- 7.2 An Hfr strain transfers genes in alphabetical order, abc. In an Hfr $a+b+c+str^S$ X F- a-b-C str-r mating, do all $b+str^R$ recombinants receive the a+ allele? Are all $b+str^R$ recombinants also a+? Why or why not?
- 7.3 Bacteriophage P1 transduction using a donor bacterial strain that has the wildtype alleles of the closely linked markers *a*, *b*, *c*, and *d* shows the following percentages of cotransduction:

a-b:29	a-c: 2	a-d: 5
b-c: 0	b-d:1	c-d: 50

What is the order of the genetic markers?

7.4 Bacterial cells of genotype *pur- pro+ his+* were transduced with PI bacteriophage grown on bacteria of genotype *pur+ pro- his-.* Transductants containing *pur+* were selected and tested for the unselected markers *pro* and *his.* The numbers of *pur+* colonies with each of four genotypes were as follows:

pro+	his+	102	
pro-	his+	25	
pro+	his-	160	
pro-	his-	1	

Is pur closer to the his gene or the pro gene?

What is the co-transduction frequency between pur and his? between pur and pro? Why can't the co-transduction frequency be determined between pro and his from this experiment?

7.5 An experiment was carried out in E. *coli* to map five genes around the chromosome using each of three different Hfr strains. The genetic markers were *bio*, *met*, *phe*, *his*, and

trp. The Hfr strains were found to transfer the genetic markers at the times indicated here. Construct a genetic map of the *E. coli* chromosome that includes all five genetic markers, the genetic distances in minutes between adjacent gene pairs, and the origin and direction of transfer of each Hfr. Complete the missing entries in the table, which are indicated by question marks.

bio	met	phe	his
26	44	66	?
phe	met	bio	trp
?	26	44	75
phe	his	?	bio
6	27	35	?
	26 phe ? phe	phe met ? 26 phe his	26 44 66 phe met bio ? 26 44 phe his ?

Problems From Chapter 10

- 10.1 Reverse transcriptase, like most enzymes that make DNA, requires a primer. Explain why, when cDNA is to be made for the purpose of cloning a eukaryotic gene, a convenient primer is a short sequence of poly(dT). Why does this method not work with a prokaryotic messenger RNA?
- 10.2 A DNA microarray is hybridized with fluorescently labeled reverse-transcribed DNA as described in the text, where the control mRNA (C) is labeled with a green fluor and the experimental mRNA (E) with a red fluor. Indicate what you can conclude about the relative levels of expression of a spot in the microarray that fluoresces:
- (a) Red
- (b) Green
- (c) Yellow
- (d) Orange
- (e) Lime green