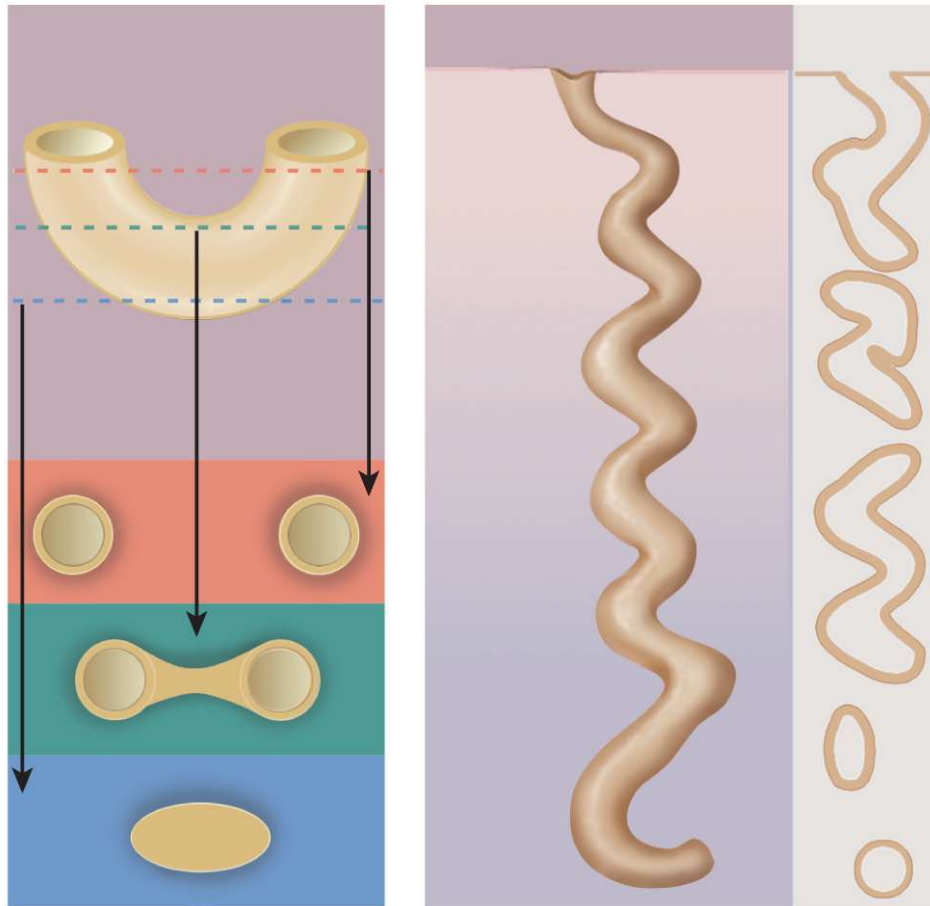


# How to Interpret Tissue Sections Using a Microscope



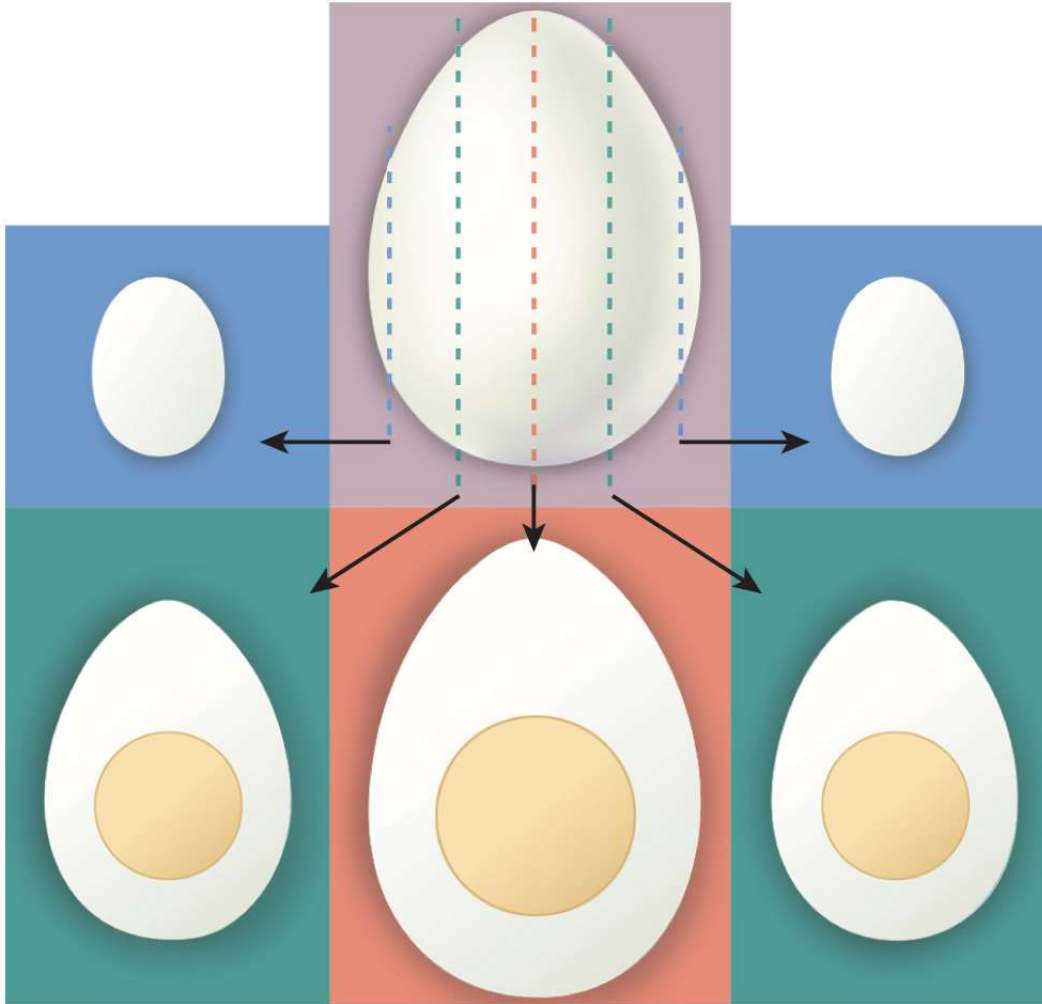
# Interpreting Tissue Sections

---

- Microscopic sectioning changes a **three-dimensional structure into a two-dimensional image**
- Preparation of histological specimens
  - fixative prevents decay (e.g. formalin) // but in process dissolves and removes the fat from the tissue
  - histological sections – tissue is sliced into thin sections one or two cells thick
  - stains – tissue is mounted on slides and artificially colored with histological stain - increases contrast // different stains may be used to bind to different cellular components to increase contrast

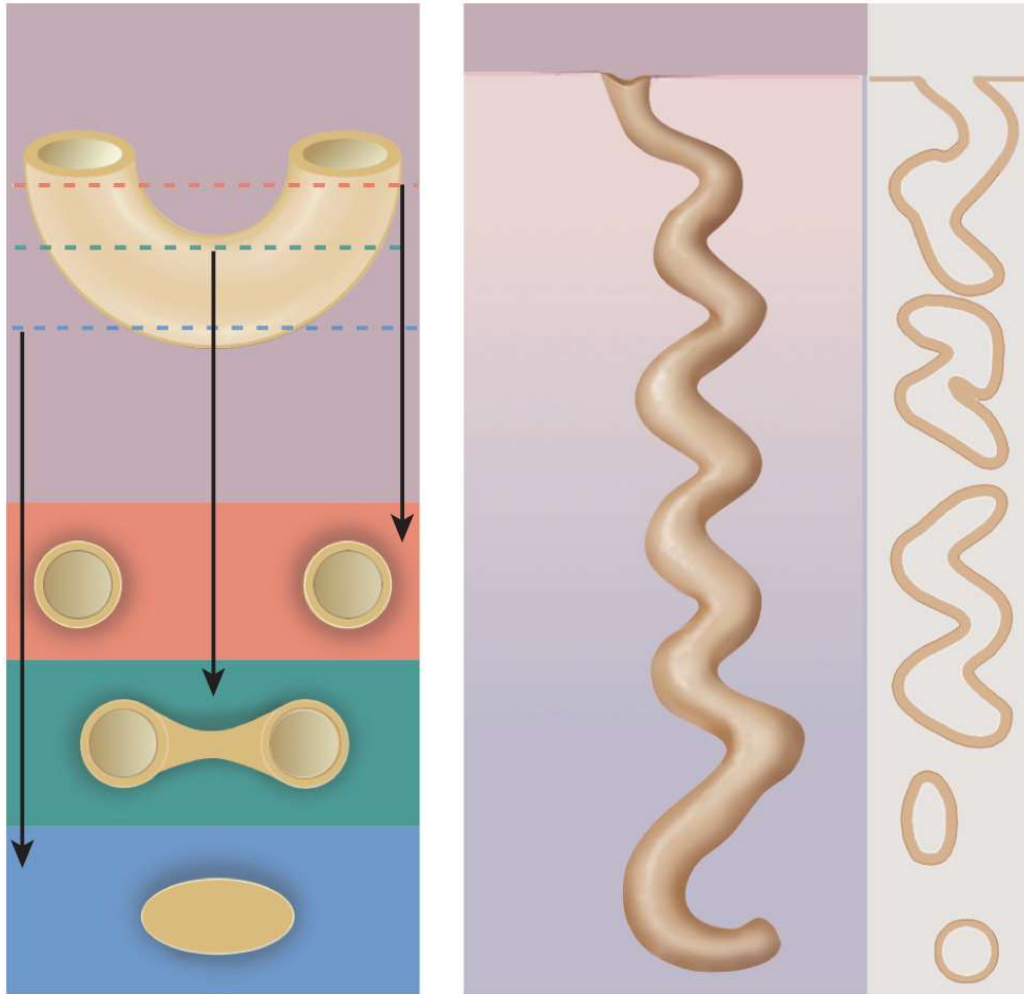
# Sectioning Solid Objects

---



- sectioning a cell with a centrally located nucleus
- some slices miss the cell nucleus
- in some sections the nucleus looks smaller than other sections

# Sectioning Hollow Structures

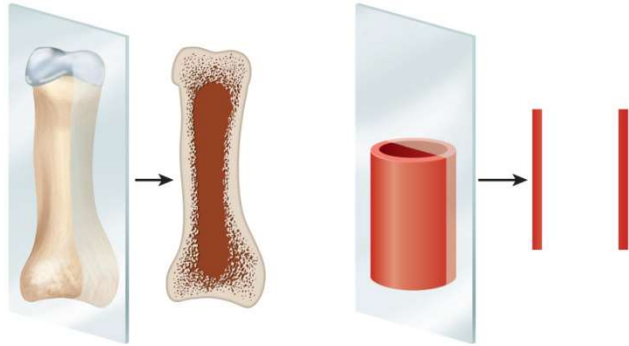


- **cross section** of blood vessel, gut, or other tubular organ.
- **longitudinal section** of a sweat gland /// notice what a single slice could look like.
- *You must use your knowledge about tissue function and your imagination to interpret what you are looking at under the microscope.*

# Types of Tissue Sections

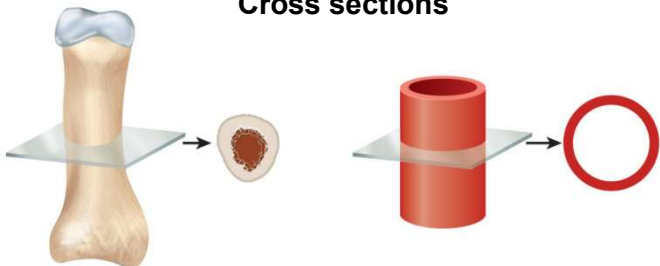
---

Longitudinal sections



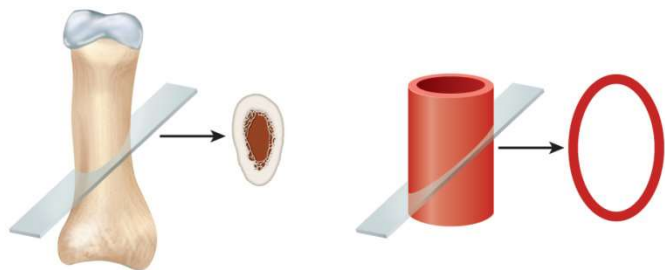
- **longitudinal section (l.s.)**
  - tissue cut along long direction of organ

Cross sections



- **cross section (c.s. or x.s.) or transverse section (t.s.)**
  - tissue cut perpendicular to length of organ

Oblique sections



- **oblique section**
  - tissue cut at angle between cross and longitudinal section

# Non-sectioned Preparation

---

- **Smear** – tissue is rubbed or spread across the slide and often a cover slip is placed over the tissue sample /// E.g. blood smear to identify formed elements
- **Spread** – cobwebby tissue is laid out on a slide without cover slip /// E.g. areolar tissue