### PLNT2530 PLANT BIOTECHNOLOGY

# FINAL EXAMINATION

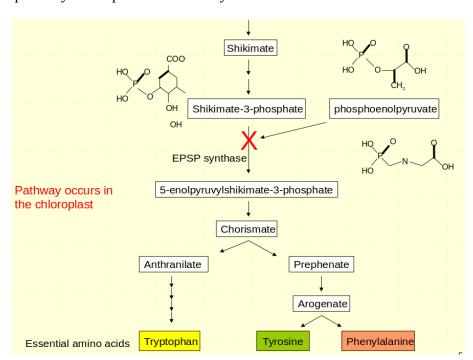
Thursday, April 12, 2018 18:00 to 20:00 Engineering E2-130, Seats 1 - 18

Answer any combination of questions totalling to <u>exactly</u> 100 points. If you answer questions totalling more than 100 points, answers will be discarded at random until the points equal 100. This exam is worth 40% of the course grade. The questions available total to 120 points.

Hand in these question sheets along with your exam book. Question sheets will be shredded.

Ways to write a readable and concise answer:

- i. Just answer the question. Save time by specifically addressing what is asked. Don't give irrelevant background if it doesn't contribute to the question that was asked.
- ii. Avoid stream of consciousness. Plan your answer by organizing your key points, and then write a concise, coherent answer. Make your point once, clearly, rather than repeating the same thing several times with no new information. iii. Point form, diagrams, tables, bar graphs, figures are welcome. Often they get the point across more clearly than a long paragraph.
- iv. Your writing must be legible. If I can't read it, I can't give you any credit.
- 1. (5 points) A diagram of the shikimic acid pathway is shown below. Give the name of the chemical that inhibits this pathway at the point indicated by the X.



2. (5 points) The Arctic non-browning apple represents what is arguably a new era in plant biotechnology. What is the significance of products such as the Arctic apple that distinguishes it from the previous generation of biotech crops, such as herbicide resistant soybean or canola, or Bt cotton or corn?

3. (10 points) Fill in the blanks.

Eukaryotes inac	ctivate	specific genes	by
methylation of	a	residues.	

By preventing <u>b</u>, methyation acts as a mechanism for selectively down-regulating genes.

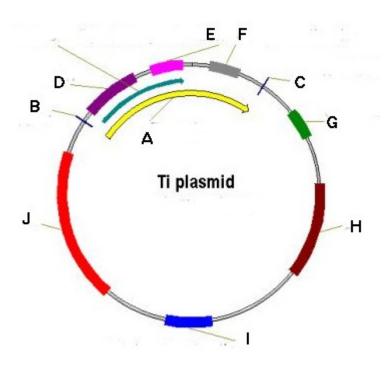
CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>	
CpG	СрG		
	cpg cpg cpg		
CH <sup>3</sup>	CH <sup>3</sup> CH <sup>3</sup> CH <sup>3</sup>		

In embryos, methylation is often

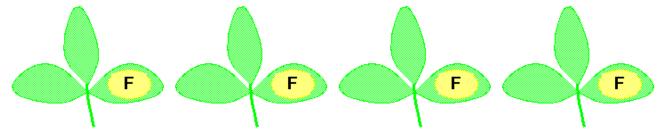
<u>c</u> . Subsequently during development, certain genes are methylated, and are d in the adult.

Methylation patterns are often \_\_\_\_\_e from one generation to the next.

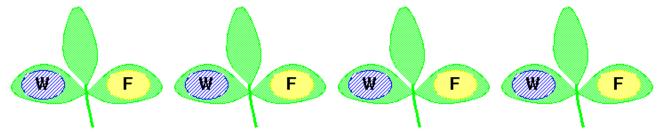
- 4. (10 points) Explain why a single gene insertion at a single locus in a transgenic plant will usually give a single band on a Southern blot, when the same gene is used as a hybridization probe. Draw a diagram that illustrates your point.
- 5. (10 points) Explain the distinction between the following terms, with reference to biotech crops: homozygous, heterozygous, hemizygous, and biallelic. Using standard genetic notation (eg. X for dominant, x for recessive and so forth) how would the genotypes for each be represented?
- 6. (5 points) Describe the distinction between transformation and transient expression.
- 7. (10 points) For the map of the octopine Ti plasmid shown below, give the name for each feature marked with a letter, A J. You do not need to describe each feature, just list its name.



- 8. (15 points) Soybeans were transformed with a pea gene (P) for resistance to fungi. You wish to determine whether the P gene will confer fungal resistance in soybean. To do this, you would inoculate leaves with a fixed concentration of fungal spores suspended in sterile water (**F**) and observe whether resistance or susceptibility is seen. One leaf is inoculated, and the other left uninoculated as a control. After observing the response, RNA is isolated from leaf tissue to observe changes in gene expression by RNAseq. Expression of genes is compared between fungal-treated leaves and control leaves.
- a) Part of the experimental design includes inoculating 4 different plants at the same time. Why are 4 plants inoculated, as opposed to 1? What would be missed if we did the experiment only once? (You can ignore the trivial reason that 4 plants would yield more RNA.)



b) An alternative design is shown below, in which control leaves, rather than remaining uninoculated, are inoculated with sterile water (**W**). Why is this a better experimental design?

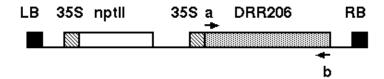


- c) There are two possible ways to extract the RNA. One is to pool inoculated leaves from all 4 plants, and do a single RNA extraction. The other is to isolate RNA independently from each leaf and sequence each RNA sample separately. This is a lot of extra work. Why is this a better experimental design? In other words, once you have the four measurements for each gene, you're going to average them anyway, so why not just pool the leaves and simplify things? What information would you be throwing away if you pooled leaves?
- 9. (10 points) Explain why mutations generated by the CRISPR-Cas9 system will be essentially indistinguishable from naturally-occuring mutations. Why is this important in light of regulation and labeling of genetically-modified crops?

10. (15 points) There are many ways of verifying transformation of a plant, each with its advantages and disadvantages. The construct shown below contains two genes, under the control of the 35S promoter, between the T-DNA left and right borders. The DRR206 gene has been shown to provide some resistance to the blackleg fungus in transgenic canola. The nptII gene encodes neomycin phosphotransferase.

The following table lists a number of possible experiments, and the types of information that might be obtained. A "yes" means that the experiment is likely to provide a particular type of information. A "no" means that the experiment will not tell us that information. Some of the table has already been filled in.

Your job is to rewrite the table in your answer book, and complete it using yes/no answers. To save time, feel free to shorten or abbreviate the column and row headings, as long as you keep them in the same order.



	What the experiment tells us:					
	Presence of DRR206 gene	Copy number of DRR206 gene	Expression of DRR206	Presence of nptII gene		
Experiment:						
PCR of genomic DNA using primers a and b	yes	no	no	no		
Southern blot using DRR206 gene as a probe						
Northern blot using DRR206 as a probe						
assay for root growth in kanamycin						
test for resistance to blackleg		no				

- 11. (5 points) What are the characteristics we should look for in choosing a plant species for production of pharmaceuticals?
- 12. (10 points) The slide below comes from the lecture material on Substantial Equivalence, in which the expression of over 51,000 genes was compared in different transgenic, mutagenized and
- a) What does comparison of gene expression between the three unmutagenized, untransformed rice lines tell us?
- b) To compare transgenics with controls, the authors used two different genes, BCBF1 and immunoglobulin. Why did they choose these two particular genes for evaluating the effects of transformation on gene expression?

Batista R,Saibo N, Lourenço T, Oliveira MM (2008) Microarray analyses reveal that plant mutagenesis may induce more transcriptomic changes than transgene Proc. Natl. Acad. Sci. USA 105:3640-3645, doi: 10.1073/pnas.070788105

<u>Hypothesis</u>: The genetic variation due to mutagenesis is greater than the genetic variation due to transformation

Oryza sativa L. spp. Japonica (rice) lines:

## Nipponbare

- unmutagenized, untransformed control
- Υ-irradiated, M1 generation ("unstable")
- Agrobacterium-transformed with BCBF1<sup>\*</sup> gene T1 generation ("unstable")

\*BCBF1 - Barley C-repeat binding factor. Activated during stress; interacts with dehydrationresponsive elements found in many stress-realted genes.

#### Bengal

- unmutagenized, untransformed control
- Agrobacterium-transformed with ScFv (immunoglobulin), T3 generation ("stable")

#### Estrella A

- · unmutagenized, untransformed control
- Y-irradiated, semi-dwarf phenotype, self-pollinated for 10 generations ("stable")

RNA from whole 12-day old seedlings Microarray: 51,279 probes ie. distinct transcripts

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13. (10 points) The following Southern blots were done during a project to transform soybean with a gene from pea for fungal resistance (**P**). Explain the results in B and C in light of what we have discussed regarding the basic genetics of transgenes.

