

Cell (biology)

The **cell** is the basic structural and functional unit of all forms of life. Every cell consists of cytoplasm enclosed within a membrane; many cells contain organelles, each with a specific function. The term comes from the Latin word *cellula* meaning 'small room'. Most cells are only visible under a microscope. Cells emerged on Earth about 4 billion years ago. All cells are capable of replication, protein synthesis, and motility.

broadly categorized into two Cells are eukaryotic cells, which possess a nucleus, and prokaryotic cells, which lack a nucleus but have a are nucleoid region. Prokaryotes single-celled organisms such as bacteria, whereas eukaryotes can be either single-celled, such as amoebae, or multicellular, such as some algae, plants, animals, and fungi. Eukaryotic cells contain organelles including mitochondria, which provide energy for cell functions; chloroplasts, which create sugars by photosynthesis, in plants; and ribosomes, which synthesise proteins.

Cells were discovered by Robert Hooke in 1665, who named them after their resemblance to cells inhabited by Christian monks in a monastery. Cell theory, developed in 1839 by Matthias Jakob Schleiden and Theodor Schwann, states that all organisms are composed of one or more cells, that cells are the fundamental unit of structure and function in all living organisms, and that all cells come from pre-existing cells.

Onion (Allium cepa) root cells in different phases of the cell cycle (drawn by E. B. Wilson, 1900) Eukaryote Prokaryote Cell Membrane Cell Wall A eukaryotic cell (left) and prokaryotic cell (right) **Identifiers** MeSH D002477 (https://meshb.nlm.nih.gov/re cord/ui?ui=D002477) TH H1.00.01.0.00001 (https://ifaa.unifr.ch/ Public/EntryPage/PDF/TH%20Chapte r%20H1.00.pdf) **FMA** 686465 (https://bioportal.bioontology.or g/ontologies/FMA/?p=classes&concepti d=http%3A%2F%2Fpurl.org%2Fsig%2 Font%2Ffma%2Ffma686465) Anatomical terminology

Cell

Cell types

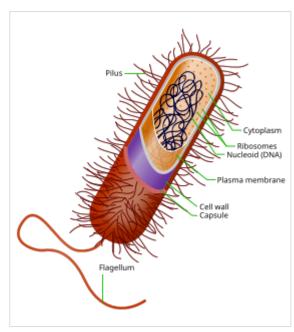
Cells are broadly categorized into two types: <u>eukaryotic cells</u>, which possess a <u>nucleus</u>, and <u>prokaryotic cells</u>, which lack a nucleus but have a nucleoid region. Prokaryotes are <u>single-celled organisms</u>, whereas eukaryotes can be either single-celled or multicellular.

Prokaryotic cells

Prokaryotes include <u>bacteria</u> and <u>archaea</u>, two of the three <u>domains</u> of <u>life</u>. Prokaryotic cells were the first form of <u>life</u> on Earth, characterized by having vital <u>biological processes</u> including <u>cell signaling</u>. They are simpler and smaller than eukaryotic cells, and lack a <u>nucleus</u>, and other membrane-bound <u>organelles</u>. The <u>DNA</u> of a prokaryotic cell consists of a single <u>circular chromosome</u> that is in direct contact with the <u>cytoplasm</u>. The nuclear region in the cytoplasm is called the <u>nucleoid</u>. Most prokaryotes are the smallest of all organisms, ranging from 0.5 to 2.0 µm in diameter. [1]

A prokaryotic cell has three regions:

Enclosing the cell is the <u>cell envelope</u>, generally consisting of a <u>plasma membrane</u> covered by a <u>cell wall</u> which, for some bacteria, may be further covered by a third layer called a <u>capsule</u>. Though most prokaryotes have both a cell



Structure of a typical prokaryotic cell

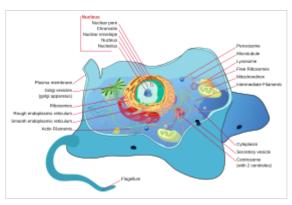
- membrane and a cell wall, there are exceptions such as <u>Mycoplasma</u> (bacteria) and <u>Thermoplasma</u> (archaea) which only possess the cell membrane layer. The envelope gives rigidity to the cell and separates the interior of the cell from its environment, serving as a protective filter. The cell wall consists of <u>peptidoglycan</u> in bacteria and acts as an additional barrier against exterior forces. It also prevents the cell from expanding and bursting (cytolysis) from <u>osmotic pressure</u> due to a <u>hypotonic</u> environment. Some eukaryotic cells (plant cells and fungal cells) also have a cell wall.
- Inside the cell is the cytoplasmic region that contains the genome (DNA), ribosomes and various sorts of inclusions. [2] The genetic material is freely found in the cytoplasm. Prokaryotes can carry extrachromosomal DNA elements called plasmids, which are usually circular. Linear bacterial plasmids have been identified in several species of spirochete bacteria, including members of the genus <u>Borrelia</u> notably <u>Borrelia burgdorferi</u>, which causes Lyme disease. [3] Though not forming a nucleus, the <u>DNA</u> is condensed in a nucleoid. Plasmids encode additional genes, such as antibiotic resistance genes.
- On the outside, some prokaryotes have <u>flagella</u> and <u>pili</u> that project from the cell's surface. These are structures made of proteins that facilitate movement and communication between cells.

Eukaryotic cells

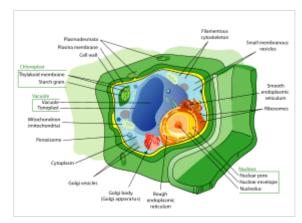
<u>Plants</u>, <u>animals</u>, <u>fungi</u>, <u>slime moulds</u>, <u>protozoa</u>, and <u>algae</u> are all <u>eukaryotic</u>. These cells are about fifteen times wider than a typical prokaryote and can be as much as a thousand times greater in volume. The main distinguishing feature of eukaryotes as compared to prokaryotes is <u>compartmentalization</u>: the presence of membrane-bound <u>organelles</u> (compartments) in which specific activities take place. Most important among these is a <u>cell nucleus</u>, <u>[2]</u> an organelle that houses the cell's <u>DNA</u>. This nucleus gives the eukaryote its name, which means "true kernel (nucleus)". Some of the other differences are:

- The plasma membrane resembles that of prokaryotes in function, with minor differences in the setup. Cell walls may or may not be present.
- The eukaryotic DNA is organized in one or more linear molecules, called <u>chromosomes</u>, which are associated with histone proteins. All chromosomal DNA is stored in the cell

- <u>nucleus</u>, separated from the cytoplasm by a membrane. [2] Some eukaryotic organelles such as mitochondria also contain some DNA.
- Many eukaryotic cells are <u>ciliated</u> with <u>primary cilia</u>. Primary cilia play important roles in chemosensation, <u>mechanosensation</u>, and <u>thermosensation</u>. Each cilium may thus be "viewed as a sensory cellular <u>antennae</u> that coordinates a large number of cellular signaling pathways, sometimes coupling the signaling to ciliary motility or alternatively to cell division and differentiation."
- Motile eukaryotes can move using motile cilia or flagella. Motile cells are absent in conifers and flowering plants. Eukaryotic flagella are more complex than those of prokaryotes.



Structure of a typical animal cell



Structure of a typical plant cell

Comparison of features of prokaryotic and eukaryotic cells

	Prokaryotes	Eukaryotes
Typical organisms	bacteria, archaea	protists, algae, fungi, plants, animals
Typical size	~ 1–5 <u>µm^[6]</u>	~ 10–100 μm ^[6]
DNA	In nucleoid region	In <u>nucleus</u> with double membrane
Chromosomes	Single, usually <u>circular</u>	Multiple paired linear chromosomes with histone proteins
RNA/protein synthesis	coupled in the cytoplasm	RNA synthesis in the nucleus protein synthesis in the cytoplasm
Ribosomes	50S and 30S	60S and 40S
Cytoplasmic structure	Microcompartments with proteins, cytoskeleton	Endomembrane system, cytoskeleton
Cell movement	flagella	flagella and cilia; lamellipodia and filopodia
Mitochondria	none	one to several thousand
Chloroplasts	none	in <u>algae</u> and <u>plants</u>
Organization	single cells, colonies, biofilms	single cells, colonies, multicellular organisms with specialized cells
Cell division	binary fission (simple division)	mitosis (fission or budding) meiosis
Membranes	cell membrane	Cell membrane and membrane-bound organelles

Many groups of eukaryotes are single-celled. Among the many-celled groups are animals and plants. The number of cells in these groups vary with species; it has been estimated that the <u>human body</u> contains around 37 trillion (3.72×10^{13}) cells, [7] and more recent studies put this number at around 30 trillion (\sim 36 trillion cells in the male, \sim 28 trillion in the female).[8]

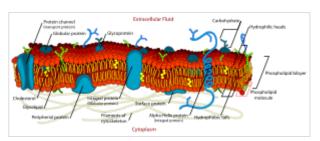
Subcellular components

All cells, whether <u>prokaryotic</u> or <u>eukaryotic</u>, have a <u>membrane</u> that envelops the cell, regulates what moves in and out (selectively permeable), and maintains the <u>electric</u> potential of the cell. Inside the membrane, the <u>cytoplasm</u> takes up most of the cell's volume. Except <u>red blood cells</u>, which lack a cell nucleus and most organelles to accommodate maximum space for <u>hemoglobin</u>, all cells possess <u>DNA</u>, the hereditary material of <u>genes</u>, and <u>RNA</u>, containing the information necessary to <u>build</u> various <u>proteins</u> such as <u>enzymes</u>, the cell's primary machinery. There are also other kinds of <u>biomolecules</u> in cells. This article lists these primary <u>cellular components</u>, then briefly describes their function.

Cell membrane

The <u>cell membrane</u>, or plasma membrane, is a selectively permeable <u>biological membrane</u> that surrounds the cytoplasm of a cell. In animals, the plasma membrane is the outer boundary of the cell, while in plants and prokaryotes it is usually covered by a <u>cell wall</u>. This membrane serves to separate and protect a cell from its surrounding environment and is made mostly from a <u>double layer of phospholipids</u>, which are amphiphilic (partly hydrophobic and partly hydrophilic). Hence, the layer is called a phospholipid

<u>bilayer</u>, or sometimes a fluid mosaic membrane. Embedded within this membrane is a macromolecular structure called the <u>porosome</u> the universal secretory portal in cells and a variety of <u>protein</u> molecules that act as channels and pumps that move different molecules into and out of the cell. The membrane is semi-permeable, and selectively permeable, in that it can either let a substance (<u>molecule</u> or <u>ion</u>) pass through freely, to a limited extent or not at all. Cell

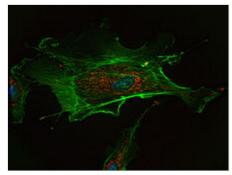


Detailed diagram of lipid bilayer of cell membrane

surface membranes also contain $\underline{\text{receptor}}$ proteins that allow cells to detect external signaling molecules such as hormones. [9]

Cytoskeleton

The cytoskeleton acts to organize and maintain the cell's shape; anchors organelles in place; helps during endocytosis, the uptake of external materials by a cell, and cytokinesis, the separation of daughter cells after cell division; and moves parts of the cell in processes of growth and mobility. The eukaryotic cytoskeleton is microtubules, composed of intermediate filaments microfilaments. In the cytoskeleton of a neuron the intermediate filaments are known as neurofilaments. There are a great number of proteins associated with them, each controlling a cell's structure by directing, bundling, and aligning filaments. [2] The prokaryotic cytoskeleton is less well-studied but is involved in the maintenance of cell shape, polarity and cytokinesis. [10] The subunit protein of microfilaments is a small, monomeric protein called actin. The subunit of microtubules is a dimeric molecule



A fluorescent image of an endothelial cell. Nuclei are stained blue, mitochondria are stained red, and microfilaments are stained green.

called <u>tubulin</u>. Intermediate filaments are heteropolymers whose subunits vary among the cell types in different tissues. Some of the subunit proteins of intermediate filaments include <u>vimentin</u>, <u>desmin</u>, <u>lamin</u> (lamins A, B and C), <u>keratin</u> (multiple acidic and basic keratins), and <u>neurofilament proteins</u> (<u>NF–L</u>, <u>NF–M</u>).

Genetic material

Two different kinds of genetic material exist: <u>deoxyribonucleic acid</u> (DNA) and <u>ribonucleic acid</u> (RNA). Cells use DNA for their long-term information storage. The biological information contained in an organism is <u>encoded</u> in its DNA sequence. RNA is used for information transport (e.g., <u>mRNA</u>) and <u>enzymatic</u> functions (e.g., <u>ribosomal</u> RNA). <u>Transfer RNA</u> (tRNA) molecules are used to add amino acids during protein <u>translation</u>.

Prokaryotic genetic material is organized in a simple <u>circular bacterial chromosome</u> in the <u>nucleoid region</u> of the cytoplasm. Eukaryotic genetic material is divided into different, [2] linear molecules called <u>chromosomes</u> inside a discrete nucleus, usually with additional genetic material in some organelles like mitochondria and chloroplasts (see endosymbiotic theory).

A human cell has genetic material contained in the cell nucleus (the nuclear genome) and in the mitochondria (the mitochondrial genome). In humans, the nuclear genome is divided into 46 linear DNA molecules called chromosomes, including 22 homologous chromosome pairs and a pair of sex chromosomes. The mitochondrial genome is a circular DNA molecule distinct from nuclear DNA. Although the mitochondrial DNA is very small compared to nuclear chromosomes. [2] it codes for 13 proteins involved in mitochondrial energy production and specific tRNAs.

Foreign genetic material (most commonly DNA) can also be artificially introduced into the cell by a process called transfection. This can be transient, if the DNA is not inserted into the cell's genome, or stable, if it is. Certain viruses also insert their genetic material into the genome.

Deoxyribonucleic acid (DNA)

Organelles

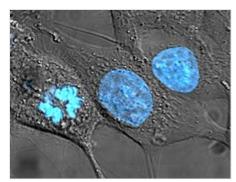
Organelles are parts of the cell that are adapted and/or specialized for carrying out one or more vital functions, analogous to the organs of the human body (such as the heart, lung, and kidney,

with each organ performing a different function). [2] Both eukaryotic and prokaryotic cells have organelles, but prokaryotic organelles are generally simpler and are not membrane-bound.

There are several types of organelles in a cell. Some (such as the nucleus and Golgi apparatus) are typically solitary, while others (such as mitochondria, chloroplasts, peroxisomes and lysosomes) can be numerous (hundreds to thousands). The cytosol is the gelatinous fluid that fills the cell and surrounds the organelles.

Eukaryotic

• **Cell nucleus**: A cell's information center, the cell nucleus is the most conspicuous organelle found in a eukaryotic cell. It houses the cell's chromosomes, and is the place where almost all DNA replication and RNA synthesis (transcription) occur. The nucleus is spherical and separated from the cytoplasm by a double membrane called the nuclear envelope, space between these two membrane is called perinuclear space. The nuclear envelope isolates and protects a cell's DNA from various molecules that could accidentally damage its structure or interfere with its processing. During processing, DNA is transcribed, or copied into a special RNA, called messenger RNA (mRNA). This mRNA is then transported out of the nucleus, where it is translated into a specific protein molecule. The nucleolus is a specialized region within the nucleus where ribosome subunits are assembled. In prokaryotes, DNA processing takes place in the cytoplasm. [2]



Human cancer cells, specifically HeLa cells, with DNA stained blue. The central and rightmost cell are in interphase, so their DNA is diffuse and the entire nuclei are labelled. The cell on the left is going through mitosis and its chromosomes have condensed.

- **Mitochondria and chloroplasts**: generate energy for the cell. <u>Mitochondria</u> are self-replicating double membrane-bound organelles that occur in various numbers, shapes, and sizes in the cytoplasm of all eukaryotic cells. <u>Pespiration</u> occurs in the cell mitochondria, which generate the cell's energy by <u>oxidative phosphorylation</u>, using <u>oxygen</u> to release energy stored in cellular nutrients (typically pertaining to <u>glucose</u>) to generate <u>ATP</u> (<u>aerobic respiration</u>). Mitochondria multiply by <u>binary fission</u>, like prokaryotes. Chloroplasts can only be found in plants and algae, and they capture the sun's energy to make carbohydrates through photosynthesis.
- Endoplasmic reticulum: The endoplasmic reticulum (ER) is a transport network for molecules targeted for certain modifications and specific destinations, as compared to molecules that float freely in the cytoplasm. The ER has two forms: the rough ER, which has ribosomes on its surface that secrete proteins into the ER, and the smooth ER, which lacks ribosomes. [2] The smooth ER plays a role in calcium sequestration and release and also helps in synthesis of lipid.
- **Golgi apparatus**: The primary function of the Golgi apparatus is to process and package the macromolecules such as proteins and lipids that are synthesized by the cell.

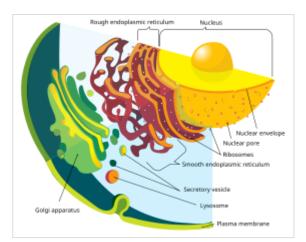


Diagram of the endomembrane system

- Lysosomes and peroxisomes: Lysosomes contain digestive enzymes (acid hydrolases). They digest excess or worn-out organelles, food particles, and engulfed viruses or bacteria. Peroxisomes have enzymes that rid the cell of toxic peroxides, Lysosomes are optimally active in an acidic environment. The cell could not house these destructive enzymes if they were not contained in a membrane-bound system. [2]
- **Centrosome**: the cytoskeleton organizer: The <u>centrosome</u> produces the <u>microtubules</u> of a cell—a key component of the <u>cytoskeleton</u>. It directs the transport through the <u>ER</u> and the <u>Golgi apparatus</u>. Centrosomes are composed of two <u>centrioles</u> which lie perpendicular to each other in which each has an organization like a <u>cartwheel</u>, which separate during <u>cell division</u> and help in the formation of the <u>mitotic spindle</u>. A single centrosome is present in the animal cells. They are also found in some fungi and algae cells.
- Vacuoles: <u>Vacuoles</u> sequester waste products and in plant cells store water. They are often described as liquid filled spaces and are surrounded by a membrane. Some cells, most notably <u>Amoeba</u>, have contractile vacuoles, which can pump water out of the cell if there is too much water. The vacuoles of plant cells and fungal cells are usually larger than those of animal cells. Vacuoles of plant cells are surrounded by a membrane which transports ions against concentration gradients.

Eukaryotic and prokaryotic

- **Ribosomes**: The <u>ribosome</u> is a large complex of <u>RNA</u> and <u>protein</u> molecules. They each consist of two subunits, and act as an assembly line where RNA from the nucleus is used to synthesise proteins from amino acids. Ribosomes can be found either floating freely or bound to a membrane (the rough endoplasmatic reticulum in eukaryotes, or the cell membrane in prokaryotes). [11]
- Plastids: Plastid are membrane-bound organelle generally found in plant cells and euglenoids and contain specific *pigments*, thus affecting the colour of the plant and organism. And these pigments also helps in food storage and tapping of light energy. There are three types of plastids based upon the specific pigments. Chloroplasts contain

<u>chlorophyll</u> and some carotenoid pigments which helps in the tapping of light energy during photosynthesis. <u>Chromoplasts</u> contain fat-soluble <u>carotenoid</u> pigments like orange carotene and yellow xanthophylls which helps in synthesis and storage. <u>Leucoplasts</u> are non-pigmented plastids and helps in storage of nutrients. [12]

Structures outside the cell membrane

Many cells also have structures which exist wholly or partially outside the cell membrane. These structures are notable because they are not protected from the external environment by the cell membrane. In order to assemble these structures, their components must be carried across the cell membrane by export processes.

Cell wall

Many types of prokaryotic and eukaryotic cells have a <u>cell wall</u>. The cell wall acts to protect the cell mechanically and chemically from its environment, and is an additional layer of protection to the cell membrane. Different types of cell have cell walls made up of different materials; plant cell walls are primarily made up of <u>cellulose</u>, fungi cell walls are made up of <u>chitin</u> and bacteria cell walls are made up of peptidoglycan.

Prokaryotic

Capsule

A gelatinous <u>capsule</u> is present in some bacteria outside the cell membrane and cell wall. The capsule may be <u>polysaccharide</u> as in <u>pneumococci</u>, <u>meningococci</u> or <u>polypeptide</u> as <u>Bacillus anthracis</u> or <u>hyaluronic acid</u> as in <u>streptococci</u>. Capsules are not marked by normal staining protocols and can be detected by <u>India ink</u> or <u>methyl blue</u>, which allows for higher contrast between the cells for observation. [13]:87

Flagella

<u>Flagella</u> are organelles for cellular mobility. The bacterial flagellum stretches from cytoplasm through the cell membrane(s) and extrudes through the cell wall. They are long and thick thread-like appendages, protein in nature. A different type of flagellum is found in archaea and a different type is found in eukaryotes.

Fimbriae

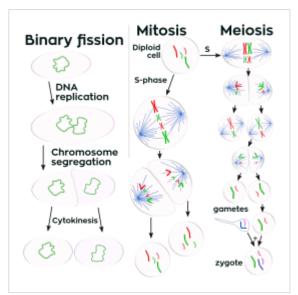
A <u>fimbria</u> (plural fimbriae also known as a <u>pilus</u>, plural pili) is a short, thin, hair-like filament found on the surface of bacteria. Fimbriae are formed of a protein called <u>pilin</u> (<u>antigenic</u>) and are responsible for the attachment of bacteria to specific receptors on human cells (<u>cell adhesion</u>). There are special types of pili involved in bacterial conjugation.

Cellular processes

Replication

Cell division involves a single cell (called a *mother cell*) dividing into two daughter cells. This leads to growth in multicellular organisms (the growth of tissue) and to procreation (vegetative reproduction) in unicellular organisms. Prokaryotic cells divide by binary fission, while eukaryotic cells usually undergo a process of nuclear division, called mitosis, followed by division of the cell, called cytokinesis. A diploid cell may also undergo meiosis to produce haploid cells, usually four. Haploid cells serve as gametes in multicellular organisms, fusing to form new diploid cells.

<u>DNA</u> replication, or the process of duplicating a cell's genome, [2] always happens when a cell divides through mitosis or binary fission. This occurs during the S phase of the cell cycle.



<u>Prokaryotes</u> divide by <u>binary fission</u>, while eukaryotes divide by mitosis or meiosis.

In meiosis, the DNA is replicated only once, while the cell divides twice. DNA replication only occurs before $\underline{\text{meiosis I}}$. DNA replication does not occur when the cells divide the second time, in $\underline{\text{meiosis II}}$. Replication, like all cellular activities, requires specialized proteins for carrying out the job. $\underline{^{[2]}}$

DNA repair

Cells of all organisms contain enzyme systems that scan their DNA for <u>damage</u> and carry out <u>repair processes</u> when it is detected. Diverse repair processes have evolved in organisms ranging from bacteria to humans. The widespread prevalence of these repair processes indicates the importance of maintaining cellular DNA in an undamaged state in order to avoid cell death or errors of replication due to damage that could lead to <u>mutation</u>. <u>E. coli</u> bacteria are a well-studied example of a cellular organism with diverse well-defined <u>DNA repair</u> processes. These include: <u>nucleotide excision repair</u>, <u>DNA mismatch repair</u>, <u>non-homologous end joining</u> of double-strand breaks, <u>recombinational repair</u> and light-dependent repair (<u>photoreactivation</u>). [15]

Growth and metabolism

Between successive cell divisions, cells grow through the functioning of cellular metabolism. Cell metabolism is the process by which individual cells process nutrient molecules. Metabolism has two distinct divisions: <u>catabolism</u>, in which the cell breaks down complex molecules to produce energy and

<u>reducing power</u>, and <u>anabolism</u>, in which the cell uses energy and reducing power to construct complex molecules and perform other biological functions.

Complex sugars can be broken down into simpler sugar molecules called <u>monosaccharides</u> such as <u>glucose</u>. Once inside the cell, glucose is broken down to make adenosine triphosphate (ATP), a molecule that possesses readily available energy, through two different pathways. In plant cells, <u>chloroplasts</u> create sugars by <u>photosynthesis</u>, using the energy of light to join molecules of water and carbon dioxide.

Protein synthesis

Cells are capable of synthesizing new proteins, which are essential for the modulation and maintenance of cellular activities. This process involves the formation of new protein molecules from <u>amino acid</u> building blocks based on information encoded in DNA/RNA. Protein synthesis generally consists of two major steps: transcription and translation.

Transcription is the process where genetic information in DNA is used to produce a complementary RNA strand. This RNA strand is then processed to give messenger RNA (mRNA), which is free to migrate through the cell. mRNA molecules bind to protein-RNA complexes called ribosomes located in the cytosol, where they are translated into polypeptide sequences. The ribosome mediates the formation of a polypeptide sequence based on the mRNA sequence. The mRNA sequence directly relates to the polypeptide sequence by binding to transfer RNA (tRNA) adapter molecules in binding pockets within the ribosome. The new polypeptide then folds into a functional three-dimensional protein molecule.

Motility

Unicellular organisms can move in order to find food or escape predators. Common mechanisms of motion include flagella and cilia.

In multicellular organisms, cells can move during processes such as wound healing, the immune response and <u>cancer metastasis</u>. For example, in wound healing in animals, white blood cells move to the wound site to kill the microorganisms that cause infection. Cell motility involves many receptors, crosslinking, bundling, binding, adhesion, motor and other proteins. The process is divided into three steps: protrusion of the leading edge of the cell, adhesion of the leading edge and de-adhesion at the cell body and rear, and cytoskeletal contraction to pull the cell forward. Each step is driven by physical forces generated by unique segments of the cytoskeleton. [17][16]

Navigation, control and communication

In August 2020, scientists described one way cells—in particular cells of a slime mold and mouse pancreatic cancer-derived cells—are able to <u>navigate</u> efficiently through a body and identify the best routes through complex mazes: generating gradients after breaking down diffused <u>chemoattractants</u> which enable them to sense upcoming maze junctions before reaching them, including around corners. [18][19][20]

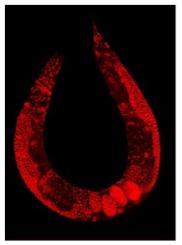
Multicellularity

Cell specialization/differentiation

Multicellular organisms are <u>organisms</u> that consist of more than one cell, in contrast to single-celled organisms. [21]

In complex multicellular organisms, cells specialize into different <u>cell</u> <u>types</u> that are adapted to particular functions. In mammals, major cell types include <u>skin cells</u>, <u>muscle cells</u>, <u>neurons</u>, <u>blood cells</u>, <u>fibroblasts</u>, <u>stem cells</u>, and others. Cell types differ both in appearance and function, yet are <u>genetically</u> identical. Cells are able to be of the same <u>genotype</u> but of different cell type due to the differential <u>expression</u> of the <u>genes</u> they contain.

Most distinct cell types arise from a single <u>totipotent</u> cell, called a <u>zygote</u>, that <u>differentiates</u> into hundreds of different cell types during the course of <u>development</u>. Differentiation of cells is driven by different environmental cues (such as cell–cell interaction) and intrinsic differences (such as those caused by the uneven distribution of molecules during division).



Staining of a <u>Caenorhabditis elegans</u> highlights the nuclei of its cells.

Origin of multicellularity

Multicellularity has evolved independently at least 25 times, [22] including in some prokaryotes, like cyanobacteria, myxobacteria, actinomycetes, or Methanosarcina. However, complex multicellular organisms evolved only in six eukaryotic groups: animals, fungi, brown algae, red algae, green algae, and plants. [23] It evolved repeatedly for plants (Chloroplastida), once or twice for animals, once for brown algae, and perhaps several times for fungi, slime molds, and red algae. [24] Multicellularity may have evolved from colonies of interdependent organisms, from cellularization, or from organisms in symbiotic relationships.

The first evidence of multicellularity is from <u>cyanobacteria</u>-like organisms that lived between 3 and 3.5 billion years ago. Other early fossils of multicellular organisms include the contested <u>Grypania</u> spiralis and the fossils of the black shales of the <u>Palaeoproterozoic</u> <u>Francevillian Group Fossil</u> B Formation in Gabon. [25]

The evolution of multicellularity from unicellular ancestors has been replicated in the laboratory, in evolution experiments using predation as the selective pressure. [22]

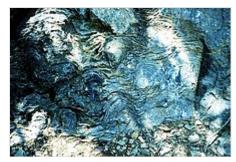
Origins

The origin of cells has to do with the origin of life, which began the history of life on Earth.

Origin of life

Small molecules needed for life may have been carried to Earth on meteorites, created at deep-sea vents, or synthesized by lightning in a reducing atmosphere. There is little experimental data defining what the first self-replicating forms were. RNA may have been the earliest self-replicating molecule, as it can both store genetic information and catalyze chemical reactions. [26]

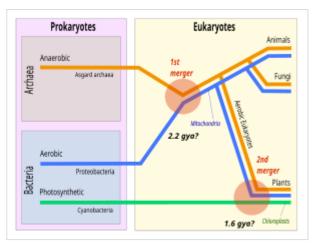
Cells emerged around 4 billion years ago. [27][28] The first cells were most likely heterotrophs. The early cell membranes were probably simpler and more permeable than modern ones, with only a single fatty acid chain per lipid. Lipids spontaneously form bilayered vesicles in water, and could have preceded RNA. [29][30]



Stromatolites are left behind by cyanobacteria, also called bluegreen algae. They are among the oldest fossils of life on Earth. This one-billion-year-old fossil is from Glacier National Park in the United States.

First eukaryotic cells

Eukaryotic cells were created some 2.2 billion years ago in a process called eukaryogenesis. This is widely agreed to have involved symbiogenesis, in which archaea and bacteria came together to create the first eukaryotic common ancestor. This cell had a new level of complexity and capability, with a nucleus^{[32][33]} and facultatively aerobic mitochondria.[31] It evolved some 2 billion years ago into a population of single-celled organisms that included the last eukaryotic common ancestor, gaining capabilities along the way, though the sequence of the steps involved has been disputed, and may not have started with symbiogenesis. It featured at least one centriole and cilium, sex (meiosis and syngamy), peroxisomes, and a dormant cyst with a cell wall of chitin and/or cellulose. [34][35] In turn, the last eukaryotic common ancestor gave rise to the eukaryotes' crown group, containing the ancestors of

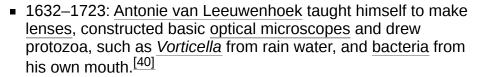


In the theory of <u>symbiogenesis</u>, a merger of an <u>archaean</u> and an aerobic bacterium created the eukaryotes, with aerobic <u>mitochondria</u>, some 2.2 billion years ago. A second merger, 1.6 billion years ago, added <u>chloroplasts</u>, creating the green plants.[31]

<u>animals</u>, <u>fungi</u>, <u>plants</u>, and a diverse range of single-celled organisms. [36][37] The plants were created around 1.6 billion years ago with a second episode of symbiogenesis that added <u>chloroplasts</u>, derived from cyanobacteria. [31]

History of research

In 1665, Robert Hooke examined a thin slice of cork under his microscope, and saw a structure of small enclosures. He wrote "I could exceeding plainly perceive it to be all perforated and porous, much like a Honey-comb, but that the pores of it were not regular". [38] To further support his theory, Matthias Schleiden and Theodor Schwann both also studied cells of both animal and plants. What they discovered were significant differences between the two types of cells. This put forth the idea that cells were not only fundamental to plants, but animals as well. [39]





Robert Hooke's drawing of cells in cork, 1665

- 1665: Robert Hooke discovered cells in cork, then in living plant tissue using an early compound microscope. He coined the term *cell* (from Latin cellula, meaning "small room" in his book Micrographia (1665). [42][40]
- 1839: Theodor Schwann [43] and Matthias Jakob Schleiden elucidated the principle that plants and animals are made of cells, concluding that cells are a common unit of structure and development, and thus founding the cell theory.
- 1855: <u>Rudolf Virchow</u> stated that new cells come from pre-existing cells by <u>cell division</u> (omnis cellula ex cellula).
- 1931: Ernst Ruska built the first transmission electron microscope (TEM) at the University of Berlin. [44] By 1935, he had built an EM with twice the resolution of a light microscope, revealing previously unresolvable organelles.
- 1981: <u>Lynn Margulis</u> published *Symbiosis in Cell Evolution* detailing how eukaryotic cells were created by symbiogenesis. [45]

See also

- Cell cortex
- Cell culture
- Cellular model
- Cytoneme
- Cytorrhysis
- Cytotoxicity
- Lipid raft
- List of distinct cell types in the adult human body
- Outline of cell biology

- Parakaryon myojinensis
- Plasmolysis
- Syncytium
- Tunneling nanotube
- Vault (organelle)

References

- 1. Black, Jacquelyn G. (2004). *Microbiology*. New York Chichester: Wiley. <u>ISBN</u> <u>978-0-471-42084-2</u>.
- 2. This article incorporates public domain material from "What Is a Cell?" (https://web.archive.org/web/20091208004309/https://web.archive.org/web/20130503014839/http://www.ncbi.nlm.nih.gov/About/primer/genetics_cell.html). Science Primer. NCBI. 30 March 2004. Archived from the original (https://web.archive.org/web/20130503014839/http://www.ncbi.nlm.nih.gov/About/primer/genetics_cell.html) on 2009-12-08. Retrieved 3 May 2013.
- 3. European Bioinformatics Institute, Karyn's Genomes: Borrelia burgdorferi (http://www.ebi.ac.uk/2can/genomes/bacteria/Borrelia_burgdorferi.html) Archived (https://web.archive.org/web/20130506040937/http://www.ebi.ac.uk/2can/genomes/bacteria/Borrelia_burgdorferi.html) 2013-05-06 at the Wayback Machine, part of 2can on the EBI-EMBL database. Retrieved 5 August 2012
- Satir, P.; Christensen, Søren T. (June 2008). "Structure and function of mammalian cilia" (htt ps://www.ncbi.nlm.nih.gov/pmc/articles/PMC2386530). Histochemistry and Cell Biology. 129 (6): 687–693. doi:10.1007/s00418-008-0416-9 (https://doi.org/10.1007%2Fs00418-008-0416-9). PMC 2386530 (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2386530). PMID 18365235 (https://pubmed.ncbi.nlm.nih.gov/18365235). 1432-119X.
- 5. Blair, D. F.; Dutcher, S. K. (October 1992). "Flagella in prokaryotes and lower eukaryotes". *Current Opinion in Genetics & Development*. **2** (5): 756–767. doi:10.1016/S0959-437X(05)80136-4 (https://doi.org/10.1016%2FS0959-437X%2805%2980136-4).

 PMID 1458024 (https://pubmed.ncbi.nlm.nih.gov/1458024).
- 6. Campbell Biology Concepts and Connections. Pearson Education. 2009. p. 320.
- 7. Bianconi, Eva; Piovesan, Allison; Facchin, Federica; Beraudi, Alina; Casadei, Raffaella; Frabetti, Flavia; Vitale, Lorenza; Pelleri, Maria Chiara; Tassani, Simone; Piva, Francesco; Perez-Amodio, Soledad (2013-11-01). "An estimation of the number of cells in the human body" (https://doi.org/10.3109%2F03014460.2013.807878). Annals of Human Biology. 40 (6): 463–471. doi:10.3109/03014460.2013.807878 (https://doi.org/10.3109%2F03014460.20 13.807878). hdl:11585/152451 (https://hdl.handle.net/11585%2F152451). ISSN 0301-4460 (https://search.worldcat.org/issn/0301-4460). PMID 23829164 (https://pubmed.ncbi.nlm.nih. gov/23829164). S2CID 16247166 (https://api.semanticscholar.org/CorpusID:16247166).
- Hatton, Ian A.; Galbraith, Eric D.; Merleau, Nono S. C.; Miettinen, Teemu P.; Smith, Benjamin McDonald; Shander, Jeffery A. (2023-09-26). "The human cell count and size distribution" (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC10523466). Proceedings of the National Academy of Sciences. 120 (39): e2303077120. Bibcode:2023PNAS..12003077H (https://ui.adsabs.harvard.edu/abs/2023PNAS..12003077H). doi:10.1073/pnas.2303077120 (https://doi.org/10.1073%2Fpnas.2303077120). ISSN 0027-8424 (https://search.worldcat.org/issn/0027-8424). PMC 10523466 (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC10523466). PMID 37722043 (https://pubmed.ncbi.nlm.nih.gov/37722043).
- 9. Guyton, Arthur C.; Hall, John E. (2016). *Guyton and Hall Textbook of Medical Physiology* (htt ps://books.google.com/books?id=3sWNCgAAQBAJ). Philadelphia: Elsevier Saunders. pp. 930–937. ISBN 978-1-4557-7005-2. OCLC 1027900365 (https://search.worldcat.org/ocl c/1027900365).

- 10. Michie, K. A.; Löwe, J. (2006). "Dynamic filaments of the bacterial cytoskeleton". *Annual Review of Biochemistry*. **75**: 467–492. doi:10.1146/annurev.biochem.75.103004.142452 (htt ps://doi.org/10.1146%2Fannurev.biochem.75.103004.142452). PMID 16756499 (https://pubmed.ncbi.nlm.nih.gov/16756499). S2CID 4550126 (https://api.semanticscholar.org/CorpusI D:4550126).
- 11. Ménétret, Jean-François; Schaletzky, Julia; Clemons, William M.; et al. (December 2007). "Ribosome binding of a single copy of the SecY complex: implications for protein translocation" (https://authors.library.caltech.edu/90566/2/1-s2.0-S1097276507008258-mmc 1.pdf) (PDF). Molecular Cell. 28 (6): 1083–1092. doi:10.1016/j.molcel.2007.10.034 (https://doi.org/10.1016%2Fj.molcel.2007.10.034). PMID 18158904 (https://pubmed.ncbi.nlm.nih.gov/18158904). Archived (https://web.archive.org/web/20210121115905/https://authors.library.caltech.edu/90566/2/1-s2.0-S1097276507008258-mmc1.pdf) (PDF) from the original on 2021-01-21. Retrieved 2020-09-01.
- 12. Sato, N. (2006). "Origin and Evolution of Plastids: Genomic View on the Unification and Diversity of Plastids". In Wise, R. R.; Hoober, J. K. (eds.). *The Structure and Function of Plastids*. Advances in Photosynthesis and Respiration. Vol. 23. Springer. pp. 75–102. doi:10.1007/978-1-4020-4061-0_4 (https://doi.org/10.1007%2F978-1-4020-4061-0_4). ISBN 978-1-4020-4060-3.
- 13. *Prokaryotes* (https://books.google.com/books?id=N2GU-DYKkk0C&q=Prokaryotic+india+ink &pg=PA87). Newnes. 1996. ISBN 978-0080984735. Archived (https://web.archive.org/web/2 0210414134256/https://books.google.com/books?id=N2GU-DYKkk0C&q=Prokaryotic+india +ink&pg=PA87) from the original on April 14, 2021. Retrieved November 9, 2020.
- 14. Campbell Biology Concepts and Connections. Pearson Education. 2009. p. 138.
- 15. Snustad, D. Peter; Simmons, Michael J. *Principles of Genetics* (5th ed.). DNA repair mechanisms, pp. 364–368.
- 16. Ananthakrishnan, R.; Ehrlicher, A. (June 2007). "The forces behind cell movement" (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1893118). International Journal of Biological Sciences. **3** (5). Biolsci.org: 303–317. doi:10.7150/ijbs.3.303 (https://doi.org/10.7150%2Fijbs.3.303). PMC 1893118 (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1893118). PMID 17589565 (https://pubmed.ncbi.nlm.nih.gov/17589565).
- 17. Alberts, Bruce (2002). *Molecular biology of the cell* (4th ed.). Garland Science. pp. 973–975. ISBN 0815340729.
- 18. Willingham, Emily. "Cells Solve an English Hedge Maze with the Same Skills They Use to Traverse the Body" (https://www.scientificamerican.com/article/cells-solve-an-english-hedge-maze-with-the-same-skills-they-use-to-traverse-the-body/). Scientific American. Archived (https://web.archive.org/web/20200904102655/https://www.scientificamerican.com/article/cells-solve-an-english-hedge-maze-with-the-same-skills-they-use-to-traverse-the-body/) from the original on 4 September 2020. Retrieved 7 September 2020.
- 19. "How cells can find their way through the human body" (https://phys.org/news/2020-08-cells human-body.html). phys.org. Archived (https://web.archive.org/web/20200903220400/https://phys.org/news/2020-08-cells-human-body.html) from the original on 3 September 2020. Retrieved 7 September 2020.
- 20. Tweedy, Luke; Thomason, Peter A.; Paschke, Peggy I.; Martin, Kirsty; Machesky, Laura M.; Zagnoni, Michele; Insall, Robert H. (August 2020). "Seeing around corners: Cells solve mazes and respond at a distance using attractant breakdown" (https://www.science.org/doi/10.1126/science.aay9792). Science. 369 (6507): eaay9792. doi:10.1126/science.aay9792 (https://doi.org/10.1126%2Fscience.aay9792). PMID 32855311 (https://pubmed.ncbi.nlm.nih.gov/32855311). S2CID 221342551 (https://api.semanticscholar.org/CorpusID:221342551). Archived (https://web.archive.org/web/20200912234645/https://science.sciencemag.org/content/369/6507/eaay9792) from the original on 2020-09-12. Retrieved 2020-09-13.
- 21. Becker, Wayne M.; et al. (2009). *The world of the cell*. Pearson Benjamin Cummings. p. 480. ISBN 978-0321554185.

- 22. Grosberg, R. K.; Strathmann, R. R. (2007). "The evolution of multicellularity: A minor major transition?" (https://web.archive.org/web/20160304121329/http://www-eve.ucdavis.edu/grosberg/Grosberg%20pdf%20papers/2007%20Grosberg%20%26%20Strathmann.AREES.pdf) (PDF). *Annu Rev Ecol Evol Syst.* **38**: 621–654. doi:10.1146/annurev.ecolsys.36.102403.114735 (https://doi.org/10.1146%2Fannurev.ecolsys.36.102403.114735). Archived from the original (http://www-eve.ucdavis.edu/grosberg/Grosberg%20pdf%20papers/2007%20Grosberg%20%26%20Strathmann.AREES.pdf) (PDF) on 2016-03-04. Retrieved 2013-12-23.
- 23. Popper, Zoë A.; Michel, Gurvan; Hervé, Cécile; et al. (2011). "Evolution and diversity of plant cell walls: from algae to flowering plants" (http://public.wsu.edu/~lange-m/Documnets/Teaching2011/Popper2011.pdf) (PDF). *Annual Review of Plant Biology*. **62**: 567–590. doi:10.1146/annurev-arplant-042110-103809 (https://doi.org/10.1146%2Fannurev-arplant-042110-103809). hdl:10379/6762 (https://hdl.handle.net/10379%2F6762). PMID 21351878 (https://pubmed.ncbi.nlm.nih.gov/21351878). S2CID 11961888 (https://api.semanticscholar.org/CorpusID:11961888). Archived (https://web.archive.org/web/20160729224035/http://public.wsu.edu/~lange-m/Documnets/Teaching2011/Popper2011.pdf) (PDF) from the original on 2016-07-29. Retrieved 2013-12-23.
- 24. Bonner, John Tyler (1998). "The Origins of Multicellularity" (https://web.archive.org/web/201 20308175112/http://courses.cit.cornell.edu/biog1101/outlines/Bonner%20-Origin%20of%20 Multicellularity.pdf) (PDF). Integrative Biology. 1 (1): 27–36. doi:10.1002/(SICI)1520-6602(1998)1:1<27::AID-INBI4>3.0.CO;2-6 (https://doi.org/10.1002%2F%28SICI%291520-6602%281998%291%3A1%3C27%3A%3AAID-INBI4%3E3.0.CO%3B2-6). ISSN 1093-4391 (https://search.worldcat.org/issn/1093-4391). Archived from the original (http://courses.cit.cornell.edu/biog1101/outlines/Bonner%20-Origin%20of%20Multicellularity.pdf) (PDF, 0.2 MB) on March 8, 2012.
- 25. Albani, Abderrazak El; Bengtson, Stefan; Canfield, Donald E.; et al. (July 2010). "Large colonial organisms with coordinated growth in oxygenated environments 2.1 Gyr ago". Nature. 466 (7302): 100–104. Bibcode:2010Natur.466..100A (https://ui.adsabs.harvard.edu/abs/2010Natur.466..100A). doi:10.1038/nature09166 (https://doi.org/10.1038%2Fnature09166). PMID 20596019 (https://pubmed.ncbi.nlm.nih.gov/20596019). S2CID 4331375 (https://api.semanticscholar.org/CorpusID:4331375).
- 26. Orgel, L. E. (December 1998). "The origin of life--a review of facts and speculations". *Trends in Biochemical Sciences*. **23** (12): 491–495. doi:10.1016/S0968-0004(98)01300-0 (https://doi.org/10.1016%2FS0968-0004%2898%2901300-0). PMID 9868373 (https://pubmed.ncbi.nlm.nih.gov/9868373).
- 27. Dodd, Matthew S.; Papineau, Dominic; Grenne, Tor; et al. (1 March 2017). "Evidence for early life in Earth's oldest hydrothermal vent precipitates" (http://eprints.whiterose.ac.uk/112 179/). Nature. 543 (7643): 60–64. Bibcode:2017Natur.543...60D (https://ui.adsabs.harvard.edu/abs/2017Natur.543...60D). doi:10.1038/nature21377 (https://doi.org/10.1038%2Fnature21377). PMID 28252057 (https://pubmed.ncbi.nlm.nih.gov/28252057). Archived (https://web.archive.org/web/20170908201821/http://eprints.whiterose.ac.uk/112179/) from the original on 8 September 2017. Retrieved 2 March 2017.
- 28. Betts, Holly C.; Puttick, Mark N.; Clark, James W.; Williams, Tom A.; Donoghue, Philip C. J.; Pisani, Davide (20 August 2018). "Integrated genomic and fossil evidence illuminates life's early evolution and eukaryote origin" (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC615291 0). Nature Ecology & Evolution. 2 (10): 1556–1562. Bibcode:2018NatEE...2.1556B (https://ui.adsabs.harvard.edu/abs/2018NatEE...2.1556B). doi:10.1038/s41559-018-0644-x (https://doi.org/10.1038%2Fs41559-018-0644-x). PMC 6152910 (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6152910). PMID 30127539 (https://pubmed.ncbi.nlm.nih.gov/30127539).
- 29. Griffiths, G. (December 2007). "Cell evolution and the problem of membrane topology" (http s://doi.org/10.1038%2Fnrm2287). Nature Reviews. Molecular Cell Biology. 8 (12): 1018—1024. doi:10.1038/nrm2287 (https://doi.org/10.1038%2Fnrm2287). PMID 17971839 (https://pubmed.ncbi.nlm.nih.gov/17971839). S2CID 31072778 (https://api.semanticscholar.org/CorpusID:31072778).

- 30. "First cells may have emerged because building blocks of proteins stabilized membranes" (https://www.sciencedaily.com/releases/2019/08/190812155502.htm). ScienceDaily. Archived (https://web.archive.org/web/20210918102211/https://www.sciencedaily.com/releases/2019/08/190812155502.htm) from the original on 2021-09-18. Retrieved 2021-09-18.
- 31. Latorre, A.; Durban, A; Moya, A.; Pereto, J. (2011). "The role of symbiosis in eukaryotic evolution" (https://books.google.com/books?id=m3oFebknu1cC&pg=PA326). In Gargaud, Muriel; López-Garcìa, Purificacion; Martin, H. (eds.). *Origins and Evolution of Life: An astrobiological perspective*. Cambridge: Cambridge University Press. pp. 326–339. ISBN 978-0-521-76131-4. Archived (https://web.archive.org/web/20190324055723/https://books.google.com/books?id=m3oFebknu1cC&pg=PA326) from the original on 24 March 2019. Retrieved 27 August 2017.
- 32. McGrath, Casey (31 May 2022). "Highlight: Unraveling the Origins of LUCA and LECA on the Tree of Life" (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC9168435). Genome Biology and Evolution. **14** (6): evac072. doi:10.1093/gbe/evac072 (https://doi.org/10.1093%2Fgbe% 2Fevac072). PMC 9168435 (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC9168435).
- 33. Weiss, Madeline C.; Sousa, F. L.; Mrnjavac, N.; et al. (2016). "The physiology and habitat of the last universal common ancestor" (http://complexityexplorer.s3.amazonaws.com/supplemental_materials/3.6+Early+Metabolisms/Weiss_et_al_Nat_Microbiol_2016.pdf) (PDF).

 Nature Microbiology. 1 (9): 16116. doi:10.1038/nmicrobiol.2016.116 (https://doi.org/10.1038/92Fnmicrobiol.2016.116). PMID 27562259 (https://pubmed.ncbi.nlm.nih.gov/27562259). S2CID 2997255 (https://api.semanticscholar.org/CorpusID:2997255).
- 34. Leander, B. S. (May 2020). "Predatory protists" (https://doi.org/10.1016%2Fj.cub.2020.03.05 2). Current Biology. **30** (10): R510 R516. doi:10.1016/j.cub.2020.03.052 (https://doi.org/1 0.1016%2Fj.cub.2020.03.052). PMID 32428491 (https://pubmed.ncbi.nlm.nih.gov/3242849 1). S2CID 218710816 (https://api.semanticscholar.org/CorpusID:218710816).
- 35. Strassert, Jürgen F. H.; Irisarri, Iker; Williams, Tom A.; Burki, Fabien (25 March 2021). "A molecular timescale for eukaryote evolution with implications for the origin of red algalderived plastids" (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7994803). Nature Communications. 12 (1): 1879. Bibcode:2021NatCo..12.1879S (https://ui.adsabs.harvard.edu/abs/2021NatCo..12.1879S). doi:10.1038/s41467-021-22044-z (https://doi.org/10.1038%2 Fs41467-021-22044-z). PMC 7994803 (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7994803). PMID 33767194 (https://pubmed.ncbi.nlm.nih.gov/33767194).
- 36. Gabaldón, T. (October 2021). "Origin and Early Evolution of the Eukaryotic Cell". *Annual Review of Microbiology.* **75** (1): 631–647. doi:10.1146/annurev-micro-090817-062213 (https://doi.org/10.1146%2Fannurev-micro-090817-062213). PMID 34343017 (https://pubmed.ncbi.nlm.nih.gov/34343017). S2CID 236916203 (https://api.semanticscholar.org/CorpusID:236916203).
- 37. Woese, C.R.; Kandler, Otto; Wheelis, Mark L. (June 1990). "Towards a natural system of organisms: proposal for the domains Archaea, Bacteria, and Eucarya" (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC54159). Proceedings of the National Academy of Sciences of the United States of America. 87 (12): 4576–4579. Bibcode:1990PNAS...87.4576W (https://ui.adsabs.harvard.edu/abs/1990PNAS...87.4576W). doi:10.1073/pnas.87.12.4576 (https://oi.org/10.1073%2Fpnas.87.12.4576). PMC 54159 (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC54159). PMID 2112744 (https://pubmed.ncbi.nlm.nih.gov/2112744).
- 38. <u>Hooke, Robert (1665)</u>. <u>"Observation 18" (https://en.wikisource.org/wiki/Micrographia/Chapter_18)</u>. *Micrographia*.
- 39. Maton, Anthea (1997). *Cells Building Blocks of Life* (https://archive.org/details/cellsbuildingbl o00mato). New Jersey: Prentice Hall. pp. 44-45 The Cell Theory. ISBN 978-0134234762.
- 40. Gest, H. (2004). "The discovery of microorganisms by Robert Hooke and Antoni Van Leeuwenhoek, fellows of the Royal Society". *Notes and Records of the Royal Society of London.* **58** (2): 187–201. doi:10.1098/rsnr.2004.0055 (https://doi.org/10.1098%2Frsnr.2004. 0055). PMID 15209075 (https://pubmed.ncbi.nlm.nih.gov/15209075). S2CID 8297229 (https://api.semanticscholar.org/CorpusID:8297229).

- 41. "The Origins Of The Word 'Cell' " (https://www.npr.org/templates/story/story.php?story Id=129934828&t=1628175572746). *National Public Radio*. September 17, 2010. Archived (https://web.archive.org/web/20210805150111/https://www.npr.org/templates/story/story.php?storyId=129934828&t=1628175572746) from the original on 2021-08-05. Retrieved 2021-08-05.
 - "cellŭla" (http://www.perseus.tufts.edu/hopper/text?doc=Perseus:text:1999.04.0059:e ntry=cellula). <u>A Latin Dictionary</u>. Charlton T. Lewis and Charles Short. 1879. ISBN 978-1999855789. Archived (https://web.archive.org/web/20210807122358/htt p://www.perseus.tufts.edu/hopper/text?doc=Perseus%3Atext%3A1999.04.0059%3A entry%3Dcellula) from the original on 7 August 2021. Retrieved 5 August 2021.
- 42. Hooke, Robert (1665). *Micrographia: ...* (https://archive.org/stream/mobot31753000817897# page/113/mode/2up) London: Royal Society of London. p. 113. "... I could exceedingly plainly perceive it to be all perforated and porous, much like a Honey-comb, but that the pores of it were not regular [...] these pores, or cells, [...] were indeed the first microscopical pores I ever saw, and perhaps, that were ever seen, for I had not met with any Writer or Person, that had made any mention of them before this ..." Hooke describing his observations on a thin slice of cork. See also: Robert Hooke (http://www.ucmp.berkeley.edu/history/hooke.html) Archived (https://web.archive.org/web/19970606013455/http://www.ucmp.berkeley.edu/history/hooke.html) 1997-06-06 at the Wayback Machine
- 43. Schwann, Theodor (1839). *Mikroskopische Untersuchungen über die Uebereinstimmung in der Struktur und dem Wachsthum der Thiere und Pflanzen* (http://www.deutschestextarchiv.de/book/show/schwann_mikroskopische_1839). Berlin: Sander.
- 44. Ernst Ruska (January 1980). *The Early Development of Electron Lenses and Electron Microscopy*. Applied Optics. Vol. 25. Translated by T. Mulvey. p. 820.

 Bibcode:1986ApOpt..25..820R (https://ui.adsabs.harvard.edu/abs/1986ApOpt..25..820R).

 doi:10.1364/AO.25.000820 (https://doi.org/10.1364%2FAO.25.000820). ISBN 978-3-7776-0364-3.
- 45. Cornish-Bowden, Athel (7 December 2017). "Lynn Margulis and the origin of the eukaryotes" (https://www.sciencedirect.com/science/article/pii/S0022519317304459). Journal of Theoretical Biology. The origin of mitosing cells: 50th anniversary of a classic paper by Lynn Sagan (Margulis). 434: 1. Bibcode:2017JThBi.434....1C (https://ui.adsabs.harvard.edu/abs/2017JThBi.434....1C). doi:10.1016/j.jtbi.2017.09.027 (https://doi.org/10.1016%2Fj.jtbi.2017.09.027). PMID 28992902 (https://pubmed.ncbi.nlm.nih.gov/28992902).

Further reading

- Alberts, Bruce; Johnson, Alexander; Lewis, Julian; Morgan, David; Raff, Martin; Roberts, Keith; Walter, Peter (2015). Molecular Biology of the Cell (6th ed.). Garland Science. p. 2. ISBN 978-0815344322.
- Alberts, B.; et al. (2014). Molecular Biology of the Cell (https://web.archive.org/web/2014071 4210549/http://www.garlandscience.com/product/isbn/9780815344322) (6th ed.). Garland. ISBN 978-0815344322. Archived from the original (http://www.garlandscience.com/product/isbn/9780815344322) on 2014-07-14. Retrieved 2016-07-06.; The fourth edition is freely available (https://www.ncbi.nlm.nih.gov/books/bv.fcgi?rid=mboc4.TOC&depth=2) Archived (https://web.archive.org/web/20091011113848/http://www.ncbi.nlm.nih.gov/books/bv.fcgi?rid=mboc4.TOC&depth=2) 2009-10-11 at the Wayback Machine from National Center for Biotechnology Information Bookshelf.
- Lodish, Harvey; et al. (2004). *Molecular Cell Biology* (https://archive.org/details/molecularcel lbio00harv) (5th ed.). New York: WH Freeman. ISBN 978-0716743668.
- Cooper, G. M. (2000). The cell: a molecular approach (https://www.ncbi.nlm.nih.gov/books/bv.fcgi?rid=cooper.TOC&depth=2) (2nd ed.). Washington, D.C.: ASM Press. ISBN 978-0878931026. Archived (https://web.archive.org/web/20090630030426/http://www.ncbi.nlm.ni

<u>h.gov/books/bv.fcgi?rid=cooper.TOC&depth=2)</u> from the original on 2009-06-30. Retrieved 2017-08-30.

External links

- MBInfo Descriptions on Cellular Functions and Processes (http://www.mechanobio.info/)
- Inside the Cell (http://publications.nigms.nih.gov/insidethecell/) Archived (https://web.archive.org/web/20170720164847/http://publications.nigms.nih.gov/insidethecell/) 2017-07-20 at the Wayback Machine a science education booklet by National Institutes of Health, in PDF and ePub.
- Cell Biology (http://www.biology.arizona.edu/cell_bio/cell_bio.html) in "The Biology Project" of University of Arizona.
- Centre of the Cell online (http://www.centreofthecell.org/)
- The Image & Video Library of The American Society for Cell Biology (http://cellimages.ascb.org/) Archived (https://web.archive.org/web/20110610012208/http://cellimages.ascb.org/) 2011-06-10 at the Wayback Machine, a collection of peer-reviewed still images, video clips and digital books that illustrate the structure, function and biology of the cell.
- WormWeb.org: Interactive Visualization of the *C. elegans* Cell lineage (http://wormweb.org/c elllineage) Visualize the entire cell lineage tree of the nematode *C. elegans*

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