

BIOLOGY ASSIGNMENT-02

REPORT TOPIC :Scanning Electron Microscopy (SEM)

PICTURE:



Introduction

Scanning Electron Microscopy (SEM) is a type of electron microscopy that produces images by scanning a specimen with a focused beam of electrons. These electrons interact with atoms in the sample to produce various signals that provide information about the sample's surface topography, composition, and other properties. SEM is extensively used in the field of biology for analyzing the surface structures of biological specimens in high detail and at very high magnifications.

In biology, SEM helps visualize complex surface architectures of cells, tissues, microorganisms, and plant structures that are otherwise difficult to analyze using conventional optical microscopes. SEM is valuable for both research and diagnostic purposes.

Principle of SEM

SEM works by directing a beam of high-energy electrons onto the surface of a specimen. The beam is generated by an electron gun and focused into a narrow spot using electromagnetic lenses. As the electron beam scans across the surface, electrons interact with the sample, resulting in the emission of various signals such as secondary electrons, backscattered electrons, and X-rays.

Secondary electrons (SE) are primarily used to generate SEM images. These are lowenergy electrons emitted from the surface atoms of the specimen and provide detailed topographical information. Backscattered electrons (BSE), on the other hand, are highenergy electrons reflected from the sample and provide compositional contrast.

Sample Preparation for Biological SEM

Preparing biological specimens for SEM imaging is crucial, as they are often soft, moist, and non-conductive. Improper preparation can damage the specimen or produce poorquality images. The preparation process involves several steps:

1. Fixation: Biological tissues are fixed using aldehydes like glutaraldehyde to preserve cell-structure.
2. Post-fixation: Osmium tetroxide is often used to stabilize lipid components and improve-electroncontrast.
3. Dehydration: Gradual replacement of water with ethanol or acetone through a graded series-(30%-100%).
4. Drying: Critical point drying or freeze-drying methods are employed to preserve structural-integrity.
5. Coating: A conductive coating (gold, platinum, or carbon) is applied using sputter coating to prevent charging under the electron beam.

Applications of SEM in Biology

SEM is widely used in biological sciences and has numerous applications, including:

- *Cell Biology: Examines cellular surfaces, including organelles like cilia, flagella, and membrane.
- *Microbiology: Helps in studying the morphology and surface features of bacteria, viruses.
- *Histology: Provides 3D views of tissue structures for detailed analysis.
- *Entomology: Allows the study of the exoskeleton and sensory structures in insects.
- *Botany: Used for analyzing stomata, trichomes, pollen grains, and seed coats.
- *Medical Research: Helps in analyzing pathological tissues and understanding disease mechanisms.
- *Biomaterials: Evaluates the biocompatibility and structural features of scaffolds, implants, and prosthetics.

Advantages of SEM in Biology

- *High Resolution: SEM can resolve fine details up to 1-2 nanometers, much higher than light.
- *Three-Dimensional Imaging: Provides depth and perspective, enhancing understanding morphology.
- *Elemental Analysis: Integration with Energy-Dispersive X-ray Spectroscopy (EDS) analysis.

*Versatility: Applicable to a wide range of biological samples including cells, tissues, and biomaterials.

Limitations of SEM

*Non-Living Samples: SEM requires vacuum conditions, so biological samples must be dehydrated and cannot be alive during imaging. *Complex Sample Preparation: Time-consuming and can introduce artifacts. *Cost and Maintenance: SEM instruments are expensive to purchase and maintain. *Conductive Coating: The need for a metal coating can obscure fine surface details or interfere with some analyses.

Recent Developments and Future Perspectives

Recent advancements in SEM technology have significantly improved its capabilities in biological research. Environmental SEM (ESEM) allows imaging of hydrated and uncoated biological samples in low-vacuum or variable pressure conditions. Cryo-SEM, which involves freezing specimens rapidly, enables visualization of biological structures in their near-native states without extensive dehydration or coating.

Future improvements in SEM aim to reduce preparation time, enhance imaging speed, and increase resolution. Integration with AI and automated image analysis tools is also transforming the way SEM data is interpreted in biological research.

Conclusion

Scanning Electron Microscopy is a vital technique in biology, offering unparalleled insights into the ultrastructure of cells and tissues. While it has limitations, technological advancements continue to expand its applications, making SEM an indispensable tool in both academic and clinical biological research.

How SEM Helps Humans

Scanning Electron Microscopy (SEM) plays a significant role in enhancing human life, particularly in healthcare, biotechnology, and environmental research. It helps researchers understand the microscopic structures of tissues, cells, bacteria, and viruses, aiding in disease diagnosis and the development of treatments. For example:

*Medical Diagnostics: SEM is used to examine biopsy tissues and identify abnormalities in cellular structure, helping diagnose diseases such as cancer. *Infectious Disease Research: It allows detailed visualization of pathogens like viruses and bacteria, which helps in developing vaccines and antimicrobial drugs. *Pharmaceuticals: SEM ensures the quality and structure of drug particles, aiding in drug formulation.

*Forensics and Pathology: Used in forensic science to analyze tissues and samples investigations.

*Environmental Monitoring: Helps identify pollutants and bio-contaminants in ecosystems, contributing to public health.

How SEM Works

The working principle of SEM involves several key steps:

1. Electron Beam Generation: An electron gun generates a beam of high-energy electrons.
2. Focusing: Electromagnetic lenses focus the beam into a fine point.
3. Scanning: The electron beam is rastered across the surface of the specimen.
4. Interaction: The electrons interact with atoms in the sample, producing secondary electrons.
5. Detection: Detectors collect these signals, and they are processed to form an image. The image displayed represents the surface structure of the sample with high magnification and resolution. By adjusting parameters like beam voltage and detector settings, researchers can gather detailed information about the texture, composition, and structure of biological specimens.

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