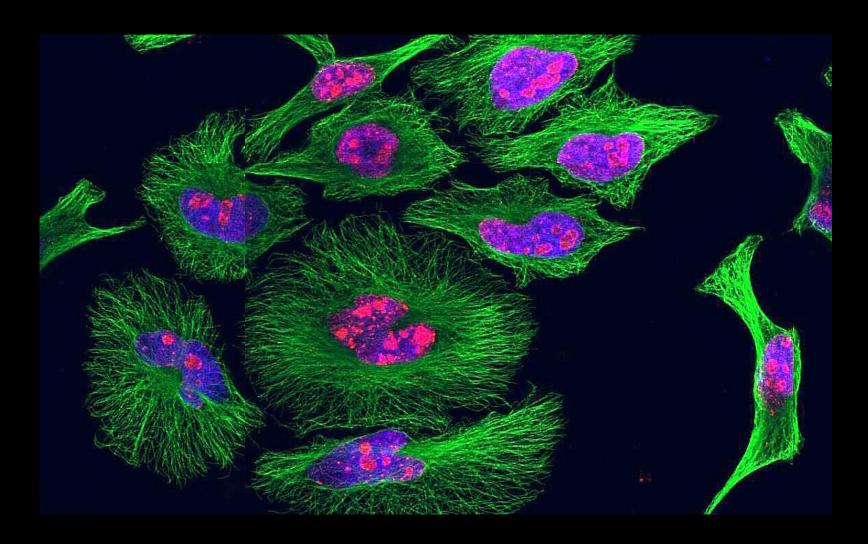
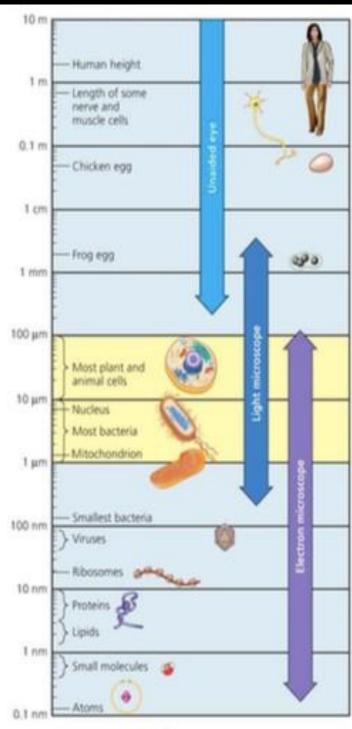
Biologi Sel Umi Baroroh, S.Si., M.Biotek.





TOUR THE CELL

seberapa kecilkah sel?

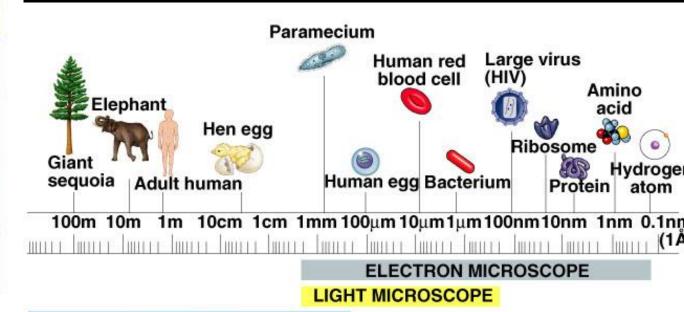


1 centimeter (cm) = 10^{-2} meter (m) = 0.4 inch 1 millimeter (mm) = 10^{-3} m

I micrometer (µm) = 10-3 mm = 10-6 m

1 nanometer (nm) = 10⁻³ µm = 10⁻⁹ m

Visualisasi sel



HUMAN EYE



berapa banyak sel dalam tubuh kita?

KONSEP KUNCI

- untuk mempelajari sel, scientist menggunakan mikroskop & alat biokimia
- 2. sel eukariot memiliki membran internal yang mengakomodasi fungsinya
- 3. instruksi genetik eukariot sel terletak di inti sel dan dibawa keluar oleh ribosom
- 4. sistem endomembran meregulasi jalur protein dan menampilkan fungsi metabolisme dalam sel
- 5. mitokondria dan kloroplas mengubah energi dari satu bentuk ke yang lainnya
- 6. sitoskeleton adalah jaringan serat yang mengorganisasi struktur dan aktivitas sel
- 7. komponen ekstraseluler dan koneksi antara sel membantu koordinasi aktivitas seluler



1. UNTUK MEMPELAJARI SEL, ILMUAN MENGGUNAKAN MIKROSKOP DAN ALAT BIOKIMIA

T Figure 6.3 Research Method

Light Microscopy

TECHNIQUE

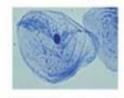
(a) Brightfield (unstained specimen). Passes light directly through specimen. Unless cell is naturally pigmented or artificially stained, image has little contrast. [Parts (a)-(d) show a human cheek epithelial cell.]



RESULTS

50 µm





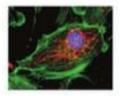
(c) Phase-contrast. Enhances contrast in unstained cells by amplifying variations in density within specimen; especially useful for examining living, unpigmented cells.



(d) Differential-interferencecontrast (Nomarski). Like phase-contrast microscopy, uses optical modifications to exaggerate differences in density, making the image appear almost 3-D.

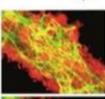


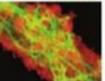
(e) Fluorescence. Shows the locations of specific molecules in the cell by tagging the molecules with fluorescent dyes or antibodies. These fluorescent substances absorbultraviolet radiation and emit visible light, as shown here in a cell from an artery.



50 µm

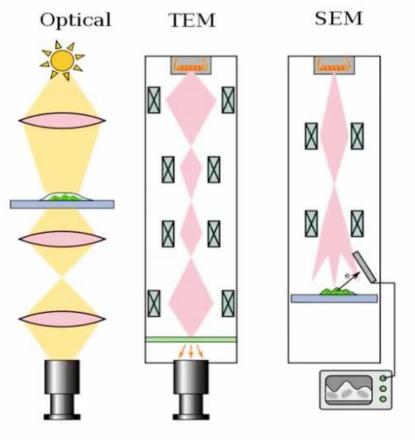
(f) Confocal. A fluorescent "optical. sectioning" technique that uses a pinhole aperture to eliminate out-of-focus light from a thick sample, creating a single plane of fluorescence in the image. By capturing sharp images at many different planes, a 3-D reconstruction can be created. At the right are confocal (top) and standard fluorescent micrographs of stained nervous tissue, where nerve cells are green, support cells are red, and regions of overlap are yellow. The standard image is blurry because the out-of-focus light is not excluded.





50 µm

MIKROSKOP

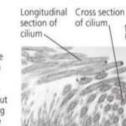


v Figure 6.4 Research Method

Electron Microscopy

TECHNIQUE

(a) Scanning electron microscopy (SEM). Micrographs taken with a scanning electron microscope show a 3-D image of the surface of a specimen. This SEM shows the surface of a cell from a rabbit trachea (windpipe) covered with motile organelles called cilia. Beating of the cilia helps move inhaled debris upward toward the throat.



RESULTS

(b) Transmission electron microscopy (TEM). A

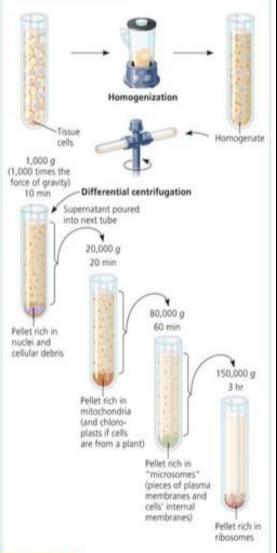
transmission electron microscope profiles a thin section of a specimen. Here we see a section through a tracheal cell, revealing its ultrastructure. In preparing the TEM, some cilia were cut along their lengths, creating longitudinal sections, while other cilia were cut straight across, creating cross sections.

▼ Figure 6.5 Research Method

Cell Fractionation

APPLICATION Cell fractionation is used to isolate (fractionate) cell components based on size and density.

TECHNIQUE First, cells are homogenized in a blender to break them up. The resulting mixture (cell homogenate) is then centrifuged at various speeds and durations to fractionate the cell components, forming a series of pellets, overlaid by the remaining homogenate (supernatant).

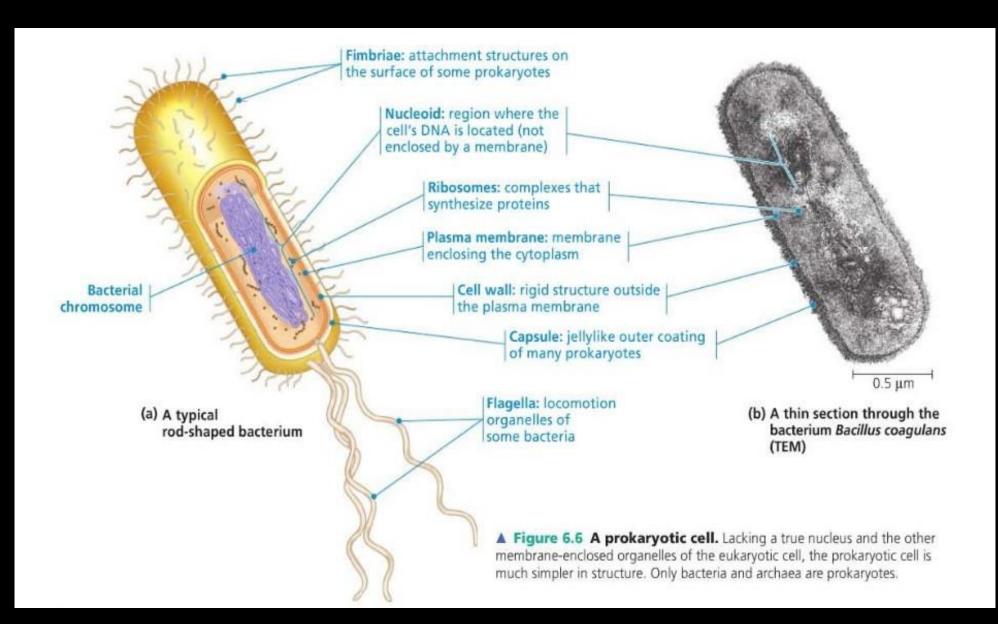


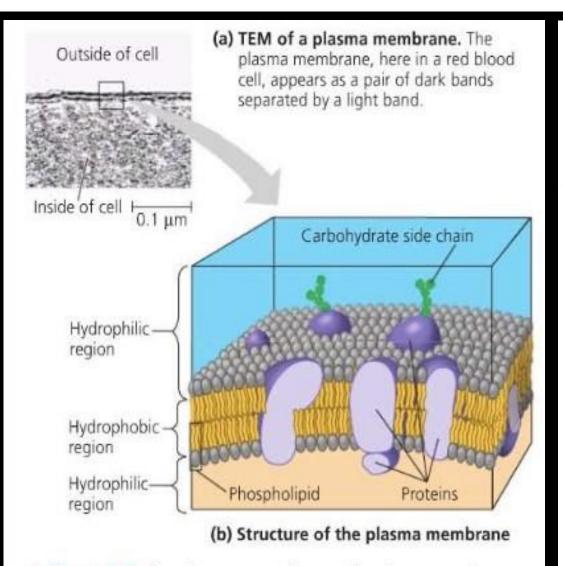
RESULTS In early experiments, researchers used microscopy to identify the organelles in each pellet and biochemical methods to determine their metabolic functions. These identifications established a baseline for this method, enabling today's researchers to know which cell fraction they should collect in order to isolate and study particular organelles.

FRAKSINASI SEL

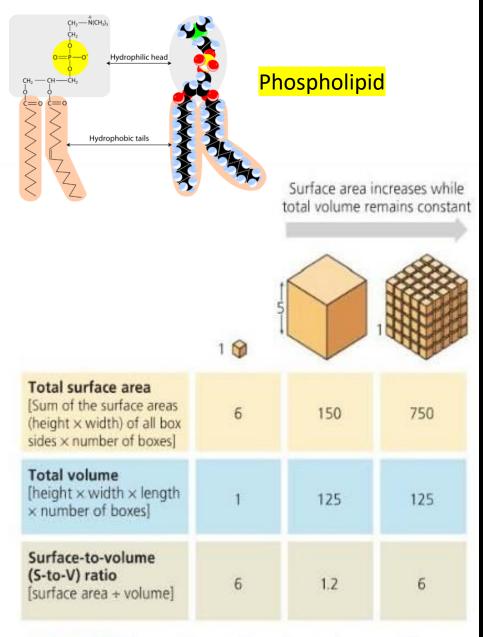
2. SEL EUKARIOT MEMILIKI MEMBRAN INTERNAL YANG MEWADAHI FUNGSINYA

Sel Prokariot





▲ Figure 6.7 The plasma membrane. The plasma membrane and the membranes of organelles consist of a double layer (bilayer) of phospholipids with various proteins attached to or embedded in it. In the interior of a membrane, the phospholipid tails are hydrophobic, as are the interior portions of membrane proteins in contact with them. The phospholipid heads are hydrophilic, as are proteins or parts of proteins in contact with the aqueous solution on either side of the membrane. (Channels through certain proteins are also hydrophilic.) Carbohydrate side chains are found only attached to proteins or lipids on the outer surface of the plasma membrane.



▲ Figure 6.8 Geometric relationships between surface area and volume. In this diagram, cells are represented as boxes. Using arbitrary units of length, we can calculate the cell's surface area (in square units, or units²), volume (in cubic units, or units³), and ratio of surface area to volume. A high surface-to-volume ratio facilitates the exchange of materials between a cell and its environment.

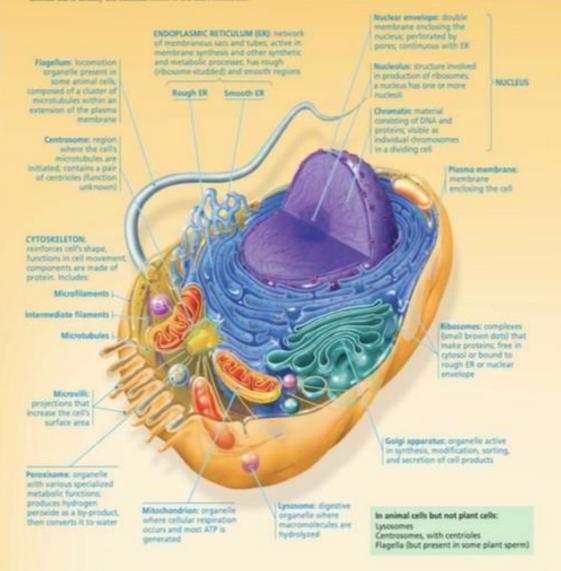
Pemandangan Sel Eukariot

Exploring Animal and Plant Cells

Animal Cell

This drawing of a generalized animal cell incorporates the most common structures of animal cells (no cell actually looks just like this). As shown by this cutaway view, the cell has a variety of components, including organelles ("little organs"), which are bounded by membranes. The most prominent organelle in an animal cell is usually the nucleus. Most of the cell's metabolic

activities occur in the cytoplasm, the entire region between the nucleus and the plasma membrane. The cytoplasm contains many organelles and other cell components suspended in a semifluid medium, the cytosol. Pervading much of the cytoplasm is a labyrinth of membranes called the endoplasmic reticulum (ER).



Plant Cell

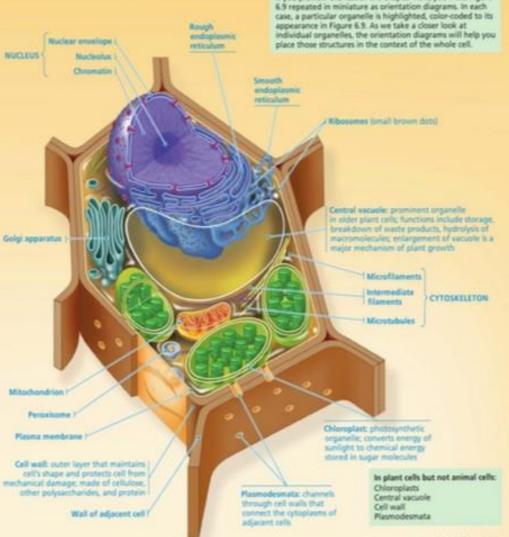
This drawing of a generalized plant cell reveals the similarities and differences between an animal cell and a plant cell. In addition to most of the features seen in an animal cell, a plant cell has organelles called plastid in the chloroplast, which carries out photosynthesis. Many plant cells have a large central vaccole; some may have one or more smaller vacuoles. Among other tasks, vaccoles carry out functions

performed by hysosomes in animal cells. Outside a plant cell's plasma membrane is a thick cell wall, perfocated by channels called plasmodesmata.



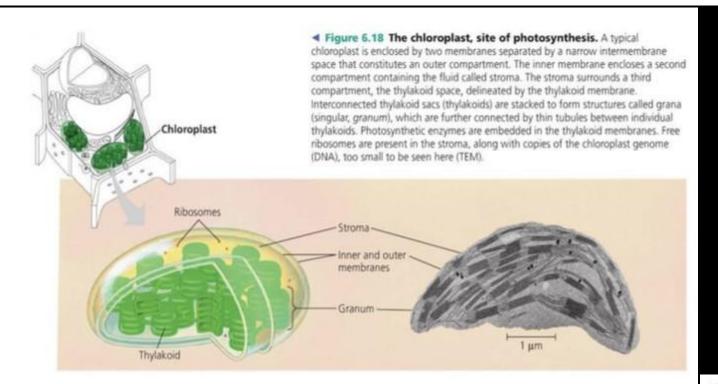
BioFlix Vist the Study Area at www.masteringbio.com for the BioFlix 3-O Ammations called Tour of an Ammal Cell and Tour of a Plant Cell.

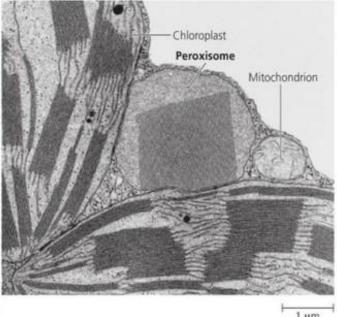
If you preview the rest of the chapter nov., you'll see Figure



Let's Tour The Cell (3-6)

| | Cell Component | Structure | Function |
|---|--|--|---|
| Concept 6.3 The eukaryotic cell's genetic instructions are housed in the nucleus and carried out by the ribosomes (pp. 102–104) MEDIA Activity Role of the Nucleus and Ribosomes in Protein Synthesis | Nucleus (ER) | Sumounded by nuclear envelope (double membrane) perforated by nuclear pores. The nuclear envelope is continuous with the endoplasmic reticulum (ER). | Houses chromosomes, made of chromatin (DNA, the genetic material, and proteins); contains nucleoil, where ribosomal subunits are made. Pores regulate entry and exit of materials. |
| | Ribosome 🚇 | Two subunits made of ribosomal RNA and proteins; can be free in cytosol or bound to ER | Protein synthesis |
| Concept 6.4 The endomembrane system regulates protein traffic and performs metabolic functions in the cell (pp. 104–108) MEDIA Activity The Endomembrane System | Endoplasmic reticulum (Nuclear envelope) | Extensive network of membrane-bounded tubules and sacs; membrane separates lumen from cytosol; continuous with the nuclear envelope | Smooth ER: synthesis of lipids, metabolism of carbohy- drates, Ca ^{2*} storage, detoxifica- tion of drugs and poisons Rough ER: Aids in synthesis of secretory and other proteins from bound ribosomes; adds carbohydrates to glycoproteins; produces new membrane |
| | Golgi apparatus | Stacks of flattened membranous sacs; has polarity (cis and trans faces) | Modification of proteins, carbo- hydrates on proteins, and phos- pholipids; synthesis of many poly- saccharides; sorting of Golgi products, which are then released in vesicles |
| | tysosome | Membranous sac of hydrolytic enzymes (in animal cells) | Breakdown of ingested substances, cell macromolecules, and damaged organelles for recycling |
| | Vacuole | Large membrane-bounded vesicle in plants | Digestion, storage, waste disposal, water balance, cell growth, and protection |
| Concept 6.5 Mitochondria and chloro- plasts change energy from one form to another (pp. 109-111) | Mitochondrion | Bounded by double membrane; inner membrane has infoldings (cristae) | Cellular respiration |
| MEDIA Activity Build a Chloroplant and a Mitochondrion | Chloroplast | Typically two membranes around fluid stroma, which contains membranous thylakoids stacked into grana (in plants) | Photosynthesis |
| | Peroxisome | Specialized metabolic compart- ment bounded by a single membrane | Contains enzymes that transfer hydrogen to water, producing hydrogen peroxide (H ₂ O ₂) as a by-product, which is converted to water by other enzymes in the peroxisome |





▲ Figure 6.19 A peroxisome. Peroxisomes are roughly spherical and often have a granular or crystalline core that is thought to be a dense collection of enzyme molecules. This peroxisome is in a leaf cell (TEM), Notice its proximity to two chloroplasts and a mitochondrion. These organelles cooperate with peroxisomes in certain metabolic functions.

Cytoskeleton

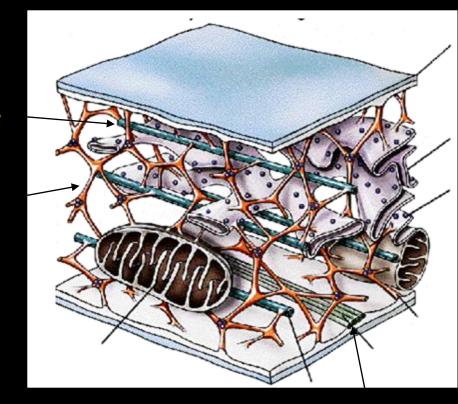
Network of protein fibers supporting cell shape and

anchoring organelles

Actin filaments cell movement

Microtubules

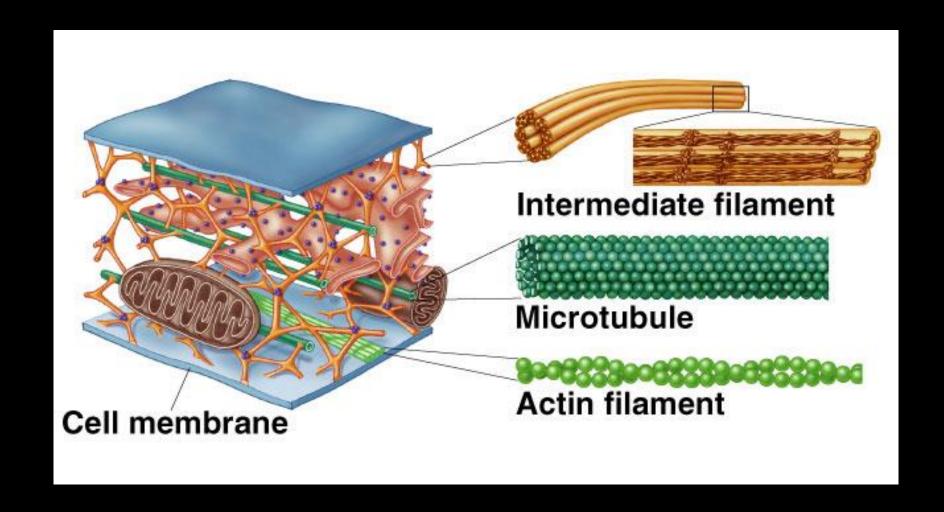
Microtubules
Hollow tubes
Facilitate cell movement
Centrioles – barrel shaped
organelles occur in pairs –
help assemble animal cell's
microtubules



Actin

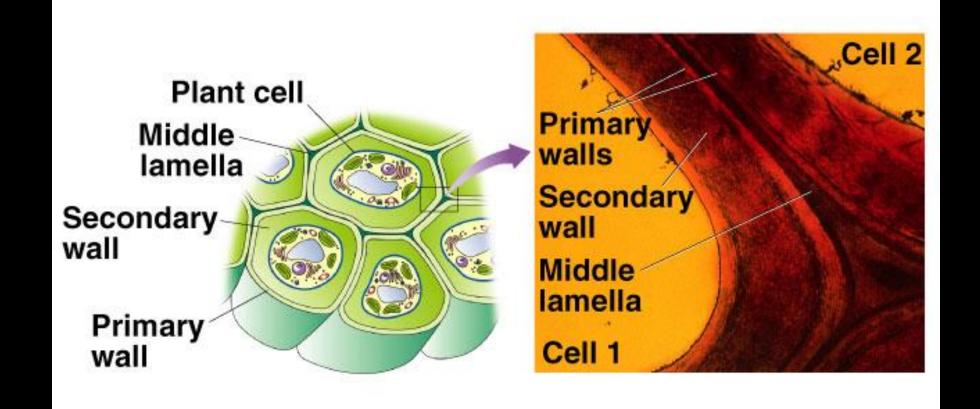
Intermediate filaments
Stable - don't break down

Cytoskeleton

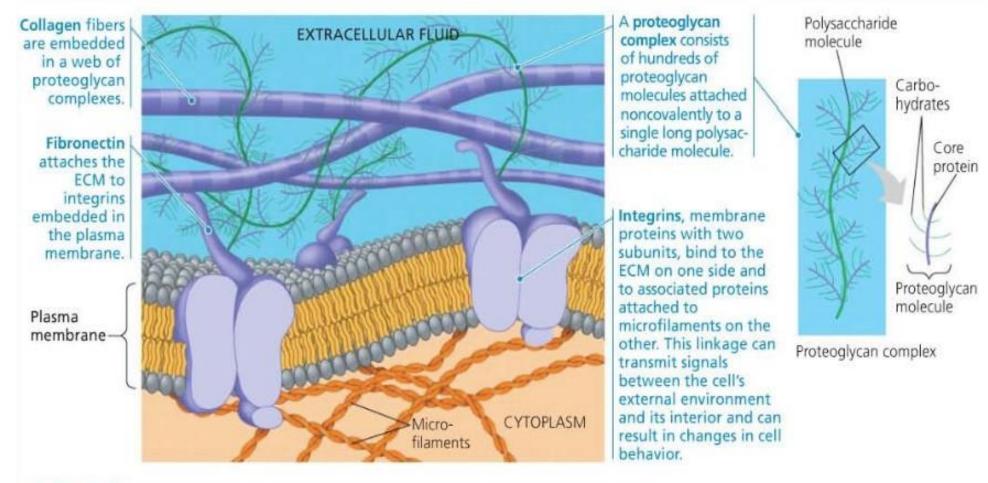


7. KOMPONEN EKSTRASELULAR & KONEKSI ANTARA SEL MEMBANTU KOORDINAT AKTIVITAS SEL

Dinding Sel Tumbuhan



Matriks Ekstraselular Sel Hewan



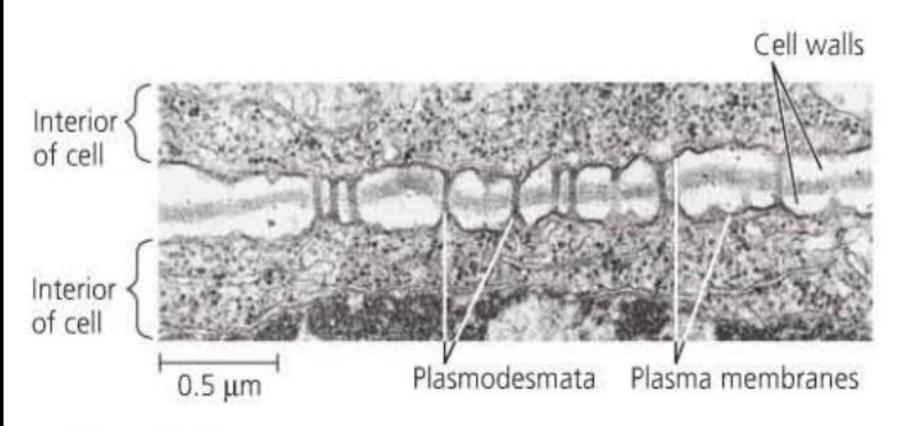
▲ Figure 6.30 Extracellular matrix (ECM) of an animal cell. The molecular composition and structure of the ECM varies from one cell type to another. In this example, three different types of glycoproteins are present: proteoglycans, collagen, and fibronectin.

Copyright © The McGraw-Hill Companies, Inc. Permission required for reproduction or display.

Table 5.3 A Comparison of Prokaryotic, Animal, and Plant Cells

| | a | 20 | ~~ | | | |
|---------------------|----------------------|----------------|-------------------------------|---|--|--|
| | Prokaryote | 3 | Animal | Plant | | |
| Exterior Structures | | | | | | |
| Cell wall | Present (protein-po | olysaccharide) | Absent | Present (cellulose) | | |
| Cell membrane | Present | | Present | Present | | |
| Flagella/cilia | May be present (sin | gle strand) | May be present | Absent except in sperm of a few species | | |
| Interior Structures | | | | | | |
| ER | Absent | | Usually present | Usually present | | |
| Ribosomes | Present | | Present | Present | | |
| Microtubules | Absent | | Present | Present | | |
| Centrioles | Absent | | Present | Absent | | |
| Golgi apparatus | Absent | | Present | Present | | |
| Nucleus | Absent | | Present | Present | | |
| Mitochondria | Absent | | Present | Present | | |
| Chloroplasts | Absent | | Absent | Present | | |
| Chromosomes | A single circle of D | NA | Multiple; DNA-protein complex | Multiple; DNA-protein complex | | |
| Lysosomes | Absent | | Usually present | Present | | |
| Vacuoles | Absent | | Absent or small | Usually a large single vacuole | | |

Plasmodesmata pada Sel Tumbuhan

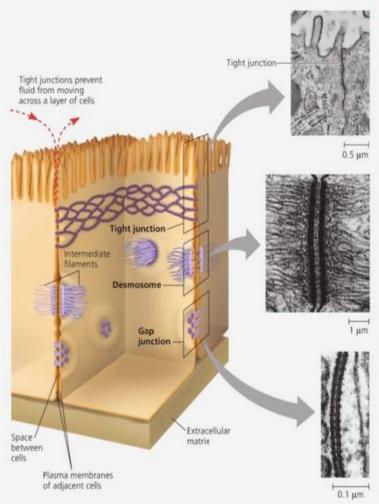


▲ Figure 6.31 Plasmodesmata between plant cells. The cytoplasm of one plant cell is continuous with the cytoplasm of its neighbors via plasmodesmata, channels through the cell walls (TEM).

Tight Junction, Desmosomes, Gap Juction pada Sel Hewan

▼ Figure 6.32

Exploring Intercellular Junctions in Animal Tissues



Tight Junctions

At tight junctions, the plasma membranes of neighboring cells are very tightly pressed against each other, bound together by specific proteins (purple). Forming continuous seals around the cells, tight junctions prevent leakage of extracellular fluid across a layer of epithelial cells. For example, tight junctions between skin cells make us watertight by preventing leakage between cells in our sweat glands.

Desmosomes

Desmosomes (also called anchoring junctions) function like rivets, fastening cells together into strong sheets. Intermediate filaments made of sturdy keratin proteins anchor desmosomes in the cytoplasm. Desmosomes attach muscle cells to each other in a muscle. Some "muscle tears" involve the rupture of desmosomes.

Gap Junctions

Gap junctions (also called communicating junctions) provide cytoplasmic channels from one cell to an adjacent cell and in this way are similar in their function to the plasmodesmata in plants. Gap junctions consist of membrane proteins that surround a pore through which ions, sugars, amino acids, and other small molecules may pass. Gap junctions are necessary for communication between cells in many types of tissues, including heart muscle, and in animal embryos.

Sel: Satuan Kehidupan yang Lebih Besar dari bagian-bagiannya

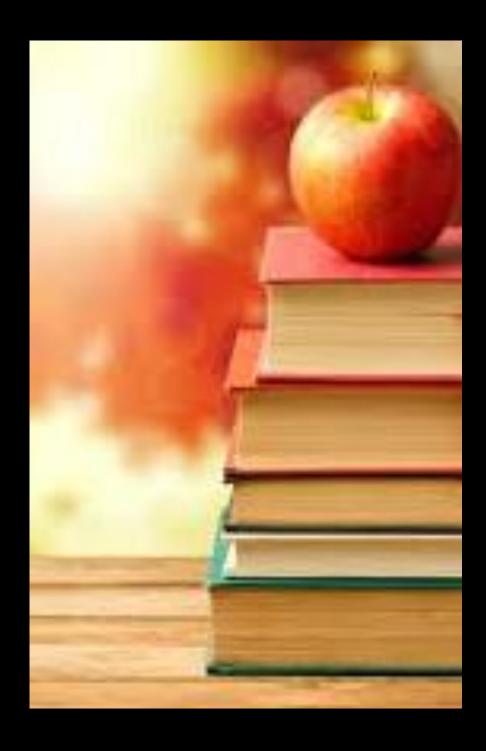
5 µm



▲ Figure 6.33 The emergence of cellular functions. The ability of this macrophage (brown) to recognize, apprehend, and destroy bacteria (yellow) is a coordinated activity of the whole cell. Its cytoskeleton, lysosomes, and plasma membrane are among the components that function in phagocytosis (colorized SEM).

DAFTAR PUSTAKA

- Campbell, Neil A., Jane B. Reece. 2008. Biology. 8th ed. Pearson Benjamin Cummings, San Francisco.
- -https://www.youtube.com/watch?v=zcUIQ14iUqA
- -https://www.youtube.com/watch?v=d4TJ4NY1IA0
- https://www.youtube.com/watch?v=gaElp0M3NZw



TUGAS

MENGGAMBAR STRUKTUR SEL BAKTERI/HEWAN/TUMBUHAN PADA KERTAS HVS/BERGAMBAR DAN DILENGKAPI DENGAN PENJELASAN SINGKAT DARI BAGIAN-BAGIAN SEL

BUAT SEKREATIF MUNGKIN. GAMBAR YANG MENARIK INSYAALLAH AKAN DIUPLOAD KE MEDIA SOSIAL STFI

DEADLINE: 1 MINGGU SETELAH PERKULIAHAN