## 2019-2020 Written Genetic PT

1.	PCR is a DNA-based test that yields a positive or negative result for					
	the presence of GM.					
2.	Seed Calc can be used to determine the of pools needed based on					
	detection level of the	test used and the G	BMO acce <sub>l</sub>	ptance lev	rel.	
3.	Quantitative PCR yie	ds a specific	of GN	M present	in a given sample.	
4.	DNA is made up of _	h	eld togeth	ner with	bonds.	
5.	The basic principle o	f PCR is that of an ex	kponential	l	_ in the amount of target	
6.	Quantitative testing r	nethods typically ha	ave softwa	ire that ca	lculates the percent	
	based on the standards used in the assay.					
7.	True or False	Ouantitative lateral	flow strip	s are an in	nmunnassay that vields	
	True or False Quantitative lateral flow strips are an immunoassay that yields only a positive or negative result.					
8.	True or False	ELISA and Herbicide	e Bioassay	ys are not	typically used in AP/LLP	
	testing but they are an option in certain circumstances.					
9.	Ture or False	Results from each t	est should	d be consi	dered based on upper	
	limits of detection, technologies assayed, and seed lot sampling.					
			quire the	use of 4 o	r more known standards	
	to enable quantification.					
11.	True or False	A sample must be re	epresenta	tive of the	seed lot to provide an	
	accurate result reflecting the true contamination level.					
12.	True or False	-	flow strips	s can be le	ess expensive and labor	
	intensive than other DNA methods.					
13.		_	•	•	ein from different traits	
	express the same in different tissue types (seed, leaf) so its not important to understand what levels can be detected in the test.					

## 2019-2020 Written Genetic PT

2019-2020 Written Genetic PI					
14. When choosing technologies for AP tests, what should be considered?					
a. Size of seed lot					
b. Detection limit of the test and the acceptable limit of AP					
c. The event in question					
d. All of the Above					
15. For quantitative methods to provide an accurate result which of the below is					
recommended:					
a. Using two controls.					
b. Testing multiple reps.					
c. Using only assays to look for promoters.					
d. All of the above.					
16. The four bases in DNA are:					
a. Adenine, Ribose, Guanine, and Cytosine					
b. Adenine, Uracil, Guanine, and Cytosine					
c. Ribose, Uracil, Thymine, and Adenine					
d. Adenine, Thymine, Guanine, and Cytosine					
17. Adventitious presence is the presence of another seed variety					
genetic material.					
a. Intended					
b. Unintended					
c. Quantitative					
d. Qualitative					
18. A negative result in AP/LLP testing means the contamination:					
a. May be lower than the LOD					
b. Is zero					
c. Both A and B					
d. Neither A or B					

- 19. What is one of the biggest issues with AP/LLP testing?
  - a. Determining what technology to use
  - b. Interpreting results
  - c. Sampling
  - d. Extraction

## 2019-2020 Written Genetic PT

20. What is the purpose of AP or LLP event testing?
21. Name two scenarios where AP testing would be used.
22. Name two different technologies used to AP test.
23. Name two advantages of qualitative PCR.
24. What type of controls should be included in an AP test for comparisons?
25. What is a false error rate and why do labs need them?
26. Name two advantages to quantitative PCR testing.
27. Name two things that a lab should consider ensuring they are obtaining expected and accurate results when AP testing.
28. In Ct Target, what does Ct stand for?
29. What would you do if you performed a PCR on a traited sample and had inconsistent results?
30. List a main source of contamination in a seed crop.