



AP BIOLOGY

Survival Guide

2026-2027

Student: _____

When life
gives you
mold, make
Penicillin!



Mrs. Turney

This coursebook contains material specifically authorized for non-commercial, single one-time use in educational settings only. The teachers of this course have designed this collective work and are making the material in this coursebook available to our currently enrolled students under sole intent to advance the understanding of course material in our Biology classroom. Publication of this coursebook on open access sites is strictly prohibited. Distribution of this coursebook is limited to one-time use per student and constitutes a 'fair use' of any such copyrighted material as provided for in section 107 of the US Copyright Law. In accordance with Title 17 U.S.C. Section 107, the material in this coursebook is distributed without profit to those students currently enrolled in this course who are receiving the included information for research and educational purposes only. If you wish to use material from this coursebook that is beyond 'fair use', you must obtain permission directly from the copyright owners.

Task Verbs Used in Free-Response Questions

The following **task verbs** are commonly used in the free-response questions:

Calculate: Perform mathematical steps to arrive at a final answer, including algebraic expressions, properly substituted numbers, and correct labeling of units and significant figures.

Construct/Draw: Create a diagram, graph, representation, or model that illustrates or explains relationships or phenomena. Labels may or may not be required.

Describe: Provide relevant characteristics of a specified topic.

Determine: Decide or conclude after reasoning, observation, or applying mathematical routines (calculations).

Evaluate: Judge or determine the significance or importance of information, or the quality or accuracy of a claim.

Explain: Provide information about how or why a relationship, process, pattern, position, situation, or outcome occurs, using evidence and/or reasoning to support or qualify a claim. Explain "how" typically requires analyzing the relationship, process, pattern, position, situation, or outcome; whereas explain "why" typically requires analysis of motivations or reasons for the relationship, process, pattern, position, situation, or outcome.

Identify: Indicate or provide information about a specified topic, without elaboration or explanation.

Justify: Provide evidence to support, qualify, or defend a claim, and/or provide reasoning to explain how that evidence supports or qualifies the claim.

Make a claim: Make an assertion that is based on evidence or knowledge.

Predict/Make a prediction: Predict the causes or effects of a change in, or disruption to, one or more components in a relationship, pattern, process, or system.

Represent: Use appropriate graphs, symbols, words, illustrations, and/or tables of numerical values to describe biological concepts, characteristics, and/or relationships.

State (the null/alternative hypothesis): Indicate or provide a hypothesis to support or defend a claim about a scientifically testable question.

Support a claim: Provide reasoning to explain how evidence supports or qualifies a claim.

Rate and Growth		Water Potential (Ψ)
Rate $\frac{dY}{dt}$	dY = amount of change dt = change in time	$\Psi = \Psi_p + \Psi_s$ Ψ_p = pressure potential Ψ_s = solute potential The water potential will be equal to the solute potential of a solution in an open container because the pressure potential of the solution in an open container is zero.
Population Growth $\frac{dN}{dt} = B - D$	B = birth rate D = death rate	The Solute Potential of a Solution $\Psi_s = -iCRT$ i = ionization constant (1.0 for sucrose because sucrose does not ionize in water) C = molar concentration R = pressure constant ($R = 0.0831$ liter bars/mole K) T = temperature in Kelvin ($^{\circ}\text{C} + 273$)
Exponential Growth $\frac{dN}{dt} = r_{\max} N$	N = population size K = carrying capacity $r_{\max}$ = maximum per capita growth rate of population	
Logistic Growth $\frac{dN}{dt} = r_{\max} N \left(\frac{K - N}{K} \right)$		
Simpson's Diversity Index Diversity Index = $1 - \sum \left(\frac{n}{N} \right)^2$ n = total number of organisms of a particular species N = total number of organisms of all species		pH = $-\log[\text{H}^+]$

Surface Area and Volume

Surface Area of a Sphere

$$SA = 4\pi r^2$$

Surface Area of a Rectangular Solid

$$SA = 2lh + 2lw + 2wh$$

Surface Area of a Cylinder

$$SA = 2\pi rh + 2\pi r^2$$

Surface Area of a Cube

$$SA = 6s^2$$

Volume of a Sphere

$$V = \frac{4}{3}\pi r^3$$

Volume of a Rectangular Solid

$$V = lwh$$

Volume of a Cylinder

$$V = \pi r^2 h$$

Volume of a Cube

$$V = s^3$$

r = radius

l = length

h = height

w = width

s = length of one side of a cube

SA = surface area

V = volume

The AP Biology Exam

FORMAT

Total Time: *Three hours duration and consisting of two sections (90 minutes each)*
 Multiple Choice: *60 multiple choice questions (some individual & some in sets of 4-5)*
 Free-Response Questions: *2 long questions and 4 short answer questions*
 Distribution by section: *Multiple Choice= 50% of your total score*
 Free Response= 50% of your total score
 Calculators are allowed on the exam (be sure to check AP central for type)

The exam also assesses each of the eight units of the course with the following exam weightings on the multiple-choice section of the AP Exam:

Unit	Exam Weighting
1: Chemistry of Life	8–11%
2: Cell Structure and Function	10–13%
3: Cellular Energetics	12–16%
4: Cell Communication and Cell Cycle	10–15%
5: Heredity	8–11%
6: Gene Expression and Regulation	12–16%
7: Natural Selection	13–20%
8: Ecology	10–15%

<https://apstudents.collegeboard.org/courses/ap-biology/assessment>

TRENDS

Results in 2025: Earning a 3 or higher (pass rate)= 71.0% (Total # students= 287,232)
 Earning a 5= 19.0% of test takers.
 Earning a 4: 24.0% of test takers.
 Earning a 3: 28.0% of test takers.
 Earning a 1: 8.0% of test takers.

Results in 2024: Earning a 3 or higher (pass rate)= 68.0% (Total # students= 260,062)
 Earning a 5= 16.0% of test takers.
 Earning a 4: 23.0% of test takers.
 Earning a 3: 29.0% of test takers.
 Earning a 1: 10.0% of test takers.

Results in 2023: Earning a 3 or higher (pass rate)= 64% (Total # students=239,470)
 Earning a 5= 14.0% of test takers.
 Earning a 4: 23.0% of test takers.
 Earning a 3: 27.0% of test takers.
 Earning a 1: 12.0% of test takers.

While "passing" is a 3, a 4 or 5 is generally required to stand out or skip introductory college courses.

<https://www.totalregistration.net/AP-Exam-Registration-Service/Compare-Score-Distributions.php?id=18&year=2021>

About the Exam

The AP Biology Exam will test your understanding of the biological concepts covered in the course units, as well as your ability to utilize the scientific method and analyze data.

This is a hybrid digital exam. You'll complete multiple-choice questions and view free-response questions in the Bluebook testing app. You'll handwrite your free-response answers in paper exam booklets.

Calculators are permitted for this exam. See [AP Exams Calculator Policy](#) for the list of calculators approved for this course.

Reference materials, such as equations sheets and reference tables, are available for this course. Go to [Reference Information for Specific AP Exams](#) to find reference materials for AP Biology and exam day details.

UNIT CONVERSIONS:

Customary Units	Metric Units
1 inch	2.54 centimeters
1 foot	30.48 centimeters or 0.3048 meter
1 yard	0.914 meter
1 mile	1.609 kilometers
1 ounce	28.350 grams
1 pound	454 grams or 0.454 kilogram
1 fluid ounce	29.574 milliliters
1 quart	0.946 liter
1 gallon	3.785 liters

Metric Units	Customary Units
1 centimeter	0.394 inch
1 meter	3.281 feet or 1.093 yards
1 kilometer	0.621 mile
1 gram	0.035 ounce
1 kilogram	2.205 pounds
1 milliliter	0.034 fluid ounce
1 liter	1.057 quart or 0.264 gallon

Energy Units Conversion Table

Btu	kWh	Therm	Dkt
1	0.00029	0.00001	.000001
3,412	1	29.308	293.08
100,000	29.308	1	10
1,000,000	293.08	10	1
3,412,000	1,000	34.12	3.412

Power Units Conversion Table

Btu/hour	Watt	HP	kW
1	0.293	0.00039	0.00029
3.413	1	0.00134	0.001
2546.10	746	1	0.746
3413	1000	1.341	1

Temperature Conversions

C	F	C	F	C	F	C	F
$^{\circ}\text{C} = \frac{5}{9} (^{\circ}\text{F} - 32)$ $^{\circ}\text{F} = \frac{9}{5} ^{\circ}\text{C} + 32$		30.5	86.9	32.8	91.0	34.2	93.6
		31.0	87.8	32.9	91.2	34.3	93.7
		31.5	88.7	33.0	91.4	34.4	93.9
		31.6	88.9	33.0	91.4	34.5	94.1
27.0 80.6 27.5 81.5 28.0 82.4 28.5 83.3 29.0 84.2 29.5 85.1 30.0 86.0		31.7	89.1	33.1	91.6	34.6	94.3
		31.8	89.2	33.2	91.8	34.7	94.5
		31.9	89.4	33.3	91.9	34.8	94.6
		32.0	89.6	33.4	92.1	34.9	94.8
		32.1	89.8	33.5	92.3	35.0	95.0
		32.2	90.0	33.6	92.5	35.1	95.2
		32.3	90.1	33.7	92.7	35.2	95.4
		32.4	90.3	33.8	92.8	35.3	95.5
		32.5	90.5	33.9	93.0	35.4	95.7
		32.6	90.7	34.0	93.2	35.5	95.9
		32.7	90.9	34.1	93.4	35.6	96.1

STATISTICAL METHODS FOR AP BIOLOGY

The Independent T-test analyzes the difference between the means of two groups (the control and the experimental groups) and takes into account the standard deviations of those two groups. It basically asks: Is there more variance **WITHIN** a single group than there is **BETWEEN** the two groups? If so, then the variable being applied to the experimental group cannot be determined to have been the cause of the difference between the group. BUT...if the difference between the subjects of the control group and the subjects of the experimental group is **GREATER** than the difference within one group, a significant amount of that difference can be attributed to the independent variable in the experiment.

$$t = \frac{(X_1 - X_2)}{\sqrt{\frac{(s_1)^2}{n_1} + \frac{(s_2)^2}{n_2}}}$$

Where:

x_1 is the mean of sample 1

s_1 is the standard deviation of sample 1

n_1 is the sample size of sample 1

x_2 is the mean of sample 2

s_2 is the standard deviation of sample 2

n_2 is the sample size in sample 2

How to Calculate T:

1. Calculate the mean (X) of each sample
2. Find the *absolute value* of the difference between the means
3. Calculate the [standard deviation](#) for each sample
4. Square the standard deviation for each sample
5. Divide each squared standard deviation by the sample size of that group.
6. Add these two values
7. Take the square root of the number to find the "standard error of the difference. 8. Divide the difference in the means (step 2) by the standard error of the difference (step 7). The answer is your "calculated T-value."
9. Determine the degrees of freedom (df) for the test. In the t-test, the degrees of freedom is the sum of the sample sizes of both groups minus 2.
10. Determine the "Critical T-value" in a table by triangulating your DF and the "p value" of 0.05.

A p-value is the probability of concluding there is a significant difference between the group's result and the null hypothesis is true (meaning, the probability of making the **WRONG** conclusion). In Biology, we use a standard "p-value" of 0.05. This means that five times out of a hundred you would find a statistically significant difference between the means even if there was none.

Draw your conclusion:

- If your calculated t value is greater than the critical T-value from the table, you can conclude that the difference between the means for the two groups is significantly different. We reject the null hypothesis and conclude that the alternative hypothesis is correct.
- If your calculated t value is lower than the critical T-value from the table, you can conclude that the difference between the means for the two groups is NOT significantly different. We accept the null hypothesis.

Confidence Interval							
	80%	90%	95%	98%	99%	99.8%	99.9%
df	0.2	0.1	0.05	0.02	0.01	0.002	0.001
1	3.078	6.314	12.706	31.821	63.657	318.313	636.589
2	1.886	2.920	4.303	6.965	9.925	22.327	31.598
3	1.638	2.353	3.182	4.541	5.841	10.215	12.924
4	1.533	2.132	2.776	3.747	4.604	7.173	8.610
5	1.476	2.015	2.571	3.365	4.032	5.893	6.869
6	1.440	1.943	2.447	3.143	3.707	5.208	5.959
7	1.415	1.895	2.365	2.998	3.499	4.785	5.408
8	1.397	1.860	2.306	2.896	3.355	4.501	5.041
9	1.383	1.833	2.262	2.821	3.250	4.297	4.781
10	1.372	1.812	2.228	2.764	3.169	4.144	4.587

Chi-Square

A chi-square analysis looks at the observed results compared to the expected results. It is typically used when you are looking at distinct groups or outcomes that should have been selected equally if there was no other variable affecting the selection. For example, if you expect that hamsters randomly select which corner of a cage to sit in, then their locations in each of the four corners over several trials should be the same. Given 40 runs, the hamster should have gone to each of the 4 corners a total of 10 times.

$$\chi^2 = \sum \frac{(O - E)^2}{E}$$

Chi-Square Test (χ^2) $\chi^2 = \sum_{i=1}^r \sum_{j=1}^c \frac{(O_{ij} - E_{ij})^2}{E_{ij}}$

Computing D_f (degrees of freedom)

Degrees of freedom for chi square test of independence is equal to:

(rows-1)*(columns-1) = D_f
 (2-1) = 1 * (2-1) = 1 * 1 = D_f
 $D_f = 1$

We then get our chi critical value by getting Chi table and locating significance level 0.05 and D_f of 1.

Critical Values of Chi Square

df	0.05	0.01
1	3.84	6.64
2	5.99	9.21
3	7.82	11.34
4	9.49	13.28
5	11.07	15.09
6	12.59	16.81
7	14.07	18.48
8	15.51	20.09
9	16.92	21.67
10	18.31	23.21
11	19.68	24.72
12	21.03	26.22
13	22.36	27.69
14	23.68	29.14
15	25.00	30.58

df	0.05	0.01
16	26.30	32.00
17	27.59	33.41
18	28.87	34.80
19	30.14	36.19
20	31.41	37.57
21	32.67	38.93
22	33.92	40.29
23	35.17	41.64
24	36.42	42.98
25	37.65	44.31
26	38.88	45.64
27	40.11	46.96
28	41.34	48.28
29	42.56	49.59
30	43.77	50.89

Source: Adapted from <http://www.statsoft.com/Textbook/ChiSquare>

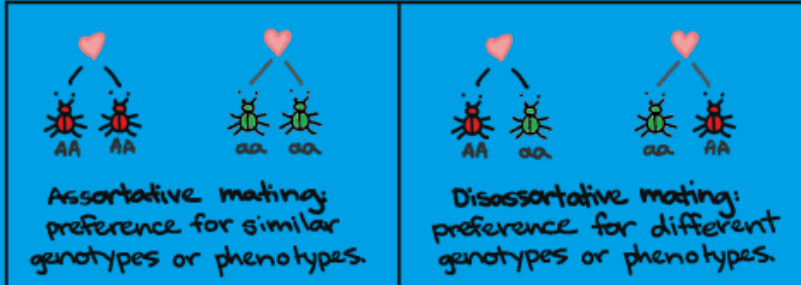
Degrees of Freedom = # Possible Observations – 1

Level of Significance (alpha) in Science is set at 0.05.

Below Critical Value = Accept. Above Critical Value = Reject

If the null hypothesis is accepted, then there probably is no relationship between the two variables and the experimental results were probably due to chance alone. If the null hypothesis is rejected, then there probably is a relationship between the two variables, and the experimental results are probably not due to chance.

The Hardy-Weinberg Theorem

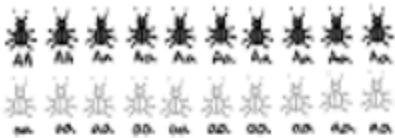


$$p^2 = (0.3)^2 = 0.09$$

$$2pq = 2(0.3)(0.7) = 0.42$$

$$q^2 = (0.7)^2 = 0.49$$

Small population of 20 beetles:



ALLELE FREQUENCIES:

Freq. of allele A = $p = 12/40 = 0.3$

Freq. of allele a = $q = 28/40 = 0.7$

GENOTYPE FREQUENCIES:

Freq. of AA = $2/20 = 0.1$

Freq. of Aa = $8/20 = 0.4$

Freq. of aa = $10/20 = 0.5$

Ah...but do you
know WHY I roll
balls of dung?!?





The Four Main Ideas in AP Biology from AP Central and the College Board:

Big Idea 1: The process of evolution drives the diversity & unity of life.

Big Idea 2: Biological systems use energy and molecular building blocks to grow, reproduce, and maintain dynamic homeostasis.

Big Idea 3: Living systems store, retrieve, transmit, and respond to information essential to life processes.

Big Idea 4: Biological systems interact, and these systems and their interactions possess complex properties.

The application of each of these “big ideas” is paramount to understanding Biology at a depth that is at the introductory college level. As such, each one of these ideas is based on a core principle on which science is founded, and thus are found in each and every one of the units we cover in AP Biology. In addition, many of the test items (not only within this class, but also on the national AP Biology Exam) address these ideas, including free-response items.

Lab Report Scoring Template

Student Name:				
Title of Experiment:				
Scientific Method	Categories	Point Value	Notes	Earned Points
Title		2		
Introduction	Background	4		
	Problem Statement	2		
	Hypothesis	4		
	Null hypothesis	2		
Materials and Methods	Materials	4		
	Independent Variable	2		
	Dependant Variable	2		
	Confounding Variable	2		
	Sample Size	2		
	Controls	2		
	Methods/ Procedure	12		
	Statistical Methods	2		
Results	Data Organization	8		
	Data Analysis	10		
Conclusions		5		
Bibliography		5		
		70 possible points	Total points earned:	

Confounding Variable: is a third variable that influences both the independent and dependent variables. How to control them: You can decrease the impact of confounding variables on your research through: **restriction, matching, statistical control and randomization**. In restriction, you restrict your sample by only including certain subjects that have the same values of potential confounding variables.

Experimental Design Tips

What variable will you manipulate? _____
(independent variable)IV- what you are changing

What variable is responding to the one you are manipulating? _____
(dependent variable)DV- what you are measuring- be specific

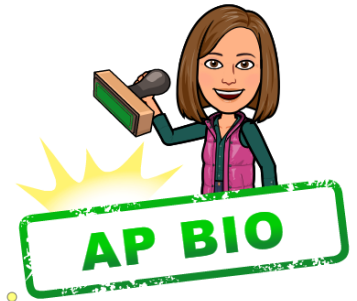
Write a title for your experiment using the following as a guide.

Title: The effect of _____ IV _____ on the _____ DV _____.

Write a hypothesis- What are you changing and how will that affect your results? Use the following as a guide.

Hypothesis: If the _____ IV _____ is _____
(describe how you changed it)

Then the _____ DV _____ will _____ because _____.
(describe the effect-what you are measuring) (give detail)



What is the **null hypothesis**?

- H₀: There is no difference between bees and butterflies in the number of seeds produced by the flowers they pollinate.



significant



Graphing for Labs

Mean: average of the data set

Median: the middle number in an ordered data set.

Mode: the most frequent number in your data set

Range= Largest - smallest

Setting up a graph don't forget:

Independent Variable goes on the X axis

Dependent Variable goes on the Y axis

What are the 7 steps to drawing a line graph in science?

1. Step 1: Identify the variables. ... Independent and dependent
2. Step 2: Determine the variable range. ...
3. Step 3: Determine the scale of the graph. ...
4. Step 4: Number and label each axis and title the graph.
5. Step 5: Determine the data points and plot on the graph. ...
6. Step 6: Draw the graph.
7. Step 7: Label the axis and title the graph (describes the experiment)

Elements of a good graph:

1. Title that describes the experiment
2. Uses the space provided.
3. Axis should be labeled with quantity being measured and the units of measurement
4. Create a consistent scale
5. Each point should be plotted accurately
6. Use a line of best fit if needed.

How to create a consistent scale:

Calculate the range (Largest # minus the smallest #)

Count the number of items (or number of boxes/lines) you have on the axis

Divide the calculated range by the number of items (Range/# of items)

Use the answer as a guide, but be consistent. Another helpful tip take your answer and multiple by 2, choose a number between the first calculated # and the multiplied by 2 answer for your grid (be sure it is divisible by 10)

A graphing example

Data to be graphed:

X Axis	Y Axis
0	0
1.0	3.4
2.0	6.9
2.9	10.5
3.9	13.3
4.9	16.9
5.9	20.8

Y axis: Range= $20.8-0=20.8$, $20.8/10= 2.1$

Multiply by 2= 4.2

-choose a # between 2.1 and 4.2 that is divisible by 10

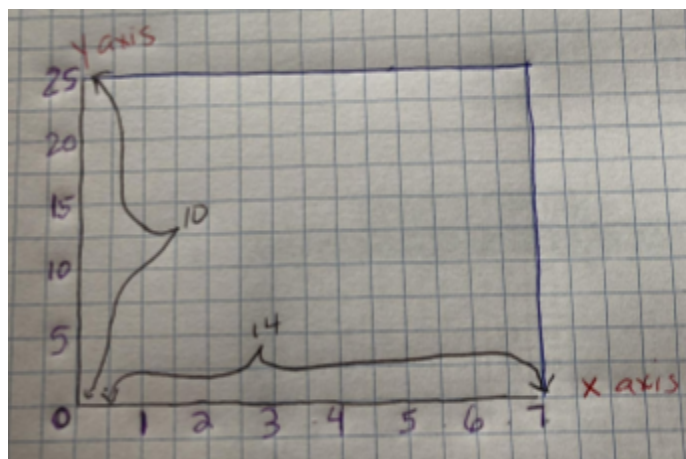
Best choice= 2.5

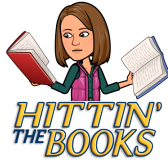
X axis: Range= $5.9-0=5.9$, $5.9/14= .42$

Multiply by 2= .84

-choose a # between .42 and .84 that is divisible by 10

Best choice= 0.5





AP Biology Reading by Units

SEMESTER ONE

Summer Assignment: Jump Start & Ecology (Quiz upon returning) Packet due 1st day of class

Unit Zero: Basics of AP Science (2 weeks)

Unit One: Biochemistry (3 weeks)

Textbook readings from chapters 1, 2, 3, 4, 5 and Sections 8.1, 8.2, 8.4, 8.5.

Unit Two: Cell Biology (3 weeks)

Textbook readings from chapters 6, 7 and 11.

Unit Three: Cellular Energetics (4 weeks)

Textbook readings from chapters 8, 9, 10.

Unit Four: Cell Communication and Cell Cycle (2 weeks)

Textbook readings from chapter 11-12

SEMESTER TWO

Unit Five: Mendelian and Human Genetics (4 weeks)

Textbook readings from chapters (sections on 12 and 13) 14, 15, 16 and 17.

Unit Six: Molecular Genetics and DNA Technology (4 weeks)

Textbook readings from chapters 18, 19, 20 and 21.

Unit Seven: Natural Selection & Evolution (4 weeks)

Textbook readings from chapters 22, 23, 24, 25 and 26.

Unit Eight: Ecology (2 Weeks Review only) -Part Self Study: Begin over Summer

Textbook readings from chapters 52, 53, 54 and 55, 56 if time.

**** Schedule is subject to change**

Statistics and Graphing Choice Charts and Guidelines

		Normally Distributed Data (Parametric)	Nonparametric	Frequency counts
Descriptive Statistics		Mean, Standard Deviation, Standard Error, 95% CI	Median, Quartiles, Interquartile Range	Percent
Comparative Statistics	Graph Type	Bar Graph	Box and Whiskers Plot	Stacked Bar Graph or Pie Chart
	Test Types	*T-Test & *ANOVA		Chi-Square (Goodness of Fit)
Association Statistics	Graph Type	Scatter Plot Line Graph—Linear Time	Scatter Plot Line Graph—Linear Time	*Mosaic Chart
	Test for Association	*Pearson Correlation		*Chi Square Test for Association
	Linear Relationship	*Linear Regression		

**These are tests or graph types that you do NOT need to be able to calculate or draw but you might need to be able to interpret them.*

Graphing

One of the best ways to **communicate the results** of a scientific investigation is graphing or creating an effective **visual representation** of the data. Investigators often can easily see **patterns** in a carefully crafted visual display that may not be readily apparent in a data table of numbers. Visual displays also can clarify how two measured variables **affect** each other.

In a scientific investigation one of the first steps in data analysis is to create graphs. Effective graphs convey **summary or descriptive statistics** as part of the display. This means that a certain amount of data analysis is required to produce appropriate graphical representations of experimental results.

When you collect data in AP Biology, it will usually fall into three categories:

1. Normal or Parametric data

- Measurement data that fit a normal curve or distribution
- Usually data are in decimal form
- Examples include plant height, body temperature, skin color etc.

2. Nonparametric data

- Data do not fit normal distribution
- May include large outliers
- May be count data that can be ordered

3. Frequency or count data

- Usually generated by counting how many items fit into a category.
- Examples: results of a genetic cross; data collected as percentages.

The typical questions asked in an AP Biology Lab can be divided into two groups:

1. Questions that compare phenomena, events or populations.
 - a. Is A different from B?
2. Questions that look for associations between variables.
 - a. How are A and B correlated?

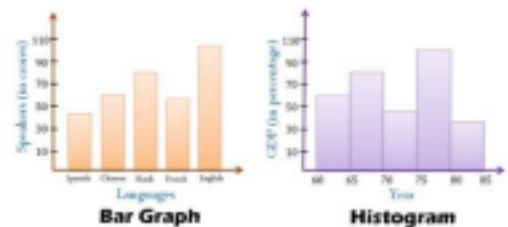
You should take time to think through **what kind of question is being asked** and **what kind of data will be required** to answer that question **prior** to starting an investigation and during the design process.

General Graphing Guidelines

1. Use the information provided here to select the appropriate graph type **PRIOR** to collecting data.
2. Every graph presented should have a **good title** that tells what the graph is about. A good title tells exactly what information the author is trying to present with the graph and usually includes both the **dependent and independent** variable.
3. Axes should be carefully **labeled** with numbers, labels and **units**. Intervals marked on line graphs should be equal in size and the scale should appropriately fit data.
4. Only plot points that have been measured. Plot points accurately!

Histograms vs. Bar Graphs

- In order to determine whether data is parametric or nonparametric you should initially make a **histogram** to look at the distribution of quantitative data.
- A histogram is different than a bar graph and is typically used for the purpose of determining distribution of data. A bar graph shows categories of data while a histogram shows a grouping of data on the x axis.



Basis for Comparison	Histogram	Bar Graph
Meaning	A graphical representation that displays data by way of bars to show the frequency of numerical data. Can be used to determine whether data is parametric or nonparametric.	A pictorial representation of data that uses bars to compare different categories of data.
Indicates	Distribution of non-discrete variables	Comparison of discrete variables
Presents	Quantitative data	Categorical data
Spaces	Bars touch	Bars do not touch
Elements	Elements are grouped together in “bins” that represent ranges of data	Elements are taken as individual entities
Can bars be reordered?	No	Yes

Guidelines for Histograms

- When an investigation involves **measurement data**, one of the first steps is to construct a histogram to represent the data's distribution to see if it approximates a **normal distribution**.
- Plots of **continuous data** and are often used to represent frequency distributions
- The data are numerical and continuous (i.e. height or weight) so the **bars touch**.
- The x axis usually records the class interval. The y axis usually records the number of individuals in each class interval (frequency).
- Requires setting up of **bins**—uniform range intervals that cover the entire range of the data.

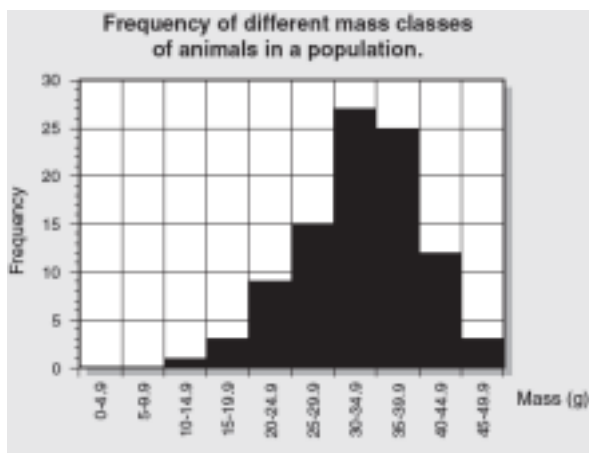
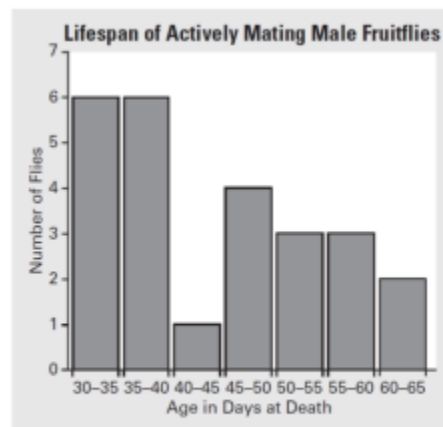


Figure 5. Histogram Showing Nonparametric Data [Source: Hanley, James A., and Stanley H. Shapiro. "Sexual Activity and the Lifespan of Male Fruitflies: A Dataset That Gets Attention." *Journal of Statistics Education* 2, no. 1 (1994).]

Guidelines for Bar Graphs

Bar graphs are appropriate for data that are **non-numerical and discrete** for at least one variable, i.e. they are grouped into separate categories.

- Data is **discontinuous**, non-numerical categories (i.e. place, color and species)—**bars do not touch** • Data values may be entered on or above bars
- Multiple sets of data can be displayed side by side for direct comparison (i.e. *males and females in the same age group*).
- Axes may be reversed so that the categories are on the x axis, i.e. the bars can be vertical or horizontal. • **Examples** of situations when using a bar graph is appropriate: *Are the spines on fish in one lake without predators shorter than the spines on fish in another lake with predators*
- **Sample standard error bars** (or sample error of the sample mean) is a notation at the top of each shaded bar that shows the sample standard error. Most of the time, bar graphs should include standard error rather than standard deviation. The SEM bars provide more information about how different the two means may be from each other.

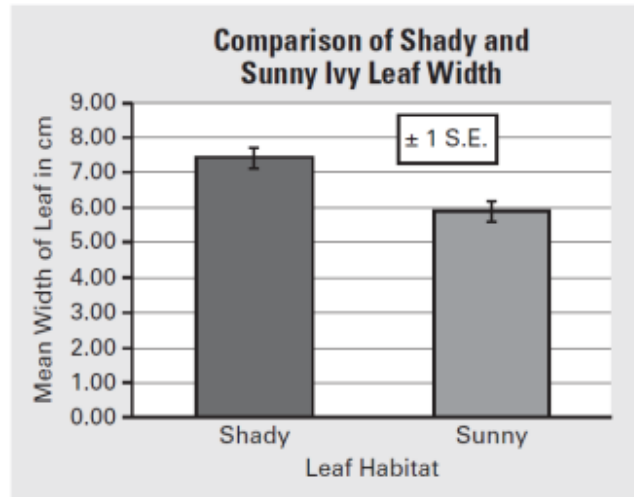
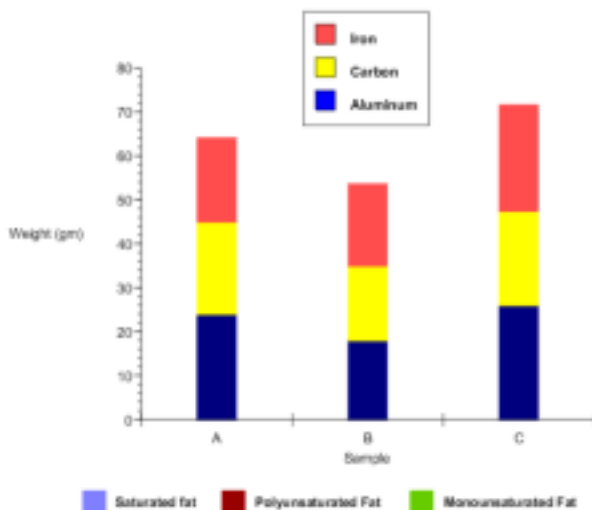


Figure 1. A Bar Graph

Stacked Bar Graph or Weighted Bar Graph

A standard **bar** chart compares individual data points with each other. In a **stacked bar** chart, parts of the data are adjacent (in the case of horizontal **bars**) or **stacked** (in the case of vertical **bars**, aka columns); each **bar** displays a total amount, broken down into sub-amounts.

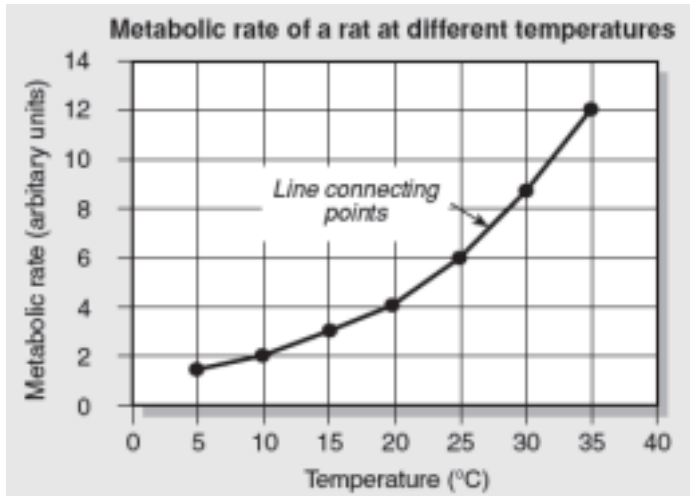
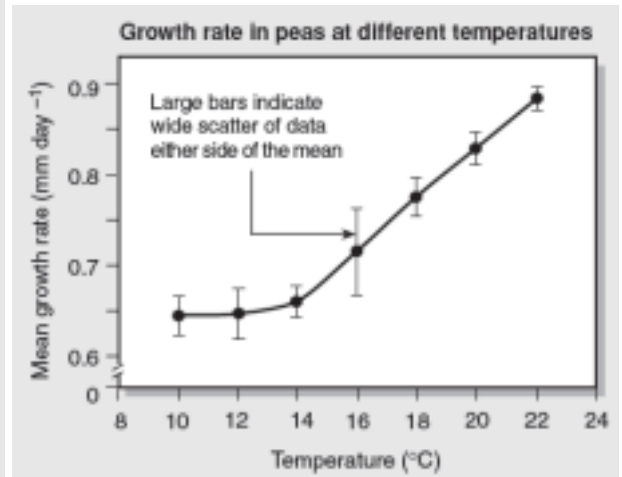


Guidelines for Line Graphs

Line graphs used when one variable (the independent variable) affects another, the dependent variable.

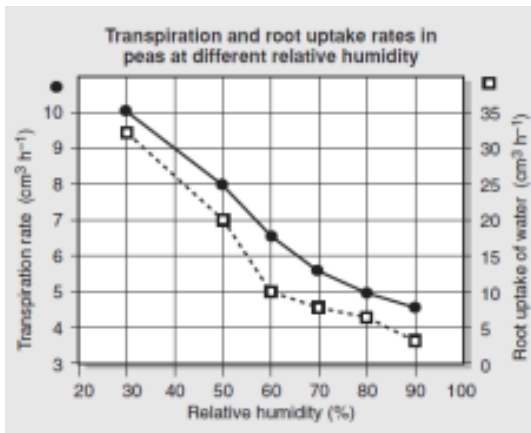
- Data must be continuous for both variables.
- The dependent variable is usually the biological response.
- The independent variable is often time or the experimental treatment.
- In cases where there is an implied trend (e.g. one variable increases with the other), a line of best fit is usually plotted through the data points to show the relationship.
- If fluctuations in the data are likely to be important (e.g. with climate and other environmental data) the data points are usually connected directly (point to point).

- Take care to **NOT** extend the line past the last data point; if the line extends past the last data point, it should be a dashed line or have an arrow point on the end to indicate that data is being extrapolated.
- Can be drawn with **measure of error**. The data are presented as points (the calculated means), with bars above and below, indicating a measure of variability or spread in the data (e.g. standard error, standard deviation or 95% confidence intervals).
- Where no error value has been calculated, the scatter can be shown by plotting the individual data points vertically above and below the mean. By convention, bars are not used to indicate the range of raw values in a data set.



Plotting Multiple Data Sets

- A single figure can be used to show two or more data sets, *i.e. more than one curve can be plotted per set of axes*. This type of presentation is useful when you want to **visually compare the trends** for two or more treatments, or the **response of one species against the response of another**.
- If the two data sets use the **same measurement units** and a similar range of values for the independent variable, one scale on the Y axis is used
- If the two data sets use **different units and/or have a very different range of values** for the independent variable, two scales for the y axis are used. The scales can be adjusted if necessary to avoid overlapping plots.



a. The two curves must be distinguished with **a key**.

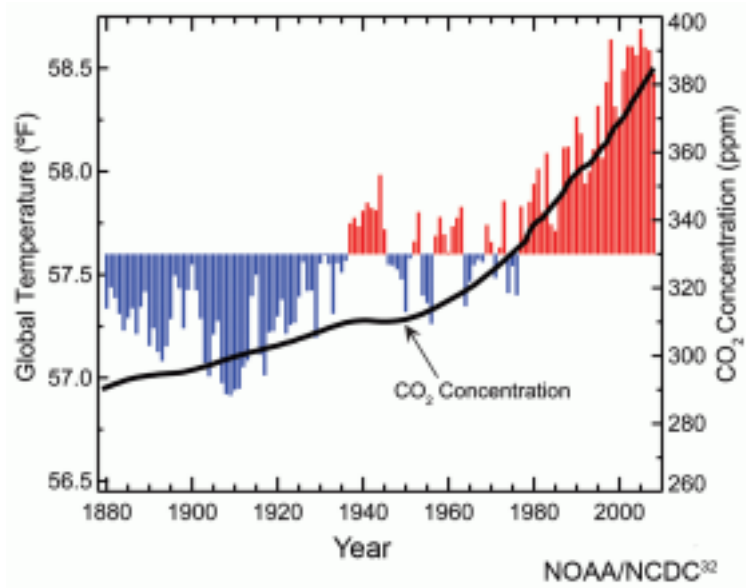
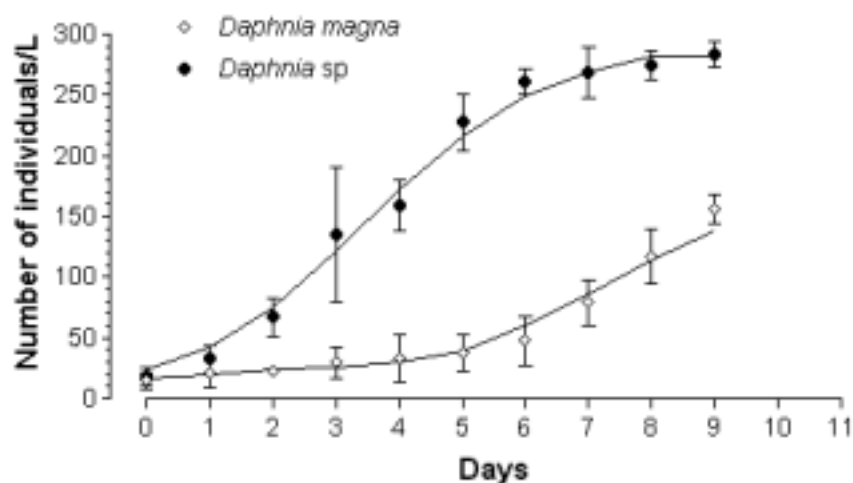
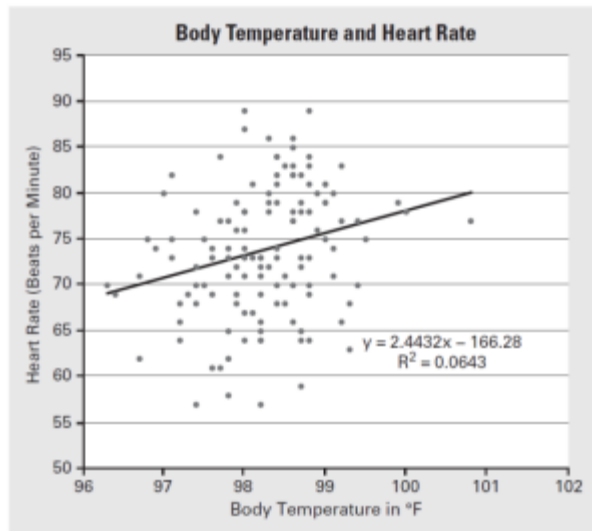
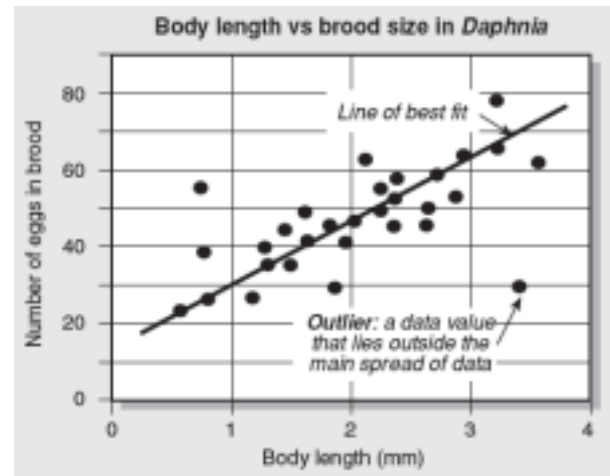


Figure 2. Mean population density (± 1 standard deviation) of two species of *Daphnia* following artificial eutrophication of a small farm pond by application of organic fertilizer. Six replicate 1 L water samples were drawn from 50 cm depth at 1100 hr each day.

Guidelines for Scatter Graphs

- Use when comparing one measured variable against another—looking for **trends or associations**. • If the relationship appears to be linear, a **linear regression line** can be calculated & plotted to help filter out the pattern that isn't always apparent. In the example shown, **the value r** can be used to help determine if there is a statistical correlation between the x & y variables to infer the possibility of causal mechanisms. • The data for this graph must be **continuous** for both variables.

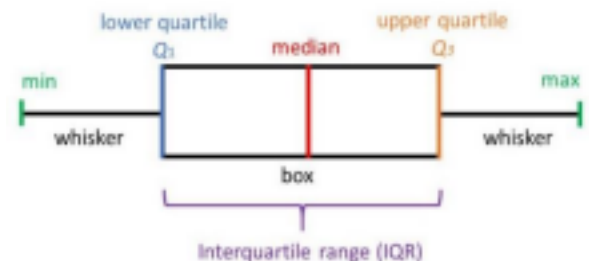


Guidelines for Box and Whisker Plot

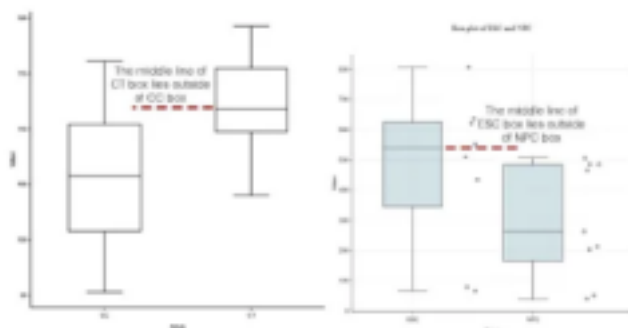
- The data should be nonparametric
- Descriptive statistics for this data should be medians and quartiles.
- Ticks at the tops and bottoms of vertical lines show highest and lowest values in the data set.
- Top of each box shows upper quartile; bottom of each box shows lower quartile
- The horizontal line represents the median.

Use to show

- skewness of data
- distribution of data
- the range of data
- outliers



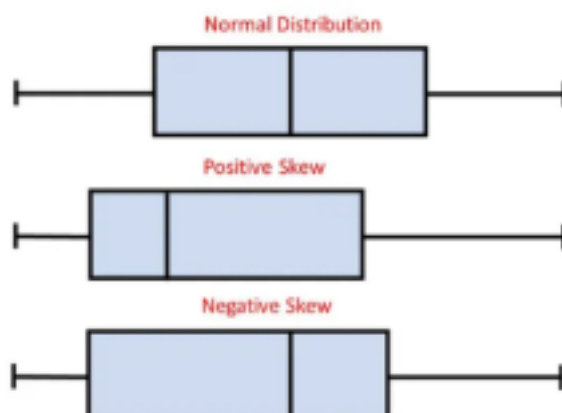
Compare the respective medians of each box plot. If the median line of a box plot lies outside of the box of a comparison box plot, then there is likely to be a difference between the two groups.



Source: <https://blog.biostatistics.com/2018/05/22/how-to-compare-box-plots/>

Box plots are useful as they show the skewness of a data set

The box plot shape will show if a statistical data set is normally distributed or skewed.

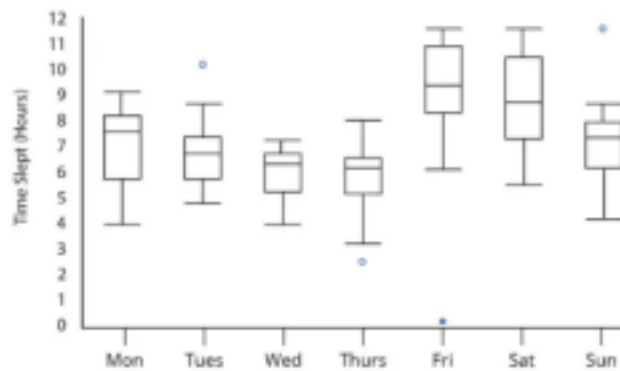
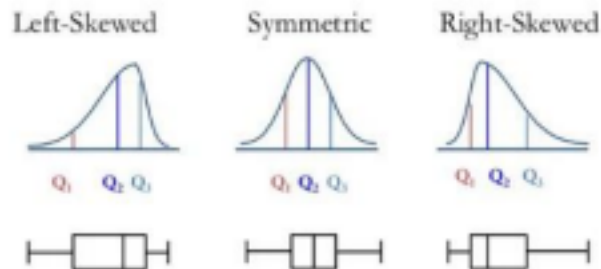


When the median is in the middle of the box, and the whiskers are about the same on both sides of the box, then the distribution is symmetric.

When the median is closer to the bottom of the box, and if the whisker is shorter on the lower end of the box, then the distribution is positively skewed (skewed right).

When the median is closer to the top of the box, and if the whisker is shorter on the upper end of the box, then the distribution is negatively skewed (skewed left).

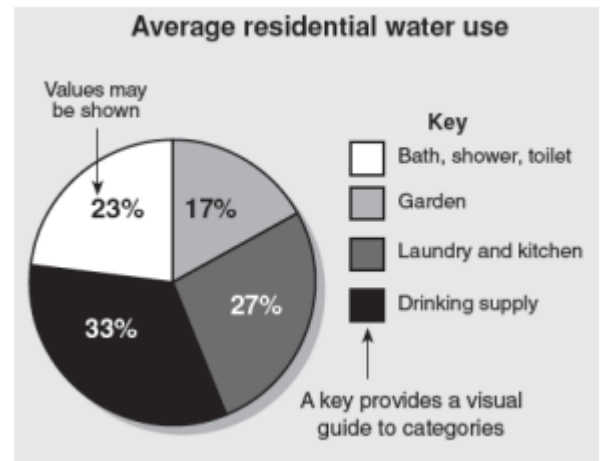
Compare the interquartile ranges (that is, the box lengths), to examine how the data is dispersed between each sample. The longer the box the more dispersed the data. The smaller the less dispersed the data.



Guidelines for Circle Graphs

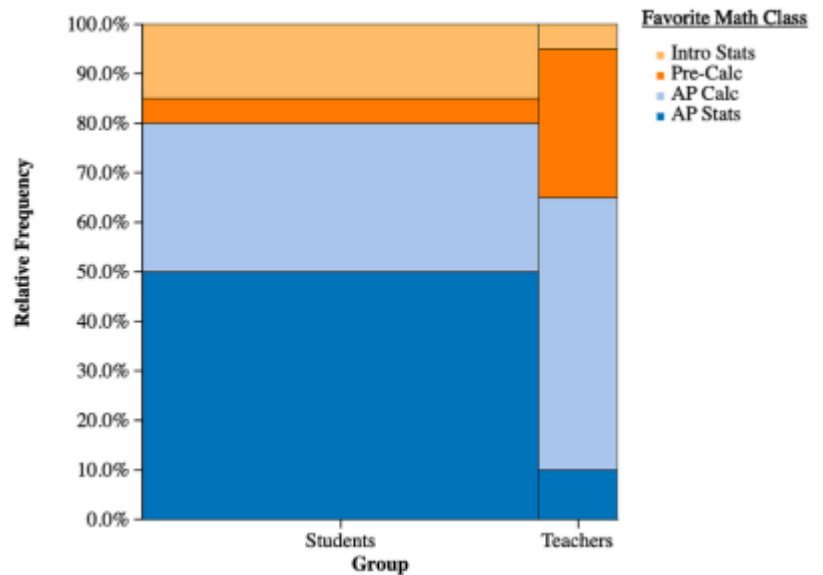
Circle graphs can be used instead of bar graphs, generally in cases where there are six or fewer categories involved. A circle graph provides strong visual impact of the **relative proportions** in each category, particularly where one of the categories is very dominant.

- Data for one variable are discontinuous (non-numerical or categories)
- Data for the dependent variable are usually in the form of **counts, proportions or percentages.**



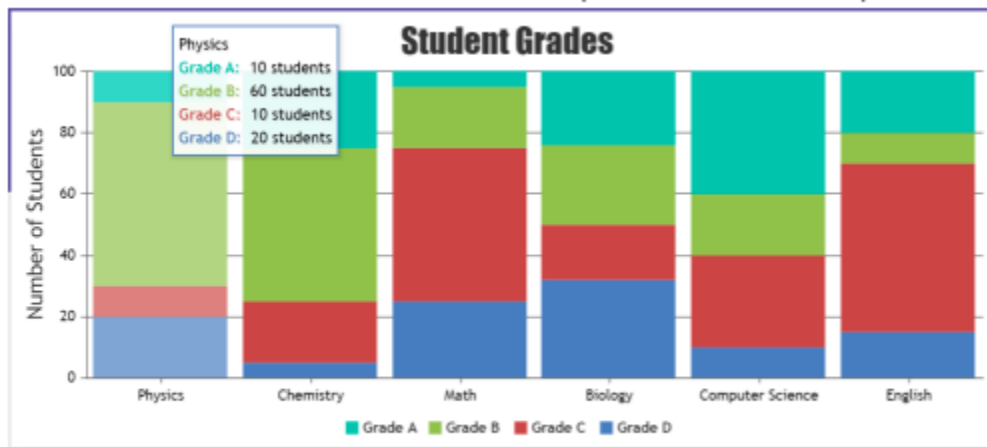
Example of Mosaic Plot

- similar to a stacked bar graph but the x axis is scaled to represent the relative size of each group.
- Students are not expected to be able to make this graph type but might need to interpret the data.



Stacked Column Charts

- Works well when the focus of the chart is to compare the totals and one part of the totals.



Key Terms for AP Biology Units

1-Basics of Biochemistry

Readings from chapters 2, 3, 4, 5 and 8

anion	isotope	acid precipitation
atom	mass number	adhesion
atomic mass	matter	aqueous solution
atomic nucleus	molecular formula	base
atomic number	molecule	buffer
cation	neutron	calorie
chemical bond	nonpolar covalent bond	Celsius scale
chemical equilibrium	orbital	cohesion
chemical reaction	periodic table of elements	colloid
compound	polar covalent bond	evaporative cooling
covalent bond	potential energy	heat
dalton	product	heat of vaporization
double bond	proton	hydration shell
electron	radioactive isotopes	hydrogen ion
electron shell	reactant	hydrophilic
electronegativity	salt	hydrophobic
element	single bond	hydroxide ion joule (j)
energy	structural formula	kilocalorie (kcal)
energy level	trace element	kinetic energy
hydrogen bond	valence	molarity
ion	valence shell	mole (mol)
ionic bond	Van der Waals interactions	pH
ionic compound	acid	polar molecule
		solute

solution	amino group	hydrocarbon
solvent	carbonyl group	hydroxyl group isomer
specific heat	carboxyl group	organic chemistry
surface tension	enantiomer	phosphate group
temperature	functional group	structural isomer
ATP	geometric isomer	sulfhydryl group
alpha helix	condensation reaction	fatty acid
amino acid	dehydration reaction	gene
antiparallel	denaturation	glycogen
beta pleated sheet	DNA	glycosidic linkage
carbohydrate	deoxyribose	hydrolysis
catalyst	disaccharide	hydrophobic interaction
cellulose	disulfide bridge	lipid
chaperonin	double helix	macromolecule
chitin	enzyme	monomer
cholesterol	fat	

2-Cell Biology

Readings from chapters 6, 7 and 11

monosaccharide	purine	unsaturated fatty acid
nucleic acid	pyrimidine	x-ray crystallography
nucleotide	quaternary structure RNA	activation energy
peptide bond	ribose	active site
phospholipid	saturated fatty acid	allosteric regulation
polymer	secondary structure	anabolic pathway
polynucleotide	starch	bioenergetics
polypeptide	steroid	catabolic pathway
polysaccharide	tertiary structure	catalyst
primary structure	triacylglycerol	chemical energy
protein		coenzyme

cofactor	first law of	noncompetitive behavior
competitive inhibitor	thermodynamics	phosphorylated
cooperativity	free energy	potential energy
endergonic reaction	free energy of activation	second law of
energy coupling	heat	thermodynamics
entropy	induced fit	substrate
enzyme-substrate complex	kinetic energy	thermal energy
exergonic reaction	metabolic pathway	thermodynamics
feedback inhibition	metabolism	
actin	microscope	mitochondrial matrix
basal body	endomembrane	mitochondrion
cell fractionation	system	myosin
cell wall	endoplasmic	nuclear envelope
central vacuole	reticulum	nuclear lamina
centriole		nucleoid
centrosome	eukaryotic cell	nucleolus
chloroplast	extracellular matrix	nucleus
chromatin	(ECM) fibronectin	organelle
chromosome	flagellum	peroxisome
cilium	food vacuole	plasma membrane
collagen	gap junction	plasmodesma
contractile vacuole	glycoprotein	plastid
crista	golgi apparatus	primary cell wall
cytoplasm	granum	prokaryotic cell
cytoplasmic	integrin	proteoglycan
streaming	intermediate filament	pseudopodium
cytoskeleton	light microscope (LM)	ribosome
cytosol	lysosome	rough ER
desmosome	microfilament	scanning electron
dynein	microtubule	microscope (SEM)
electron	middle lamella	secondary cell wall

smooth ER	aquaporin	glycolipid
stroma	concentration gradient	glycoprotein
thylakoid	cotransport	hypertonic
tight junction	diffusion	hypotonic
tonoplast	electrochemical gradient	integral protein
transmission electron microscope (TEM)	endocytosis	ion channel
transport vesicle	exocytosis	isotonic
ultracentrifuge	facilitated diffusion	ligand
vesicle	flaccid	membrane potential
active transport	fluid mosaic model	osmoregulation
amphipathic molecule	gated protein	osmosis
peripheral protein	turgid	passive transport
phagocytosis	adenylyl cyclase	local regulator
pinocytosis	cyclic amp	protein kinase
plasmolysis	diacylglycerol (DAG)	protein phosphatase
proton pump	G protein	reception
receptor-mediated	G-protein-linked	receptor tyrosine
endocytosis	receptor hormone	kinase response
selective permeability	inositol	scaffolding protein
sodium-potassium	trisphosphate (IP ₃)	second messenger
pump	ligand	signal transduction
tonicity	ligand-gated ion	pathway
transport protein	channel	transduction

3- Energy Transformation: Photosynthesis

Readings from chapters 10,35,36 & 37

absorption spectrum	C ₃ plant	carbon fixation
action spectrum	C ₄ plant	carotenoid
autotroph	Calvin cycle	chlorophyll
bundle-sheath cell	CAM plant	chlorophyll <i>a</i>

chlorophyll <i>b</i>	pep carboxylase	visible light
crassulacean acid metabolism (CAM)	photon	wavelength
cyclic electron flow	photophosphorylation	abc model
electromagnetic spectrum	photorespiration	adventitious
glyceraldehyde-3-phosphate (G3P)	photosynthesis	annual
heterotroph	photosystem	apical dominance
light reactions	photosystem I	apical meristem
light-harvesting complex	photosystem II	axillary bud
mesophyll	primary electron acceptor	bark
mesophyll cell	reaction center	biennial
NADP ⁺	RuBisCo	blade
noncyclic electron flow	spectrophotometer stoma	bundle sheath
	stroma	companion cell cork
	thylakoid	cambium
cortex	genesis	root system
cuticle	morphol	secondary growth
endodermis	ogy	secondary plant body
epidermis	perennial	shoot system
fiber	petiole	sieve plate
fibrous root	phase change	sieve-tube member
system	phloem	spongy mesophyll
guard cells	pith	active transport
herbaceous	plasticity	aquaporin
lateral	polarity	bulk flow
meristem	primary	Casparian strip
lateral	growth	chemiosmosis
root	primary	circadian rhythm
leaf	plant	endodermis
meristem		flaccid
mesophyll	body root	guttation
morpho	root hair	

membrane potential	transpiration	humus
mycorrhizae	transport protein	hydroponic culture loam
osmosis	turgid	macronutrient
osmotic potential	turgor pressure	micronutrient
passive transport	vacuolar membrane	mineral nutrient
pressure potential	water potential	mycorrhizae
proton pump	wilting	nitrogen fixation
root pressure	xerophytes	nitrogen-fixing bacteria
solute potential	cation exchange	nodule
sugar sink	crop rotation	Phytoremediation
translocation	mycorrhizae	sustainable
	epiphyte	agriculture topsoil
	essential element horizon	

4-Respiration: Anaerobic & Aerobic

Readings from chapter 9

acetyl CoA	ATP synthase	electron transport chain
aerobic	beta oxidation	facultative
alcohol	cellular respiration	anaerobe fermentation
fermentation	Chemiosmosis	glycolysis
anaerobic	citric acid cycle	lactic acid fermentation
	cytochrome	
NAD+	oxidative phosphorylation	proton-motive force redox
oxidation	oxidizing agent	reaction

5- Cell Reproduction

Readings from chapters 12 & 13

anchorage dependence	binary fission	cell division
aster	cell cycle	cell plate
benign tumor	cell cycle control system	centromere

centrosome	metastasis	gamete
checkpoint	mitosis	gametophyte
chromatin	mitotic (m) phase	gene
chromosome	mitotic spindle	genetics
cleavage	mpf	haploid cell
cleavage furrow	origin of replication	homologous chromosomes
cyclin	prometaphase	karyotype
Cyclin-dependent kinase (Cdk)	S phase	life cycle
cytokinesis	sister chromatids	locus
density-dependent inhibition	somatic cell	meiosis
G ₀ phase	telophase	meiosis I
G ₁ phase	transformation	meiosis II
G ₂ phase	alternation of generations	recombinant chromosome
gamete	asexual reproduction	sex chromosome
genome	autosome	sexual reproduction
growth factor	Chiasma	somatic cell
interphase	clone	spore
kinetochore	reducing agent	sporophyte
M phase	reduction	synapsis
malignant tumor	substrate-level phosphorylation	tetrad
meiosis	Crossing over	variation
metaphase	diploid cell	zygote
metaphase plate	fertilization	

6-Mendelian & Human Genetics

Readings from chapters 12 & 13

allele	sampling (cvs)	dominant allele
amniocentesis	codominance	epistasis
carrier	complete dominance	F ₁ generation
character	cystic fibrosis	F ₂ generation
chorionic villus	dihybrid	genotype

heterozygous	Barr body	wild type
homozygous	chromosome theory of inheritance	bacteriophage
Huntington's disease	crossing over	DNA ligase
hybridization	cytogenetic map	DNA polymerase
incomplete (intermediate) dominance	deletion	double helix
law of independent assortment	Duchenne muscular dystrophy	helicase
law of segregation	duplication	lagging strand
monohybrid	genetic map	leading strand
multifactorial	genetic recombination	nuclease
norm of reaction	genomic imprinting	nucleotide excision repair
P generation	hemophilia	Okazaki fragment
pedigree	inversion	origin of replication
phenotype	linkage map	phage
pleiotropy	linked genes	primase
5' cap	map unit	primer
a site	monosomic	replication fork
polygenic inheritance	nondisjunction	semiconservative model
Punnett square	parental type	single-strand binding protein
quantitative character	alternative RNA splicing	telomerase
recessive allele	aminoacyl-tRNA synthetase	telomere
sickle-cell anemia	polyploidy	topoisomerase
Tay-Sachs disease	recombinant	transformation
testcross	sex-linked gene	
trait	translocation	anticodon
true-breeding	trisomic	base-pair substitution
aneuploidy		
codon	exon	intron
deletion	frameshift mutation	messenger RNA
domain	insertion	(mRNA) missense
e site		

mutation	transcript	tata box
mutagen	promoter	template strand
mutation	reading frame	terminator
nonsense mutation	ribosomal RNA	transcription
one gene-one polypeptide hypothesis	(rRNA) ribosome	transcription factor
p site	ribozyme	transcription initiation complex
point mutation	RNA polymerase	transcription unit
poly-a tail	RNA processing	transfer RNA (tRNA)
polyribosome	RNA splicing	translation
(polysome) primary	signal peptide	triplet code
	signal-recognition particle (srp)	wobble (third base)
	spliceosome	

7-Molecular Genetics & DNA Technology

Readings from chapters 18-21

activator	microRNA (miRNA)	transposon
alternative RNA	multigene family	tumor-suppressor gene
splicing cell	nucleosome	bacterial artificial chromosome (bac)
differentiation	oncogene	biotechnology
chromatin	p53 gene	cDNA library
control element	proteasome	clone
differential gene	proto-oncogene	cloning vector
expression enhancer	pseudogene	complementary DNA
epigenetic	<i>ras</i> gene	(cDNA) denaturation
inheritance	repetitive DNA	DNA fingerprint
euchromatin	repressor	DNA ligase
genomic imprinting	retrotransposon	DNA microarray assay
heterochromatin	small interfering RNA	electroporation
histone	(siRNA) transcription factor	expression vector
histone acetylation		
gel electrophoresis	gene cloning	gene therapy

genetic engineering
genetically modified organism (GMO)
genomic library
genomics

human genome project
in vitro mutagenesis
linkage map
nucleic acid hybridization
physical map

polymerase chain reaction
(pcr) proteomics
recombinant DNA
restriction enzyme

8-Evolution

Readings from chapters 22-26

restriction fragment
restriction fragment length polymorphism (RFLP)
restriction site
RNA interference (RNAi)
single nucleotide polymorphism (snp)
southern blotting
sticky end
transgenic
yeast artificial chromosome (yac)
apical meristem

apoptosis
cell differentiation
cell lineage
chimera
clone
cytoplasmic determinants
egg-polarity gene
embryonic lethal
homeobox
homeotic gene
induction

maternal effect gene model
organism
morphogen
morphogenesis
organ identity gene
pluripotent
positional information
segmentation gene stem cell
totipotent
CRISPR

artificial selection
biogeography
catastrophism
endemic
evolution
evolutionary
adaptation fossil
gradualism
homologous structures
homology
natural selection

paleontology
sedimentary rock
taxonomy
uniformitarianism
vestigial organ
descent with modification
average heterozygosity
balanced polymorphism
balancing selection

bottleneck effect
cline
directional selection
disruptive selection
duplication
fitness
founder effect
frequency-dependent selection
gene flow
gene pool

genetic drift	geographic variation	heterozygote advantage
genetic polymorphism	hardy-weinberg theorem	intersexual selection
intrasexual selection	polyploidy	paralogous genes
microevolution	postzygotic barrier prezygotic	paraphyletic
modern synthesis	barrier	phylogenetic tree
mutation	punctuated equilibrium	phylogeny
neutral variation	reproductive isolation speciation	phylogram
phenotypic polymorphism	species	phylum
population	species selection	polyphyletic
population genetics	sympatric speciation analogy	shared derived character
pseudogene	binomial	shared primitive character
relative fitness	clade	specific epithet
sexual dimorphism	cladistics	systematic
sexual selection	cladogram	taxon
stabilizing selection	class	taxonomy
adaptive radiation	domain	ultrametric tree
allometric growth	family	colony
allopatric speciation	fossil record	genetic annealing
allopolyploid	genus	geologic record
autopolyploid	homoplasy	half-life
biological species concept	ingroup	magnetic reversal
ecological species concept	kingdom	pangaea
heterochrony	maximum parsimony molecular	radiometric dating
homeotic gene	clock	ribozyme
macroevolution	molecular systematic	serial endosymbiosis snowball
macroevolution	monophyletic	earth hypothesis
morphological species concept	neutral theory	stromatolite
paedomorphosis	order	three-domain system
paleontological species concept	orthologous genes	
phylogenetic species concept	outgroup	

9-Diversity of Life I: Prokaryotes, Protists, Fungi, and Plants

Readings from chapters 27, 28, 29, 30 and 31

anaerobic respiration	capsule	Commensalism
biofilm	chemoautotroph	decomposer
bioremediation	chemoheterotroph	endospore
endotoxin	symbiont	oomycete
exotoxin	symbiosis	parabasalid
extreme halophile	taxis	plasmodial slime
extreme	alternation of	mold plasmodium
thermophile	generations	protest
extremophile	amoeba	pseudopodium
facultative	apicomplexan	radiolarian
anaerobe	blade	red alga
fimbria	brown alga	secondary endosymbiosis
gram stain	cellular slime mold	sporozoite
gram-positive	ciliate	stipe
gram-negative	conjugation	thallus
host	diatom	lycophyte
methanogen	dinoflagellate	megaphyll
mutualism	diplomonad	megaspore
nitrogen fixation	euglenid	microphyll
nucleoid region	food vacuole	microspore
obligate aerobe	foraminifera	moss
obligate anaerobe	golden alga	peat
parasitism	green alga	peristome
peptidoglycan	heteromorphic	phloem
photoautotroph	holdfast	phragmoplast
photoheterotroph	isomorphic	placental transfer cell
pilus	kinetoplastid	pterophyta
plasmid	mixotroph	rhizoid

root	anther	Integument
seed	carpel	ovary
seedless vascular plants seta	conifer	ovule
sorus	cotyledon	pericarp
sporangium	cross-pollination	petal
spore	dicot vs. monocot	pollen grain
stoma	endosperm	sepal
tracheid	filament	stamen
vascular plant	flower	stigma
vascular tissue	fruit	style
xylem		ascomycete
basidiomycete	hypha	mycorrhizae
chitin	imperfect fungi	yeast
deuteromycete	lichen	zoospore
ectomycorrhizal fungi	mold	zygosporangium
endomycorrhizal fungi	mycelium	

10-Diversity of Life II: Invertebrates and Vertebrates

Readings from chapters 32, 33, 34, 40, 41, 42, 43, 44 and 47

acoelomate	deuterostome development	larva
anterior vs. posterior	diploblastic	metamorphosis
archenteron	dorsal vs. ventral	protostome development
blastopore	ectoderm, mesoderm,	pseudocoelomate
morula, blastula, gastrula	endoderm	radial vs. spiral cleavage
	ediacaran fauna	radial symmetry
cambrian explosion	gastrulation	schizocoelous
cephalization	germ layers	triploblastic
coelom	grade	trochophore larva
coelomate	indeterminate cleavage	alimentary canal
determinate cleavage		ammonite

arthropod	gastrovascular cavity	trilobite
book lung	hermaphrodite	trochophore
brachiopod	incomplete metamorphosis	tube foot
chelicerae	invertebrate	visceral mass
closed vs. open circulation	isopod	water vascular system
complete metamorphosis	mandible	amniote
copepod	mantle	amphibian
crustacean	medusa	chondrichthyan
cuticle	molting	chordate
decapods	nematocyst	cloaca
echinoderm	osculum	ectothermic
ectoproct	parthenogenesis	endothermic
exoskeleton	polyp	lateral line system
endoskeleton	radula	mammal
foot	suspension feeder	marsupial
	torsion	
monotreme	osteichthyan	ratite
mosaic evolution	oviparous	ray & lobe-finned fish
neural crest	ovoviviparous	reptile
notochord	paleoanthropology	somites
operculum	pharyngeal clefts & slits	swim bladder
opposable thumb	placenta	tetrapod

11-Plant & Animal Behavior

Readings from chapters 39, 40 and 51

phototropism	thigmonasty	abscisic acid
heliotropism	auxin	ethylene
geotropism	gibberellins	cytokinins

photomorphogenesis	circadian rhythms	vaccination
germination	photoperiodism	nephron
tunicate	vernalization	afferent & efferent arterioles
vertebrate	gravitropism	antidiuretic hormone (ADH)
viviparous	statoliths	osmoregulation
connective tissue	thigmomorphogenesis	negative & positive feedback systems
muscle tissue	virulent and avirulent	daily torpor
nerve tissue	systemic acquired resistance	ethology
epithelial tissue	negative feedback system	proximate & ultimate
estivation	positive feedback system	questions
metabolic rate	vasodilation	imprinting
thermoregulation	vasoconstriction	kinesis vs. taxis
torpor	erythrocytes	rheotaxis
vasoconstriction	leukocytes	pheromones
vasodilation	platelets (thrombocytes)	innate behaviors
omnivore, carnivore,	antigen & antibody	cognitive maps
herbivore	apoptosis	associative learning
countercurrent exchange	major histocompatibility complex (MHC)	classical conditioning
stem cell	interferon	operant conditioning
atrium & ventricle		
hemoglobin		
optimal foraging theory		
game theory		
agonistic behavior		
altruism		

12-Ecology

Readings from chapters 50, 51, 52, 53, 54 and 55

coefficient of relatedness

Hamilton's rule

Kin selection		
Vicarious learning		
biogeography	neritic zone	classical conditioning
ecology	pelagic biome	
environmentalism	organismal ecology	coefficient of relatedness
biotic vs. abiotic	patchiness	cognition
aphotic vs. photic zones	permafrost	cognitive ethology
	population	cognitive map
benthos/benthic zone	population ecology	communication
canopy	precautionary principle	ethology
biota	savanna	fixed action pattern
chaparral	temperate broadleaf (deciduous) forest	
climate		foraging
climograph	temperate grassland	game theory
community ecology	thermocline	habituation
coniferous forest	tropical rainforest	imprinting
coral reef	tropics	inclusive fitness
deep-sea hydrothermal vents	tundra	innate behavior
	turnover	kin selection
desert	wetland	kinesis
detritus	imprinting	landmark
dispersal	behavior	mate choice
ecosystem ecology	fixed action patterns	
ecotone	polygyny	copying monogamous
estuary	polyandry	operant conditioning
eutrophic vs. oligotrophic lakes	Hamilton's rule & kin selection	optimal foraging
		theory pheromone
intertidal zone	social learning & culture	proximate question
landscape ecology	agonistic behavior	reciprocal altruism
limnetic zone	altruism	
littoral zone	associative learning behavior	sensitive period
macroclimate vs. microclimate	behavioral ecology	sign stimulus
signal	social learning	spatial learning

sociobiology	reproduction	herbivory
taxis	r-selection	host
ultimate question	semelparity	individualistic hypothesis
cohort	survivorship curve	integrated hypothesis
demographic	territoriality	intermediate disturbance hypothesis
transition	zero population growth	interspecific competition
demography		
density	aposematic	intraspecific competition
density dependent	coloration	interspecific interaction
density	Batesian	
independent	mimicry	invasive species
dispersion	biomass	keystone species
ecological capacity	bottom-up model	Mullerian mimicry
ecological footprint	coevolution	mutualism
emigration	commensalism	Nonequilibrium
exponential growth	community	model parasitism
immigration	competitive exclusion	pathogen
infant mortality	cryptic coloration	predation
iteroparity	disturbance	primary succession
k-selection	dominant species	redundancy model
logistic population growth	dynamic stability hypothesis	relative abundance
	ecological niche	resource partitioning
mark-recapture	ecological succession	secondary succession
method population	ectoparasite	species diversity
population	endoparasite	species richness
dynamics	energetic hypothesis	species-area curve
population	evapotranspiration	top-down model
ecology	facilitator	trophic structure
repeated	food chain	biogeochemical cycle
	food web	

biological magnification

critical load

decomposer

Detrivore

detritus

ecosystem

eutrophication

greenhouse effect

green world hypothesis

gross primary production

limiting nutrient

net primary production

primary consumer

production efficiency

secondary consumer

secondary production

tertiary consumer

trophic efficiency

turnover time

biological augmentation

bioremediation

conservation biology

endangered species

threatened species

extinction vortex

minimum viable

population movement

corridor

population viability

analysis restoration

ecology

sustainable development



BIOLOGY IS FUN!

Mrs. Turney

MV Room A- 103