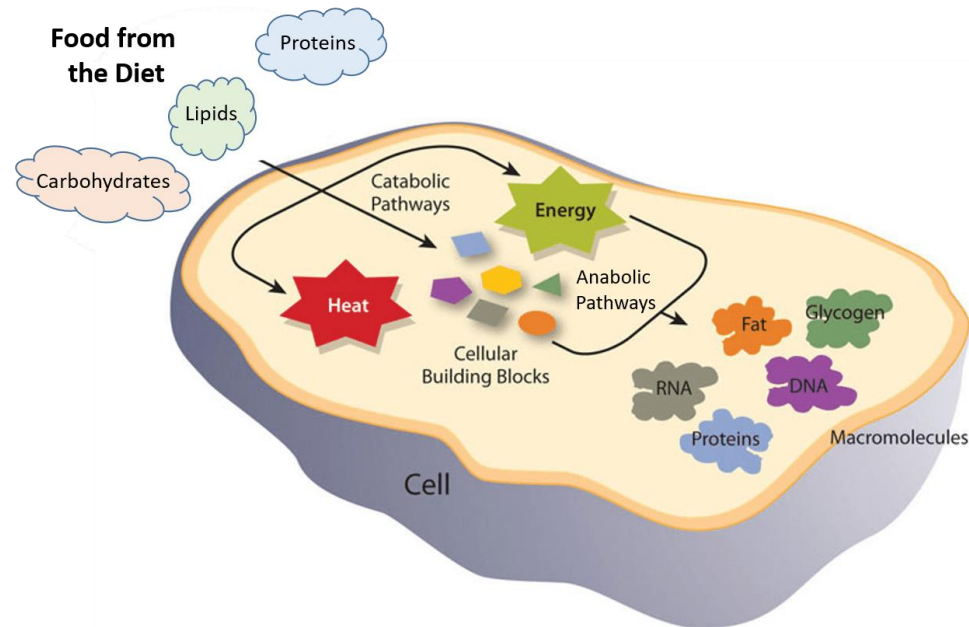


Lesson 5

Cellular Chemistry, Reaction Thermodynamics and Enzymes

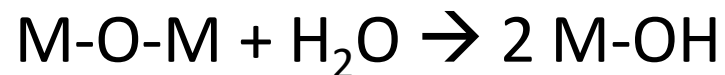


Metabolism

- The cell is an ideal continuous chemical reactor
- The ensemble of cellular chemical reactions = **metabolism**
- **Anabolic reactions** = build up molecules (*e.g.*, condensation)

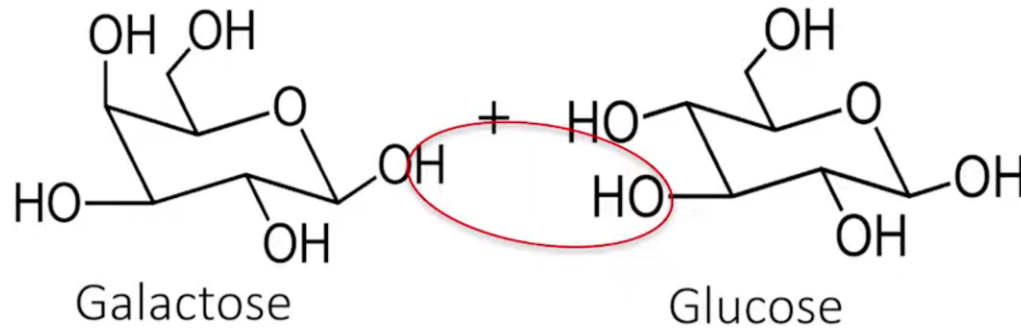


- **Catabolic reactions** = break down molecules (*e.g.*, hydrolysis)



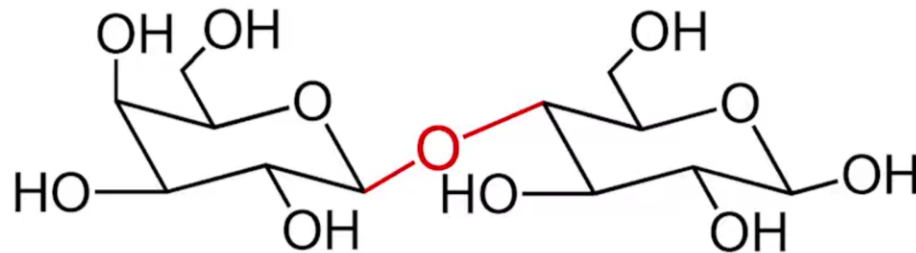
Factory, Maurice Utrillo, 1923

Condensation & hydrolysis



Condensation:

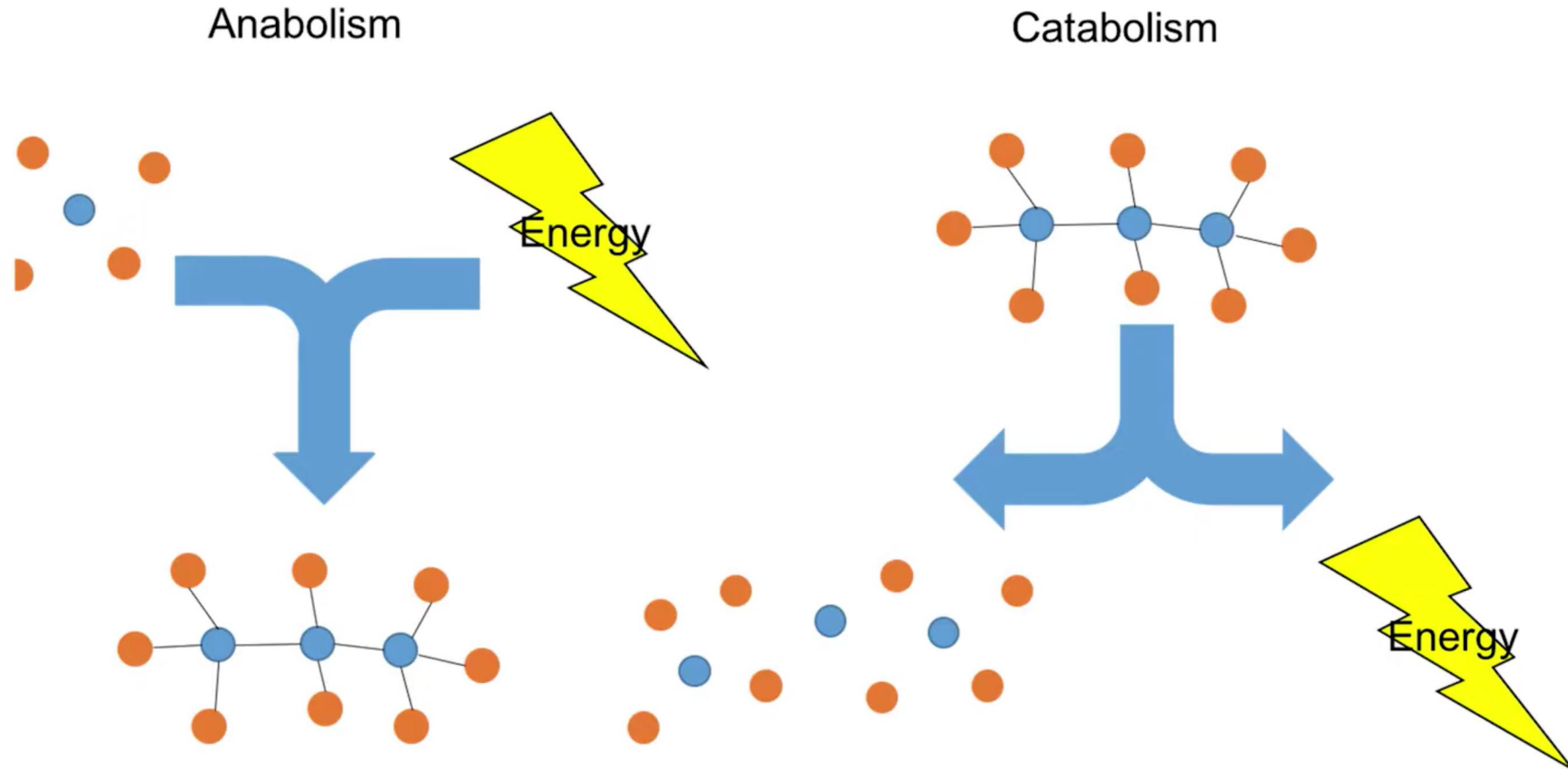
Covalent bonds form
with loss of water
Build molecules
Anabolic



Hydrolysis:

Covalent bonds break
with addition of water
Break down molecules
Catabolic

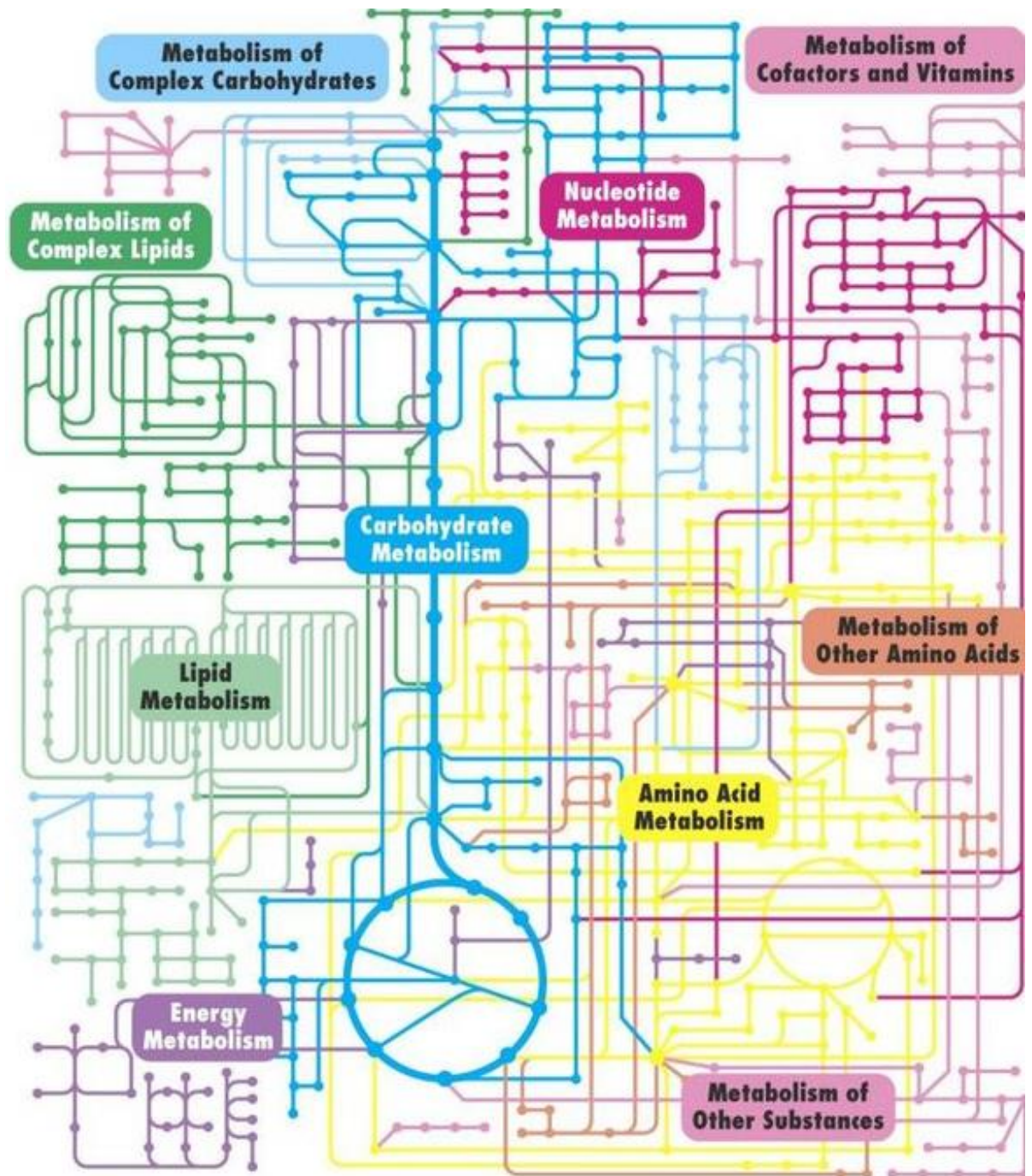
Metabolism & Energy



The metabolic map

Dots = molecules

Lines = chemical reactions
making up the metabolism of
the cell



Free energy difference

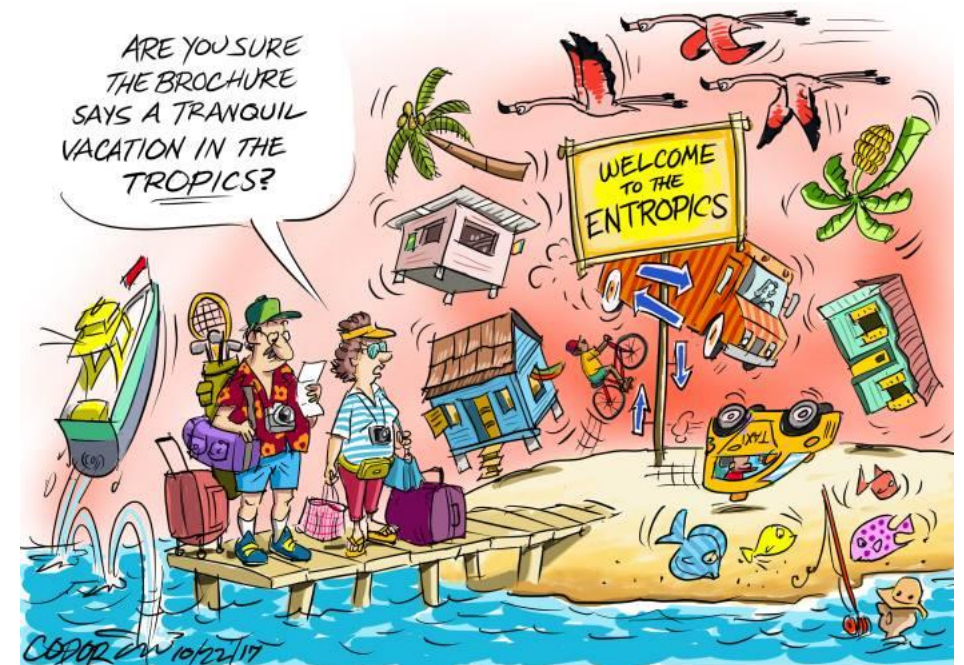
- Reactions are governed by **free energy** (*usable energy*) **G**

Reagents $\rightleftharpoons$ Products

- What really matters is the **free energy difference** $\Delta G = \sum G_p - \sum G_R$
- ΔG stems from a fundamental law of thermodynamics

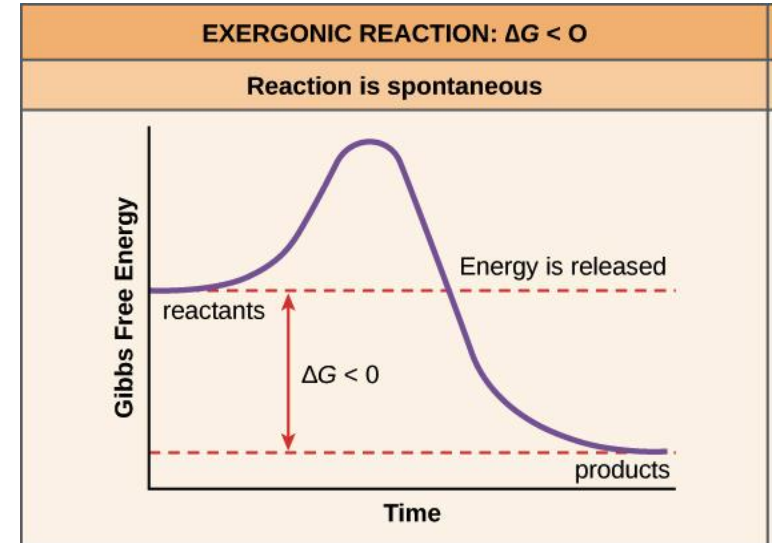
$$\Delta G = \Delta H - T\Delta S$$

- H** = **Enthalpy** = *total energy*
- T** = **Temperature**
- S** = **Entropy** = *useless energy*



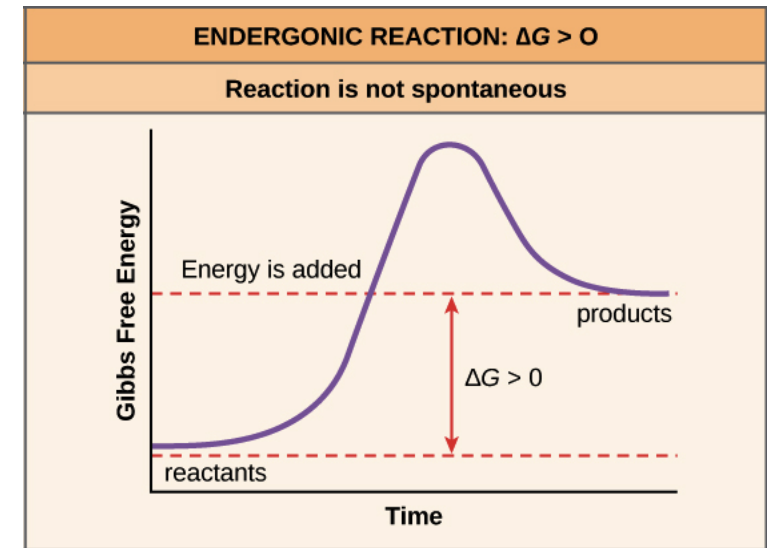
Free energy difference

- $\Delta G < 0$ ($G_P < G_R$) $\rightarrow$ Energy released, reaction proceeds (thermodynamically spontaneous, exergonic)



Free energy difference

- $\Delta G < 0$ ($G_P < G_R$) $\rightarrow$ Energy released, reaction proceeds (thermodynamically spontaneous, exergonic)
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Free energy difference

- $\Delta G < 0$ ($G_P < G_R$) $\rightarrow$ Energy released, reaction proceeds (thermodynamically spontaneous, exergonic)
- $\Delta G > 0$ ($G_P > G_R$) $\rightarrow$ Reactions requires energy to proceed (thermodynamically non-spontaneous, endergonic)
- **$\Delta G = 0$** ($G_P = G_R$) $\rightarrow$ Chemical equilibrium ($R \rightarrow P = P \rightarrow R$)



$$K_{eq} = [B]/[A]$$

$$\Delta G = -RT \ln [B]/[A]$$

┆

$$\Delta G = -RT \ln K_{eq}$$

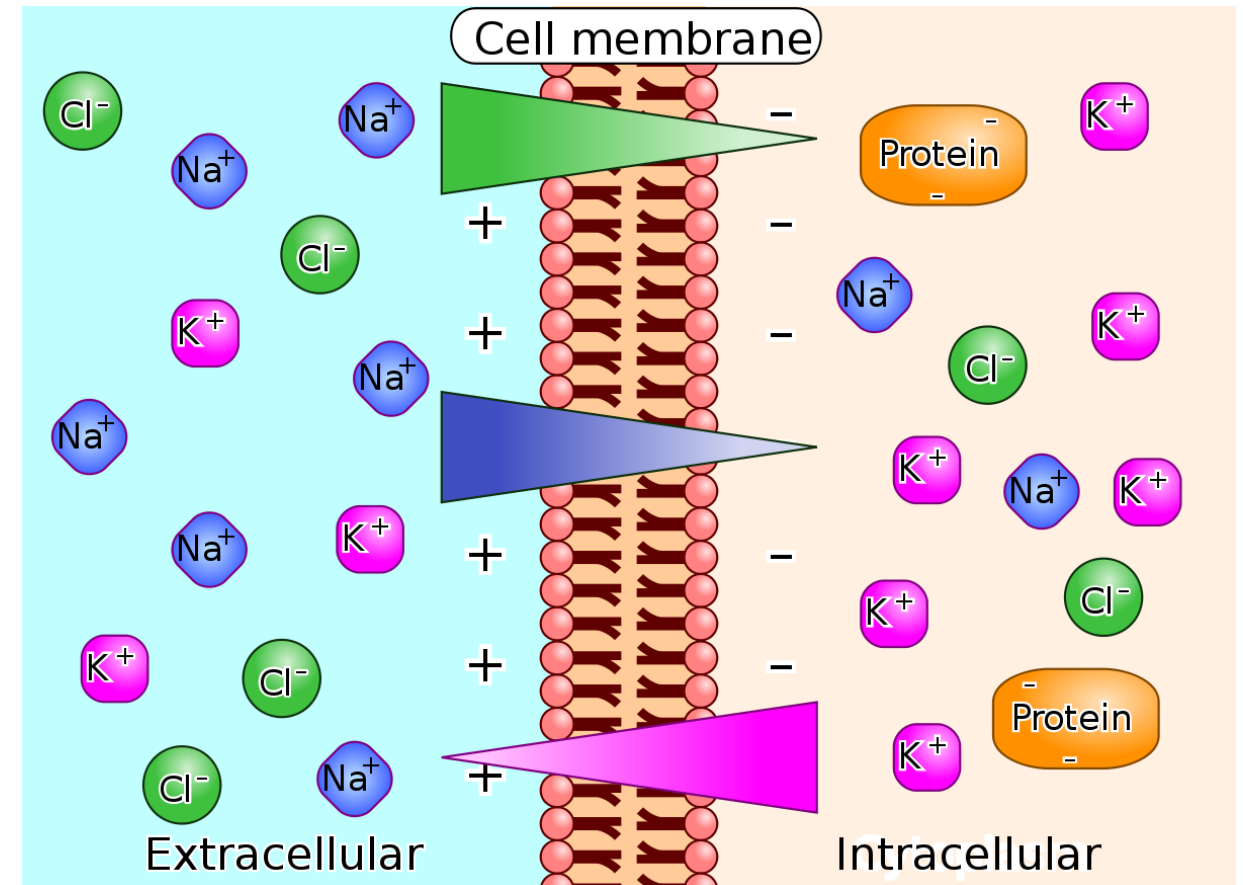
$$R = 8.314 \text{ J/(mol K)} = 1.987 \text{ cal/(mol K)}$$

Going to work in the cell factory

- Cells transfer energy from food molecules to run their processes
 - **Synthesis**
 - Complex molecules (DNA, proteins)
 - Organized structures (organelles)
 - **Organization**
 - Generation of specialized compartments to store and organize materials for specific tasks

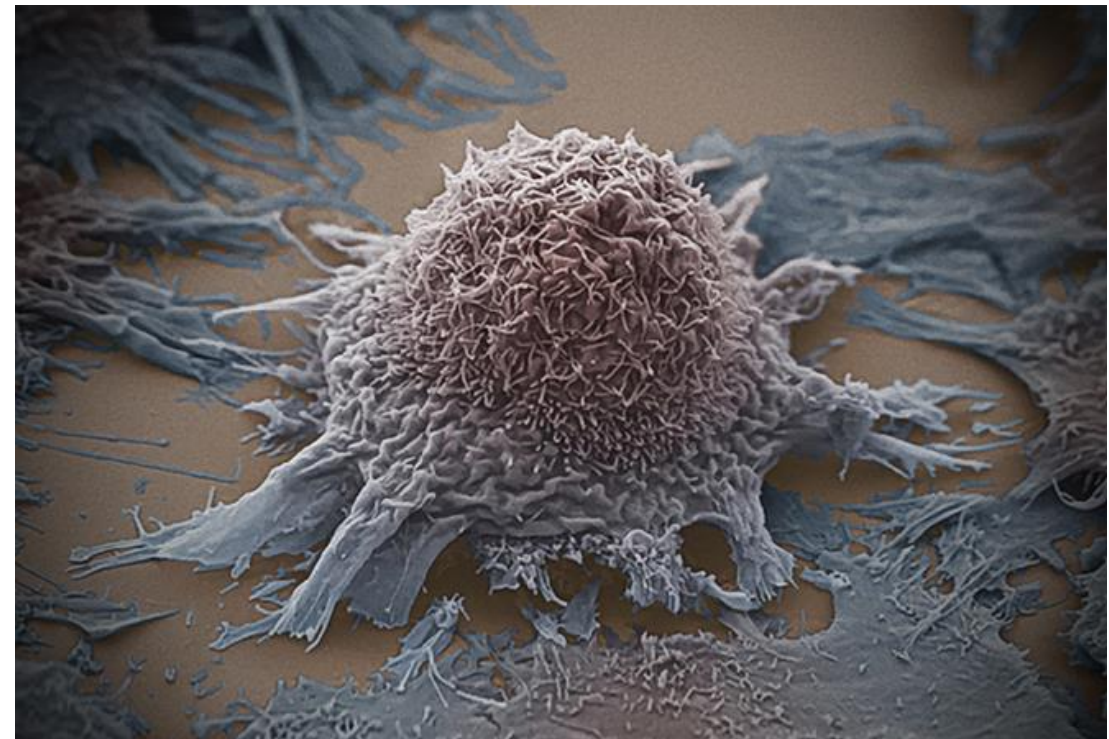
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- **Creation of electrochemical gradients**
 - Cells organize ions on either cell membrane sides to create electrochemical potentials
 - Mechanism that controls signaling and transfer of materials across the membranes



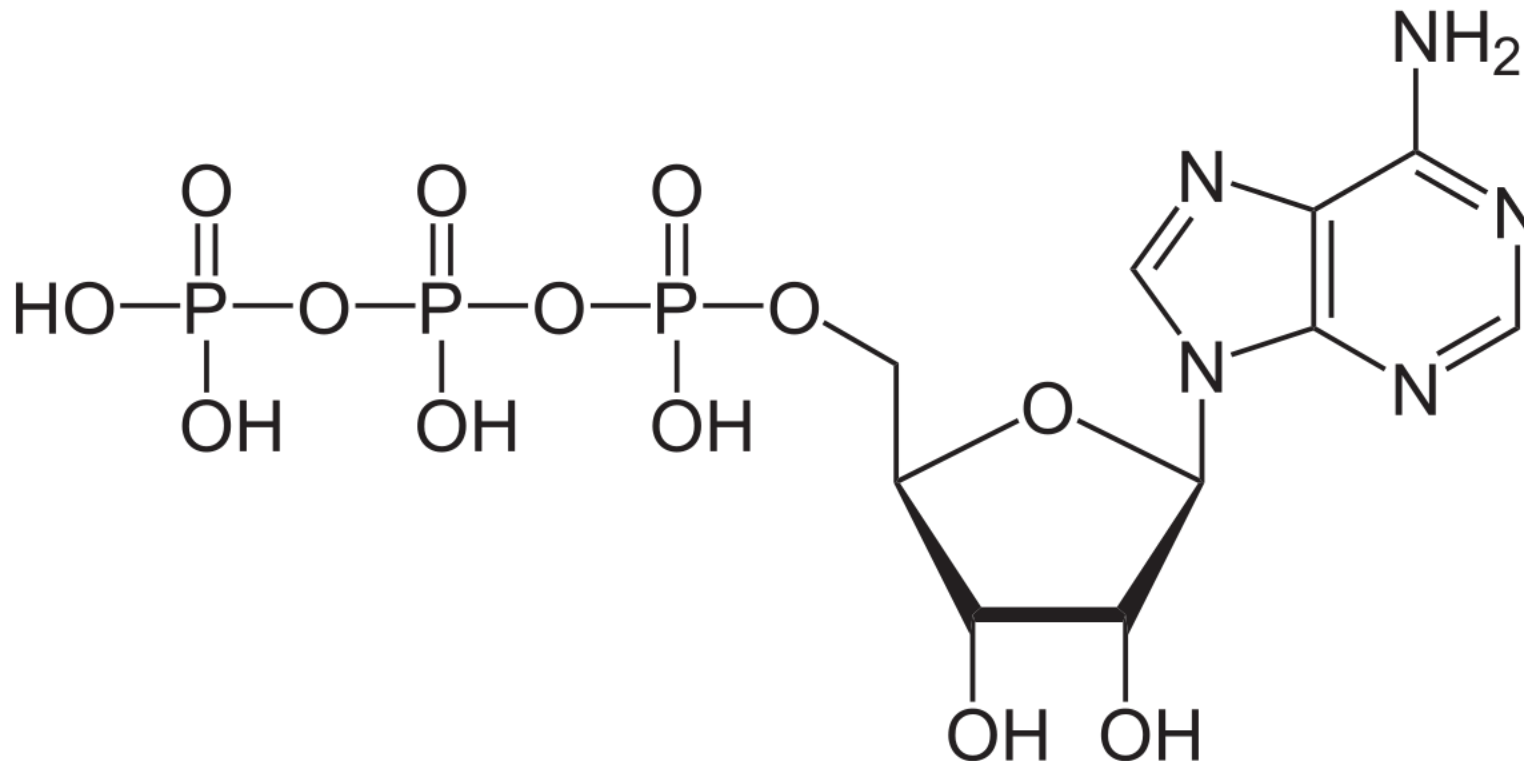
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 - Mechanism that controls signaling and transfer of materials across the membranes
- **Transport and movement**
 - Cells transport molecules from one side of the cell to another
 - Some cells swim or crawl using special motility proteins



Where does cell energy really come from?

- Cells do not use the energy from food directly
- Adenosine triphosphate, the energy middleman



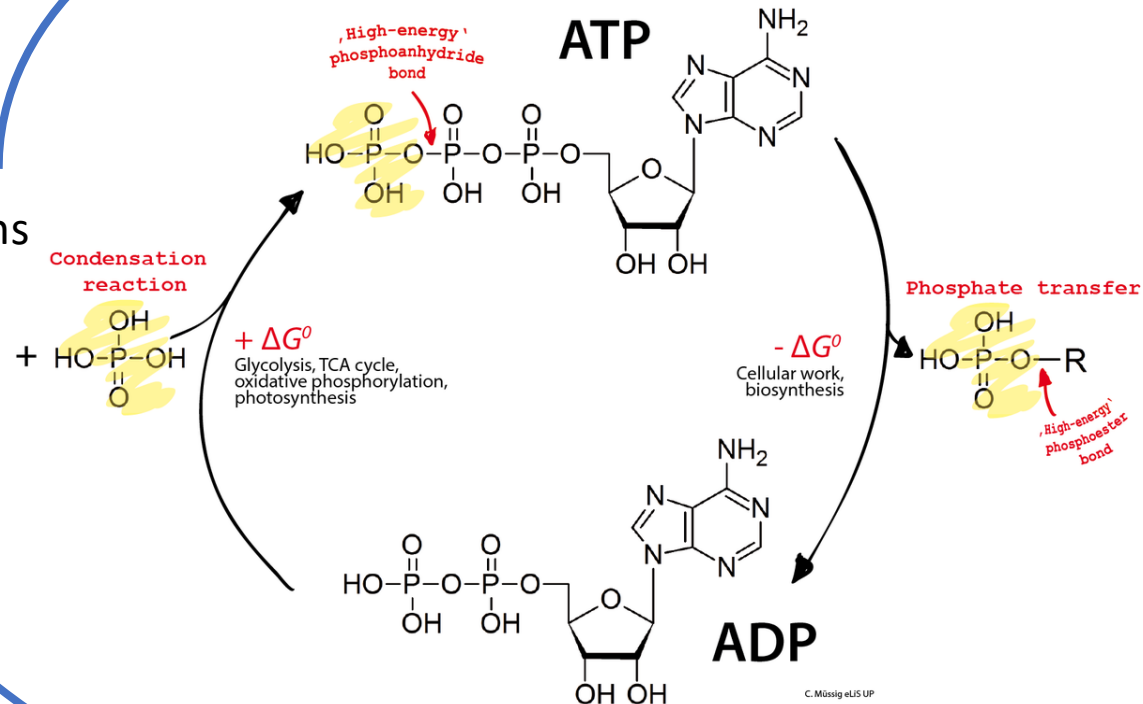
The ATP/ADP cycle

Simple molecules
glucose, amino acids, fatty acids

Catabolism – Exergonic reactions transfer energy from complex molecules to ATP

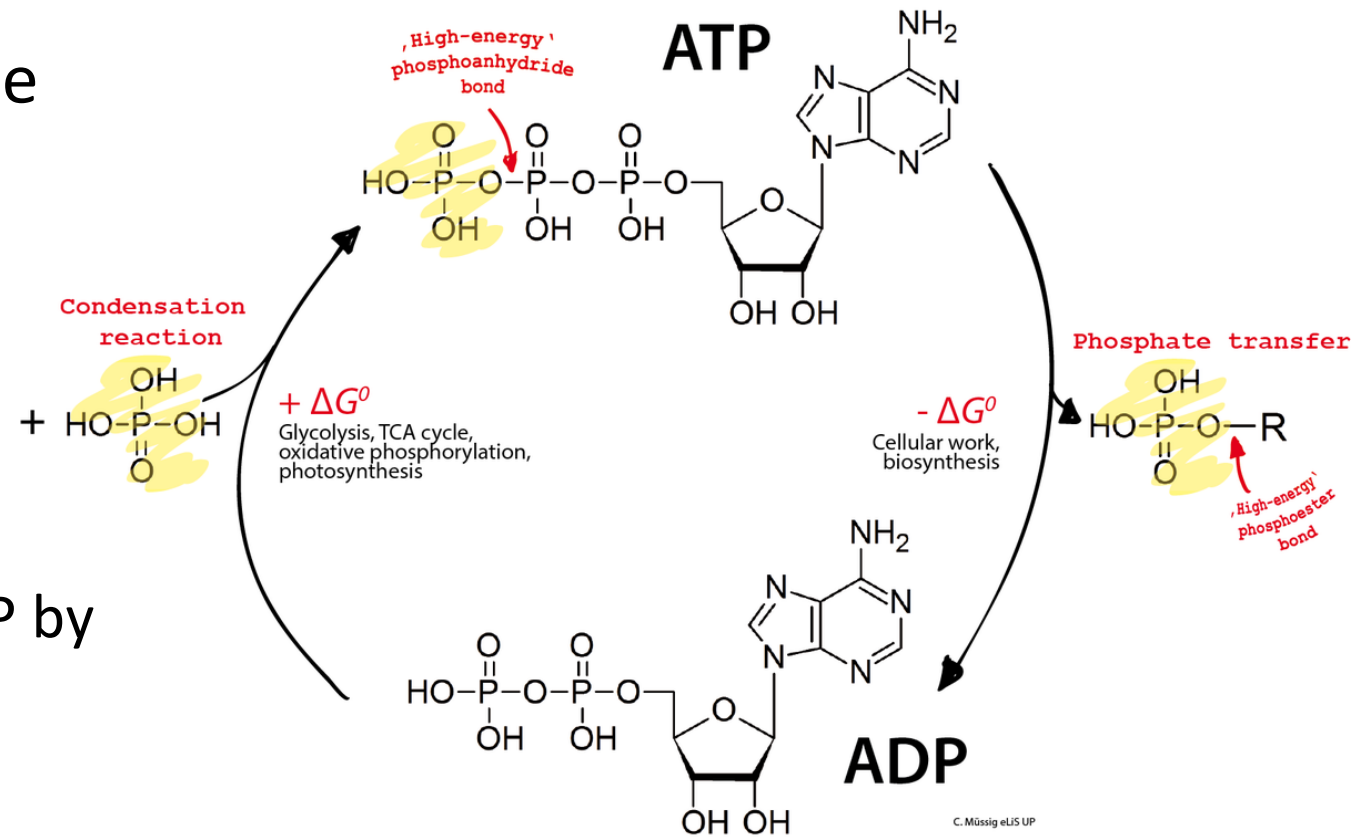
Complex molecules
polycarbs, proteins, lipids

Anabolism – Endergonic reactions transfer energy from ATP to complex molecules

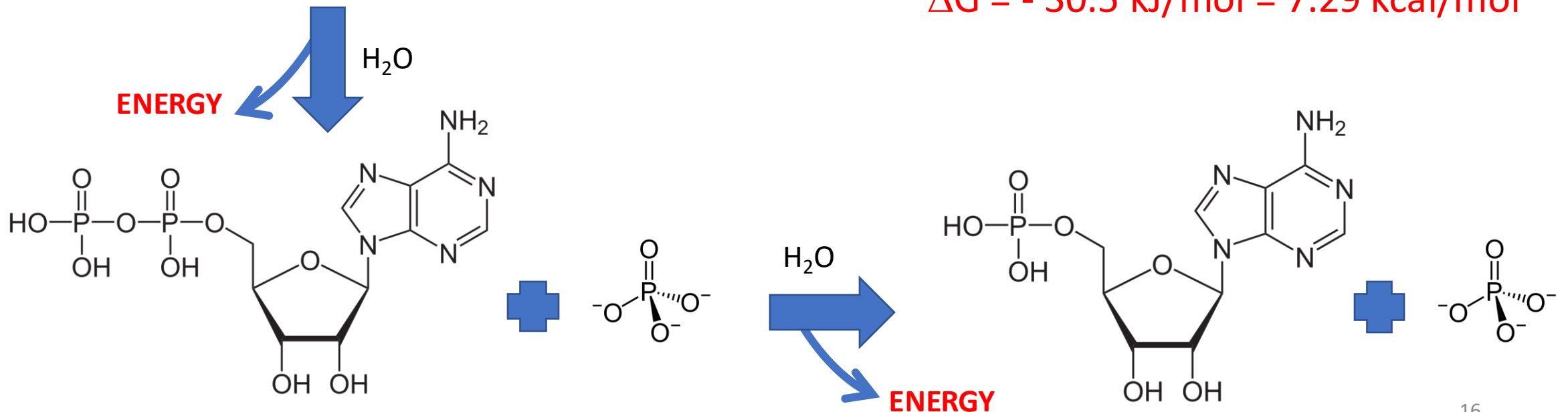
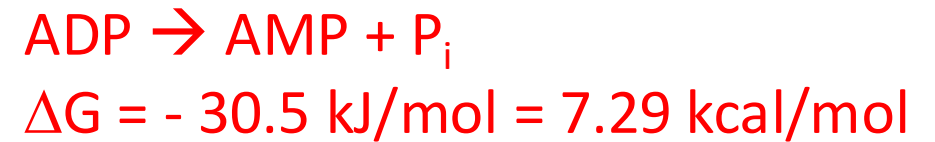
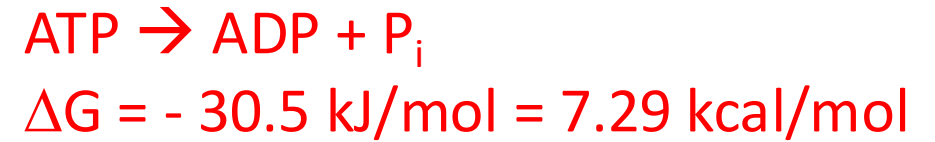
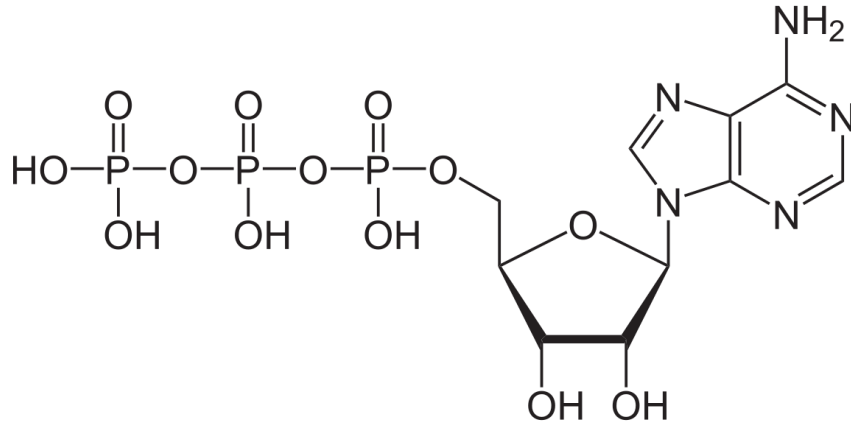


The ATP/ADP cycle – energy coupling

- Cells make and break ATP all the time
- In endergonic reactions
 - Energy is provided by ATP hydrolysis to ADP and P_i
- In exergonic reactions
 - Energy is captured to restore ATP by ADP and P_i condensation

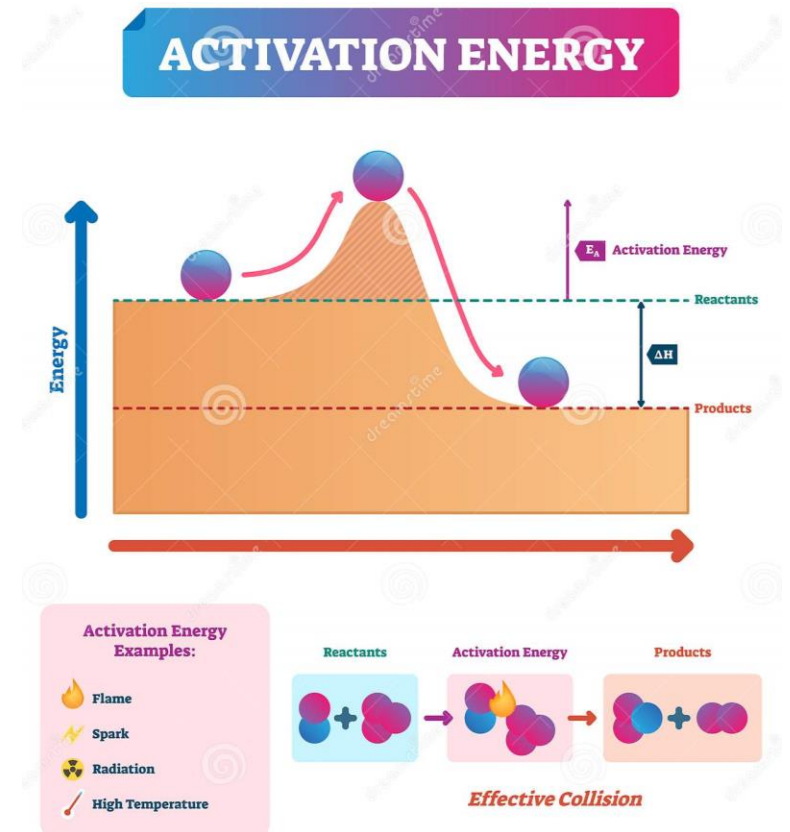


As easy as 1,2,3 – ATP, ADP and AMP



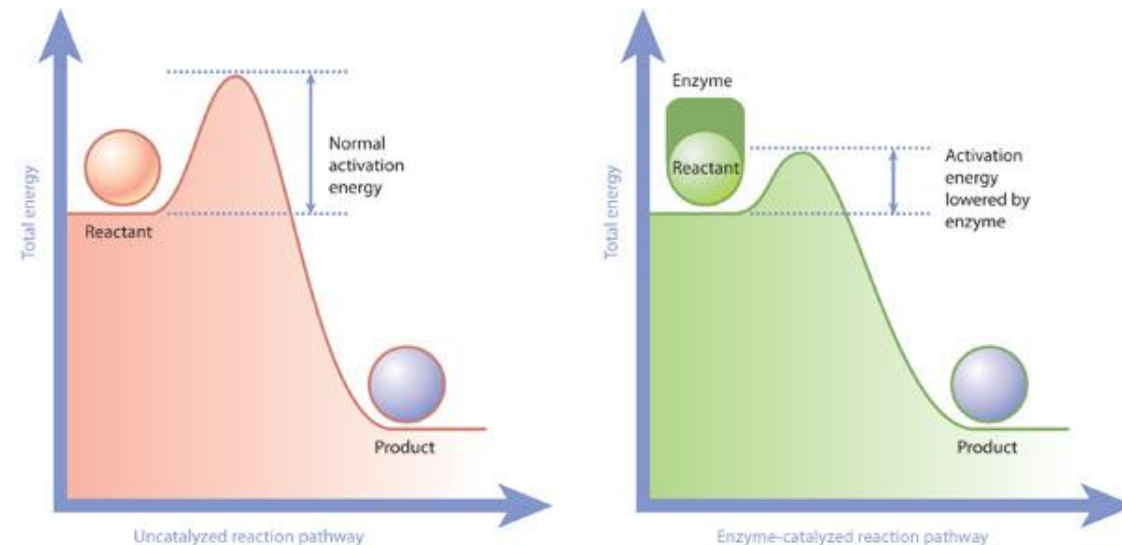
Free energy difference

- Even if a reaction is thermodynamically spontaneous ($\Delta G < 0$), it may not occur
 - It needs an “energetical push”
- **Activation energy E_a = energy barrier**



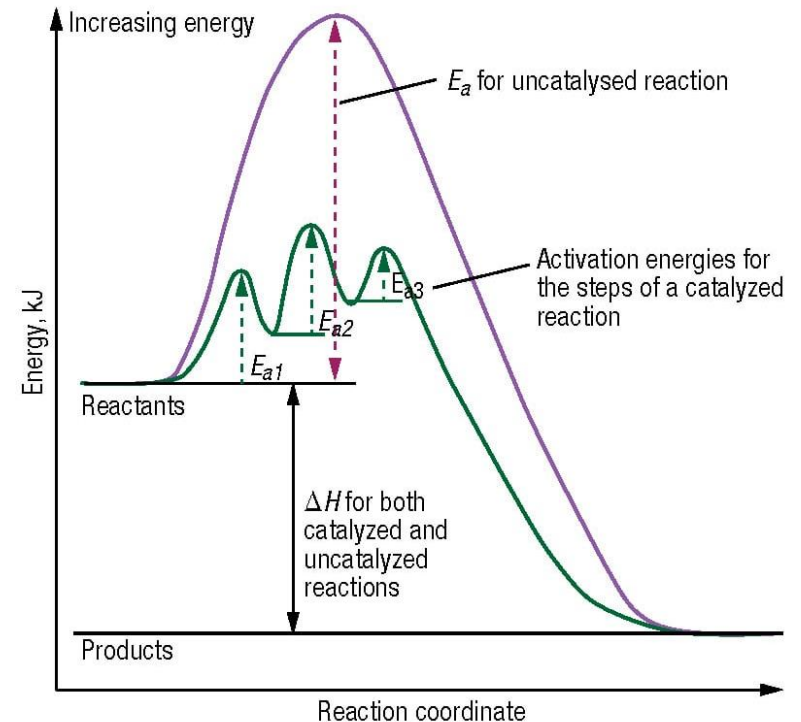
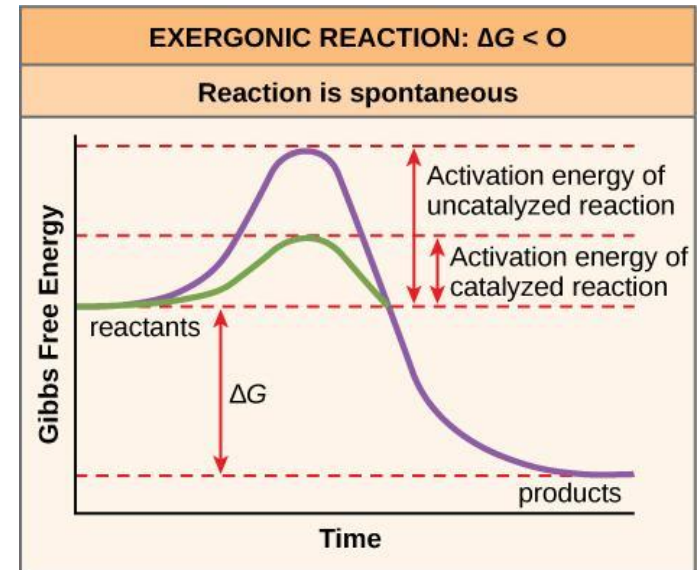
Enzymes

- Even if a reaction is thermodynamically spontaneous ($\Delta G < 0$), it may not occur
 - It needs an “energetical push”
- **Activation energy** E_a = energy barrier
- **Catalysts** = particular class of chemical substances that lower E_a and promote reactions
- **Enzymes** = biological catalysts (mostly proteins)

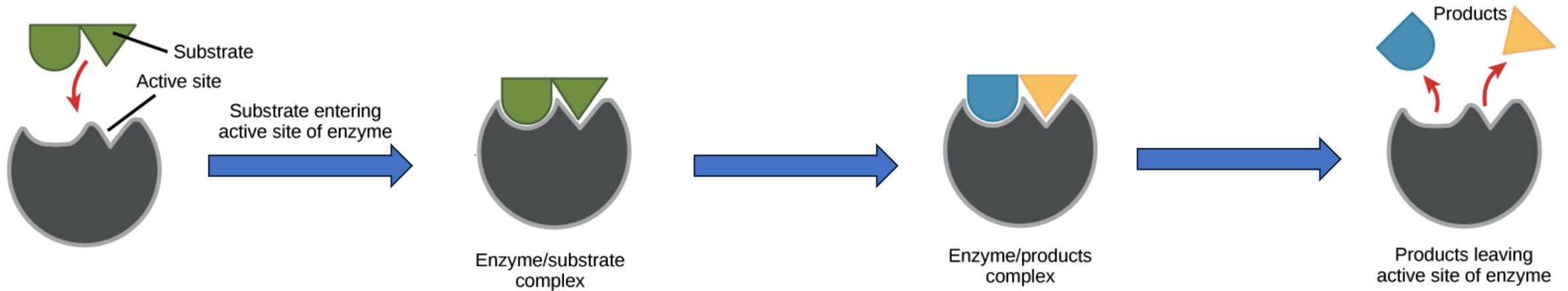


Enzymes

- There are approximately 1300 different enzymes found in the human cell
- Each enzyme catalyzes a specific chemical reaction
- **ENZYMES DO NOT CHANGE THE ΔG OF A REACTION BUT JUST SPEED UP THE REACTION RATE**
 - By breaking down the reaction into different steps, each of which has a low E_a



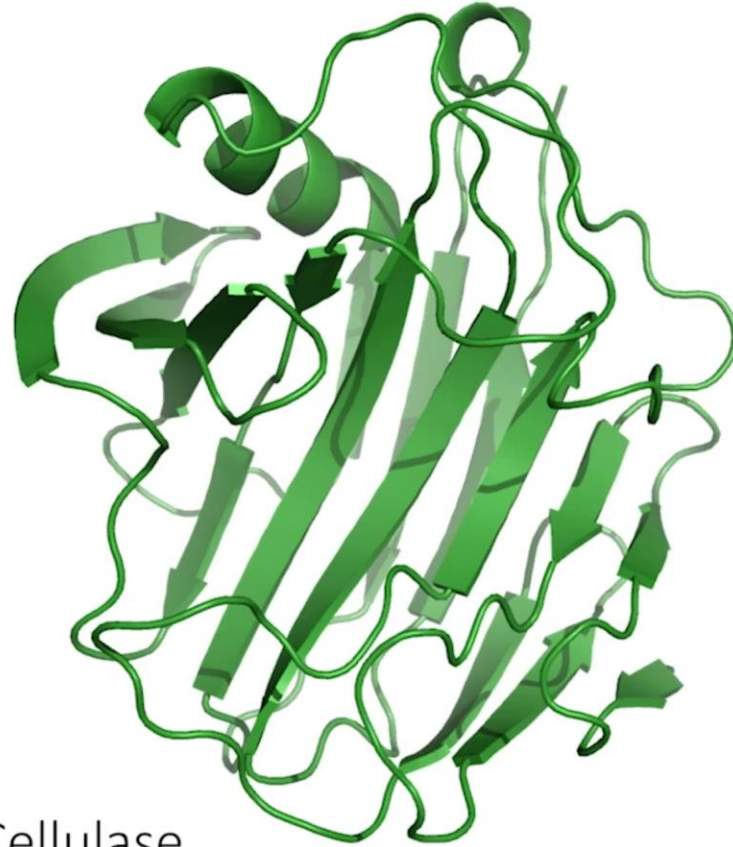
Enzymes



ENZYME CYCLE
(reversible)

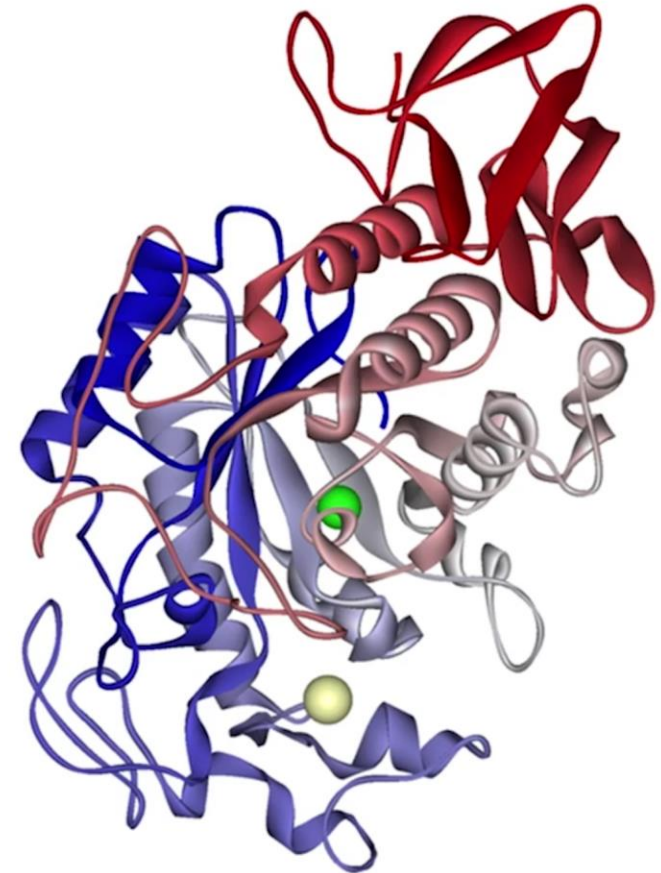
Enzyme specificity

Cellulose and starch
are both glucose polymers



Cellulase
breaks β -1,4 bonds in cellulose
People do not have this enzyme
so we cannot digest grass!

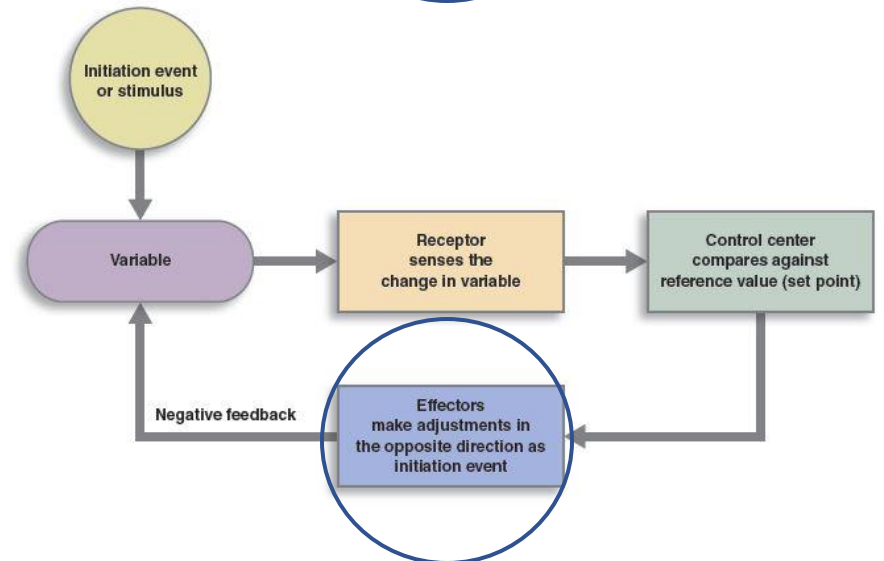
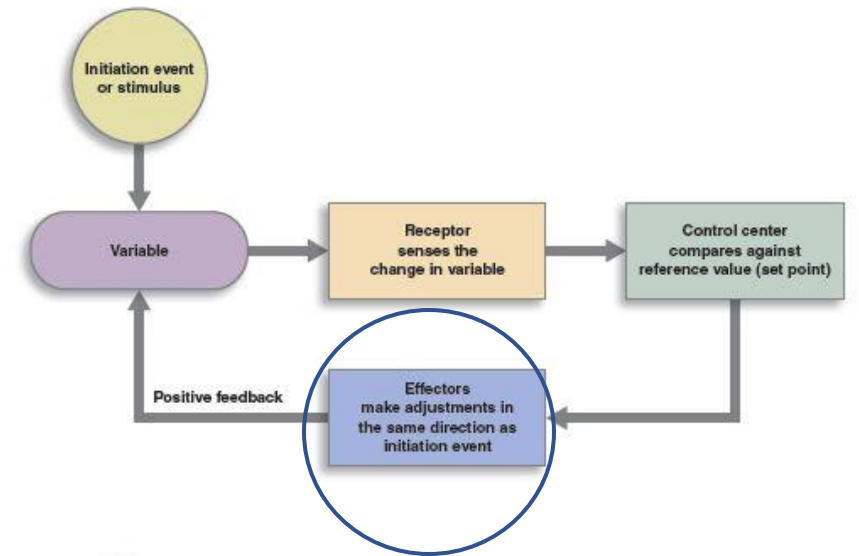
Amylase
breaks α -1,4 bonds in starch
People have this enzyme!



ENZYME SPECIFICITY

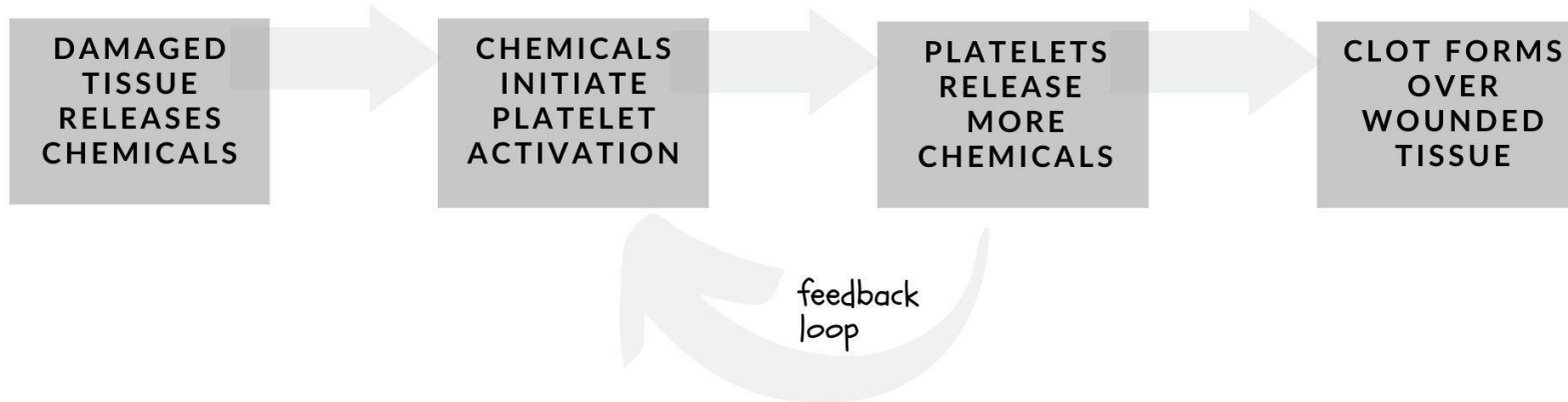
Metabolic pathways

- Cellular chemical reactions all catalyzed by enzymes
- Cellular chemical reactions are often linked into **pathways**
 - Ordered sequence of chemical reactions
- Pathways = the “cell production line”
 - Starting from point A → land up with a particular product that the cell really needs
- Reactions are organized into multistep pathways
- Cellular pathways are governed by internal feedback mechanisms
 - **Positive feedback** = make more product(s) along that particular pathway
 - **Negative feedback** = make less and/or stop producing product(s) along that particular pathways
- Cellular pathways can also be mastered by external control signals

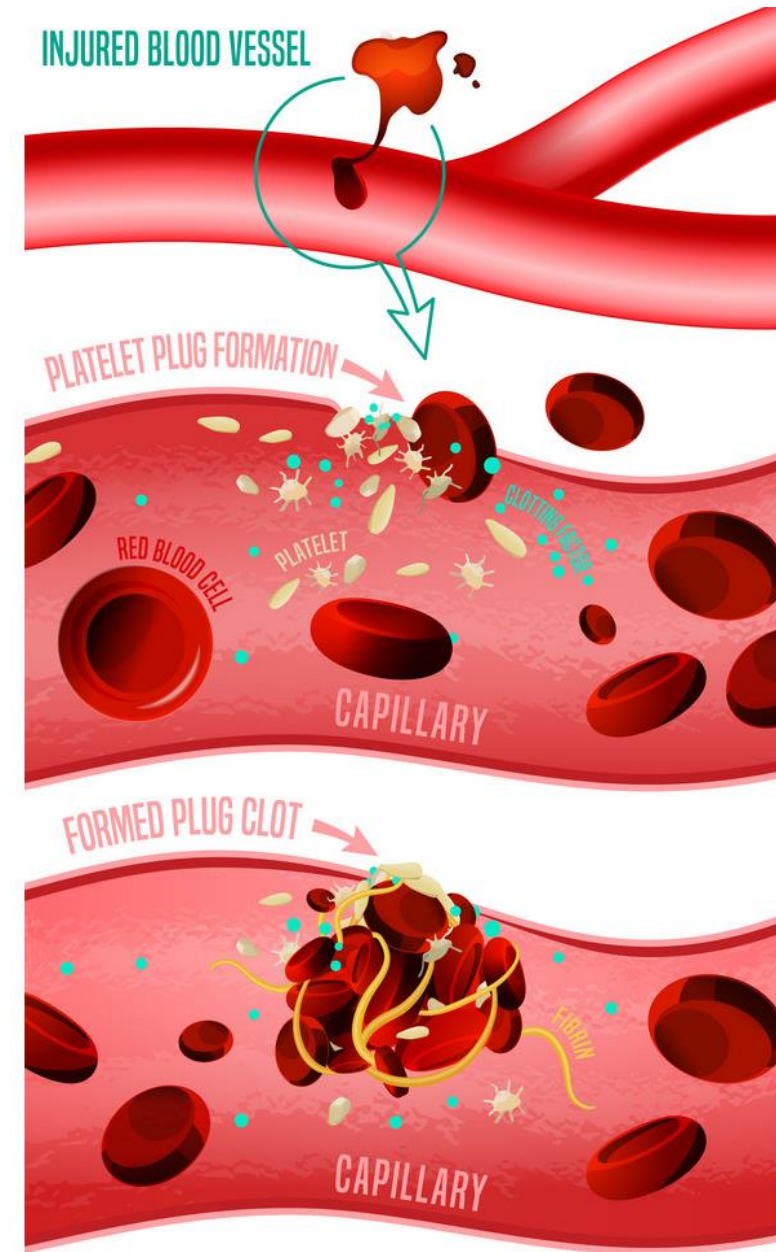


Positive feedback loops

Blood clot formation

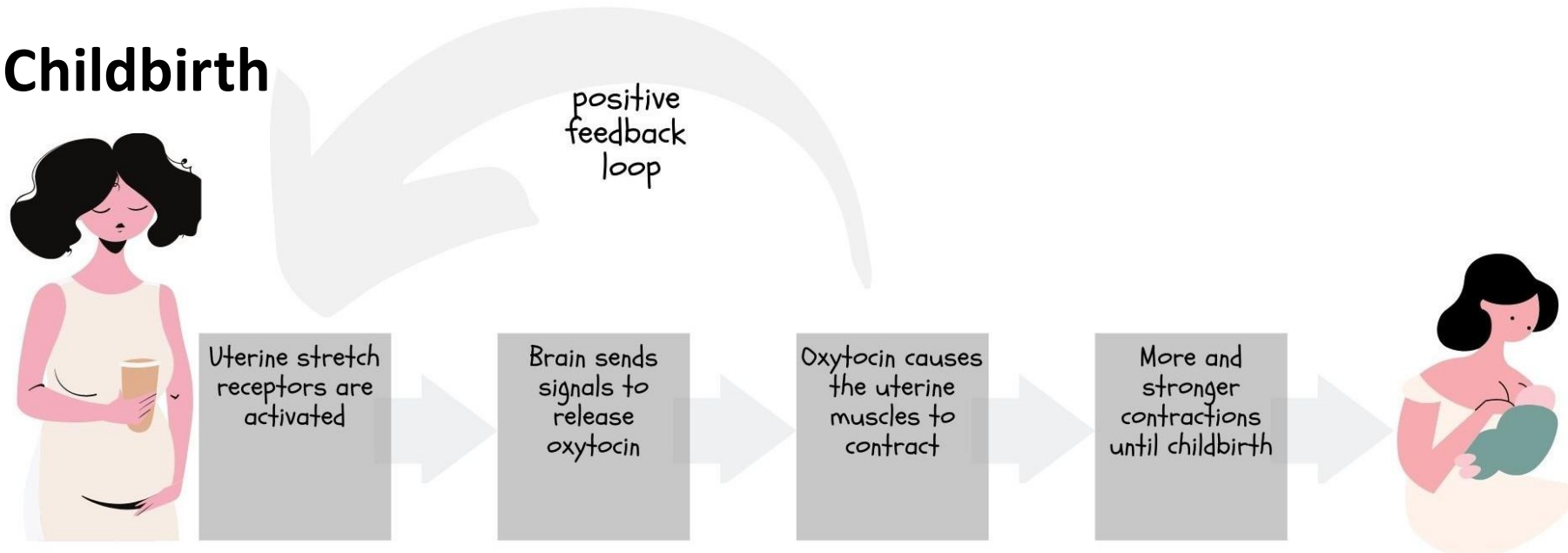


- When the body gets injured the major threat to life is excessive loss of blood
- Blood pressure and blood flow at the site of injury are reduced
- At the site of injury, blood clotting factors are released to initiate blood clotting
 - Clotting factors are responsible for the formation of a clot in the injured or wounded area
- Once the process begins it promotes the clotting process further → overall, the process of sealing the injured site is speeded up
- **This is one of the life-saving examples of positive feedback**

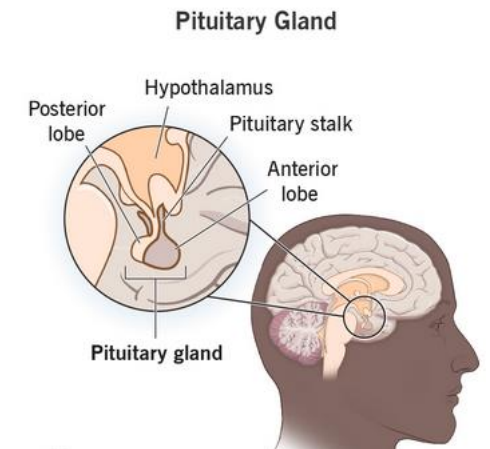


Positive feedback loops

Childbirth

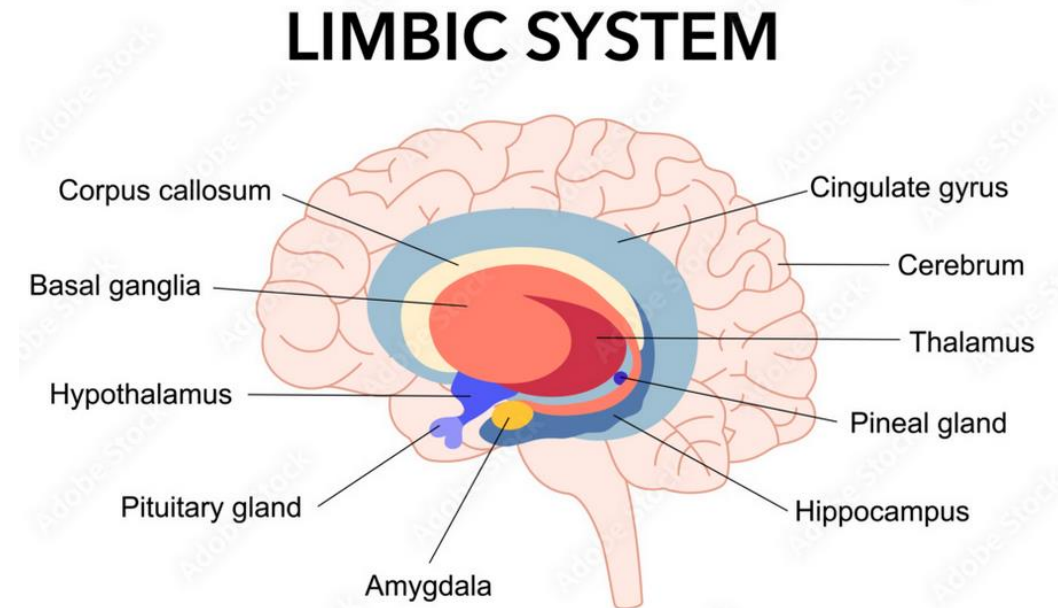


- The onset of contractions in childbirth is also known as the *Ferguson reflex*
- The baby pushes against cervix, causing it to stretch
- Stretching of cervix causes nerve impulses to be sent to the brain
- Brains stimulates the posterior pituitary gland to release oxytocin
- Oxytocin stimulates the uterine muscle to contract, initiating the birth process
- The fetus responds to uterine contractions by releasing prostaglandins, which triggers further uterine contractions



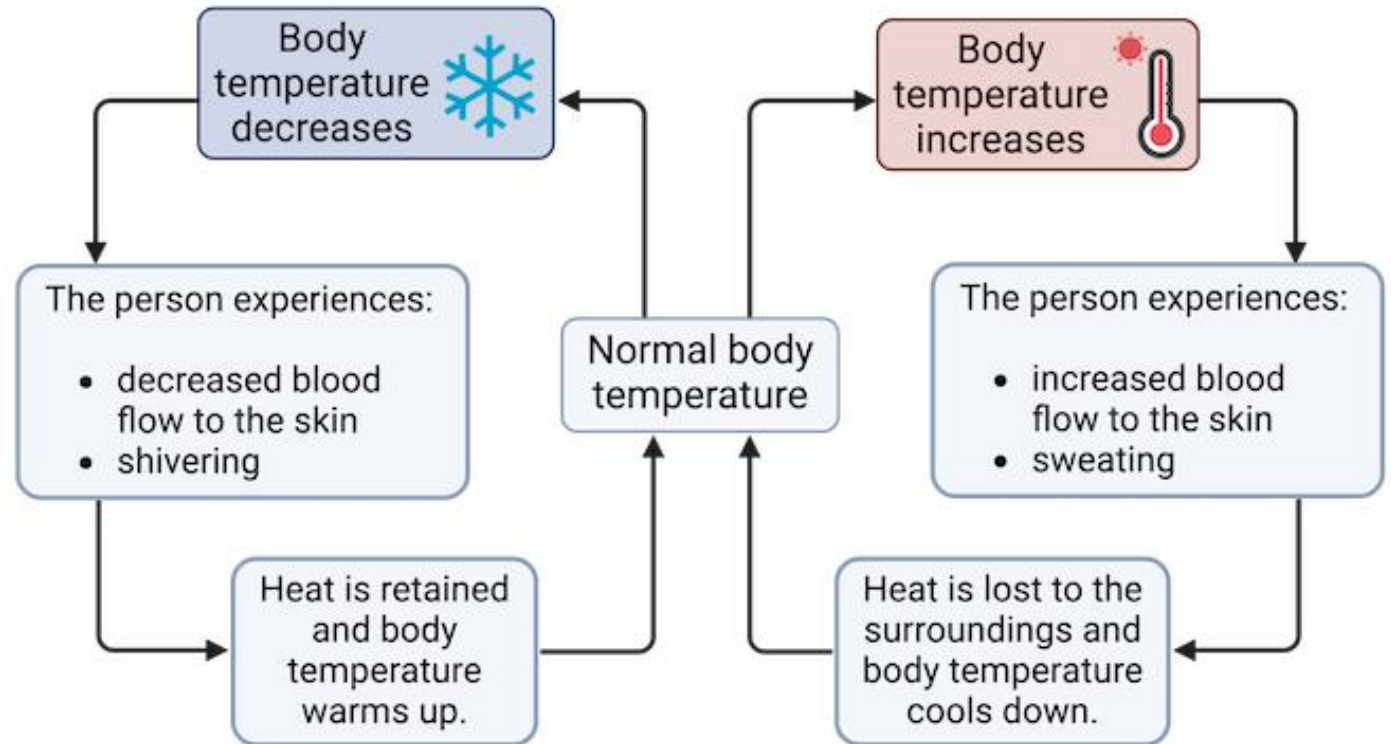
Negative feedback loops maintain homeostasis in organisms

- **Homeostasis:** tendency of organisms to maintain relatively stable internal environments despite changes in external conditions, diet, or activity level
- The primary way that organisms maintain homeostasis is through **negative feedback loops**
 - These feedback loops *counteract*, or oppose, a change in the organism
- **Hypothalamus** is the region in the ventral brain that **governs homeostasis** by directly influencing your autonomic nervous system or by managing hormones



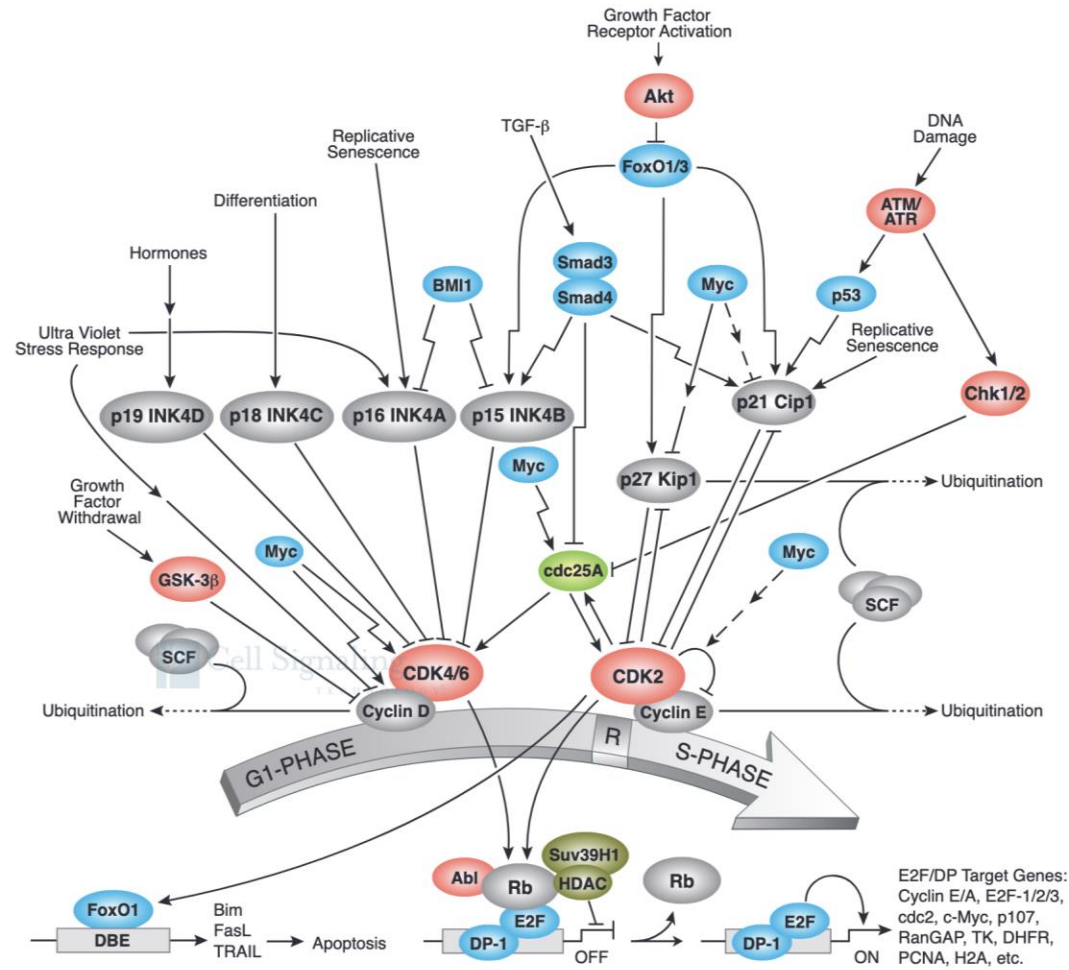
Negative feedback loops maintain homeostasis in organisms

- Nerves in the body and brain detect if you are getting too hot or too cold
- These signals are sent to the hypothalamus
- The hypothalamus then coordinates a physiological response:
 - If body temperature is too high, you begins to sweat and blood vessels dilate, increasing blood flow to the skin.
 - This increases the amount of heat lost to the surroundings
 - If body temperature is too low, you begins to shiver so that your muscles will generate more heat through cellular respiration.
 - In addition, blood vessels constrict, decreasing blood flow to the skin which limits heat loss



Cellular pathways (examples)

Cell Cycle Control: G1/S Checkpoint



Cell Cycle Control: G2/M DNA Damage Checkpoint

