Effect of sugar solution concentration on plant tissue mass:

Method:

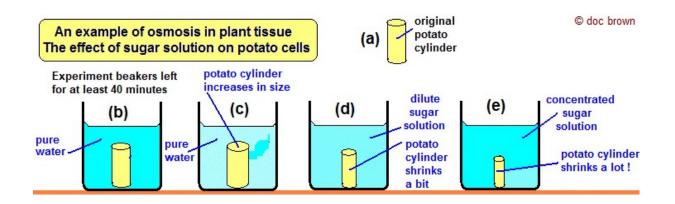
 Measure the mass of 3 identical potato cylinders
Put each in a beaker, one with pure water, one with a slightly concentrated sugar solution & one with a very concentrated sugar solution

- 3. Leave to soak for 20 hours
- 4. Take cylinders out, dry them & measure their mass

Increase in mass = water is drawn in via osmosis Decrease in mass = water has been drawn out via osmosis

Independent variable: concentration of sugar solution Dependent: affect of sugar concentration on mass Control: volume of solution, time, sugar type ect

Errors: potato isn't dry so extra mass/ water evaporates from beaker = sugar concentration changes Calculating mean percentage change reduces errors



Food tests:

Food sample prep: Break up food in pestle & mortar Transfer 2 beaker with water Stir with glass rod 2 dissolve food Filter solution through funnel

Reducing sugars via Benedicts:

1. Prep food sample

2. Add to test tube

3. Heat water bath to 75 degrees cel

4. Add benedict's solution with pipette

5. Leave test tube in bath for 5 mins

Presence of sugars= brick-red solution

Starch via Iodine: 1. Transfer food sample to test tube 2. Add iodine & shake gently Presence of starch = orange -> blue-black

Proteins via Biuret: Transfer food sample to test tube Gently shake after adding biuret solution Presence of protein= blue -> purple

Lipids via Sudan III: 1. Prep sample (don't filter it) & add to test tube, 2. Add Sudan solution & shake Presence of lipids= mixture separates into 2 layers (top is bright red) Plant growth responses:

Method:

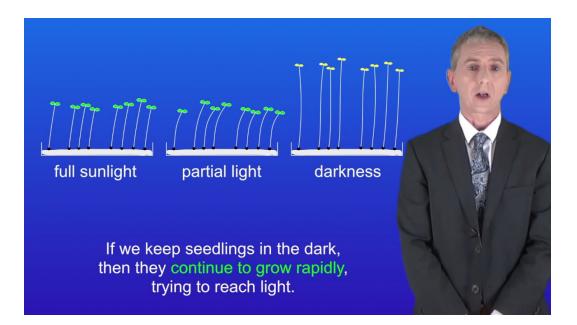
1. Put 10 crest seeds in 3 Petri dishes, lined with moist filter paper

- 2. Allow seedlings to germinate, measure their height
- 4. Place 1 dish in direct sunlight, one in partial & one in none
- 3. Leave for a week, measuring their height daily

=Seedlings in full & partial light will grow to the same height (chloroplasts are efficient at absorbing light) & the seedlings in total darkness will be the tallest with yellow leaves (energy source has ran out) = they are searching for light!

Seedlings grow towards the light = phototropism (auxin is on the side of the seedling with the least light = rapid growth)

Control variables: seed type & distance between bulb & dish (light intensity)



Quadrats:

- 1. Place 1m quadrat on a field randomly
- 2. Count organisms within the quadrat
- 3. Repeat
- 4. Work out the

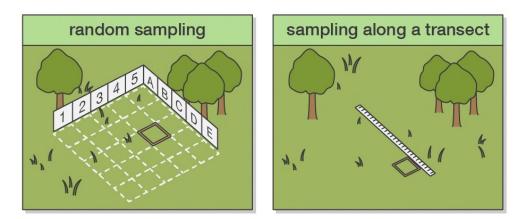
MEAN = total number of organisms / number of quadrats

Distribution can depend on light intensity Quadrats & transects = quantitative methods

Transects:

- 1. Mark out a line (with a tape measure) of an area
- 2. Count organisms touching the line
- = measures distribution from the hedge -> middle of a field

of organisms



Microscopy:

Preparing slide:

- 1. Add water to clean slide
- 2. Separate an onion's layers & peel the epidermal tissue off
- 3. Place the tissue on the water slide
- 4. Add iodine to stain the tissue
- 5. Place cover slip over the specimen

Light microscope:

- 1. Clip slide onto stage
- 2. Select lowest objective lens
- 3. Move the stage upwards with the coarse objective knob
- 4. Focus the object with the fine adjustment knob

If higher magnification needed, use higher objective lens

Draw cells with a pencil & label features

Magnification = image size / real size

There's 1000000 nano meters in 1 mm

