Fill in the Blank

- 1. Adventitious presence as the <u>unintended</u> or <u>unintentional</u> presence of another seed variety or genetic material, and/or trait(s) from another variety as a result of natural, mechanical or human means.
- 2. Lateral Flow Strips are a <u>immunoassay</u> test.
- 3. Non-herbicide tolerance traits such as insect resistance are often tested via an immunoassay utilizing either ELISA and/or lateral flow strip methods.
- 4. Molecular methods, including SNPs, may be used to test for trait purity for all defined traits including herbicide tolerance, insect resistance, and drought tolerance.
- 5. Herbicide tolerance traits can be categorized as **genetically** modified or **naturally** occurring.
- 6. Herbicide concentrations are typically expressed in parts per million (ppm) in lab test applications. Parts per million is <u>"unitless"</u> and is similar to a <u>percent</u>.
- 7. The use of check (or control) seed for each sample tested is essential to assure test accuracy and limit laboratory liability. Both negative and positive trait check samples should be included.
- 8. Immunoglobin is another name for <u>antibody</u>.
- 9. Element Specific AP testing could target the <u>promoter</u> and <u>terminator</u> sequences associated with many GMO events.
- 10. A typical ELISA kit purchased from a vendor contains the plate, <u>enzymes</u>, <u>substrate</u> and <u>detergent</u>.
- 11. SNPs are powerful genomic markers that assess a line's identity and purity.

True/False

- 12. True False Trait testing is regulated by the Federal Seed Act
- 13. True False The main advantage to herbicide bioassay is cost.
- 14. True False Lateral Flow Strips can be used to measure both AP and Trait Confirmation.

Short Answer

- 15. What four technologies can be used for trait detection?
- a. herbicide bioassay
- b. **ELISA**
- c. Lateral Flow Strips
- d. PCR
- 16. What is the difference between Quantitative PCR and Qualitative PCR?
 - Quantitative DNA-based testing methods yield a specific percentage of GM present in a given sample, while qualitative PCR is a DNA-based testing method that yields a positive or negative result for the presence of GM.
- 17. Please name two advantages of using Quantitative PCR for trait testing.
 - Highly sensitive (some assays detect as low as 0.01% GMO in a given sample).
 - Can typically be used on bulk seed, feed, and some processed food samples.
 - Recognized worldwide as the standard test method for GMO detection, and standards are easily obtained for testing.
 - Results are stored electronically (no gels or images to save).
 - Results are typically calculated automatically with software tools.
 - Can test for all GMO events with an all-purpose promoter/terminator assay.
- 18. Name two reasons why Lateral Flow strips are advantageous.
 - Less expensive to perform than DNA methods.
 - Currently provide a quantitative estimate of commercially available GMO traits, not just promoter and terminator. Useful if additional knowledge of trait is needed. (must use a quantitative strip reader).
 - Can test bulk seed and some processed seed/food products.
 - Less labor intensive and easier to perform than DNA methods.
 - Readers can save images of strip results for documentation and archiving results.
- 19. Trait impurities are often due to contaminations, caused by (name two)
 - Trait impurities are often due to contaminations of the seed source.
 - Contamination during production (outcrossing and selfing).
 - A seed mix of the seed lot.
 - Contamination during sampling or non-uniform sampling.

20. What is the difference between Polyclonal Anitbodies and Monoclonal Antibodies?

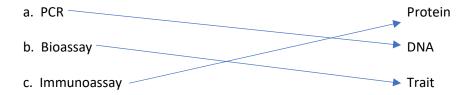
The two types of antibodies utilized in ELISA tests are polyclonal antibodies and monoclonal antibodies (Figure I.14.4). Polyclonal antibodies are produced by injecting animals, usually rabbits, with the target antigen. The animal's immune system responds to the introduction of foreign material by producing antibodies. After time, circulating antibodies are collected in the blood of the animal and subsequently purified. This antiserum contains many different antibodies for the target antigen, thus the term "poly". Monoclonal antibodies are produced by selecting antibody-producing cells from an immunized mouse and hybridizing them with cells derived from melanomas, creating "hybridoma" cells. The antibodies produced from individual hybrid cell clones are identical to one another, thus the term "mono".

- 21. In an ELISA test what would be three reasons that there is no color present in the wells at the end of the test?
 - Wrong antibodies or enzyme conjugate used.
 - Substrate was not added or the wrong substrate was used.
 - The wrong buffers were used.
 - Enzyme conjugate was not added.
 - Incorrect concentrations were used.
 - The antibodies or enzyme conjugate were inactive due to improper handling or exceeding the storage time.
 - Color development.
- 22. Name the three methods used for Herbicide Bioassay tests.
- a. Seed Soak
- b. Seedling Spray Test
- c. Substrate Imbibition
- 23. Non-Trait Monocot seedling will commonly express what symptoms?
 - Inhibition of root growth
 - Inhibition of secondary root growth
 - Shortened shoot and root growth
 - Browning of mesocotyl tissue
 - Lack of chlorophyll development
 - Clear coleoptiles with stunted plumule leaf growth
- 24. In a greenhouse spray test the plant affected by the herbicide show what symptoms?
 - Reduced seedling development (roots and shoot/hypocotyl)
 - Discoloration
 - Tissue or plant death

- 25. If your negative control shows no sensitivity to herbicide at the end of a test what would be the reason? Please name 3.
 - Incorrect herbicide may have been used.
 - Wrong formulation/amount of herbicide used
 - Herbicide stocks may be expired
 - Herbicide may be contaminated
 - May have needed an adjuvant
 - Herbicide may have been improperly mixed or prepared
 - Wrong check sample may have been used
 - Growth conditions for test may not have been appropriate
- 26. Name the four nitrogenous bases for deoxyribonucleotide triphosphates.
- a. Adenine
- b. Guanine
- c. Cytosine
- d. Thymine

Multiple Choice

- 27. False positive or negative for lateral flow strips can be caused by?
- a. wrong sample buffer
- b. Strips submerged too deep in the samples extract
- c. A and B
- d. None of the above.
- 28. Edge Effect on ELISA plates is cause by?
- a. Stacking of plates.
- b. Strong light
- c. Cold plates or reagents
- d. All of the above.
- 29. Match the technology on the left with what it detects on the right side.



Calculations

30. Use Seedcalc8 to calculate the % contamination and upper bound true % purity in a sample using the information below.

of seed pools - 30 # of seeds per pool- 1000 # of deviant pools 15 Desired confidence level 95%

31. Use Seedcalc8 to calculate the % contamination and lower bound true % purity in a sample using the information below.

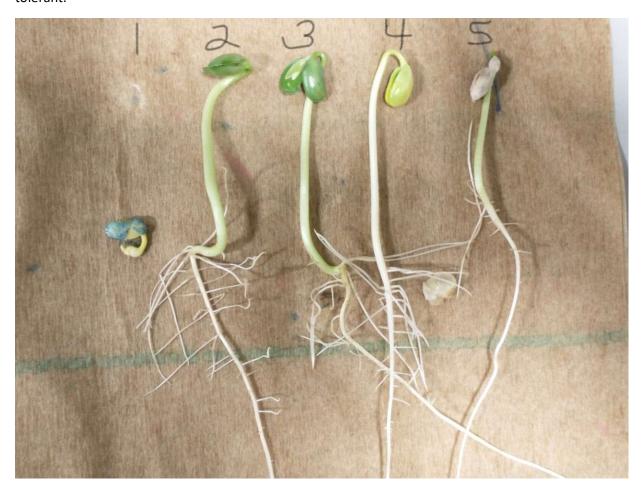
of seed pools- 20 # of seeds per pool- 500 # of deviant pools- 8 Desired confidence level 95%

0.10 % contamination 99.81 lower bound true % purity

32. Calculate the ppm for the Herbicide Concentration of Roundup Ultra 41%.

410,000 ppm

33. In this picture of the 2,4,D Soybeans rate each plant as normal or abnormal and tolerant versus non tolerant.



1 normal <mark>_X_</mark> abnormal	tolerant	X nontolerant
2. X normal abnormal	X_ tolerant	nontolerant
3. X normal abnormal	X_ tolerant	nontolerant
4. X normal abnormal	X_ tolerant	nontolerant
5. normal X abnormal	X tolerant	nontolerant

34. In this picture of Liberty Corn rate each plant as normal or abnormal and tolerant versus non tolerant.



1. X normal _	abnormal	X tolerant	nontolerant
2. X normal _	abnormal	X tolerant	nontolerant
3. X normal _	abnormal	tolerant	X nontolerant
4. X normal _	abnormal	X tolerant	nontolerant
5. X normal _	abnormal	_X_ tole	rant nontolerant

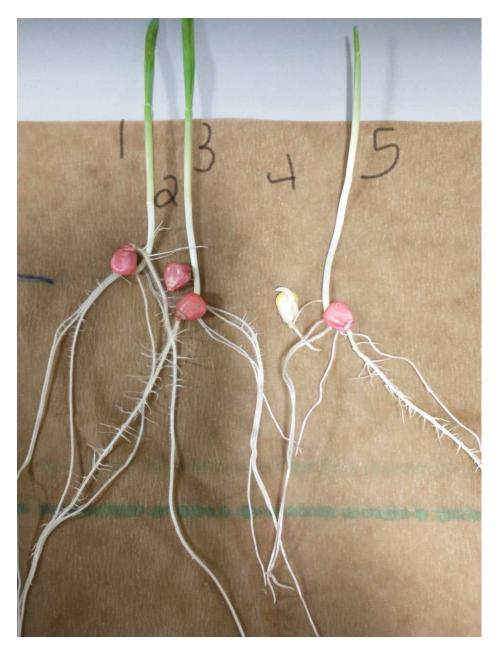
35. In this picture of Liberty Soybeans rate each plant as normal or abnormal and tolerant versus non tolerant.

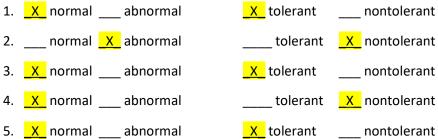


- 1. ___ normal X abnormal
- 2. X normal ___ abnormal
- 3. X normal X abnormal
- 4. X normal ___ abnormal
- 5. X normal ___ abnormal

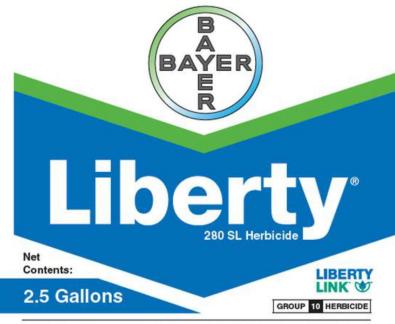
- X tolerant ___ nontolerant
- X tolerant ____ nontolerant
- X tolerant ____ nontolerant
- ____ tolerant X_ nontolerant
- X tolerant ___ nontolerant

36. In this picture of Round Up Corn rate each plant as normal or abnormal and tolerant versus non tolerant.





281,000 ppm



LIBERTY 280 SL HERBICIDE is a non-selective herbicide that provides control of a broad spectrum of broadleaf and grassy weeds.

LIBERTY 280 SL HERBICIDE is registered for use:

- as a burndown treatment prior to planting or prior to emergence of canola, corn, cotton, soybean, sugar beet, LL canola, LL corn, and LL soybean.
- post emergence weed control herbicide to be applied on all LibertyLink (LL) crops including LL canola, LL soybeans, LL com, and LL cotton
- · post emergence weed control herbicide to be applied on cotton with a hooded sprayer only

ACTIVE INGREDIENT:
Glufosinate-ammonium*
0THER INGREDIENTS:
**CAS Number 77182-82-2
**Equivalent to 2.34 pounds of active ingredient per U.S. gallon.

EPA Reg. No. 264-829

WARNING AVISO

Si usted no entiende la etiqueta, busque a alguien para que se la explique a usted en detalle. (If you do not understand the label, find someone to explain it to you in detail.)

For MEDICAL And TRANSPORTATION Emergencies ONLY Call 24 Hours A Day 1-800-334-7577 For PRODUCT USE Information Call 1-866-99BAYER (1-866-992-2937)

Please refer to booklet for additional precautionary statements and directions for use.

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