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## **Tool 3: Mathematics**



# **IB Biology - Revision Notes**

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## Applying General Mathematics in Biology

### Applying General Mathematics in Biology

- Biology often requires the use of calculations, which can include
  - **Decimals**
    - Most biological calculations use decimals, e.g. calculating the size of a bacterial cell
  - **Fractions**
    - Most scientific calculators will initially give answers as fractions
      - Make sure you know where the  $\frac{\square}{\square}$  button is so that you convert the fraction into a decimal
  - **Percentages**
    - There are many percentage calculations, including percentage change and percentage difference
  - **Ratios**
    - The most common ratio requiring understanding is that of surface area to volume ratio
  - **Proportions**
    - Proportionality can be used to understand quantity and scale and is important in biology in topics such as cell biology when creating biological drawings of cells and tissues from a microscope image or micrograph
  - **Frequencies**
    - This is most commonly used in understanding change in allele frequency
  - **Densities**
    - We often look at and examine population density in ecology or stomatal density in plant biology
  - **Approximations**
    - This is used to obtain an approximate value for example when using the magnification formula
  - **Reciprocals**
    - We frequently use reciprocals ( $1/n$ ) when dealing with concentration versus rate graphs, using  $1/T$  where  $T$  is time

### Measures of central tendency

- Measures of central tendency involve calculations of **mean, median and mode** which you should be able to apply to a range of scenarios and contexts
  - **Mean**
    - The mean is an **average of a group of numbers** calculated by totaling all values and dividing by the number of values
    - Mean is used to summarise a dataset with a single number which represents the data's typical value

- **Median**
  - This is the **middle number** which can be found by ordering all values and picking out the one in the middle
  - It helps us to understand that 50% of values have are smaller or equal to the median and 50% of values are higher or equal to the median
- **Mode**
  - This is the **most frequent value** in a dataset
  - It can be useful to understand the most common value in categorical data when the mean and median can't be used

## Measures of dispersion

- Measures of dispersion involve applying calculations of **standard deviation (SD), standard error (SE) and interquartile range (IQR)** to a range of contexts
- These ideas are also considered [here](#) with reference to the use of error bars on graph
  - **Standard Deviation**
    - The mean is a more informative statistic when it is provided alongside standard deviation
    - Standard deviation **measures the spread of data around the mean** value
      - It is very useful when comparing consistency between different data sets
    - The mean must be calculated before working out the standard deviation
  - **Standard Error**
    - Standard error of the mean **measures how far the mean of the data is likely to be from the true mean**
    - It measures the accuracy with which a sample represents a population
    - The SE is always smaller than the SD
  - **Interquartile Range**
    - This is another method of analysing dispersion of data
    - It is the **difference between the 75th and 25th percentiles of the data**
      - Quartiles are the values that divide the whole series into four equal parts

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## Scientific notation

- **Scientific notation** is also known as **standard form**
- It is a system of writing and working with very large or very small numbers
- Numbers in scientific notation are written as:

$$a \times 10^n$$

- They follow these rules:
  - **a** is a number above 1 and below 10
  - For large numbers, **n** is an integer that is greater than 0
    - i.e It shows how many times **a** is multiplied by 10
  - For small numbers, **n** is an integer that is less than 0
    - i.e It shows how many times **a** is divided by 10

- $n < 0$  for small numbers i.e how many times a is divided by 10

## Approximation and estimation

- Approximation and estimation are both methods used to **obtain values that are close to the true or accurate values**
  - While they share some similarities, they have distinct characteristics and are used in different contexts

### Approximation

- Approximation involves finding a value that is close to the actual value of a quantity
  - It may not necessarily be very precise or accurate
- It is often used when an exact calculation is challenging or time-consuming and a reasonably close value is sufficient

### Estimation

- Estimation involves making an educated guess or assessment based on available information or data
- It is used when the true value of a quantity is unknown or cannot be directly measured
  - For example biologists estimate dates of the first living cells and the last universal common ancestor or the method of estimating times by use of the "molecular clock"

## Scales of magnification

- Magnification is an important skill used widely in biology and frequently assessed in examinations
- For more information and worked examples see our revision note on [microscope skills](#)

## Rates of change

- The rate of change tells us **how something changes over time**
  - For example oxygen consumption in germinating seeds over a period of days
- To determine rates of change from tabulated data, you can use the average rate of change or gradient, if the data has been plotted as a graph
- The average rate of change between two points on a graph or in a table is:

$$\text{Rate of change} = \frac{\text{Change in the dependent variable}}{\text{Change in the independent variable}}$$

## Proportionality and correlations

- There are a number of terms that are commonly applied to trends, particularly in graphs
  - **Direct and inverse proportionality**
    - Direct proportionality applies to a trend that has a clearly **linear** relationship which means the relationship can be described as "**when one variable increases, the other increases**" or "if x doubles, then y doubles"

- Inverse proportionality means that the relationship can be described as "**when one variable increases, the other decreases**" or "if x doubles, then y halves"
- **Positive and negative correlations**
  - Positive correlations show when the gradient of the graph is positive / slopes or curves upwards and describes a relationship where as x increases, y also increases
  - Negative correlations is when the gradient of the graph is negative / slopes or curves downwards; this describes a relationship where as x increases, y decreases

## Percentage change and percentage difference

- Percentage change and percentage difference are commonly used to express **the relative change between two values**
  - They are useful for comparing experimental results, determining reaction yields and analysing other chemical data

### Percentage change

- Percentage change is used to express the relative change between an initial value and a final value
- It is calculated using the following formula:

$$\text{Percentage Change} = \frac{\text{Final value} - \text{Initial value}}{\text{Initial value}} \times 100$$

### Percentage difference

- Percentage difference is used to compare two values to determine how much they differ from each other as a percentage
- It is calculated using the following formula

$$\text{Percentage Difference} = \frac{(\text{Value 1} - \text{Value 2})}{\text{Average value 1 and value 2}} \times 100$$

## Continuous and discrete data

- **Discrete** data is quantitative
  - It consists of separate, distinct and countable values
  - For example:
    - Number of an organism in a sample
- **Continuous** data is also quantitative
  - It is based on measurements and can include decimal numbers or fractions
  - This allows for an infinite number of values
  - For example:
    - The temperature of an enzyme reaction as time progresses
    - The volume of oxygen gas produced during a photosynthesis reaction

## Statistical tests



- Statistical tests can be used to analyse a range of different data sets
- The type of test used will depend on a number of factors such as
  - The size of the sample
  - The type of data, i.e. is it discrete or continuous
  - The nature of the question being investigated

### Simpson's reciprocal index

- The Simpson's reciprocal index can be used to **measure the relative biodiversity of a given community**
- It accounts for both the number of species present (richness) and the number of individuals per species (evenness)
- A **higher index value is indicative of a greater degree of biodiversity** within the community

### The Lincoln index.

- This calculation allows an **estimate of population sizes of individual animal species**
- You can read more about the Lincoln Index [here](#)

### Chi-squared test

- A chi-square test is a statistical test that is used **to compare observed and expected results**
- Our revision notes [here](#) cover this in detail

### The t-test

- The **t-test** can be used to **compare the means of two sets of data** and determine whether they are **significantly** different or not
- The sets of data must follow a rough **normal distribution**, be **continuous** and the **standard deviations** should be approximately equal

#### Exam Tip

You will be provided with the formulae for these statistical tests in the exam, your job is to apply them to a range of contexts and data.



# Using Units, Symbols & Numerical Values in Biology

## Using Appropriate Units

- The **International System of Units** (SI) is also called the metric system
  - This is the international standard for measurement
- There are several SI base units that are used in science

SI Base Units Table

Quantity	SI base unit	Symbol
length	metre	m
mass	kilogram	kg
time	second	s
temperature	Kelvin	K
amount of substance	mole	mol
current	Ampere	A
luminous intensity	candela	cd

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- Measurements of physical quantities can require very large and very small values, for example:
  - The diameter of an atom is about  $10^{-10}$  m or 0.0000000001m
  - One mole of a substance contains  $6.02 \times 10^{23}$  or 602 000 000 000 000 000 000 000 particles
- Powers of ten are numbers that can be achieved by multiplying 10 times itself
- These come under two categories of units:
  - **Multiples** e.g.  $10^2, 10^3$
  - **Sub-multiples** e.g.  $10^{-1}, 10^{-2}$
- Each power of ten is defined by a prefix, the most common ones used in biology are listed in the table below

Table of common prefixes in biology



Prefix	Abbreviation	Power of ten
kilo-	k	$10^3$
centi-	c	$10^{-2}$
milli-	m	$10^{-3}$
micro-	$\mu$	$10^{-6}$
nano-	n	$10^{-9}$

- It essential that the correct scientific measurements are used when discussing biological experiments
- Ensure that the **correct symbols** are used in conjunction with the unit of measurement
  - E.g.  $m^3$  for cubic metres

### Units of Measurement Table

Measurement	Base unit	Symbol	Units used
Length	Metre	m	$1000\text{ m} = 1\text{ km}$ $0.01\text{ m} = 1\text{ cm}$ $0.001\text{ m} = 1\text{ mm}$ $0.000001\text{ m} = 1\mu\text{m}$
Volume	Cubic metre	$m^3$	$10^9\text{ m}^3 = 1\text{ km}^3$ $0.000001\text{ m}^3 = 1\text{ cm}^3$ $10^{-9}\text{ m}^3 = 1\text{ mm}^3$ $10^{-18}\text{ m}^3 = 1\mu\text{m}^3$
Volume	Cubic decimetre	$\text{dm}^3$	$0.001\text{ dm}^3 = 1\text{ cm}^3$
Area	Square metre	$\text{m}^2$	$10\,000\text{ m}^2 = 1\text{ ha}$ $0.0001\text{ m}^2 = 1\text{ cm}^2$
Mass	Kilogram	kg	$1000\text{ kg} = 1\text{ tonne}$ $0.001\text{ kg} = 1\text{ g}$

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			0.000001kg = 1mg $10^{-9}$ kg = 1 $\mu$ g
Time	Second	s	60 s = 1min 60 min = 1hour
Pressure	pascal	Pa	1000 Pa = 1kPa
Energy	joule	J	1000 J = 1kJ
Temperature	degree Celcius	$^{\circ}$ C	
Amount of substance	mole	mol	0.001mol = 1millimole

- $\text{cm}^3$  is the same as millilitre (ml)
- $\text{dm}^3$  is the same as litre (l)

### Exam Tip

Be careful when using the word "amount" in your answers. "Amount" has a very specific meaning in science - "mole". Instead refer to the mass, volume or concentration of a substance!

## Significant figures

- Significant figures must be used when dealing with quantitative data
- Significant figures are the digits in a number that are **reliable and absolutely necessary** to indicate the quantity of that number
- There are some important **rules** to remember for significant figures
  - All non-zero digits are significant
  - Zeros between non-zero digits are significant
    - 4107 (4.s.f.)
    - 29.009 (5.s.f.)
  - Zeros that come before all non-zero digits are not significant
    - 0.00079 (2.s.f.)
    - 0.48 (2.s.f.)
  - Zeros after non-zero digits within a number without decimals are not significant
    - 57,000 (2.s.f.)
    - 640 (2.s.f.)
  - Zeros after non-zero digits within a number with decimals are significant
    - 689.0023 (7.s.f.)



- When rounding to a certain number of significant figures:
  - Identify the significant figures within the number using the rules above
  - Count from the first significant figure to the specified number
  - Use the next number as the 'rounder decider'
  - If the decider is 5 or greater, increase the previous value by 1

### Worked example

Write 1.0478 to 3 significant figures.

**Answer:**

#### Step 1: Identify the significant figures

They are all significant figures

#### Step 2: Count to the specified number (3rd s.f.)

1.0478

#### Step 3: Round up or down

1.05

### Exam Tip

An exam question may sometimes specify how many significant figures the answer should be, make sure you keep an eye out for this!

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## Processing Uncertainties in Biology

### Processing Uncertainties in Biology

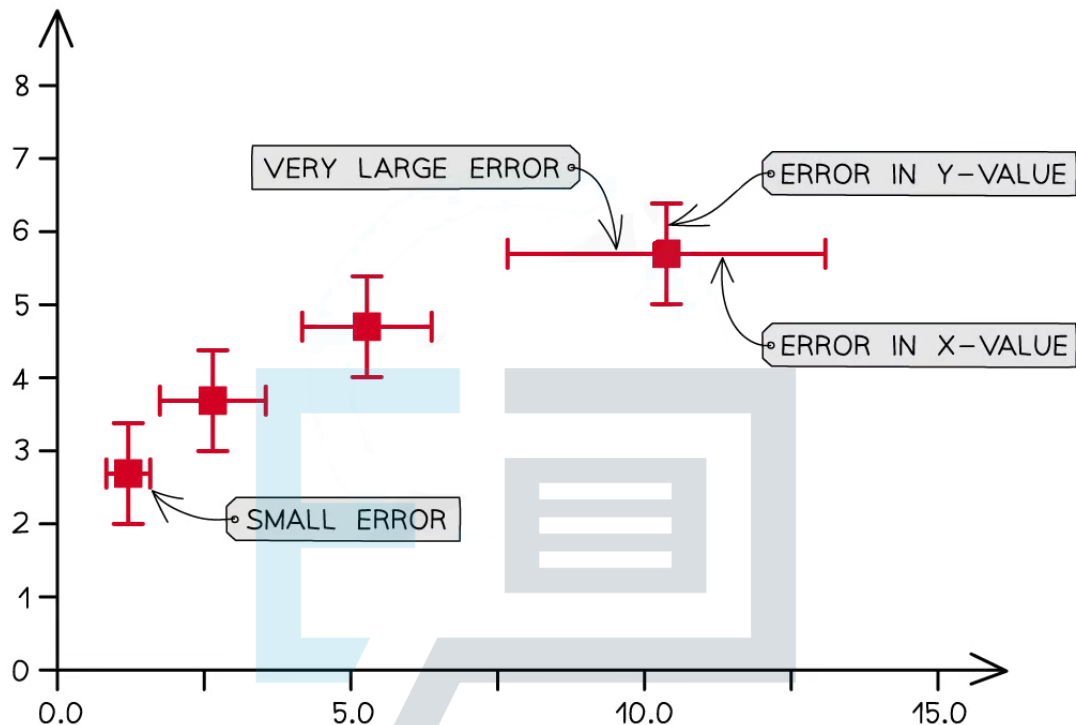
#### What is uncertainty?

- Uncertainty is **quantitative** indication of the quality of numerical results
  - It is the **difference between the actual measurement**, resulting from the **equipment** or techniques used to collect data, **and the true value**
  - It is a **range of values** around a measurement within which the true value is expected to lie
- Uncertainties are **not** the same as errors
  - Errors arise from equipment or practical techniques that cause a reading to be different from the true value
- Uncertainties in measurements are **recorded as a range ( $\pm$ )** to an appropriate level of precision, e.g.
  - If a balance that measures mass shows scale graduations of 10 g, then mass is measured to the nearest 10 g (this is known as the margin of error)
    - The true value could be 5 g higher or lower than the measured value, so the uncertainty would be  $\pm 5$  g
  - If a pipette shows scale graduations every  $0.1 \text{ cm}^3$ , then volume is measured to the nearest  $0.1 \text{ cm}^3$ 
    - The true value could be  $0.05 \text{ cm}^3$  more or less than this, so the uncertainty would be  $\pm 0.05 \text{ cm}^3$

#### Error bars

- The uncertainty in a measurement can be shown on a graph as an **error bar**
  - This bar is drawn above and below the point (or from side to side) and shows the **uncertainty** in that measurement
- Usually, error bars will be in the vertical direction, for y-values, but can also be plotted horizontally, for x-values
- **Range, degree of precision, standard error** and **standard deviation**; can be expressed on a graph using error bars
  - Range = the difference between the lowest and highest value
  - Degree of precision = how close a set of data points are to each other
  - Standard error = an estimate of the reliability of the mean
  - Standard deviation = the spread of data around the mean
- Note that it is important that you know **what is represented** by error bars on a graph, e.g. whether they represent standard deviation or standard error; in an exam this information would be provided in the question
  - Error bars that represent standard deviation can be used to assess whether or not two data sets are **significantly different** to each other

- Overlapping error bars indicate that two sets of data are not significantly different
- Error bars are used in the specification when [measuring osmotic concentration](#)



*Error bars on a graph can be used to show uncertainty*

### Level of precision

- Measurements and processed uncertainties must be expressed to an appropriate level of precision
  - E.g. number of **decimal places**
- This may depend on the sensitivity of the apparatus used to collect data; the level of precision used to express the data should not exceed the level of precision at which the data is initially measured
- Values in a raw data set should all be expressed to the same level of precision

### The coefficient of determination, $R^2$

- The **coefficient of determination** is a **measure of fit** that can be applied to lines and curves on graphs
- The coefficient of determination is written as  $R^2$
- It is used to evaluate the fit of a trend line / curve with its data set:
  - $R^2 = 0$



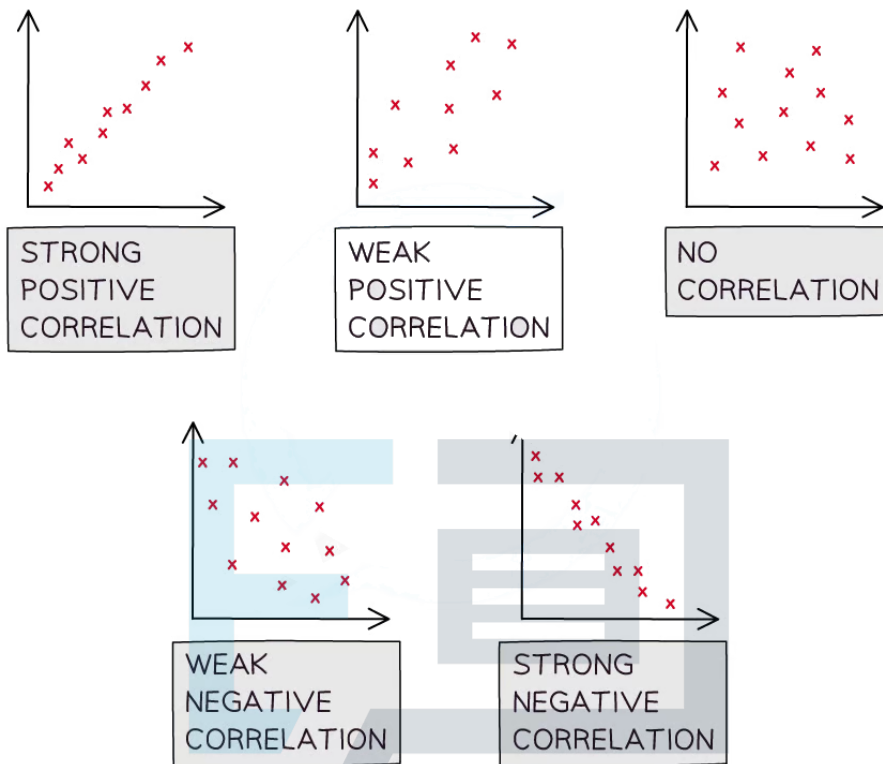
- The dependent variable cannot be predicted from the independent variable.
- $R^2$  is usually greater than or equal to zero
- **$R^2$  between 0 and 1**
  - The dependent variable can be predicted from the independent variable, although the degree of success depends on the value of  $R^2$
  - The closer to 1, the better the fit of the trend line / curve
- **$R^2 = 1$** 
  - The dependent variable can be predicted from the independent variable
  - The trend line / curve is a perfect fit
  - **Note:** This does not guarantee that the trend line / curve is a good model for the relationship between the dependent and independent variables
- Coefficient of determination is used in the specification when [comparing the speed of nerve impulse transmission](#)

## Correlation

- Correlation is an **association**, or relationship, between variables
  - Note that there is a clear distinction between correlation and causation: correlation **does not necessarily indicate a causal relationship**
  - Causation occurs when one variable has an influence or is influenced by another
- Correlation can be **positive** or **negative**
  - Positive correlation: as variable A increases, variable B increases
  - Negative correlation: as variable A increases, variable B decreases
- The **correlation coefficient (r)** can be calculated to determine **whether a linear relationship exists** between variables and **how strong** that relationship is
  - Perfect correlation occurs when all of the data points lie on a straight line; this will give a correlation coefficient of 1 or -1
    - 1 = a perfect positive correlation
    - -1 = a perfect negative correlation
  - A less-than perfect correlation will give a correlation coefficient between 1 and 0, or between 0 and -1
    - The closer to 1, or -1, the coefficient is, the stronger the correlation
  - If there is no correlation between variables the correlation coefficient will be 0
- Correlation coefficients are used in the specification when [evaluating data on coronary heart disease](#)

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*A strong correlation will have a correlation coefficient close to 1, a weak correlation will have a correlation coefficient close to 0, while a lack of any correlation will give a correlation coefficient of 0*

## Statistical tests

- Statistical tests are used to assess whether or not a data set **supports a particular hypothesis**.

e.g.

- A **null hypothesis** will state that there is **no** significant difference, or association, between two variables
- An **alternative hypothesis** will state that there **is** a significant difference, or association, between two variables
- Statistical analysis allows researchers to **accept** or **reject** the null hypothesis
- If a statistical test shows that there is no significant difference, or association, between variables, then it is said that any visible difference is **due to chance** alone
- Different statistical tests** are used for **different types of data set**, e.g.
  - A t-test determines whether the means of two data sets differ significantly
  - A correlation test determines the presence and strength of a correlation
  - A chi-squared test determines whether the difference between observed and expected values is significant
- You should be able to **select** and **apply** the correct statistical test
- The chi-squared test is used in the specification as follows:
  - To test for difference between observed and expected [outcomes of a genetic cross](#)
  - To test for [association between species](#)

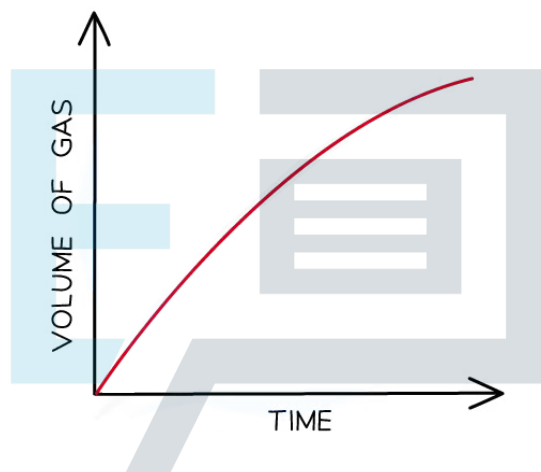
## Graphing in Biology

### Graphing in Biology

#### Sketch graphs

- **Sketch graphs** are a way to represent qualitative trends where the variables shown are often **proportional** or **inversely proportional**

#### A simple sketch graph



A sketch graph of the relationship between time and volume of gas given off, these two variables show a proportional relationship trend

#### General guidance on drawing graphs

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- The types of graphs that students are expected to be able to draw include:
  - Bar charts
  - Histograms
  - Scatter graphs
  - Line / curve graphs
  - [Logarithmic graphs](#)
  - Pie charts
  - [Box-and-whisker plots](#)

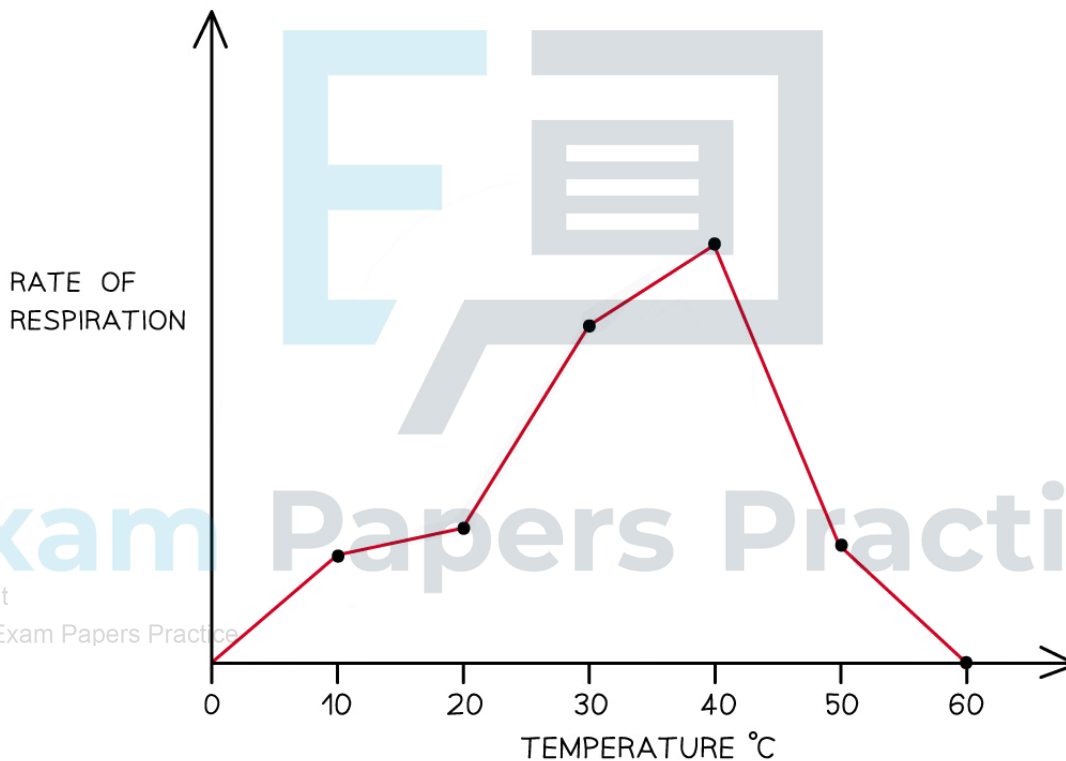
#### Tips for plotting data

- Whatever type of graph you use, remember the following:
  - The data should be plotted with the **independent** variable on the **x-axis** and the **dependent** variable on the **y-axis**
  - Plot data points **accurately**



- Use **appropriate** linear **scales** on axes
- Choose scales that enable **all** data points to be plotted **within** the **graph area**
- **Label axes**, with **units** included
- Make graphs that **fill the space** the exam paper gives you
- Draw a **line of best fit**. This may be **straight** or **curved** depending on the **trend** shown by the data. If the line of best fit is a curve make sure it is drawn **smoothly**. A line of best-fit should have a **balance of data points** above and below the line
- In some cases, the line or curve of best fit should be drawn through the **origin** (but only if the data and trend allow it)

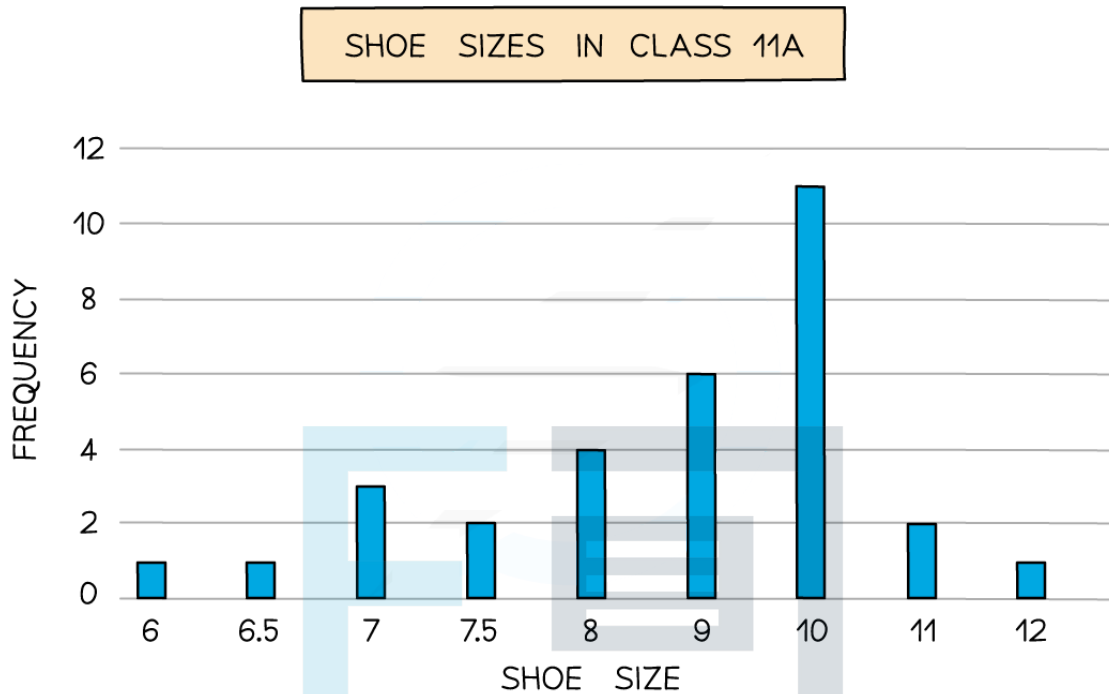
### Continuous data represented in a line graph



### Discontinuous data represented in a bar chart

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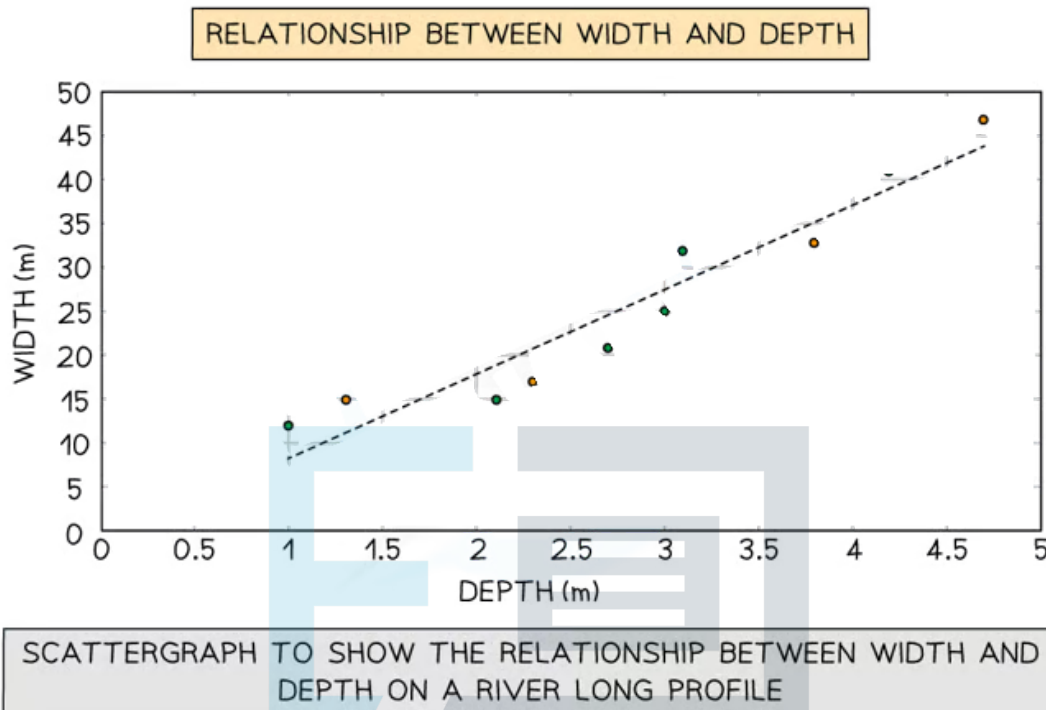
*The line graph has been used to display continuous data over time while the bar chart has been used to display grouped data*

- **Remember:** The independent variable is the one you control or manipulate and the dependent variable is the one that changes as a result of your manipulation
- Always draw data points in pencil as it makes it easier to make corrections and adjustments

### Best fit lines

- Students often confuse the term **lines of best fit** with straight lines
- **Lines of best fit** can be straight lines or curves and:
  - They show the trend of the data
    - It does not have to go through all the points, but shows the general trend
  - They must go through the majority of the points
  - Where the data is scattered the points should be evenly distributed on either side of the **best fit line**

### Graph to show use of a best fit line



### Other features of graphs

#### Using a tangent to find the initial rate of a reaction

- For linear graphs (i.e. graphs with a straight-line), the gradient is the same throughout

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- This makes it easy to calculate the rate of change (rate of change = change ÷ time)
- However, many **enzyme rate experiments** produce non-linear graphs (i.e. graphs with a **curved line**), meaning they have an **ever-changing gradient**
  - They are shaped this way because the **reaction rate** is **changing over time**
- In these cases, a **tangent** can be used to find the **reaction rate** at any **one point** on the graph:
  - A tangent is a **straight line** that is drawn so it just **touches** the curve at a **single point**
  - The **slope** of this tangent **matches** the slope of the **curve** at just that point
  - You then simply find the **gradient** of the straight line (tangent) you have drawn
- The **initial** rate of reaction is the rate of reaction at the **start** of the reaction (i.e. where **time = 0**)