The Biology Concepts Instrument (BCI): previously known as the Biology Concepts Inventory Champagne Queloz et al (2017). Diagnostic of students' misconceptions using the Biological Concepts Instrument (BCI): A method for conducting an educational needs assessment. PLoS One. 2017 May 11;12(5):e0176906.

Klymkowsky et al., 2010. Biological Concepts Instrument (BCI): A diagnostic tool for revealing student thinking. <a href="https://arxiv.org/abs/1012.4501">https://arxiv.org/abs/1012.4501</a>. \* underlined times have been identified as difficult!

Question 6: Natural selection produces evolutionary change by
$\square$ changing the frequency of various versions of genes.
☐ reducing the number of new mutations.
$\square$ producing genes needed for new environments.
$\square$ reducing the effects of detrimental versions of genes.
Question 7: If two parents display distinct forms of a trait and all their offspring (of which there are hundreds) display the same new form of the trait, you would be justified in
concluding that
□ both parents were heterozygous for the gene that controls the trait.
□ both parents were homozygous for the gene that controls the trait.
one parent was heterozygous, the other was homozygous for the gene that controls the trait.
$\ \square$ a recombination event has occurred in one or both parents.
Question 8: You are doing experiments to test whether a specific type of acupuncture works. This type of acupuncture holds that specific needle insertion points influence specific parts of the body. As part of your experimental design, you randomize your treatments so that some people get acupuncture needles inserted into the "correct" sites and others into "incorrect" sites. What is the point of inserting needles into incorrect places?  It serves as a negative control.
☐ It controls for whether the person can feel the needle.
☐ It controls for whether needles are necessary.
Question 9: As part of your experiments on the scientific validity of this particular type of acupuncture, it would be important to  \[ \text{test only people who believe in acupuncture.} \]
test only people without opinions, pro or con, about acupuncture.
$\square$ have the study performed by researchers who believe in this form of acupuncture.
$\square$ determine whether placing needles in different places produces different results.
Question 10: What makes DNA a good place to store information?
☐ The hydrogen bonds that hold it together are very stable and difficult to break.
☐ The bases always bind to their correct partner.
☐ The sequence of bases does not greatly influence the structure of the molecule.
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$\square$ The overall shape of the molecule reflects the information stored in it.

Question 11: What is it about nucleic acids that makes copying genetic information
straightforward?
☐ Hydrogen bonds are easily broken.
☐ The binding of bases to one another is specific.
☐ The sequence of bases encodes information.
☐ The shape of the molecule is determined by the information it contains.
Question 12: It is often the case that a structure (such as a functional eye) is lost during the course of evolution. This is because
☐ It is no longer actively used.
☐ Mutations accumulate that disrupt its function.
☐ It interferes with other traits and functions.
☐ The cost of maintaining it is not justified by the benefits it brings.
<ul> <li>Question 13: When we want to know whether a specific molecule will pass through a biological membrane, we need to consider</li> <li>The specific types of lipids present in the membrane.</li> <li>The degree to which the molecule is water soluble.</li> </ul>
☐ Whether the molecule is actively repelled by the lipid layer.
☐ Whether the molecule is harmful to the cell.
<ul> <li>Question 14: How might a mutation be creative?</li> <li>It could not be; all naturally occurring mutations are destructive.</li> <li>If the mutation inactivated a gene that was harmful.</li> <li>If the mutation altered the gene product's activity.</li> <li>If the mutation had no effect on the activity of the gene product.</li> </ul>
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Question 17: How does a molecule bind to its correct partner and avoid "incorrect"
<ul> <li>interactions?</li> <li>☐ The two molecules send signals to each other.</li> <li>☐ The molecules have sensors that check for incorrect bindings.</li> <li>☐ Correct binding results in lower energy than incorrect binding.</li> <li>☐ Correctly bound molecules fit perfectly, like puzzle pieces.</li> </ul>
Question 18: Once two molecules bind to one another, how could they come back apart again?
<ul> <li>☐ A chemical reaction must change the structure of one of the molecules.</li> <li>☐ Collisions with other molecules could knock them apart.</li> <li>☐ The complex will need to be degraded.</li> <li>☐ They would need to bind to yet another molecule.</li> </ul>
<ul> <li>Question 19: Why is double-stranded DNA not a good catalyst?</li> <li>☐ It is stable and does not bind to other molecules.</li> <li>☐ It isn't very flexible and can't fold into different shapes.</li> <li>☐ It easily binds to other molecules.</li> <li>☐ It is located in the nucleus.</li> </ul>
<ul> <li>Question 20: Lipids can form structures like micelles and bilayers because of</li> <li>☐ their inability to bond with water molecules.</li> <li>☐ their inability to interact with other molecules.</li> <li>☐ their ability to bind specifically to other lipid molecules.</li> <li>☐ the ability of parts of lipid molecules to interact strongly with water.</li> </ul>
<ul> <li>Question 21: A mutation leads to a dominant trait; what can you conclude about the mutation's effect?</li> <li>It results in an overactive gene product.</li> <li>It results in a normal gene product that accumulates to higher levels than normal.</li> <li>It results in a gene product with a new function.</li> </ul>
☐ It depends upon the nature of the gene product and the mutation.  Question 22: How similar is your genetic information to that of your parents?
<ul> <li>□ For each gene, one of your alleles is from one parent and the other is from the other parent.</li> <li>□ You have a set of genes similar to those your parents inherited from their parents.</li> <li>□ You contain the same genetic information as each of your parents, just half as much.</li> <li>□ Depending on how much crossing over happens, you could have a lot of one parent's genetic information and little of the other parent's genetic information.</li> </ul>

controls each trait. You examine A's offspring, of which there are hundreds, and find that most display either the same two traits displayed by A, or neither trait. There are, however, rare offspring that display one or the other trait, but not both.  The genes controlling the two traits are located on different chromosomes.  The genes controlling the two traits are located close together on a single chromosome.  The genes controlling the two traits are located at opposite ends of the same chromosome.
<ul> <li>Question 24: A mutation leads to a recessive trait; what can you conclude about the mutation's effect?</li> <li>It results in a non-functional gene product.</li> <li>It results in a normal gene product that accumulates to lower levels than normal.</li> <li>It results in a gene product with a new function.</li> <li>It depends upon the nature of the gene product and the mutation.</li> </ul>
<ul> <li>Question 25: Imagine an ADP molecule inside a bacterial cell. Which best describes how it would manage to "find" an ATP synthase so that it could become an ATP molecule?</li> <li>☐ It would follow the hydrogen ion flow.</li> <li>☐ The ATP synthase would grab it.</li> <li>☐ Its electronegativity would attract it to the ATP synthase.</li> <li>☐ It would actively be pumped to the right area.</li> <li>☐ Random movements would bring it to the ATP synthase.</li> </ul>
Question 26: You follow the frequency of a particular version of a gene in a population of <a href="mailto:asexual">asexual</a> organisms. Over time, you find that this version of the gene disappears from the population. Its disappearance is presumably due to  genetic drift.  its effects on reproductive success.  its mutation.  the randomness of survival.
<ul> <li>Question 27: Consider a diploid organism that is homozygous for a particular gene. How might the deletion of this gene from one of the two chromosomes produce a phenotype?</li> <li>If the gene encodes a multifunctional protein.</li> <li>If one copy of the gene did not produce enough gene product.</li> <li>If the deleted allele were dominant.</li> <li>If the gene encoded a transcription factor.</li> </ul>

Question 26. Gene A and gene B are located on the same chromosome. Consider the
following cross: AB/ab X ab/ab. Under what conditions would you expect to find 25% of
the individuals with an Ab genotype.
☐ It cannot happen because the A and B genes are linked.
☐ It will always occur, because of independent assortment.
☐ It will occur only when the genes are far away from one another.
☐ It will occur only when the genes are close enough for recombination to occur between
them.
Question 29: Sexual reproduction leads to genetic drift because
☐ there is randomness associated with finding a mate.
not all alleles are passed from parent to offspring.
it is associated with an increase in mutation rate.
☐ it produces new combinations of alleles.
Question 30: How is genetic drift like molecular diffusion?
☐ Both are the result of directed movements.
☐ Both involve passing through a barrier.
Both involve random events without regard to ultimate outcome.
☐ They are not alike. Genetic drift is random; diffusion typically has a direction.