

# Introduction to Bioinformatics

## Structure of Cells

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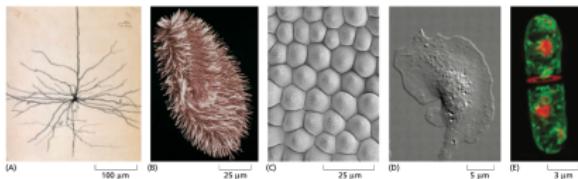
Fall 1404

# Presentation Overview

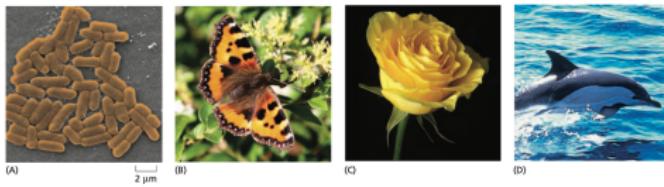
- ① Cells: The Fundamental Units of Life
- ② Cells & HeLa Cells
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- ④ Kingdoms/Domains & Prokaryotes
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# Cells: The Fundamental Units of Life

- **What is life?** Petunias, people, and pond scum are living; stones and sand are not.
- **Cell theory:** All living things are built from cells that grow and divide from pre-existing cells.

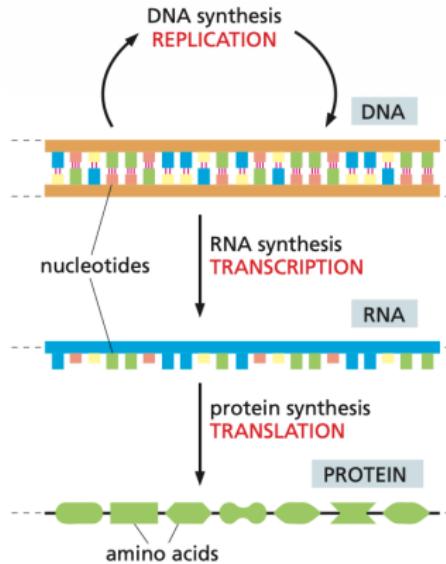


- **Why cells?** They are the basic units of structure and function; communities of specialized cells form tissues and organisms.



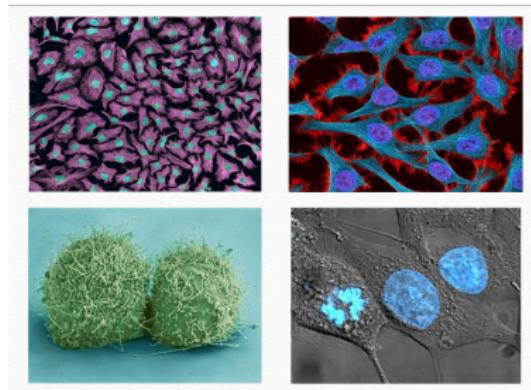
# Cells

- Small, membrane-enclosed units with aqueous interior that can grow and divide.
- Sizes:  $\sim 1\text{--}1000 \mu\text{m}$  (bacteria  $\rightarrow$  frog egg).
- Shapes reflect function: neurons, ciliated protozoa, macrophages, yeasts.
- **Common chemistry:** DNA  $\rightarrow$  RNA  $\rightarrow$  Protein (central dogma).
- **Autocatalysis:** DNA/RNA store information; proteins catalyze making DNA, RNA, and proteins.



# HeLa Cells

- Immortal human epithelial cell line from Henrietta Lacks (1951, cervical cancer).
- First robust human cells in culture; enabled vaccines, virology, cancer biology, drug screens.



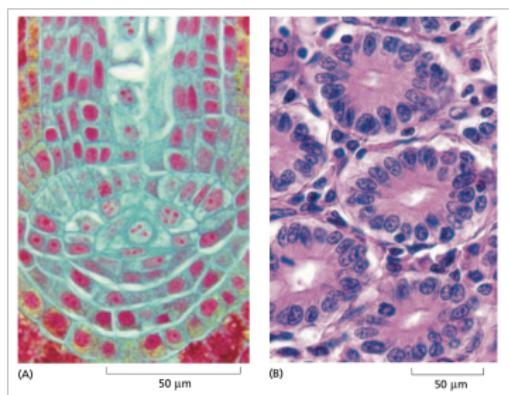
# Cells in Embryos

- Same genome, different fates: cells proliferate, migrate, and **differentiate**.
- Cell-cell signals and environment → selective gene expression.
- One fertilized egg → many specialized cell types (muscle, neuron, skin).



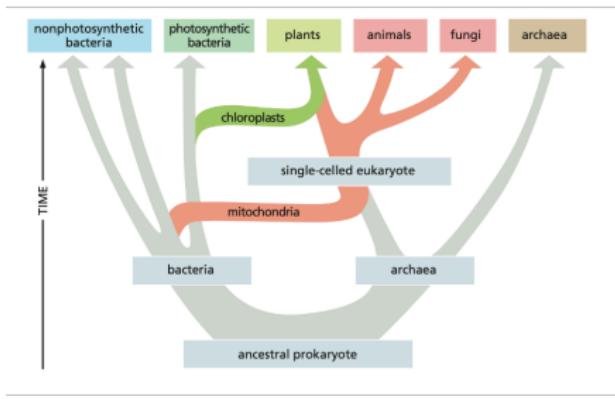
# Tissues

- Cooperative groups of specialized cells + extracellular matrix (ECM).
- Examples: epithelial sheets, muscle, connective tissue, nervous tissue, plant epidermis.
- Division of labor enables specialization (contraction, secretion); ECM provides structural and signaling context.



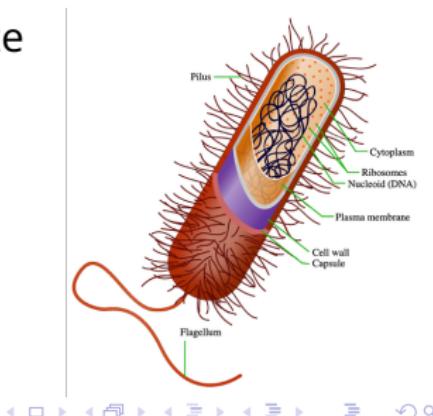
# Kingdoms of Life vs Domains

- **Classic Kingdoms (historical):** Animals, Plants, Fungi, Protists, (Monera).
- **Modern view:** Three Domains by molecular phylogeny: **Bacteria, Archaea, Eukarya.**
- Eukarya includes animals, plants, fungi, and protists; Archaea occur in extreme and common habitats.



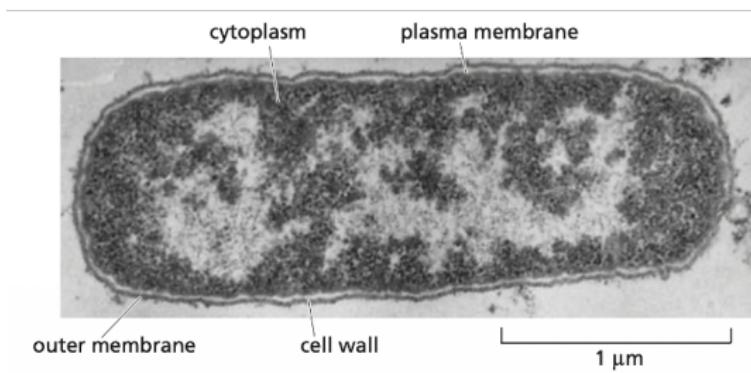
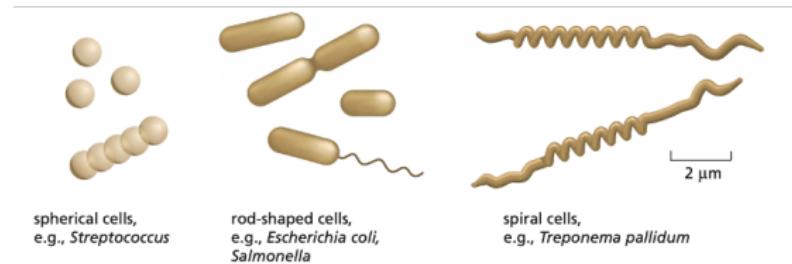
# Prokaryotes

- Cells **without** a nucleus; include **Bacteria** and **Archaea**.
- Small (few  $\mu\text{m}$ ), fast division (as little as  $\sim 20$  min), diverse metabolisms.
- Single compartment: cytoplasm + DNA; ribosomes present; no membrane-bound organelles.
- Most are single-celled; some form chains/clusters; chemically the most diverse class of cells.
- Some are aerobic, using oxygen to oxidize food molecules. Some are strictly anaerobic and are killed by the slightest exposure to oxygen



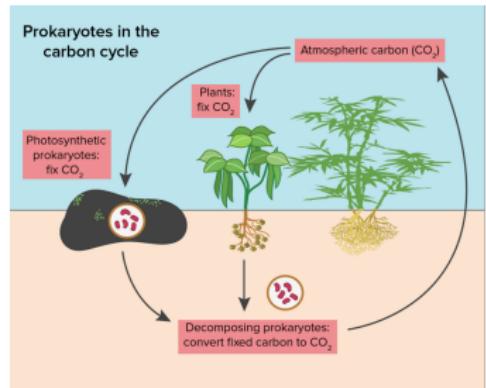
# Shapes of Bacteria

- **Coccus** (spherical)
- **Bacillus** (rod-shaped)
- **Spirillum/Spirochete** (spiral/corkscrew)



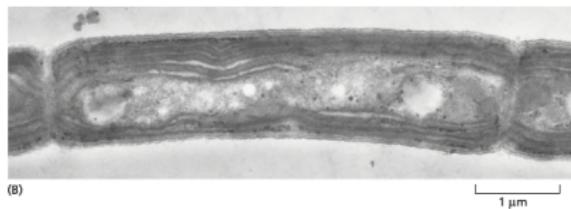
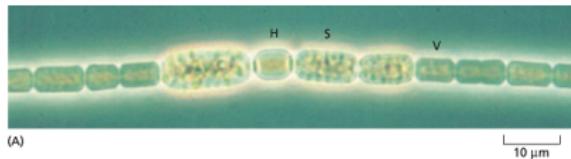
# Prokaryotic Metabolism

- **Aerobic vs anaerobic** ( $O_2$  required vs toxic).
- **Carbon sources:**  $CO_2$  (autotroph) vs organic carbon (heterotroph).
- **Nitrogen:** some fix  $N_2$ ; others use ammonia, nitrate, etc.
- Specialists: sulfur oxidizers (e.g., *Beggiatoa*), iron oxidizers, methanogens.
- Some prokaryotes can live entirely on inorganic substances ( $CO_2$ ,  $N_2$ ,  $O_2$ ,  $H$ ,  $S$ ,  $P$ ).



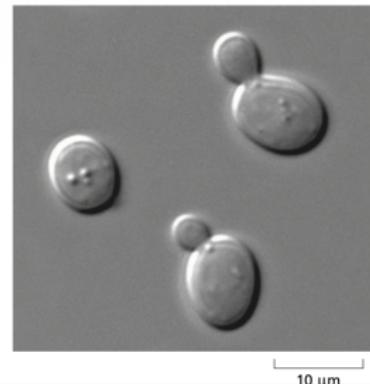
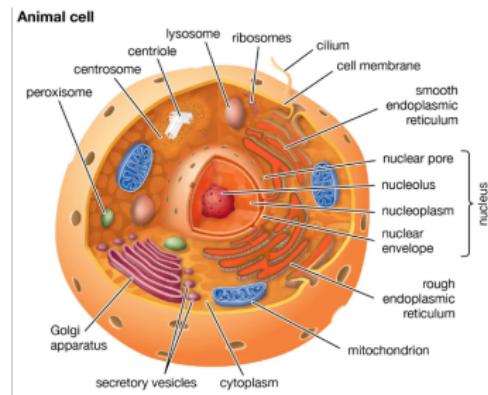
# Prokaryotic Sources of Energy

- **Phototrophs:** capture light (photosynthesis), e.g., cyanobacteria.
- **Chemolithotrophs:** energy from inorganic chemicals ( $H_2S$ ,  $Fe^{2+}$ ,  $NH_3$ ).
- **Chemoorganotrophs:** energy from organic molecules (sugars, fats).



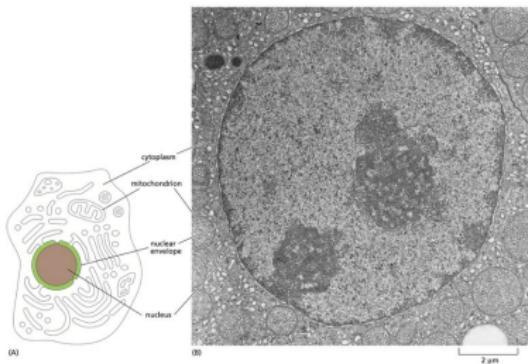
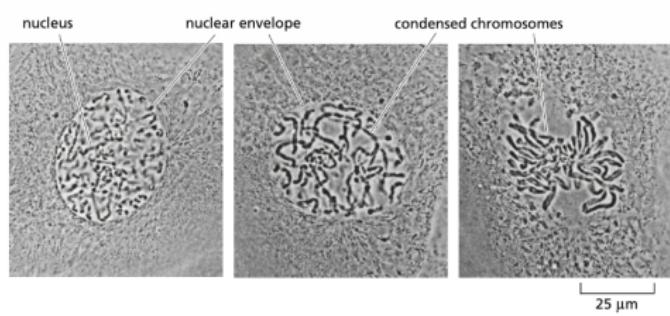
# Eukaryotes

- Cells with a **nucleus** and **membrane-bound organelles**.
- Single-celled (amoebae, yeasts) and multicellular (animals, plants, fungi).
- Typically larger; internal compartmentalization enables complex regulation.



# Nucleus: Information Store of the Cell

- **Nuclear envelope:** double membrane; nuclear pores regulate traffic.
- **DNA packaged into chromosomes;** condense visibly during division.
- **Nucleolus:** rRNA synthesis and ribosome assembly.



# Mitochondria

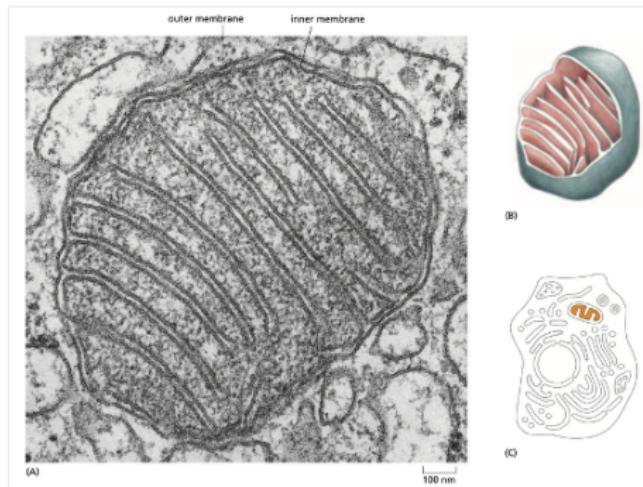
- Present in essentially all eukaryotic cells; conspicuous in cytoplasm.
- Contain their own DNA and divide by fission.
- Double membrane with folded inner membrane (cristae).



10 μm

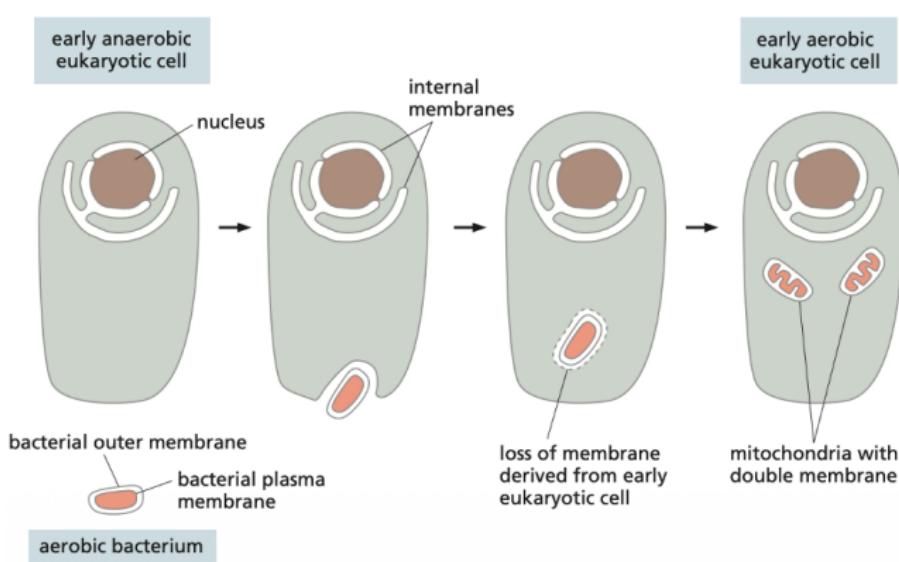
# Mitochondria: Function

- Generators of chemical energy (ATP) by oxidizing food molecules: **cellular respiration**.
- Consume O<sub>2</sub> and release CO<sub>2</sub>; central to energy metabolism.
- Functional discovery via cell fractionation and testing of purified mitochondria.
- Some anaerobic eukaryotes (e.g., *Giardia*) lack typical mitochondria.



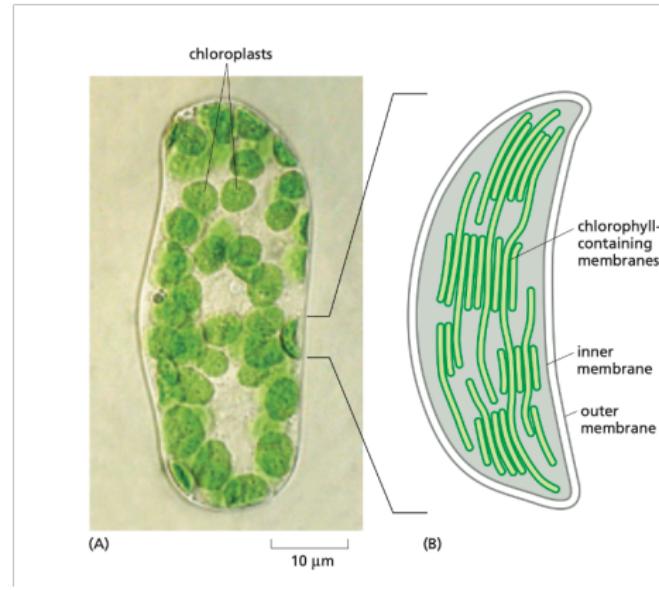
# Mitochondria: Origin (Endosymbiosis)

- Derived from engulfed aerobic bacteria → symbiosis with early eukaryote.
- Evidence: double membrane, bacterial-like ribosomes, circular DNA, division by fission.



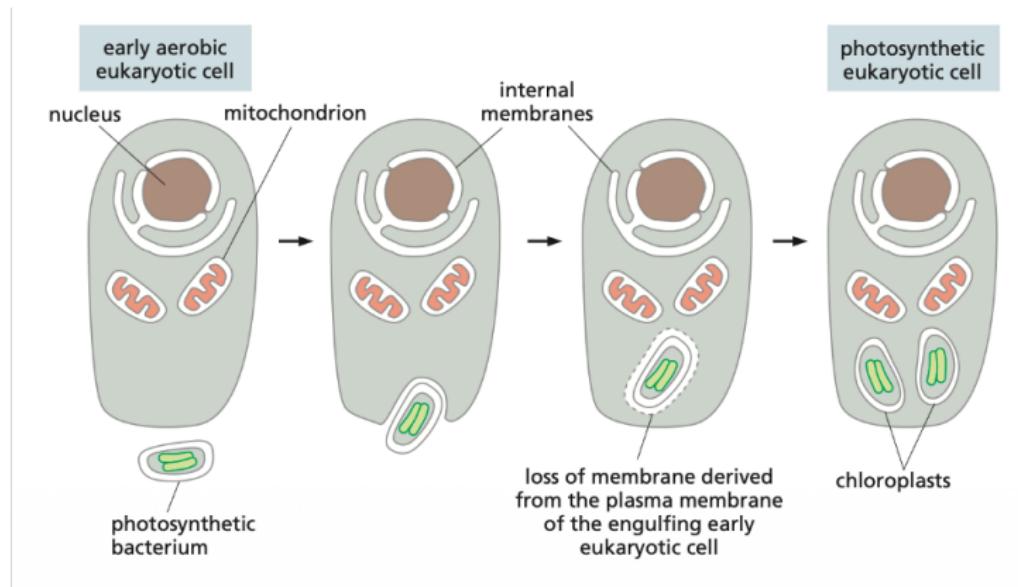
# Chloroplasts: Function

- **Photosynthesis:** light → chemical energy; fixes CO<sub>2</sub> into sugars; releases O<sub>2</sub>.
- Internal thylakoid membranes with chlorophyll.
- Plant cells oxidize sugars in mitochondria to make ATP as needed.
- Large, green organelles in plants and algae.



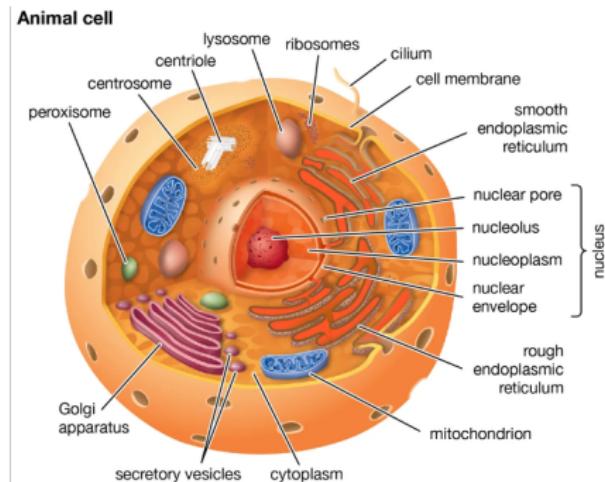
# Chloroplasts: Origin (Endosymbiosis)

- From engulfed photosynthetic bacteria (cyanobacteria-like).
- Own DNA, bacterial ribosomes; divide by fission; double membrane.



# Other Organelles

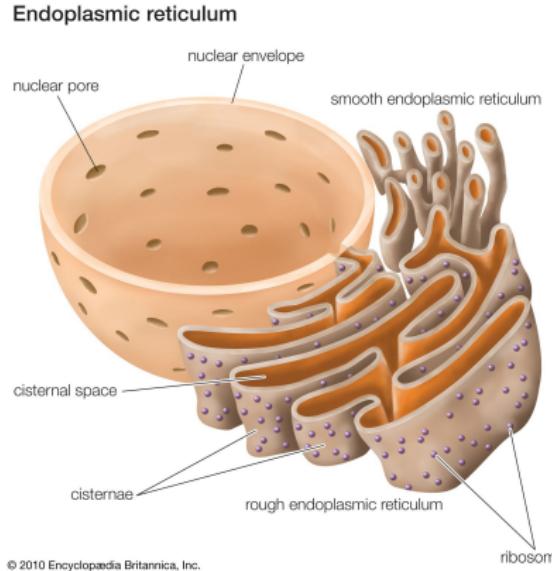
- **Endoplasmic Reticulum (ER)**: lipid/protein synthesis (RER with ribosomes; SER without).
- **Golgi apparatus**: processing, modification, and sorting.
- **Lysosomes**: intracellular digestion and recycling.
- **Peroxisomes**: detox reactions using  $H_2O_2$ .
- **Endosomes**: sorting centers after endocytosis.
- **Cytosol**: concentrated aqueous gel for metabolism and protein synthesis.



# Endoplasmic Reticulum (ER)

- Network of **interconnected membrane-bound sacs and tubules**.
- Synthesizes **lipids** and **membrane proteins**.
- **Rough ER** has ribosomes attached → primary site of **protein synthesis**.
- Continuous with the **nuclear envelope**, allowing direct transport of proteins into the ER.
- Key for maintaining cell structure and intracellular transport.

# Endoplasmic Reticulum Image



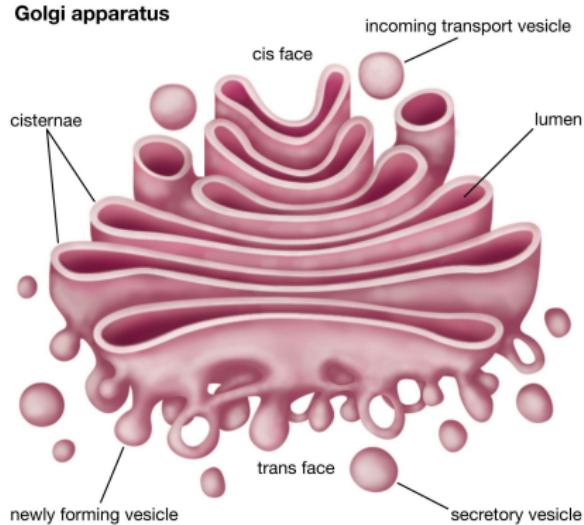
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**Figure:** Rough and smooth ER network. Ribosome-studded areas are rough ER. Source: Britannica

# Golgi Apparatus

- Composed of **stacked, flattened membrane sacs.**
- Receives proteins and lipids from the ER for further modification.
- Performs chemical modifications: **glycosylation, phosphorylation, sulfation.**
- Sorts and packages molecules into vesicles for transport to other organelles or the plasma membrane.
- Essential for secretion and membrane composition regulation.

# Golgi Apparatus Image



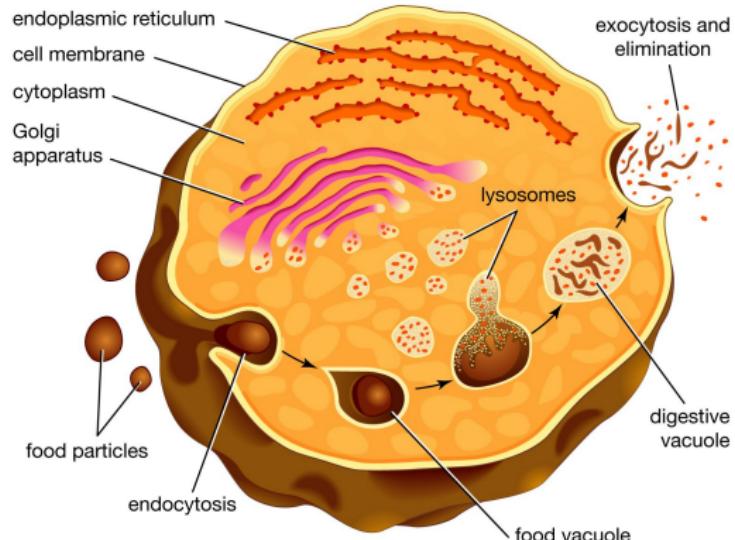
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**Figure:** Golgi apparatus; site of protein modification and vesicle packaging. Source: Britannica

# Lysosomes

- Small, membrane-bound organelles containing **digestive enzymes**.
- Break down macromolecules and recycle cell components.
- Formed from **endosomes** after endocytosis.
- Protect the cell by safely digesting waste and foreign material.
- Critical for autophagy, immune response, and nutrient recycling.

# Lysosome Image



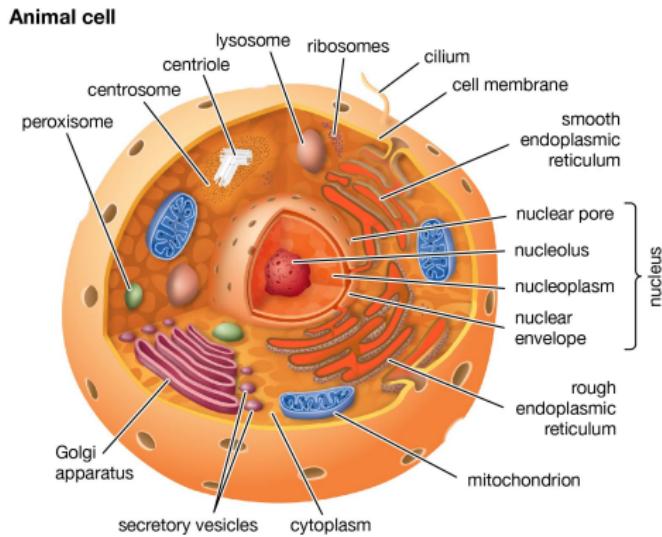
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Figure: Lysosomes digest internalized material. Source: Britannica

# Peroxisomes

- Membrane-enclosed vesicles containing **oxidative enzymes**.
- Metabolize hydrogen peroxide and other reactive oxygen species.
- Detoxify harmful molecules and protect the cell from oxidative damage.
- Important in lipid metabolism and liver function.

# Peroxisome Image



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Figure: Peroxisome containing oxidative enzymes. Source: Britannica

# Vesicles

- Small, membrane-bound sacs that transport cargo between organelles or to the cell surface.
- Can form from budding of ER or Golgi and fuse with other membranes.
- Analogy: **tiny transport vehicles** for proteins and molecules.
- Crucial for cellular logistics and secretion.

# Vesicle Image

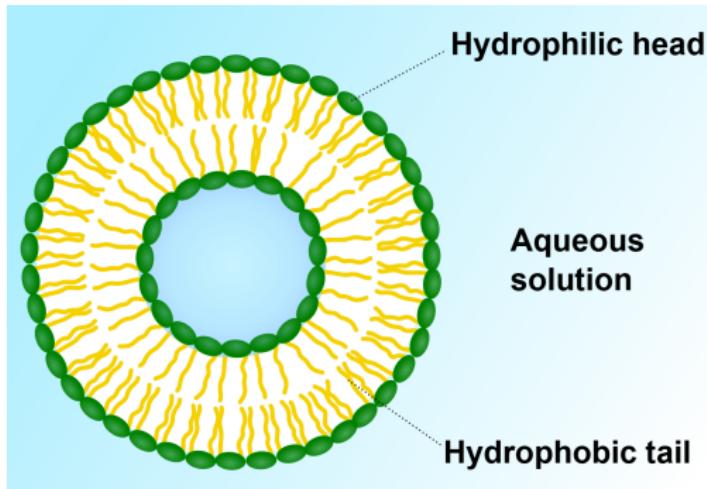


Figure: Vesicle structure. Source: Wikipedia

# Endocytosis

- Plasma membrane folds inward to form vesicles.
- Enables cells to engulf large particles, fluids, or even other cells.
- Important in nutrient uptake, immune defense, and signaling.

# Endocytosis Image

## Endocytosis

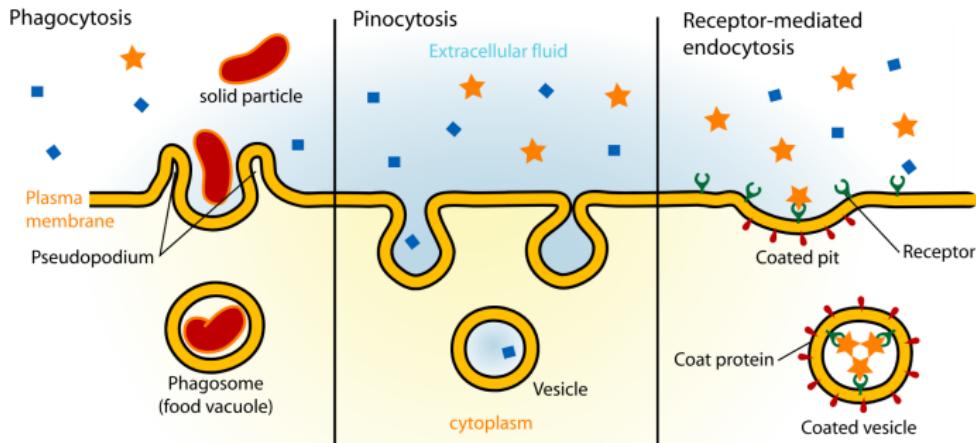
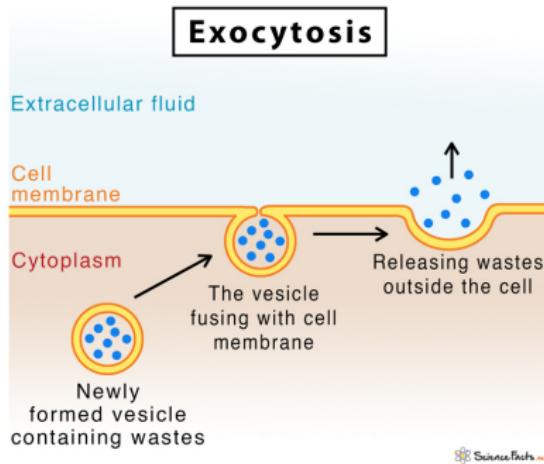


Figure: Types of endocytosis. Source: Wikipedia

# Exocytosis

- Vesicles fuse with the plasma membrane to release contents outside the cell.
- Secretes hormones, neurotransmitters, enzymes, and signaling molecules.
- Balances membrane composition and supports intercellular communication.

# Exocytosis Image

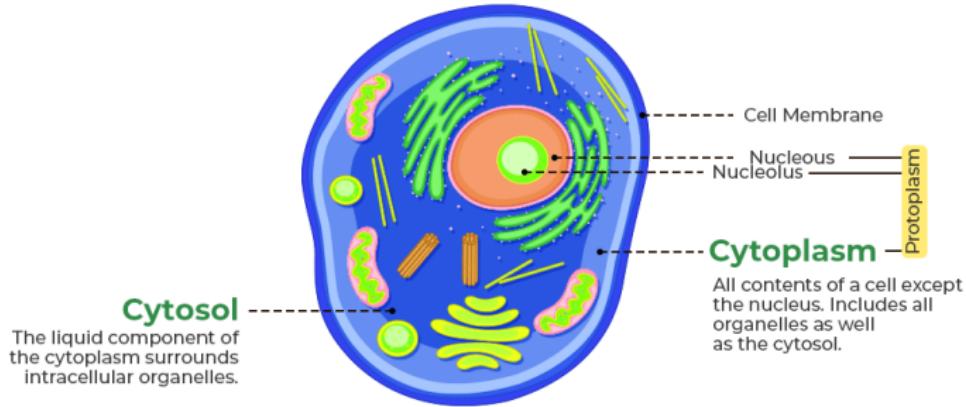


**Figure:** Exocytosis process. Source: ScienceFacts

# Cytosol

- Fluid portion of the cytoplasm where organelles are suspended.
- Contains ions, small molecules, and macromolecules.
- Site for many **metabolic pathways**, such as glycolysis and protein synthesis.
- Supports intracellular transport and acts as a medium for signaling molecules.

# Cytosol Image



**Figure:** Cytosol surrounds organelles and facilitates biochemical reactions. Source: GeeksforGeeks

# Ribosomes

- Molecular machines responsible for **protein synthesis**.
- Found **free in cytosol** or **bound to rough ER**.
- Translate mRNA sequences into polypeptide chains.
- Fundamental for cell growth, division, and enzyme production.

# Ribosome Images

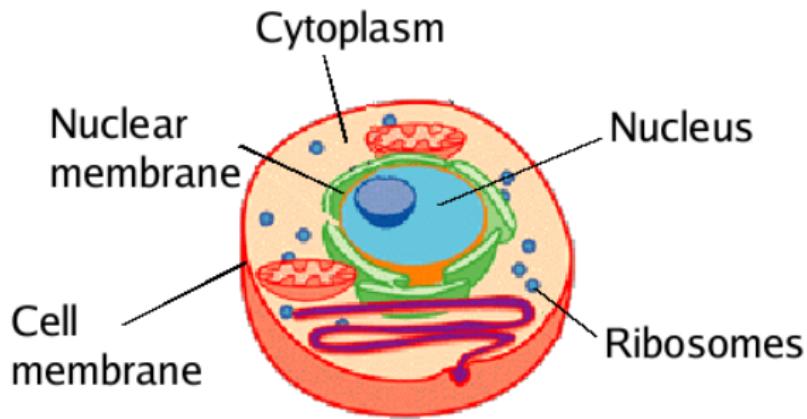
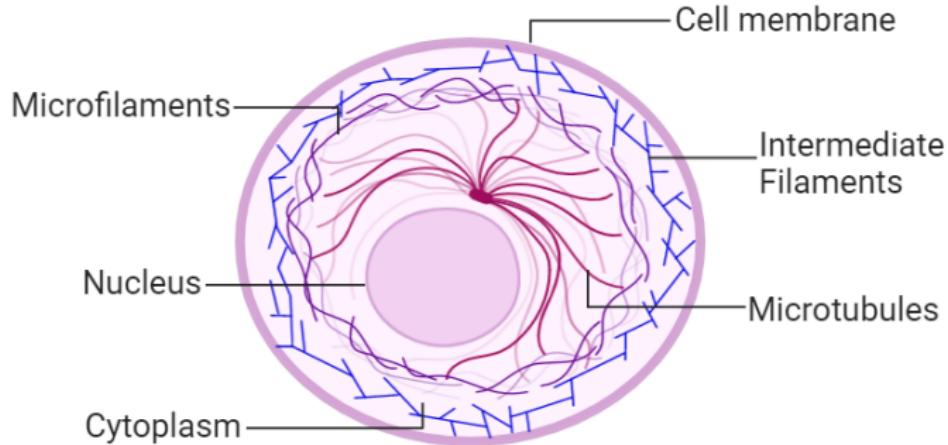


Figure: Ribosome during protein synthesis. Source: News-Medical

# Cytoskeleton Overview

- Network of **protein filaments**: actin filaments, microtubules, and intermediate filaments.
- Provides **mechanical support, maintains cell shape, and organizes internal structures**.
- Drives **cell movement, division, and intracellular transport**.

# Cytoskeleton Image

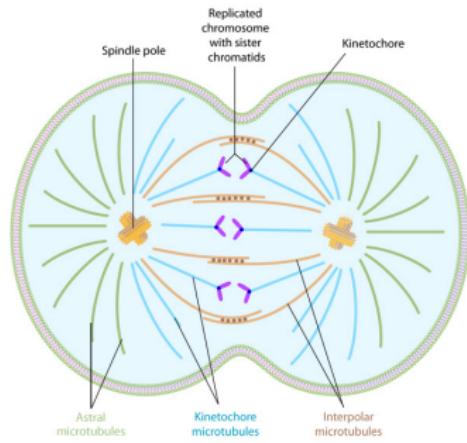
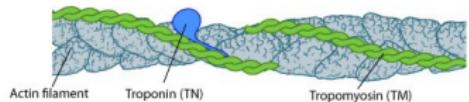


**Figure:** Cytoskeleton network maintaining structure and transport.  
Source: Ahmad Coaching

# Actin Filaments and Microtubules

- **Actin filaments:** thin, flexible fibers; generate contractile forces; abundant in cell cortex and muscle.
- **Microtubules:** thick, hollow tubes; organize organelles and guide chromosome movement during division.
- **Intermediate filaments:** provide tensile strength and maintain nuclear positioning.

# Actin and Microtubule Images

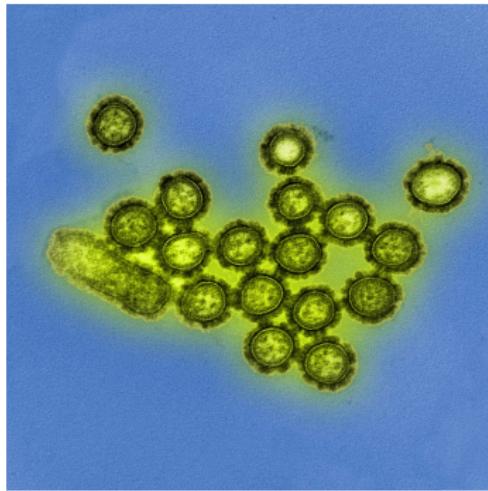


**Figure:** Actin filaments (left) and microtubules (right). Sources: MBI NUS, Nature

# Observing Cells

- **Light microscopy:** allows visualization of large organelles (> 200 nm).
- **Electron microscopy:** provides detailed ultrastructure at nanometer scale.
- **Fluorescence microscopy:** highlights specific organelles or molecules using dyes or tags.
- Techniques complement each other to study structure and function in cells.

# Cells Under Microscope



**Figure:** Surface proteins of virus particles and cells. Source: Britannica

# Cell Membrane Thickness

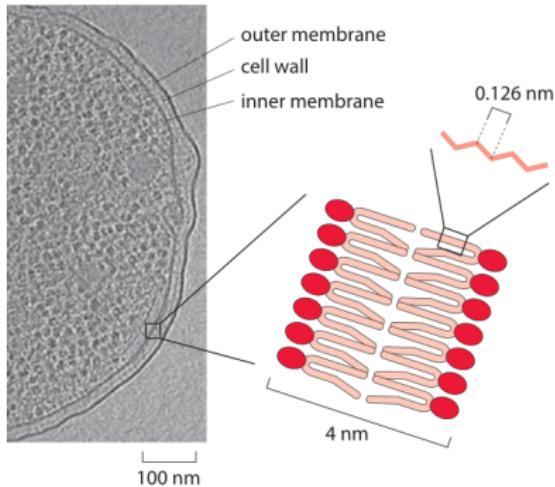


Figure: Cell membrane thickness ( 5–10 nm). Source: BioNumbers

# Signs of Life in Cells

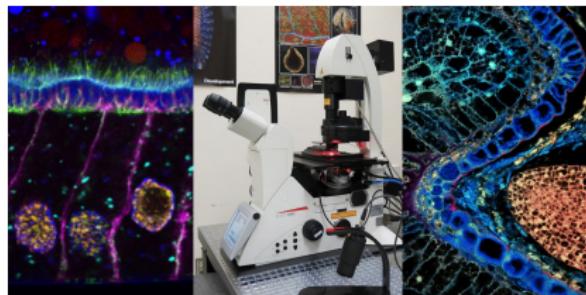
- Cells exhibit internal motion: particles, vesicles, organelles move around.
- Cells change shape, grow, divide — these dynamics signal life.
- Without staining, many organelles are transparent and difficult to see.
- To observe them, we use contrast-enhancing techniques and specialized microscopy.

# Light Microscopy

- Uses visible light passed through or reflected from specimens.
- Typically can resolve structures down to 200 nm.
- Useful for observing whole cells, larger organelles, live cells.
- Often combined with dyes or stains to highlight structures.

# Confocal Microscopy

- A laser scans a focused point and rejects out-of-focus light via pinholes.
- Produces sharper optical “slices” through cells (optical sectioning).
- Can reconstruct 3D images from stacked slices.

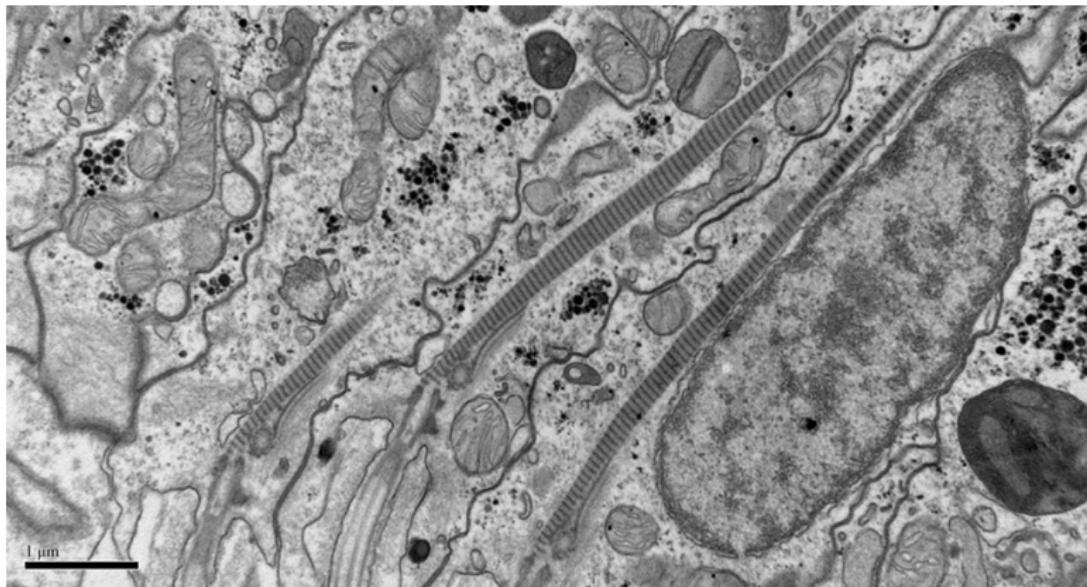


**Figure:** Confocal microscopy image (optical section of a cell). Source: Illinois State Univ.

# Transmission Electron Microscopy (TEM)

- TEM passes an electron beam through an ultra-thin section of the specimen.
- Reveals detailed **internal ultrastructure**: membranes, organelles, macromolecular complexes.
- Resolution is extremely high (on the order of nanometers), far beyond light microscopes.
- Requires specimen preparation (sectioning, staining with heavy metals) and vacuum environment.

# TEM Image



**Figure:** TEM micrograph showing internal cellular structure (e.g. organelle membranes). Source: University of Exeter Bioimaging TEM

# Scanning Electron Microscopy (SEM)

- SEM scans a focused electron beam over the specimen's surface.
- Creates a 3D-like image of surface topography and external morphology.
- Lower resolution for internal details but excellent for surface structures (microvilli, cilia, cell edges).
- Specimens often must be coated in a conductive material (e.g. gold) and placed in vacuum.

# SEM Image

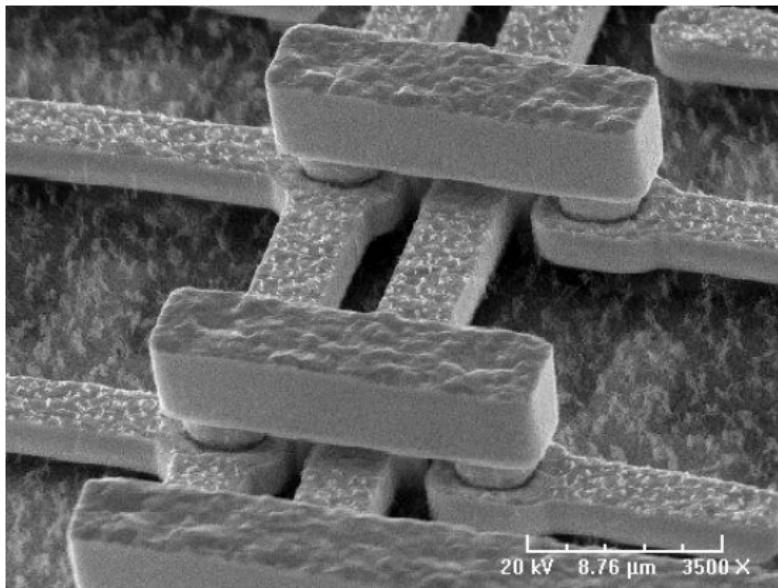
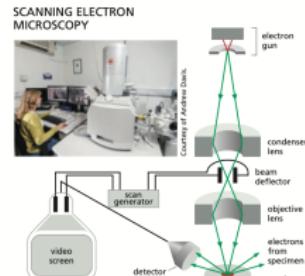
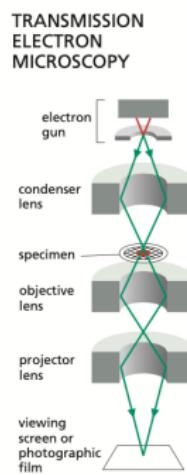
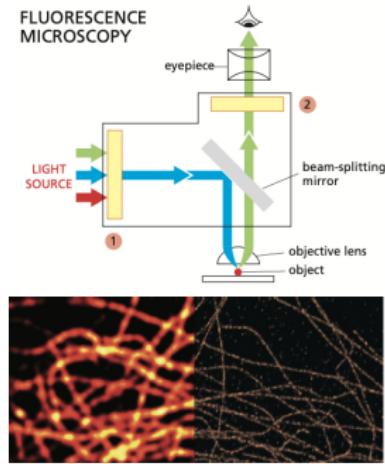
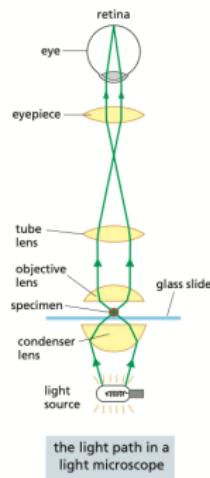


Figure: SEM image of cell surface morphology. Source: ORSlabs SEM

# Observing Cells: Microscopy (Brief)

- **Light microscopy:** brightfield + stains; phase/DIC for live cells.
- **Fluorescence/confocal:** specific labeling; optical sections; 3D.
- **Super-resolution:** ~20 nm resolution.
- **Electron microscopy:** TEM (internal details), SEM (surface).



# The End

Questions? Comments?