



## مقرر

### علم الحيوان العام (أجنة وخلية وأنسجة وفسولوجي)

الفرقة الأولى شعبة العلوم البيولوجية والجيولوجية باللغة الإنجليزية (101 عل ح)

أستاذ المقرر

د/ رانا عبد الستار علي (جزء الفسيولوجي)

د/ نادية سمير (جزء الخلية والأنسجة)

د/ سهام علي مبارك (جزء الأجنة)

قسم علم الحيوان - كلية العلوم بقنا

العام الجامعي  
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نموذج توزيع المحتوى العلمي لمقرر (الأجنة والخلية والأنسجة والفسولوجي Zoo 101) للعام الدراسي ٢٠٢٢/٢٠٢٣  
اولا:- بيانات المقرر

كود المقرر:-	اسم المقرر:- الأجنة والخلية والأنسجة والفسولوجي	الفرقة/الشعبة:- الفرقة الأولى بكلية التربية شعبة العلوم البيولوجية والجيولوجية باللغة الإنجليزية	الكلية:- كلية التربية
الفصل الدراسي:-	عدد الساعات اسبوعيا	محادثة: ٤	محادثة:- أم د/ رانا عبد الستار- د/ نادية سمير- د/ سهام علي
	معمل: ٤	القائمين بالتدريس	معمل:- م ايمان جمال + م عبد الرحمن

ثانيا:- موضوعات المقرر

ملاحظات	موضوع المحاضرة/ المعمل	اسابيع الدراسة
	An introduction of general physiology ( <b>Physiology</b> ) Introduction Types of microscopes ( <b>Cell and Histology</b> ) Introduction of Embryology: Study of basic concepts of embryology ( <b>Embryology</b> )	الاول محاضرة
	<ul style="list-style-type: none"> <li>مقدمه عن المقرر العملي التعرف على تركيب الميكروسكوب الضوئي وكيفية التعامل مع كافة انواعه -</li> <li>(الميكروسكوب ذو العدسة العينية الواحدة الميكروسكوب ذو العدستين العينيتين ميكروسكوب التشريح)</li> <li>مقدمة عامة عن المنهج (علم الأجنة وأهم تعريفاته) (جزء الأجنة)</li> </ul>	معمل
	Homeostasis ( <b>Physiology</b> ) Cell components and ultra-structure of the animal cell Organisms and Cells ( <b>Cell and Histology</b> ) Gametogenesis & fertilization ( <b>Embryology</b> )	الثاني محاضرة
	<ul style="list-style-type: none"> <li>تشريح الأحشاء العامة للضفدعة</li> <li>رسم قطاع عرضي في خصية الفأر وقطاع عرضي في مبيض القطة (جزء الأجنة)</li> </ul>	معمل

صادر القسم ( )	قسم علم الحيوان	رؤية كلية العلوم
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المرفقات	الرمز البريدي 83523 قنا. جمهورية مصر العربية. تليفاكس:- +20963213383 داخل:- رئيس القسم 1367، سكرتارية 1523. البريد الإلكتروني:- zoology@sci.svu.edu.eg	رؤية قسم علم الحيوان خريجون متميزون علميا وبحثيا محليا ودوليا خدمة للمجتمع وتنمية للبيئة

	The skeletal system ( <b>Physiology</b> )	محاضرة	الثالث
	Cell structure and function ( <b>Cell and Histology</b> )		
	Pattern of cleavage and embryonic membranes of vertebrate species ( <b>Embryology</b> )		
	<ul style="list-style-type: none"> <li>تشريح الجهاز البولي التناسلي للضفدعة</li> <li>رسم مراحل تكوين الحيوانات المنوية والبويضات (<b>جزء الأجنة</b>)</li> </ul>	معمل	
	The skeletal system ( <b>Physiology</b> )	محاضرة	الرابع
	Cell Organelles structure and function Part 1 ( <b>Cell and Histology</b> )		
	Embryonic development of vertebrates (amphioxus) ( <b>Embryology</b> )		
	<ul style="list-style-type: none"> <li>تشريح الجهاز الوريدي البالي للضفدعة</li> <li>أنواع البويضات مع رسم أمثلة (<b>جزء الأجنة</b>)</li> </ul>	معمل	
	The Digestive System ( <b>Physiology</b> )	محاضرة	الخامس
	Cell Organelles structure and function Part 2 ( <b>Cell and Histology</b> )		
	Embryonic development of vertebrates (amphioxus) ( <b>Embryology</b> )		
	<ul style="list-style-type: none"> <li>النسيج الضام الفجوي و النسيج الضام الدهني</li> <li>الغضروف الزجاجي و ق.ط. من عظم كثيف</li> <li>أنواع ومستويات التقلج مع رسم أمثلة (<b>جزء الأجنة</b>)</li> </ul>	معمل	
	The Digestive System ( <b>Physiology</b> )	محاضرة	السادس
	Cell Organelles structure and function Part 3 ( <b>Cell and Histology</b> )		
	Embryonic development of vertebrates (tadpole) ( <b>Embryology</b> )		
	<ul style="list-style-type: none"> <li>سحبة من دم الضفدعة سحبة من دم الإنسان</li> <li>ق.ع. من الحبل الشوكي للأرنب</li> <li>مراحل التكوين الجنيني للرأس حبليات (جنين السهيم) حتى طور الجاسترولا (<b>جزء الأجنة</b>)</li> </ul>	معمل	
	<b>Midterm</b>	محاضرة	السابع

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المرفقات		

		معمل	
	The Circulatory System (The Heart, Blood Vessels, Blood Types) <b>(Physiology)</b>	محاضرة	الثامن
	Cell cycle (Mitosis) <b>(Cell and Histology)</b>		
	Embryonic development of vertebrates (tadpole) <b>(Embryology)</b>		
	ق.ر. من جلد الضفدعة	معمل	
	• مراحل التكوين الجنيني للرأس حبلليات (جنين السهيم) أطوار ما بعد الجاسترولا (جزء الأجنة)		
	The Circulatory System (The Heart, Blood Vessels, Blood Types) <b>(Physiology)</b>	محاضرة	التاسع
	Cell cycle (Meiosis) and Cell death <b>(Cell and Histology)</b>		
	Embryonic development of vertebrates (chicken) <b>(Embryology)</b>		
	ق.ع. من مرئ الضفدعة	معمل	
	• مراحل التكوين الجنيني للبرمائيات (جنين الضفدعة) حتى طور البلاستولا (جزء الأجنة)		
	The Neuromuscular System 1- The Muscular System <b>(Physiology)</b>	محاضرة	العاشر
	DNA & genes <b>(Cell and Histology)</b>		
	Embryonic development of vertebrates (chicken) <b>(Embryology)</b>		
	ق.ع. من معدة الضفدعة	معمل	

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المرفقات		

	<ul style="list-style-type: none"> <li>• مراحل التكوين الجنيني للبرمائيات (جنين الضفدعة) حتى طور الجاسترولا (جزء الأجنة)</li> </ul>		
	2- The Nervous System (Physiology)	محاضرة	الحادي عشر
	Types of animal tissues (Epithelial tissue) (Cell and Histology)		
	Embryonic development of vertebrates (mammal) (Embryology)		
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	Urinogenital system (Physiology)	محاضرة	الثاني عشر
	Types of animal tissues (Connective tissue) (Cell and Histology)		
	Embryonic development of vertebrates (mammal) (Embryology)		
	<ul style="list-style-type: none"> <li>• قطاع من كبد الضفدعة ق.ع. من كلية الضفدعة</li> <li>• مراحل التكوين الجنيني للطيور (جنين الكتكوت) حتى طور البلستولا (جزء الأجنة)</li> </ul>	معمل	
	Urinogenital system (Physiology)	محاضرة	الثالث عشر
	Types of animal tissues (Muscular tissue) (Cell and Histology)		
	Embryonic membranes (Embryology)		
	<ul style="list-style-type: none"> <li>• ق.ع. من رئة الضفدعة</li> <li>• مراحل التكوين الجنيني للثدييات من الزيجوت حتى طور البلستولا (جزء الأجنة)</li> </ul>	معمل	
	Types of animal tissues (Nervous tissue) (Cell and Histology)	محاضرة	الرابع عشر

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قسم علم الحيوان



جامعة جنوب الوادي

Placenta (Embryology)		
مراجعة عامة	معمل	

استاذ المقرر      رئيس مجلس القسم      وكيل الكلية لشئون التعليم و الطلاب      عميد الكلية

أ م د/ رانا عبد الستار      د/ عبد الناصر أحمد حسين      د/ جمال عبد الله أحمد      د/ خالد بن الوليد عبد الفتاح  
د/ نادية سمير  
د/ سهام علي

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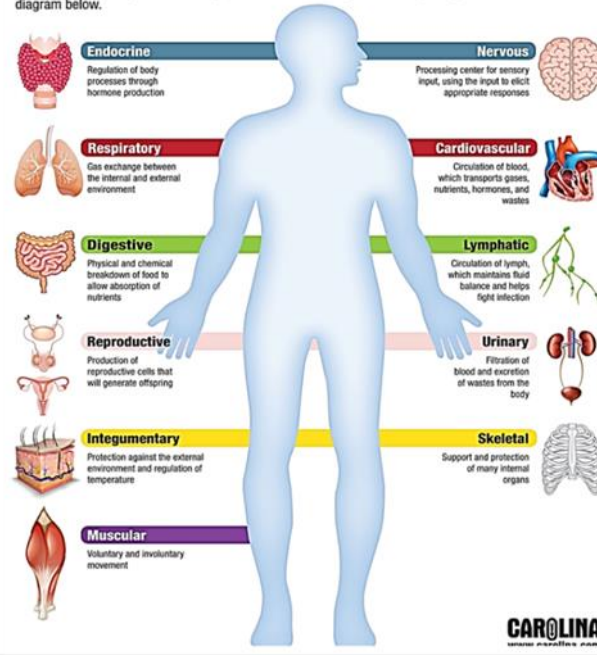
مقرر (علم الحيوان Zoo 101– Zoology I "جزء الفسيولوجي") لطلاب الفرقة الأولى  
شعبة العلوم البيولوجية والجيولوجية باللغة الإنجليزية (101 عل ح)

أستاذ المقرر

د/ رانا عبد الستار علي (جزء الفسيولوجي)

# Human Body Systems

There are 11 main systems that keep our bodies functioning. Learn the primary roles of each in the diagram below.





## Physiology content:

- An introduction of general physiology.
- Nutrition and Digestion.
- Absorption.
- Metabolism.
- Excretion.
- Respiration.
- Circulation system, Blood and Lymph.
- Reproductive system.
- Nervous System.
- Endocrine System and hormone's function.

### Physiology and life processes

Physiology tells us how our bodies work structurally and functionally.

**The most important life processes of human:**

**Metabolism:** includes catabolism and anabolism that provides energy and body components.

**Excitability:** ability to sense changes in and around us.

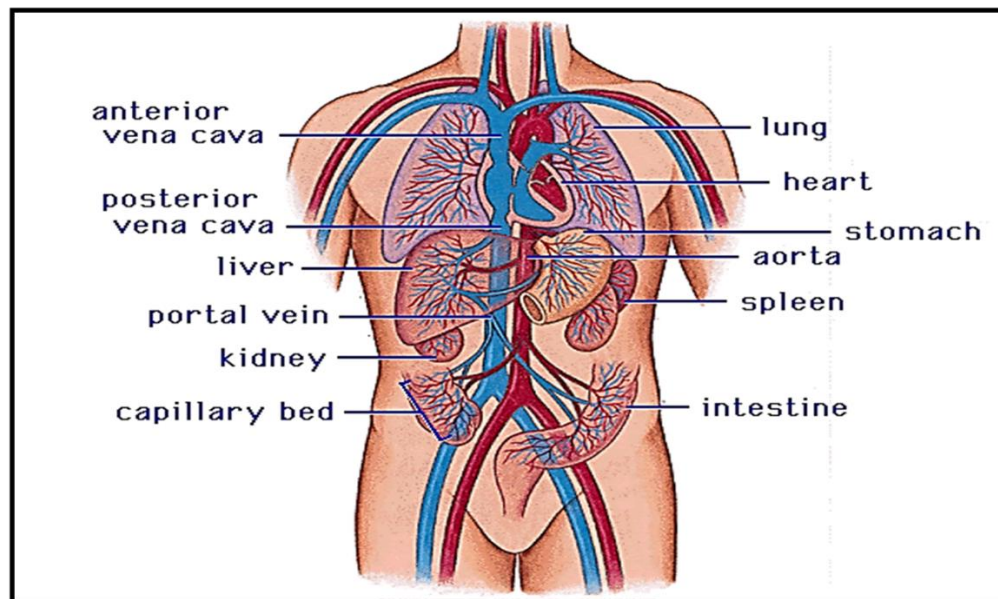
**Conductivity:** ability to carry the effects of stimulus from part of a cell to another.

**Contractility:** ability to contract in response to stimulus.

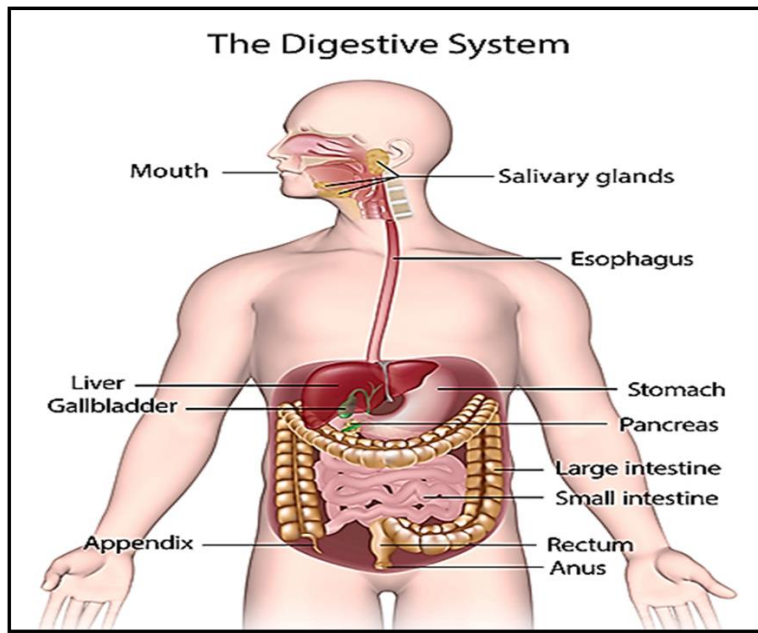
**Growth**

**Reproduction**

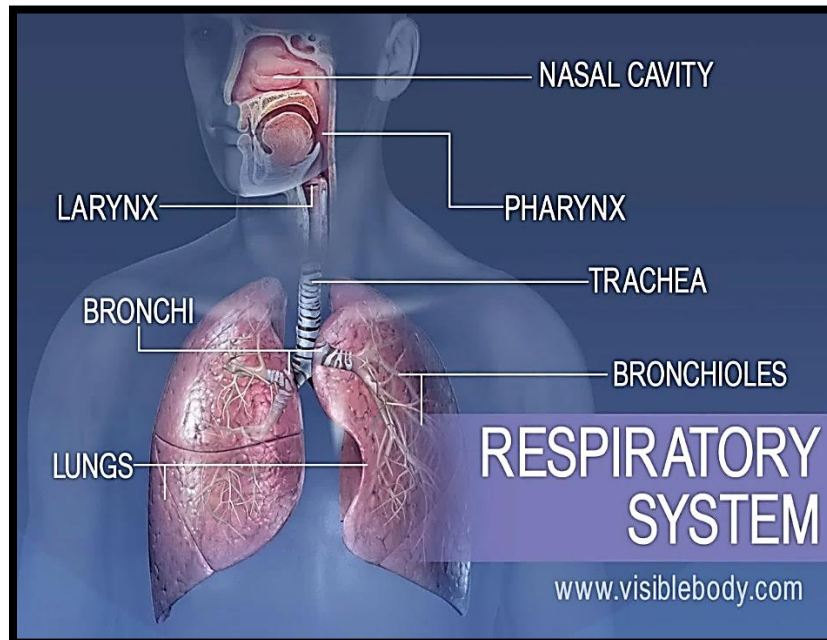
### Components of body system



### Circulatory system

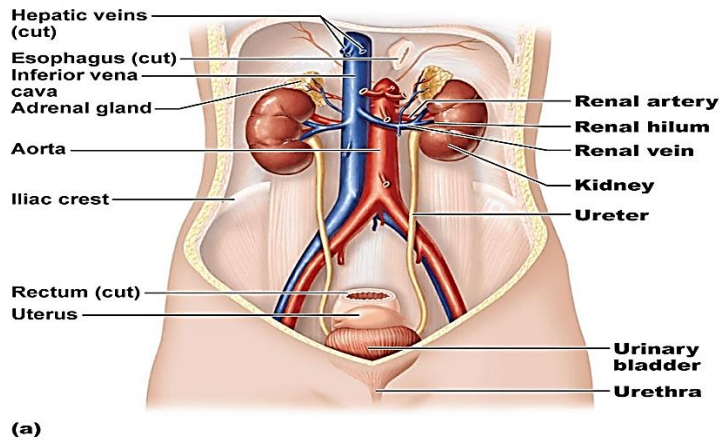


**Digestive System**



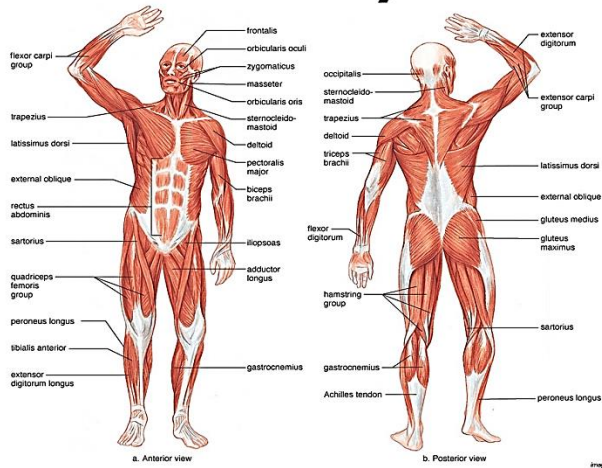
**Respiratory system**

# Urinary System

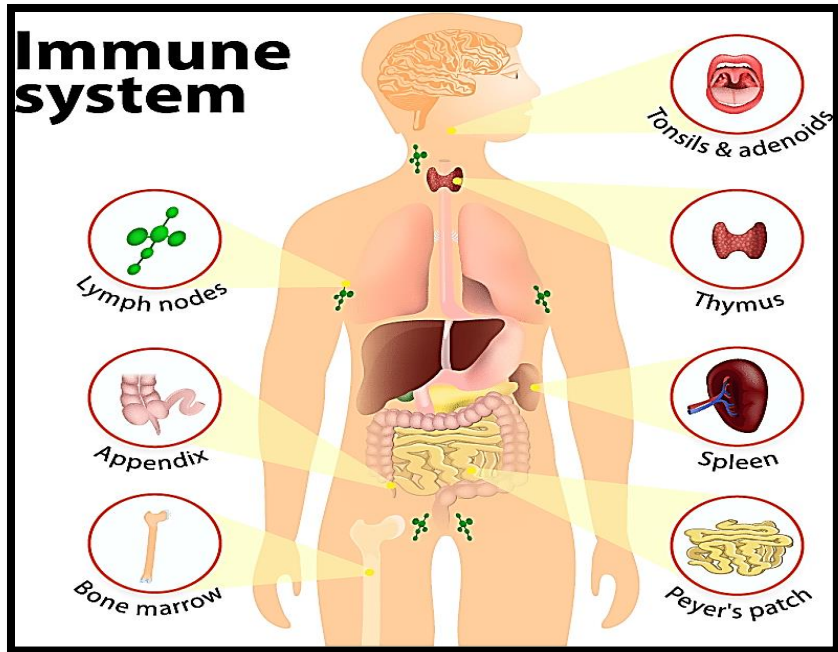


## Urinary system

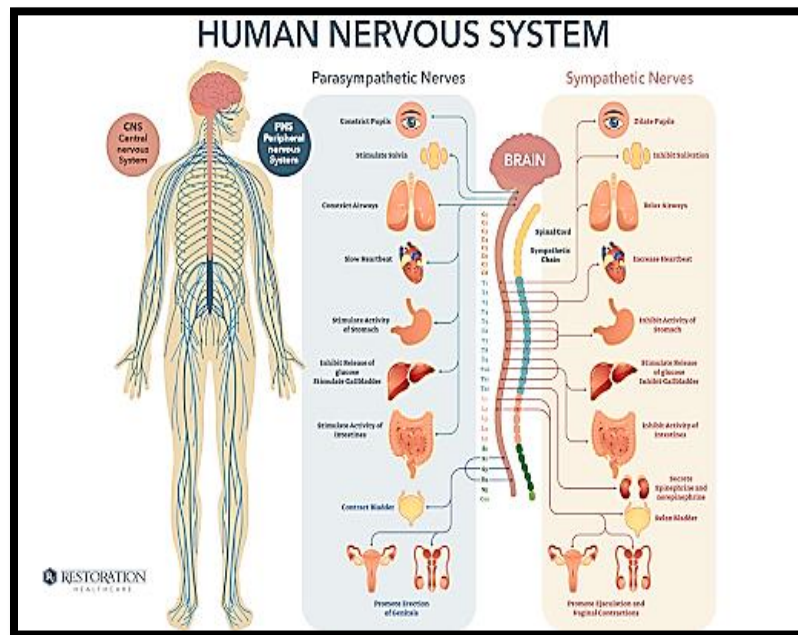
# Muscular System



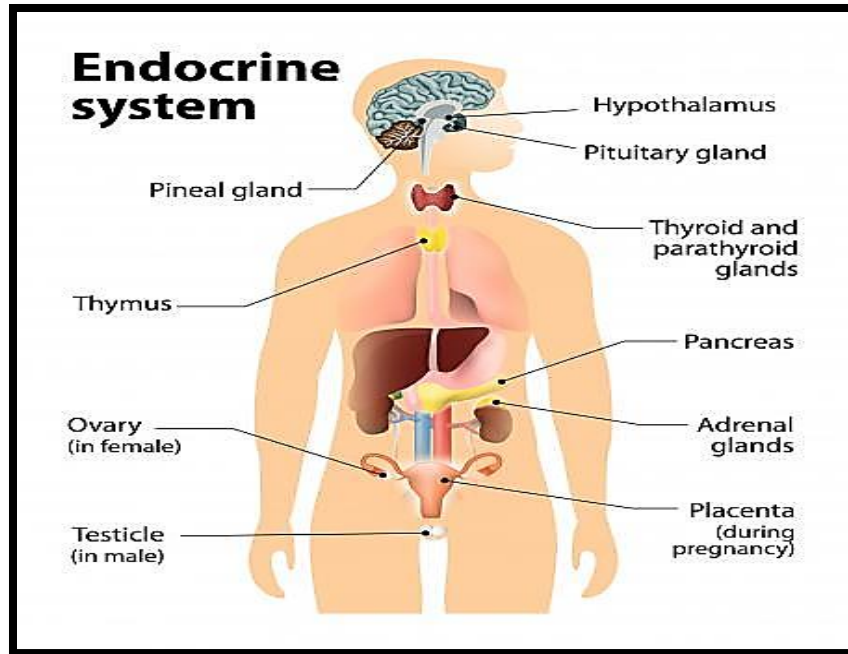
## Muscular system



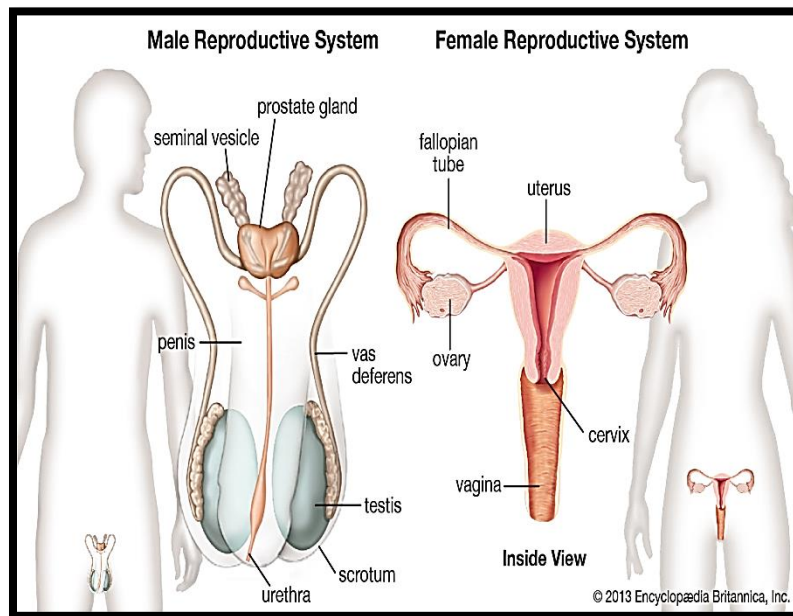
## Immune system



## Nervous system

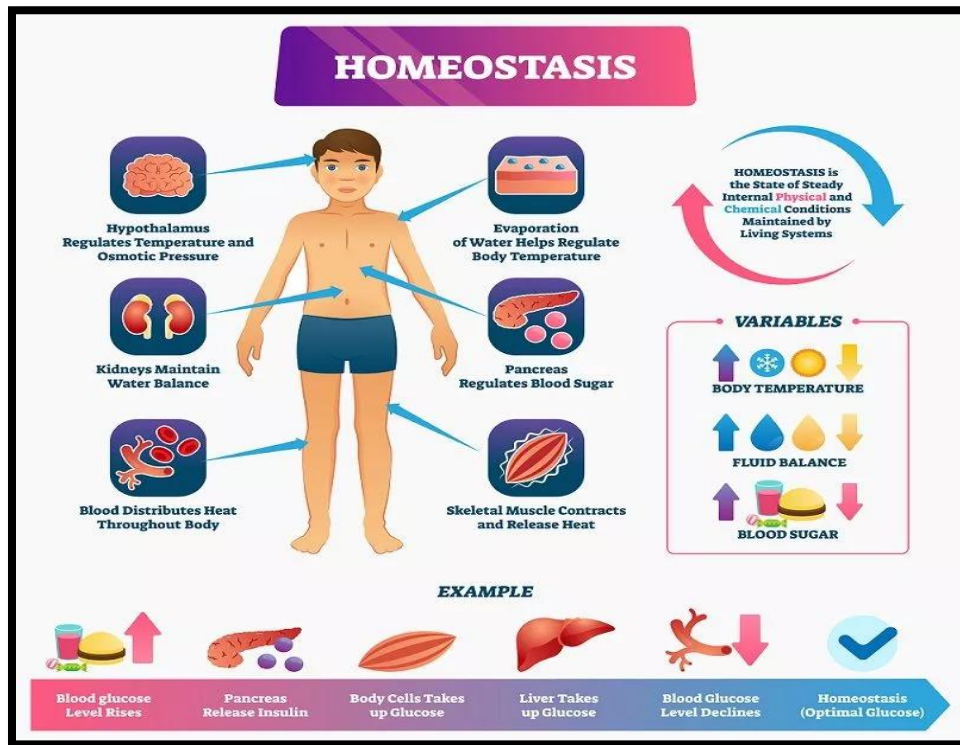


**Endocrine system**



**Reproductive system**

# Homeostasis



Role of body system in maintaining homeostasis:

## Nervous system:

Information from the external environment, also acts through electrical signals to control rapid responses for higher functions such as concentration, memory, and creativity.

## Endocrine system:

Acts by hormones secreted into the blood to control processes that require duration rather than speed, such as metabolic activity, water and electrolytes balances.

## Circulatory system:

Transports nutrients, oxygen, CO<sub>2</sub>, wastes, electrolytes and hormones through the body.

## Respiratory system:

Obtains oxygen and eliminates CO<sub>2</sub> to the external environment; helps regulate pH by adjusting the rate of removal of acid-forming carbon dioxide.

## **Urinary system:**

Important in regulating the volume, electrolyte composition, and pH of the internal environment; removes waste and excess water, salt, acid and other electrolytes from the plasma and eliminate them into the urine.

## **Digestive system:**

Obtains nutrients, water and electrolytes from the external environment and transfers them into the plasma; eliminates undigested food residues to the external environment.

## **Muscular and skeletal system:**

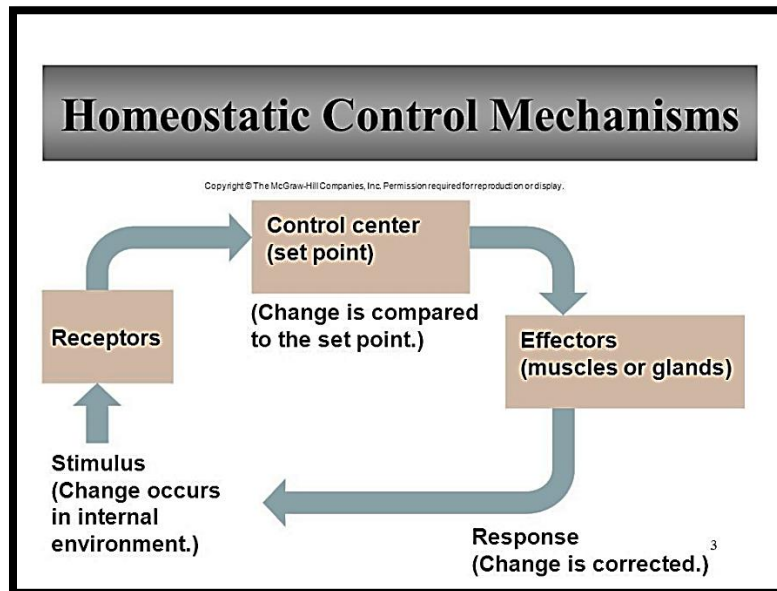
Supports and protects body parts and allows body movements, heat generated by muscular contraction are important in temperature regulation, calcium stored in the bones.

## **Immune system:**

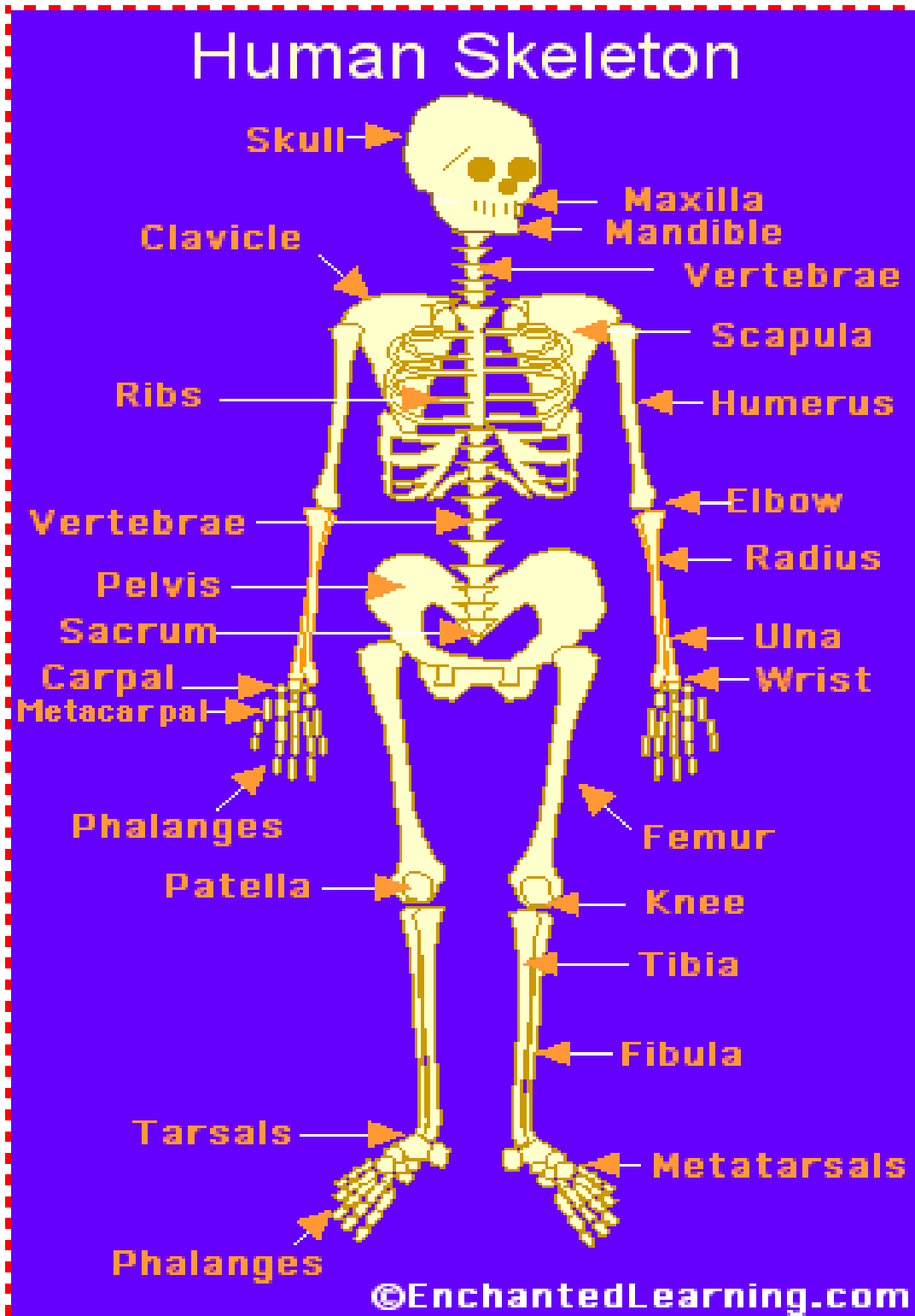
Defense against foreign invaders and cancer cells; tissue repair.

## **Integumentary system:**

Keeps internal fluids in and foreign materials out serves as a protective barrier between the external environment and the remainder of the body; temperature regulation.



# The skeletal system





The skeletal system is composed of bones and cartilage connected by ligaments to form a framework for the rest of the body tissues. There are two parts to the skeleton:

- **Axial skeleton** – bones along the axis of the body, including the skull, vertebral column and ribcage;
- **Appendicular skeleton** – appendages, such as the upper and lower limbs, pelvic girdle and shoulder girdle.

## **Function**

As well as contributing to the body's overall shape, the skeletal system has several key functions, including:

- Support and movement;
- Protection;
- Mineral homeostasis;
- Blood-cell formation;
- Triglyceride storage.

### **Support and movement**

Bones are a site of attachment for ligaments and tendons, providing a skeletal framework that can produce movement through the coordinated use of levers, muscles, tendons and ligaments. The bones act as levers, while the muscles generate the forces responsible for moving the bones.

### **Protection**

Bones provide protective boundaries for soft organs: the cranium around the brain, the vertebral column surrounding the spinal cord, the ribcage containing the heart and lungs, and the pelvis protecting the urogenital organs.

### **Mineral homeostasis**

As the main reservoirs for minerals in the body, bones contain approximately 99% of the body's calcium, 85% of its phosphate and 50% of its magnesium (Bartl and Bartl, 2017). They are essential in maintaining homeostasis of minerals in the blood with minerals stored in the bone are released in response to the body's demands, with levels maintained and regulated by hormones, such as parathyroid hormone.

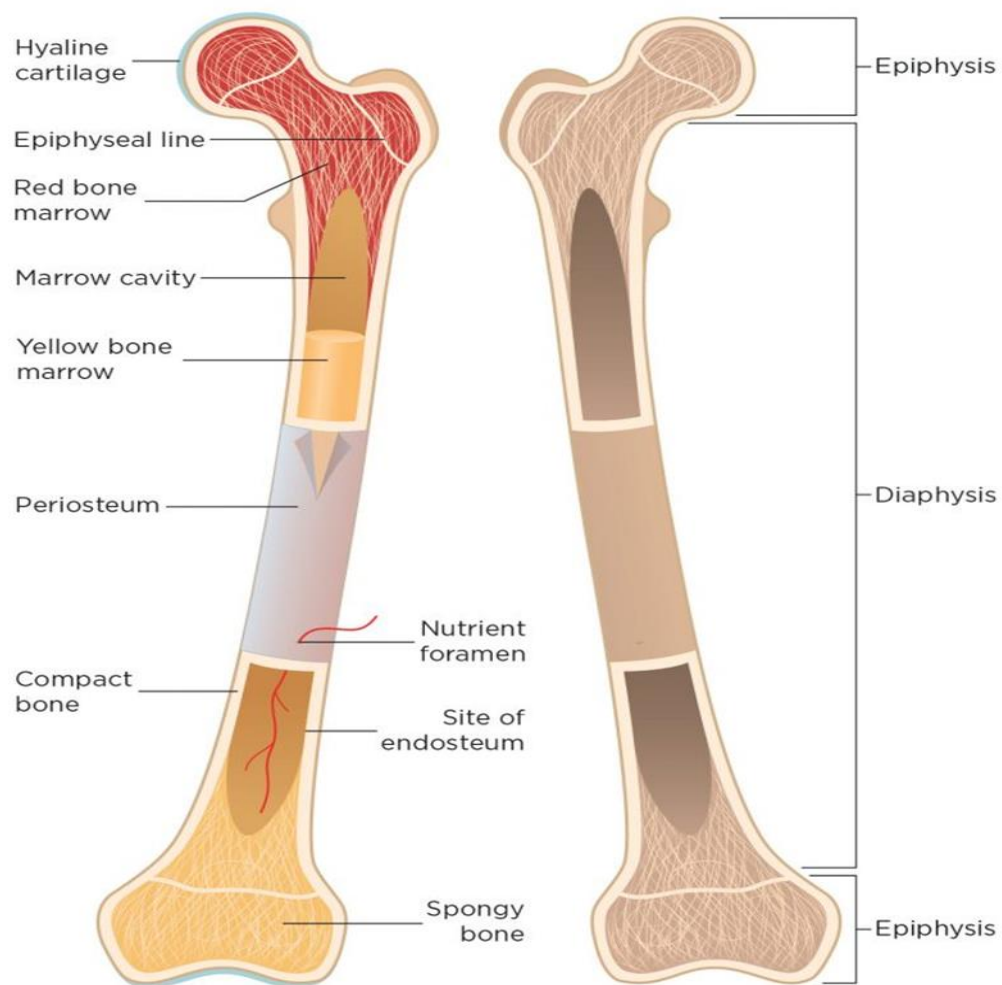
### **Blood-cell formation (haemopoiesis)**

Blood cells are formed from haemopoietic stem cells present in red bone marrow. Babies are born with only red bone marrow; over time this is replaced by yellow marrow due to a decrease in erythropoietin, the hormone responsible for stimulating the production of erythrocytes (red blood cells) in the bone marrow. By adulthood, the amount of red marrow has halved, and this reduces further to around 30% in older age (Robson and Syndercombe Court, 2018).

### **Triglyceride storage**

Yellow bone marrow (Fig 1) acts as a potential energy reserve for the body; it consists largely of adipose cells, which store triglycerides (a type of lipid that occurs naturally in the blood) (Tortora and Derrickson, 2009).

Fig 1. **Bone structure**



## **Development and structure of the skeleton**

Infants are born with about 300 separate bones, a nonprofit children's health provider. As a child grows, some of those bones fuse together until growth stops, typically by the age of 25, leaving the skeleton with 206 bones.

Our bones are separated into two categories based on the purpose and location of the bones: The axial skeleton and the appendicular skeleton, according to "Anatomy & Physiology."

The axial skeleton contains 80 bones, including the skull, spine and rib cage. It forms the central structure of the skeleton, with the function of protecting the brain, spinal cord, heart and lungs.

The remaining 126 bones make up the appendicular skeleton; they include the arms, legs, shoulder girdle and pelvic girdle. The lower portion of the appendicular skeleton protects the major organs associated with digestion and reproduction and provides

stability when a person is walking or running. The upper portion allows for a greater range of motion when lifting and carrying objects.

Bones are further classified by their shape: long, short, flat, irregular or sesamoid, according to "Anatomy & Physiology".

- Long bones are found in the arms, legs, fingers and toes. These bones are longer than they are wide and are cylindrical. They move when the muscles around them contract, and they are the most mobile parts of the skeleton.
- Short bones are found in the wrists and ankles and are about equal in their length, width and thickness.
- Flat bones make up the skull, shoulder blades, sternum and ribs. These curved, thin bones protect internal organs and provide an anchor for muscles.
- Irregular bones are those in the spinal cord and face, which, because of their unique dimension, don't fit in any of the other shape categories.
- Sesamoid bones are found in the hands, wrists, feet, ears and knees. These small, round bones are embedded in tendons and protect them from the great pressure and force they encounter.

There are some variations between male and female skeletons. For example, the female pelvis is typically more broad, thin, and round than the male pelvis, according to "Anatomy & Physiology."

### **What's inside your bones?**

All about your body's skeleton, the framework of bones that keeps you together.

Three main types of material make up every bone in your body: compact bone, spongy bone and bone marrow.

Approximately 80% of every bone is compact bone, which is the hardest and strongest type of bone and is what allows the body to support its weight. Compact bone makes up the outer layers of the bone and protects the inner parts of the bones where many vital functions occur, such as bone marrow production. Compact bone consists primarily of cells called osteocytes. Microscopic passages in between the cells to allow nerves and blood vessels to pass through.

About 20% of each bone is spongy bone, which is filled with large holes and passages. Most often found toward the ends of individual bones, the spongy bone material is filled with bone marrow, nerves and blood vessels.

Two types of bone marrow fill the pores in spongy bone. Approximately half is red bone marrow, which is found mainly within flat bones such as shoulder blades and ribs. This is where all red and white blood cells and platelets (cells that help a cut stop bleeding) are made. Infant's bones contain all red bone marrow to produce enough blood cells to keep up with the youngsters' growth.

The other half of marrow is yellow bone marrow, which is found in long bones, such as thigh bones, and consists primarily of fat. Blood vessels run through both types of bone marrow to deliver nutrients and remove waste from the bones.

There are four main types of cells within bones: Osteoblasts, osteocytes, osteoclasts and lining cells.

**Osteoblasts** are cells that create new or repair existing bone material as the bones grow or break. The cells create a flexible material called osteoid and then fortify it with minerals to harden and strengthen it. When osteoblasts successfully finish their job, they retire to become osteocytes or lining cells.

**Osteocytes**, found in the compact bone, are responsible for exchanging minerals and communicating with other cells in the vicinity. They are formed from old osteoblasts that have gotten stuck in the center of bones.

**Osteoclasts** break down existing bone material and reabsorb it. These cells often work with osteoblasts to heal and reshape bone after a break (the osteoclasts break down the extra callus formed by the healing process) to make room for new blood vessels and nerves and to make bones thicker and stronger.

**Lining cells** are flat bone cells that completely cover the outside surface of bones. Their primary function is controlling the movement of minerals, cells and other materials into and out of the bones.

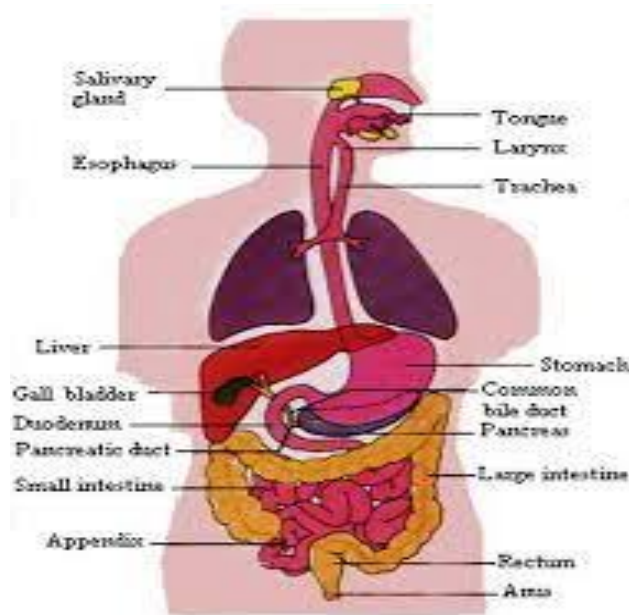
### **Diseases of the skeletal system**

As with any part of the human body, bones are susceptible to injury and disease.

Some of the most common diseases that can affect the skeletal system include:

- **Osteoporosis** is a disease that causes the density and strength of bones to decrease because bone loss occurs faster than bone growth. It can be caused by genetics or unhealthy lifestyle habits (such as lack of calcium or vitamin D, and heavy smoking or drinking with little exercise).
- **Leukemia** is a type of cancer that starts in the bone marrow and the lymphatic system. Several types of leukemia affect various blood cells and other systems of the body.
- **Osteoarthritis** is a disease that causes the breakdown of the cartilage that protects the ends of bones in joints. This lack of cartilage leads to bone-on-bone rubbing, which can cause significant pain, damage to the bones and connective tissues, inflammation of the surrounding tissue and restricted motion.

# The Digestive System



the digestive system uses mechanical and chemical activities to break food down into absorbable substances during its journey through the digestive system. Table 1 provides an overview of the basic functions of the digestive organs.

Main function of digestive system: The main function of the digestive system is to turn the food into simple sugars, amino acids, and carbohydrates. This is fuel for the human body.

**Table 1: Functions of the Digestive Organs**

Organ	Major functions	Other functions
Mouth	<ul style="list-style-type: none"> <li>• Ingests food</li> <li>• Chews and mixes food</li> <li>• Begins chemical breakdown of carbohydrates</li> <li>• Moves food into the pharynx</li> <li>• Begins breakdown of lipids via lingual lipase</li> </ul>	<ul style="list-style-type: none"> <li>• Moistens and dissolves food, allowing you to taste it</li> <li>• Cleans and lubricates the teeth and oral cavity</li> <li>• Has some antimicrobial activity</li> </ul>
Pharynx	<ul style="list-style-type: none"> <li>• Propels food from the oral cavity to the esophagus</li> </ul>	<ul style="list-style-type: none"> <li>• Lubricates food and passageways</li> </ul>
Esophagus	<ul style="list-style-type: none"> <li>• Propels food to the stomach</li> </ul>	<ul style="list-style-type: none"> <li>• Lubricates food and passageways</li> </ul>
Stomach	<ul style="list-style-type: none"> <li>• Mixes and churns food with gastric juices to form chyme</li> <li>• Begins chemical breakdown of proteins</li> <li>• Releases food into the duodenum as</li> </ul>	<ul style="list-style-type: none"> <li>• Stimulates protein-digesting enzymes</li> <li>• Secretes intrinsic factor required for vitamin B<sub>12</sub> absorption in small intestine</li> </ul>

Organ	Major functions	Other functions
	chyme <ul style="list-style-type: none"> <li>• Absorbs some fat-soluble substances (for example, alcohol, aspirin)</li> <li>• Possesses antimicrobial functions</li> </ul>	
Small intestine	<ul style="list-style-type: none"> <li>• Mixes chyme with digestive juices</li> <li>• Propels food at a rate slow enough for digestion and absorption</li> <li>• Absorbs breakdown products of carbohydrates, proteins, lipids, and nucleic acids, along with vitamins, minerals, and water</li> <li>• Performs physical digestion via segmentation</li> </ul>	<ul style="list-style-type: none"> <li>• Provides optimal medium for enzymatic activity</li> </ul>
Accessory organs	<ul style="list-style-type: none"> <li>• Liver: produces bile salts, which emulsify lipids, aiding their digestion and absorption</li> <li>• Gallbladder: stores, concentrates, and releases bile</li> <li>• Pancreas: produces digestive enzymes and bicarbonate</li> </ul>	<ul style="list-style-type: none"> <li>• Bicarbonate-rich pancreatic juices help neutralize acidic chyme and provide optimal environment for enzymatic activity</li> </ul>
Large intestine	<ul style="list-style-type: none"> <li>• Further breaks down food residues</li> <li>• Absorbs most residual water, electrolytes, and vitamins produced by enteric bacteria</li> <li>• Propels feces toward rectum</li> <li>• Eliminates feces</li> </ul>	<ul style="list-style-type: none"> <li>• Food residue is concentrated and temporarily stored prior to defecation</li> <li>• Mucus eases passage of feces through colon</li> </ul>

## Functions of the Digestive System

1- ingestion – the oral cavity allows food to enter the digestive tract and have mastication (chewing) occurs, and the resulting food bolus is swallowed.

2- Digestion:

♣ Mechanical digestion – muscular movement of the digestive tract (mainly in the oral cavity and stomach) physically break down food into smaller particles.

♣ chemical digestion – hydrolysis reactions aided by enzymes (mainly in the stomach and small intestine) chemically break down food particles into nutrient molecules, small enough to be absorbed.

♣ Secretion – enzymes and digestive fluids secreted by the digestive tract and its accessory organs facilitate chemical digestion.

♣ Absorption – passage of the end – products (nutrients) of chemical digestion from the digestive tract into blood or lymph for distribution to tissue cells.

♣ Elimination (defecation) – undigested material will be released through the rectum and anus by defecation. Copyright © 2006 Pearson Education, Inc., publishing as Benjamin Cummings

## Digestive Processes

The processes of digestion include six activities: ingestion, propulsion, mechanical or physical digestion, chemical digestion, absorption, and defecation.

The first of these processes, **ingestion**, refers to the entry of food into the alimentary canal through the mouth. There, the food is chewed and mixed with saliva, which contains enzymes that begin breaking down the carbohydrates in the food plus some lipid digestion via lingual lipase. Chewing increases the surface area of the food and allows an appropriately sized bolus to be produced.

Food leaves the mouth when the tongue and pharyngeal muscles propel it into the esophagus. This act of swallowing, the last voluntary act until defecation, is an example of propulsion, which refers to the movement of food through the digestive tract. It includes both the voluntary process of swallowing and the involuntary process of peristalsis. Peristalsis consists of sequential, alternating waves of contraction and relaxation of alimentary wall smooth muscles, which act to propel food along (Figure 1). These waves also play a role in mixing food with digestive juices. Peristalsis is so powerful that foods and liquids you swallow enter your stomach even if you are standing on your head.

### Peristalsis

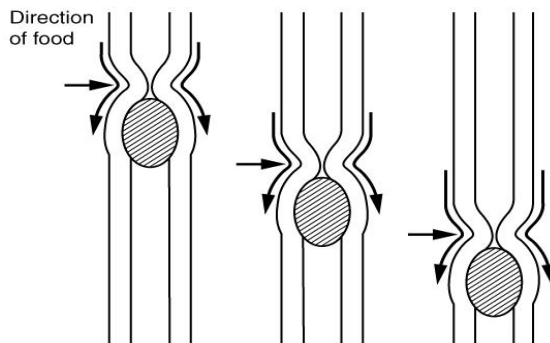


Figure 1: Peristalsis moves food through the digestive tract with alternating waves of muscle contraction and relaxation.

Digestion includes both mechanical and chemical processes.

**Mechanical digestion** is a purely physical process that does not change the chemical nature of the food. Instead, it makes the food smaller to increase both surface area and mobility. It includes mastication, or chewing, as well as tongue movements that help break food into smaller bits and mix food with saliva. Although there may be a tendency to think that mechanical digestion is limited to the first steps of the digestive process, it occurs after the food leaves the mouth, as well. The mechanical churning of food in the stomach serves to further break it apart and expose more of its surface area to digestive juices, creating an acidic “soup”

called chyme. Segmentation, which occurs mainly in the small intestine, consists of localized contractions of circular muscle of the muscularis layer of the alimentary canal. These contractions isolate small sections of the intestine, moving their contents back and forth while continuously subdividing, breaking up, and mixing the contents. By moving food back and forth in the intestinal lumen, segmentation mixes food with digestive juices and facilitates absorption.

**chemical digestion**, starting in the mouth, digestive secretions break down complex food molecules into their chemical building blocks (for example, proteins into separate amino acids). These secretions vary in composition, but typically contain water, various enzymes, acids, and salts. The process is completed in the small intestine.

Food that has been broken down is of no value to the body unless it enters the bloodstream and its nutrients are put to work. This occurs through the process of absorption, which takes place primarily within the small intestine. There, most nutrients are absorbed from the lumen of the alimentary canal into the bloodstream through the epithelial cells that make up the mucosa. Lipids are absorbed into lacteals and are transported via the lymphatic vessels to the bloodstream (the subclavian veins near the heart). The details of these processes will be discussed later.

**defecation**, the final step in digestion, undigested materials are removed from the body as feces.

in some cases, a single organ is in charge of a digestive process. For example, ingestion occurs only in the mouth and defecation only in the anus. However, most digestive processes involve the interaction of several organs and occur gradually as food moves through the alimentary canal (Figure 2).

## Digestive Processes

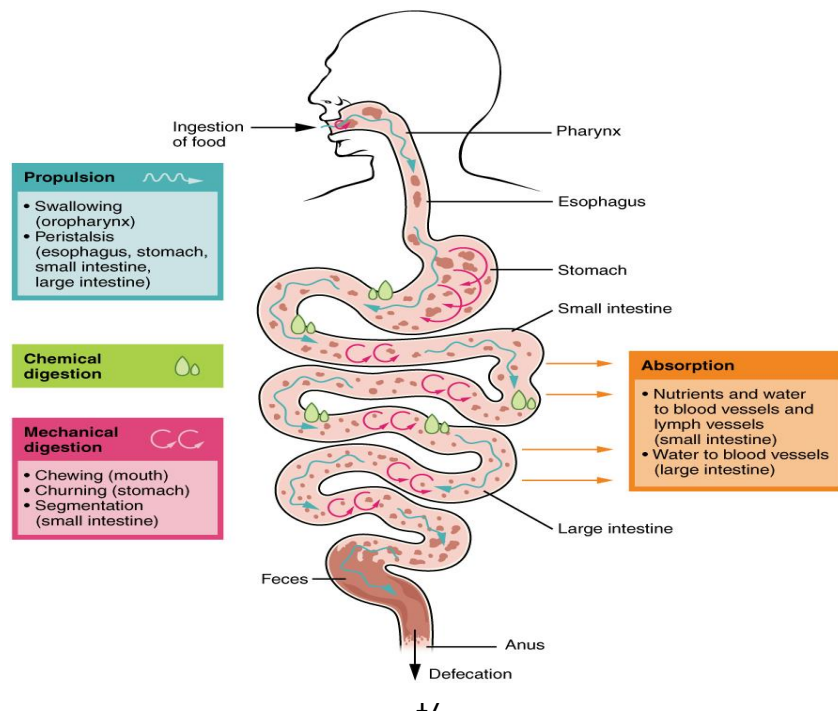




Figure 2: The digestive processes are ingestion, propulsion, mechanical digestion, chemical digestion, absorption, and defecation.

Some chemical digestion occurs in the mouth. Some absorption can occur in the mouth and stomach, for example, alcohol and aspirin.

## **Regulatory Mechanisms**

Neural and endocrine regulatory mechanisms work to maintain the optimal conditions in the lumen needed for digestion and absorption. These regulatory mechanisms, which stimulate digestive activity through mechanical and chemical activity, are controlled both extrinsically and intrinsically.

### **Neural Controls**

The walls of the alimentary canal contain a variety of sensors that help regulate digestive functions. These include mechanoreceptors, chemoreceptors, and osmoreceptors, which are capable of detecting mechanical, chemical, and osmotic stimuli, respectively. For example, these receptors can sense when the presence of food has caused the stomach to expand, whether food particles have been sufficiently broken down, how much liquid is present, and the type of nutrients in the food (lipids, carbohydrates, and/or proteins). Stimulation of these receptors provokes an appropriate reflex that furthers the process of digestion. This may entail sending a message that activates the glands that secrete digestive juices into the lumen, or it may mean the stimulation of muscles within the alimentary canal, thereby activating peristalsis and segmentation that move food along the intestinal tract.

The walls of the entire alimentary canal are embedded with nerve plexuses that interact with the central nervous system and other nerve plexuses—either within the same digestive organ or in different ones. These interactions prompt several types of reflexes. Extrinsic nerve plexuses orchestrate long reflexes, which involve the central and autonomic nervous systems and work in response to stimuli from outside the digestive system. Short reflexes, on the other hand, are orchestrated by intrinsic nerve plexuses within the alimentary canal wall. These two plexuses and their connections were introduced earlier as the enteric nervous system. Short reflexes regulate activities in one area of the digestive tract and may coordinate local peristaltic movements and stimulate digestive secretions. For example, the sight, smell, and taste of food initiate long reflexes that begin with a sensory neuron delivering a signal to the medulla oblongata. The response to the signal is to stimulate cells in the stomach to begin secreting digestive juices in preparation for incoming food. In contrast, food that distends the stomach initiates short reflexes that cause cells in the stomach wall to increase their secretion of digestive juices.

### **Hormonal Controls**

A variety of hormones are involved in the digestive process. The main digestive hormone of the stomach is gastrin, which is secreted in response to the presence of food. Gastrin stimulates the secretion of gastric acid by the parietal cells of the stomach mucosa. Other GI hormones are produced and act upon the gut and its accessory organs. Hormones produced by the duodenum

include secretin, which stimulates a watery secretion of bicarbonate by the pancreas; cholecystokinin (CCK), which stimulates the secretion of pancreatic enzymes and bile from the liver and release of bile from the gallbladder; and gastric inhibitory peptide, which inhibits gastric secretion and slows gastric emptying and motility. These GI hormones are secreted by specialized epithelial cells, called endocrinocytes, located in the mucosal epithelium of the stomach and small intestine. These hormones then enter the bloodstream, through which they can reach their target organs.

# The Circulatory System

(The Heart, Blood Vessels, Blood Types)

**Circulatory systems generally have three main features:**

- Fluid (blood or lymph) that transports materials.
- System of blood vessels.
- A heart to pump the fluid through the vessels.

**Types of circulatory systems:**

- Animals that have a circulatory system have one of two kinds:  
Open: fluid is circulated through an open body chamber.  
Closed: fluid is circulated through blood vessels.

**Function**

- Transport materials needed by cells (Oxygen- Glucose).
- Remove waste materials from cells (Carbon dioxide- Urea)

**Major Components**

**Blood:** Fluid that fills the circulatory system

**Pump (heart):** Continuously circulates blood

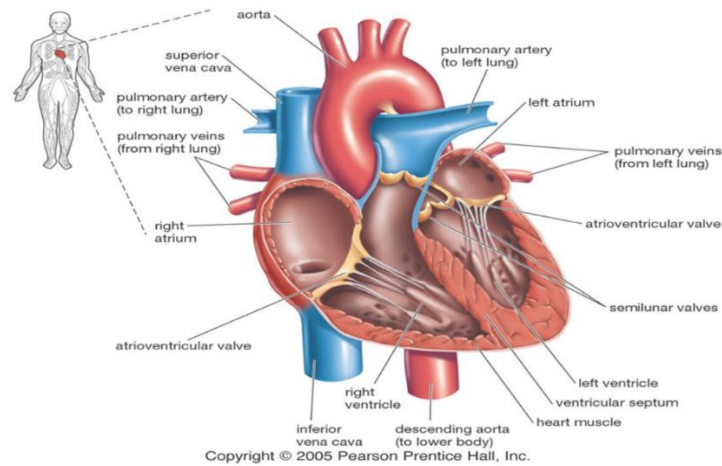
**Network of tubes** (blood vessels)

- Arteries- blood away from heart
- Veins- blood back to the heart
- Capillaries- link Arteries with Veins

**1-The Heart**

The human heart has four chambers (Left and right ventricle - Left and right atrium).

The left side of the heart pumps oxygenated blood to the body while the right side of the heart pumps deoxygenated blood to the lungs where oxygen can be absorbed by the hemoglobin carrying red blood cells.



## Functions of the Heart

- Generating blood pressure.
- Routing blood.
- Heart separates pulmonary and systemic circulations.
- Ensuring one-way blood flow.
- Heart valves ensure one-way flow.
- Regulating blood supply.

## Size, Shape, Location of the Heart

### Shape:

**Apex:** Blunt rounded point of cone.

**Base:** Flat part at opposite of end of cone.

- Size of a closed fist
- Located in thoracic cavity between two lungs

### External Anatomy

- **Four chambers**

2 atria

2 ventricles

- **Major veins**

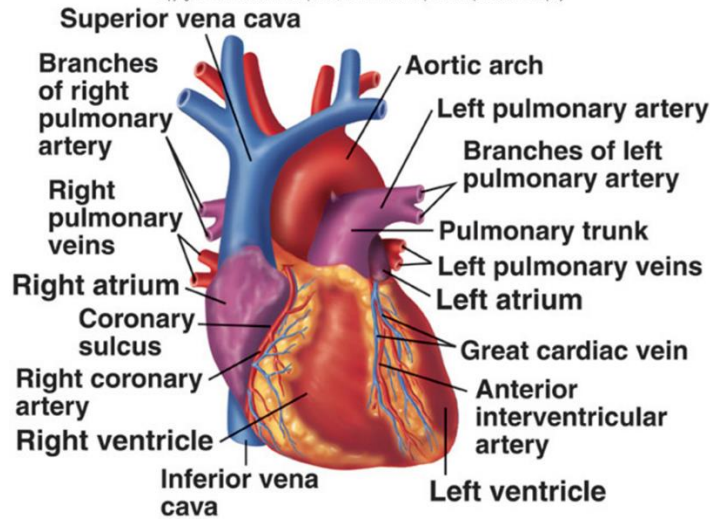
Superior and inferior vena cava

Pulmonary veins

- **Major arteries**

Aorta

Pulmonary trunk



## Heart Valves

### - Atrioventricular valves

Tricuspid

Bicuspid or mitral

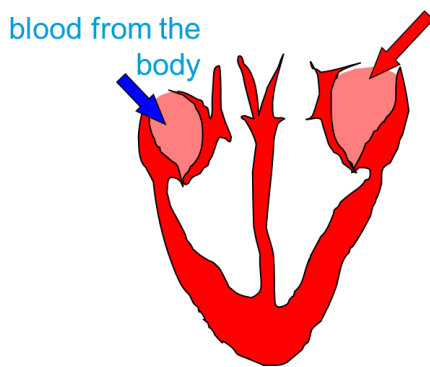
### - Semilunar valves

Aortic

Pulmonary

Prevent blood from flowing back

## How does the Heart work? Step one

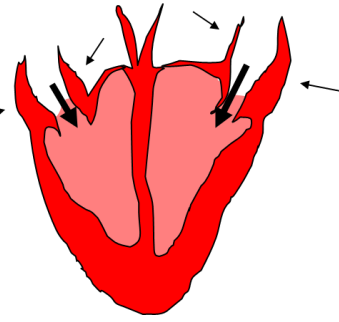


blood from the lungs

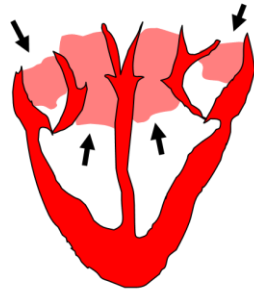
The heart beat begins when the heart muscles **relax** and blood flows into the atria.

## STEP TWO

The atria then **contract** and the valves **open** to allow blood into the ventricles.



### STEP THREE



The valves **close** to stop blood flowing backwards.

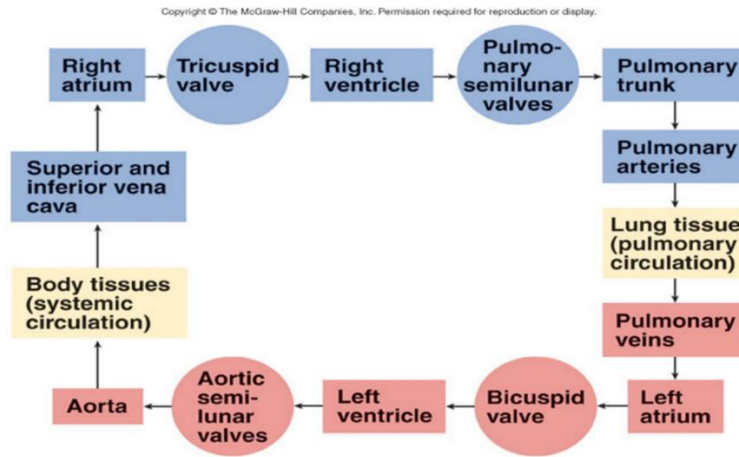
The ventricles **contract** forcing the blood to leave the heart.

At the same time, the atria are **relaxing** and once again filling with blood

The cycle then repeats itself.

### Circulation

After passing through the capillaries of the lungs, the blood which is now oxygenated returns to the heart in the pulmonary veins. The left atrium receives blood from the pulmonary vein. Blood passes through the mitral valve into the left ventricle. Contraction of the left ventricle pushes blood through the aortic semilunar valve into the aorta. Blood travels to all regions of the body where it feeds cells with oxygen picked up from the lungs and nutrients from the digestive tract. Deoxygenated blood returns from the rest of the body through the superior and inferior vena cava. Contraction of the right ventricle pushes blood through the pulmonary semilunar valve into the pulmonary arteries in which it travels to the lungs.



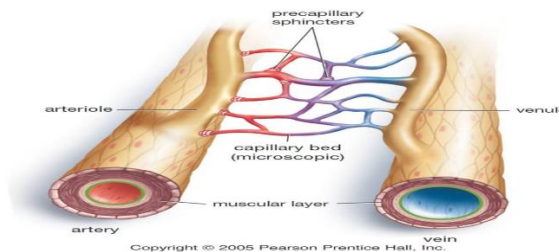
## Blood Pressure

Blood pressure is a measure of the force exerted by the blood on the wall of the arteries. An example is 120/80 (systolic pressure/diastolic pressure). Systolic pressure is the result of the contraction of the ventricles (normal 110-140). Diastolic pressure is during the ventricle relaxation (normal 70-90)

## 2-Blood Vessels:

Blood vessels fall into three major classes:

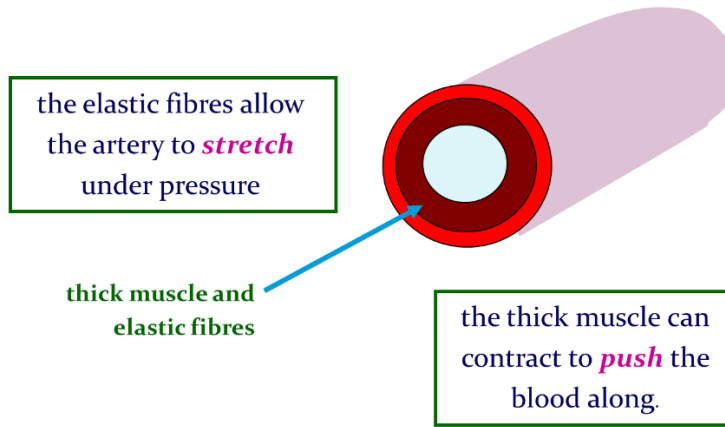
- Arteries and arterioles carry blood away from the heart.
- Veins and venules carry blood to the heart.
- Capillaries allow exchange of nutrients, wastes and gases.



## a-The ARTERY

Arteries are thick-walled and lined with smooth muscle. How does the structure of an artery help with its function?

Arteries carry blood away from the heart.

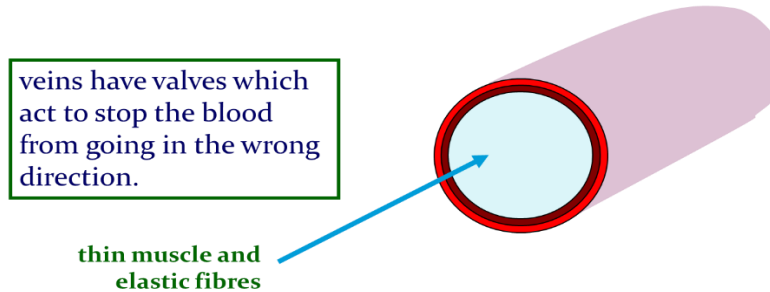


Arteries carry blood away from the heart.

### **b-The Vein**

Veins have thinner walls than arteries. Veins have fewer smooth muscle cells but do have valves. How do valves and the skeletal muscles help vein's function?

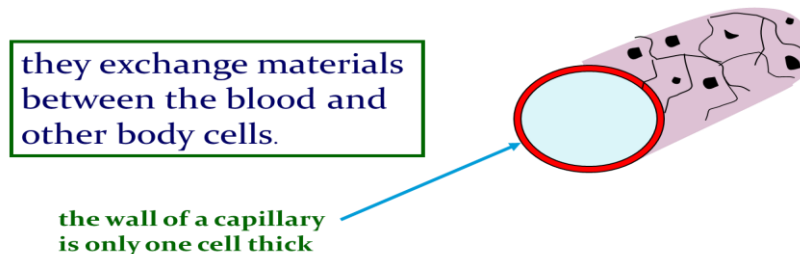
Veins carry blood towards from the heart.



### **c- The Capillary**

Body tissues contain a vast network of thin capillaries. Capillary walls are only one cell thick, allowing exchange of gases, nutrients, and wastes.

Capillaries link Arteries with Veins



### **3- Blood Components:**

Blood is made up of four major components. What do each of these do?

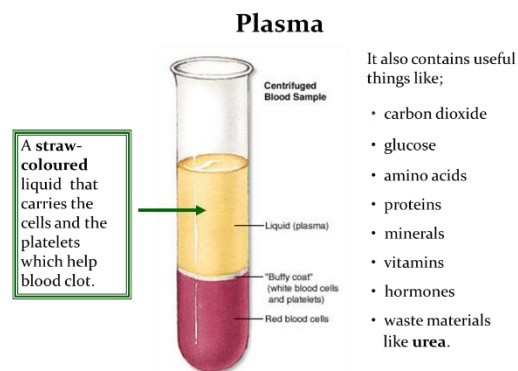
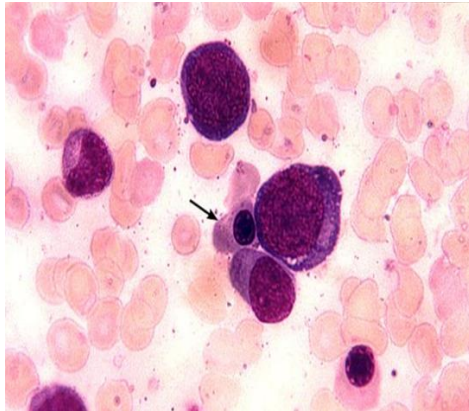
- Plasma: the liquid portion.



- Red blood cells (RBCs).
- White blood cells (WBCs).
- Platelets (PLT).

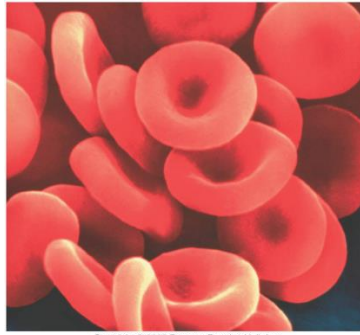
### A-) Plasma

Liquid portion of the blood. Contains clotting factors, hormones, antibodies, dissolved gases, nutrients, and waste.



### B.) Red Blood Cells

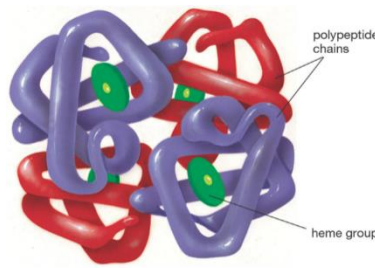
- A biconcave disc that is round and flat.
- Carry hemoglobin and oxygen. Do not have a nucleus and live only about 120 days.
- Can change shape to an amazing extent, without breaking, as it squeezes single file through the capillaries.
- Cannot repair themselves.
- Make up about 99% of the blood's cellular component.
- Red color is due to hemoglobin.



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## Hemoglobin

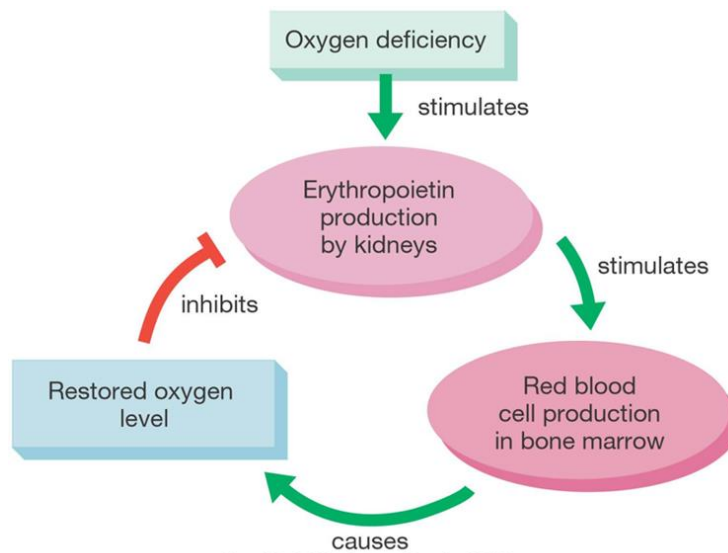
Hemoglobin is a complex protein made up of four protein strands, plus iron-rich heme groups. Each hemoglobin molecule can carry four oxygen atoms. The presence of oxygen turns hemoglobin bright red.



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## RBC lifespan

RBCs live about 4 months. Iron from hemoglobin is recycled in the liver and spleen. The hormone erythropoietin, made by the kidneys, stimulates the production of RBCs in red bone marrow.

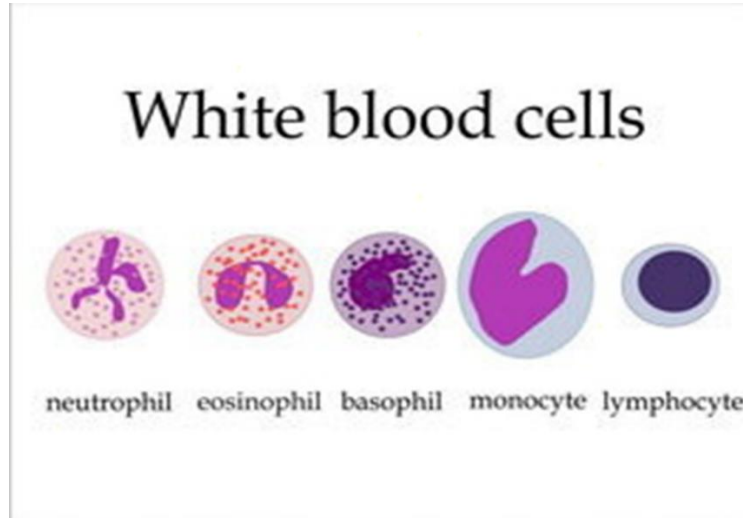


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## C.) White Blood cells

- Fight infection and are formed in the bone marrow.

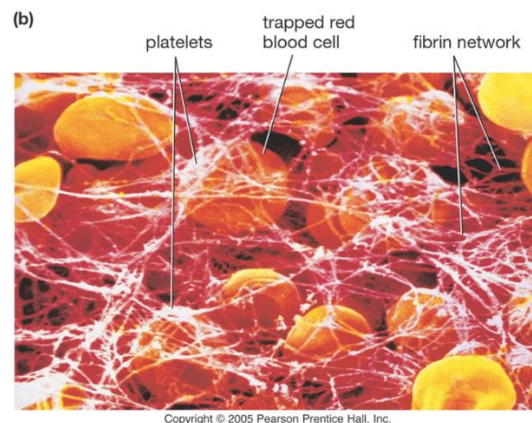
- White blood cells defend against disease by recognizing proteins that do not belong to the body.
- White cells can ooze through the walls of capillaries to patrol the tissues and reach the lymph system
- Have five types (neutrophils, lymphocytes, eosinophils, basophils, and monocytes).



the two main ones are the lymphocytes and the macrophages, macrophages 'eat' and digest micro-organisms, some lymphocytes fight disease by making antibodies to destroy invaders by dissolving them. other lymphocytes make antitoxins to break down poisons.

#### **D) Platelets**

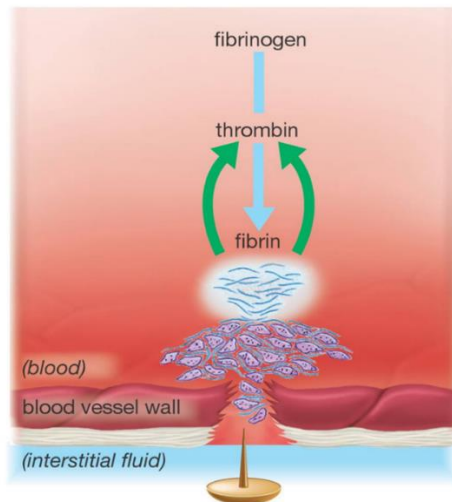
Platelets are cell fragments used in blood clotting. Platelets are derived from megakaryocytes. Because they lack a nucleus, platelets have a short lifespan, usually about 10 days.



#### **Blood clotting**

- 1- Platelets aggregate at the site of a wound.
- 2- Broken cells and platelets release chemicals to stimulate thrombin production.
- 3- Thrombin converts the protein fibrinogen into sticky fibrin, which binds the clot.

(a)



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## Disorders of the Circulatory System

- Anemia - lack of iron in the blood, low RBC count.
- Leukemia - white blood cells proliferate wildly, causing anemia.
- Hemophilia - bleeder's disease, due to lack of fibrinogen in thrombocytes.
- Heart Murmur - abnormal heartbeat, caused by valve problems.
- Heart attack - blood vessels around the heart become blocked with plaque, also called myocardial infarction.

## The Neuromuscular System

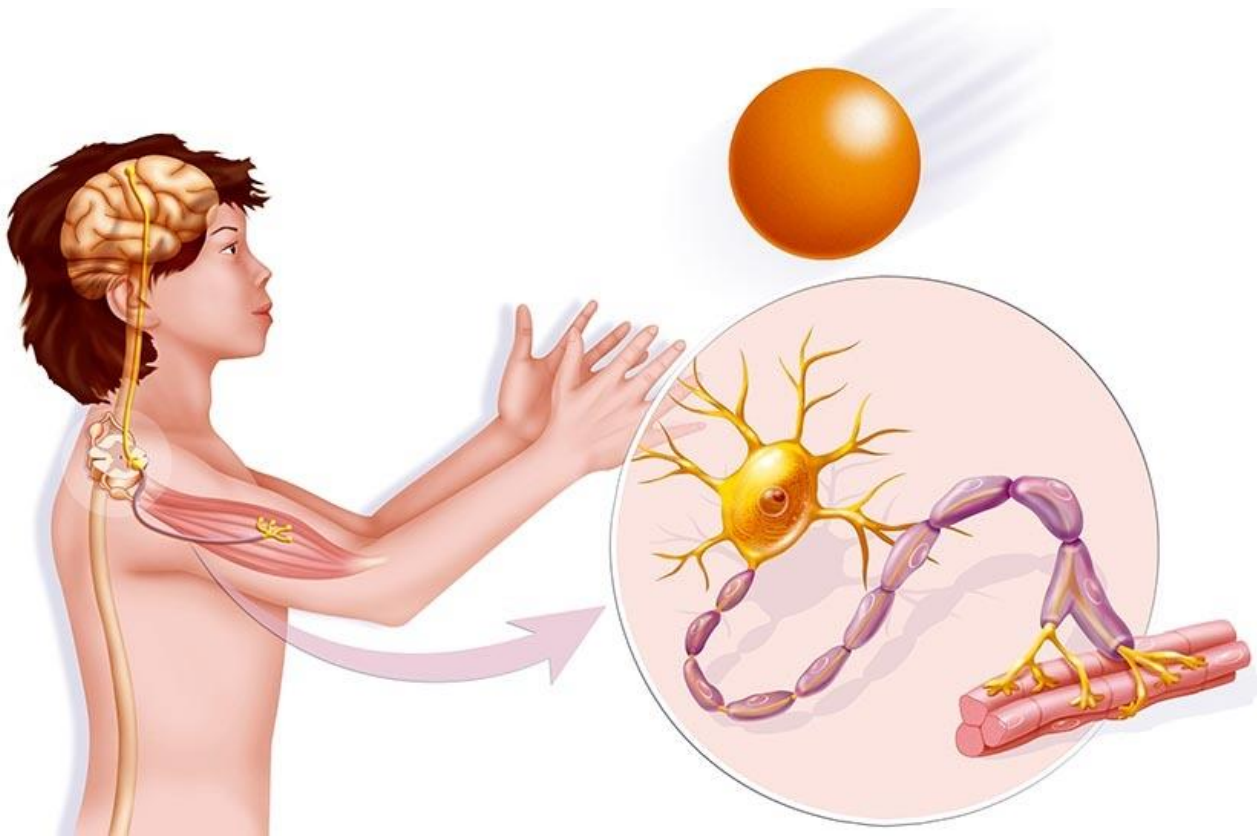
The neuromuscular system includes all the muscles in the body and the nerves serving them. Every movement the body makes requires communication between the brain and the muscles. The nervous system provides the link between thoughts and actions by relaying messages from the brain to other parts of the body.

Nerves and muscles, working together as the neuromuscular system, make the body move as you want it to and also control functions such as breathing.

### **How does the neuromuscular system work?**

Nerves have cells called neurons. Neurons carry messages from the brain via the spinal cord. The neurons that carry these messages to the muscles are called motor neurons.

Each motor neuron ending sits very close to a muscle fibre. Where they sit together is called a neuromuscular junction. The motor neurons release a chemical, which is picked up by the muscle fiber. This tells the muscle fiber to contract, which makes the muscles move.



## The Muscular System

The muscular system controls numerous functions, which is possible with the significant differentiation of muscle tissue morphology and ability.

### Key Points

- The muscular system is responsible for functions such as maintenance of posture, locomotion, and control of various circulatory systems.
- Muscle tissue can be divided functionally (voluntarily or involuntarily controlled) and morphologically (striated or non-striated).
- These classifications describe three distinct muscle types: skeletal, cardiac and smooth. Skeletal muscle is voluntary and striated, cardiac muscle is involuntary and striated, and smooth muscle is involuntary and non-striated.

### Key Terms

- **myofibril:** A fiber made up of several myofilaments that facilitates the generation of tension in a myocyte.
- **myofilament:** A filament composed of either multiple myosin or actin proteins that slide over each other to generate tension.
- **myosin:** A motor protein which forms myofilaments that interact with actin filaments to generate tension.
- **actin:** A protein which forms myofilaments that interact with myosin filaments to generate tension.
- **striated:** The striped appearance of certain muscle types in which myofibrils are aligned to produce a constant directional tension.
- **voluntary:** A muscle movement under conscious control (e.g. deciding to move the forearm).
- **involuntary:** A muscle movement not under conscious control (e.g. the beating of the heart).
- **myocyte:** A muscle cell.

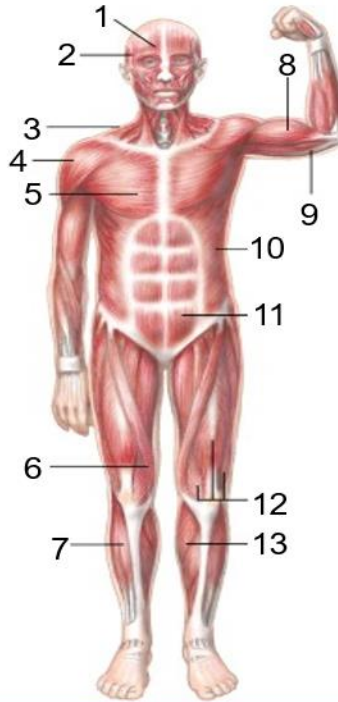
## The Musculoskeletal System

The muscular system is made up of muscle tissue and is responsible for functions such as maintenance of posture, locomotion and control of various circulatory systems. This includes the beating of the heart and the movement of food through the digestive system. The muscular system is closely associated with the skeletal system in facilitating movement. Both voluntary and involuntary muscular system functions are controlled by the nervous system.

## Section 11.2 Your Muscular System

### The Muscular System

- 1) Frontalis
- 2) Temporalis
- 3) Trapezius
- 4) Deltoid
- 5) Pectoralis major
- 6) Sartorius
- 7) Tibialis anterior



- 8) Biceps
- 9) Triceps
- 10) External oblique
- 11) Rectus abdominus
- 12) Quadriceps muscles
- 13) Gastrocnemius

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Muscle is a highly-specialized soft tissue that produces tension which results in the generation of force. Muscle cells, or myocytes, contain myofibrils comprised of actin and myosin myofilaments which slide past each other producing tension that changes the shape of the myocyte. Numerous myocytes make up muscle tissue and the controlled production of tension in these cells can generate significant force.

Muscle tissue can be classified functionally as voluntary or involuntary and morphologically as striated or non-striated. Voluntary refers to whether the muscle is under conscious control, while striation refers to the presence of visible banding within myocytes caused by the organization of myofibrils to produce constant tension.

### Types of Muscle

The above classifications describe three forms of muscle tissue that perform a wide range of diverse functions.

### Skeletal Muscle

Skeletal muscle mainly attaches to the skeletal system via tendons to maintain posture and control movement. For example, contraction of the biceps muscle, attached to the scapula and radius, will raise the forearm. Some skeletal muscle can attach directly to other muscles or to the skin, as seen in the face where numerous muscles control facial expression.

Skeletal muscle is under voluntary control, although this can be subconscious when maintaining posture or balance. Morphologically skeletal myocytes are elongated and tubular and appear striated with multiple peripheral nuclei.

### **Cardiac Muscle Tissue**

Cardiac muscle tissue is found only in the heart, where cardiac contractions pump blood throughout the body and maintain blood pressure.

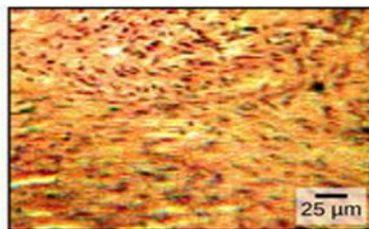
As with skeletal muscle, cardiac muscle is striated; however it is not consciously controlled and so is classified as involuntary. Cardiac muscle can be further differentiated from skeletal muscle by the presence of intercalated discs that control the synchronized contraction of cardiac tissues. Cardiac myocytes are shorter than skeletal equivalents and contain only one or two centrally located nuclei.

### **Smooth Muscle Tissue**

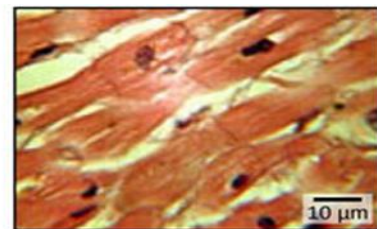
Smooth muscle tissue is associated with numerous organs and tissue systems, such as the digestive system and respiratory system. It plays an important role in the regulation of flow in such systems, such as aiding the movement of food through the digestive system via peristalsis. Smooth muscle is non-striated and involuntary. Smooth muscle myocytes are spindle shaped with a single centrally located nucleus.



Skeletal muscle



Smooth muscle



Cardiac muscle

**Types of muscle:** The body contains three types of muscle tissue: skeletal muscle, smooth muscle, and cardiac muscle, visualized here using light microscopy. Visible striations in skeletal and cardiac muscle are visible, differentiating them from the more randomized appearance of smooth muscle.

### **Function of Muscle Tissue**

The main function of the muscular system is movement. Muscles are the only tissue in the body that has the ability to contract and therefore move the other parts of the body.

Related to the function of movement is the muscular system's second function: the maintenance of posture and body position. Muscles often contract to hold the body still or in a particular position rather than to cause movement. The muscles responsible for the body's posture have the greatest endurance of all muscles in the body—they hold up the body throughout the day without becoming tired.

Another function related to movement is the movement of substances inside the body. The cardiac and visceral muscles are primarily responsible for transporting substances like blood or food from one part of the body to another.

The final function of muscle tissue is the generation of body heat. As a result of the high metabolic rate of contracting muscle, our muscular system produces a great deal of waste heat. Many small muscle contractions within the body produce our natural body heat. When we exert ourselves more than normal, the extra muscle contractions lead to a rise in body temperature and eventually to sweating.



# The Nervous System

## **Basic Structure and Function of the Nervous System**

the nervous system probably includes the brain, the nervous tissue contained within the cranium, and the spinal cord, the extension of nervous tissue within the vertebral column. That suggests it is made of two organs—and you may not even think of the spinal cord as an organ—but the nervous system is a very complex structure. Within the brain, many different and separate regions are responsible for many different and separate functions.

## **The Central and Peripheral Nervous Systems**

The nervous system can be divided into two major regions: the central and peripheral nervous systems.

The central nervous system (CNS) is the brain and spinal cord, and the peripheral nervous system (PNS) are referred to as ganglia and Nerves. The brain is contained within the cranial cavity of the skull, and the spinal cord is contained within the vertebral cavity of the vertebral column.

## **Basic Functions of the Nervous System**

The nervous system is involved in receiving information about the environment around us (sensation) and generating responses to that information (motor responses).

But there is a third function that needs to be included. Sensory input needs to be integrated with other sensations, as well as with memories, emotional state, or learning (cognition). Some regions of the nervous system are termed integration or association areas. The process of integration combines sensory perceptions and higher cognitive functions such as memories, learning, and emotion to produce a response.

### **Sensation:**

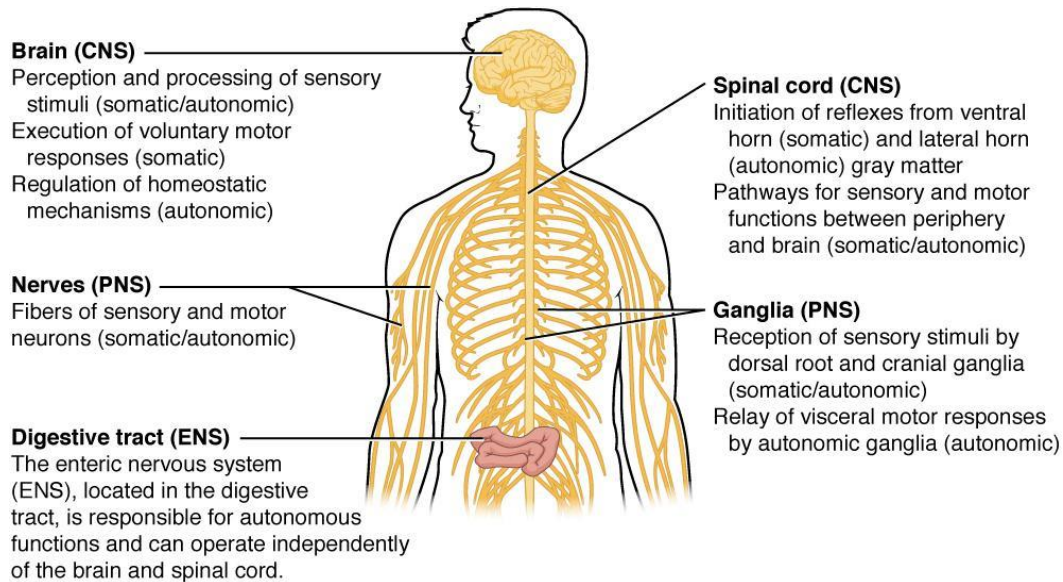
The first major function of the nervous system is sensation—receiving information about the environment to gain input about what is happening outside the body (or, sometimes, within the body). The sensory functions of the nervous system register the presence of a change from homeostasis or a particular event in the environment, known as a stimulus. The senses we think of most are the “big five”: taste, smell, touch, sight, and hearing.

### **Response:**

The nervous system produces a response based on the stimuli perceived by sensory structures. An obvious response would be the movement of muscles, such as withdrawing a hand from a hot stove.

### **Controlling the Body:**

The nervous system can be divided into two parts mostly based on a functional difference in responses. The somatic nervous system (SNS) is responsible for conscious perception and voluntary motor responses. Voluntary motor response means the contraction of skeletal muscle, The autonomic nervous system (ANS) is responsible for involuntary control of the body, usually for the sake of homeostasis (regulation of the internal environment). Sensory input for autonomic functions can be from sensory structures tuned to external or internal environmental stimuli.



**Figure:** Somatic, Autonomic, and Enteric Structures of the Nervous System Somatic structures include the spinal nerves, both motor and sensory fibers, as well as the sensory ganglia (posterior root ganglia and cranial nerve ganglia). Autonomic structures are found in the nerves also but include the sympathetic and parasympathetic ganglia. The enteric nervous system includes the nervous tissue within the organs of the digestive tract.

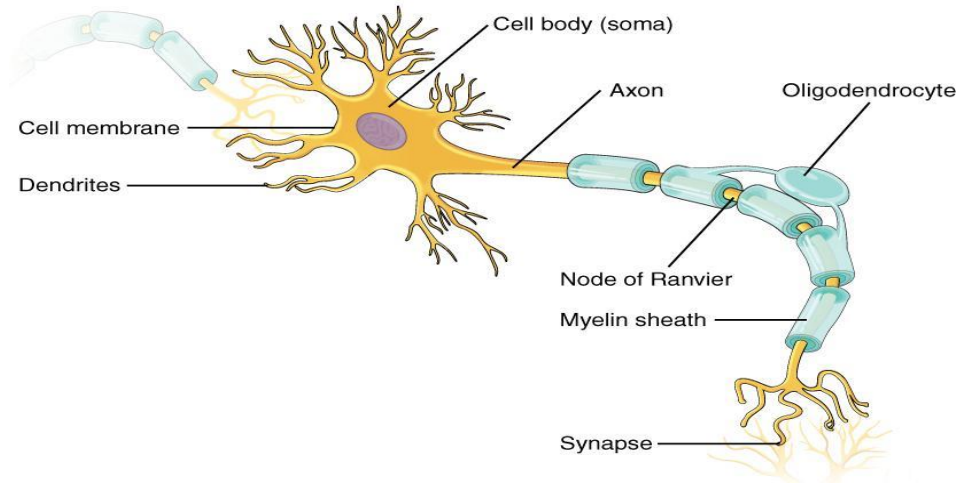
**Nervous Tissue:**

Nervous tissue is composed of two types of cells, neurons, and glial cells. Neurons are the primary type of cell that most anyone associates with the nervous system.

They are responsible for the computation and communication that the nervous system provides.

**Neurons:**

Neurons are the cells considered to be the basis of nervous tissue. They are responsible for the electrical signals that communicate information about sensations, and that produce movements in response to those stimuli.



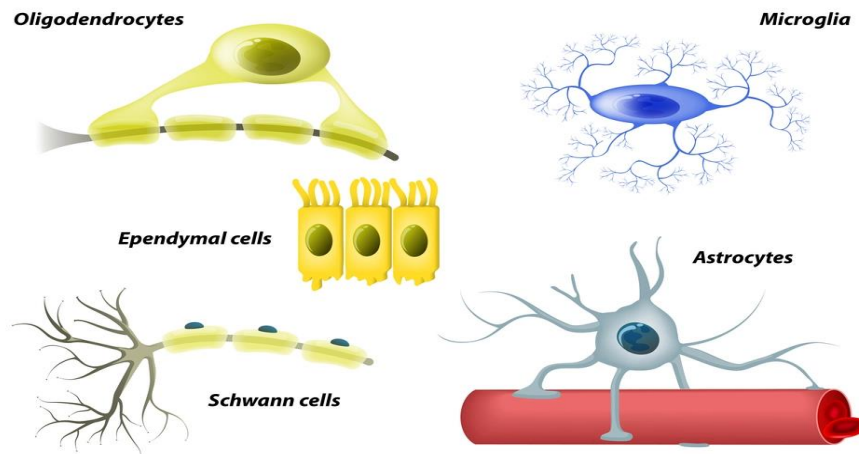
**Figure: Parts of a Neuron**

**Structure:** The major parts of the neuron are labeled on a multipolar neuron from the CNS. Where the axon emerges from the cell body, there is a special region referred to as the axon hillock. This is a tapering of the cell body toward the axon fiber. Within the axon hillock, the cytoplasm changes to a solution of limited components called axoplasm. Because the axon hillock represents the beginning of the axon, it is also referred to as the initial segment.

### Glial Cells

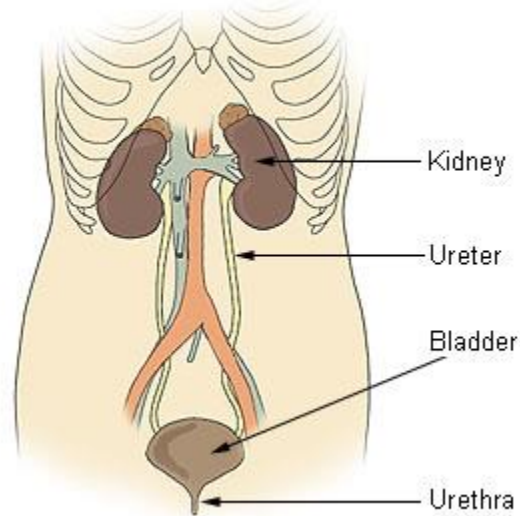
Glial cells, or neuroglia or simply glia, are the other type of cell found in nervous tissue. They are supporting cells, and many functions are directed at helping neurons complete their function for communication.

## GLIAL CELLS



# Urinogenital system

## **Components of the Urinary System**



### 1-Urinary System

The Urinary System is a group of organs in the body concerned with filtering out excess fluid and other substances from the bloodstream. The substances are filtered out from the body in the form of urine. Urine is a liquid produced by the kidneys, collected in the bladder and excreted through the urethra. Urine is used to extract excess minerals or vitamins as well as blood corpuscles from the body.

The Urinary organs include the kidneys, ureters, bladder, and urethra. The Urinary system works with the other systems of the body to help maintain homeostasis. The kidneys are the main organs of homeostasis because they maintain the acid base balance and the water salt balance of the blood.

### Functions of the Urinary System

One of the major functions of the Urinary system is the process of excretion.

1-Excretion is the process of eliminating, from an organism, waste products of metabolism and other materials that are of no use.

2-The urinary system maintains an appropriate fluid volume by regulating the amount of water that is excreted in the urine. Other aspects of its function include

3-regulating the concentrations of various electrolytes in the body fluids and maintaining normal pH of the blood.

Several body organs carry out excretion, but the kidneys are the most important excretory organ. The primary function of the kidneys is to maintain a stable internal environment (homeostasis) for optimal cell and tissue metabolism. They do this by separating urea, mineral salts, toxins, and other waste products from the blood. They also do the job of

conserving water, salts, and electrolytes. At least one kidney must function properly for life to be maintained.

Six important roles of the kidneys are:

**Regulation of plasma ionic composition.** Ions such as sodium, potassium, calcium, magnesium, chloride, bicarbonate, and phosphates are regulated by the amount that the kidney excretes. Regulation of plasma osmolarity. The kidneys regulate osmolarity because they have direct control over how many ions and how much water a person excretes.

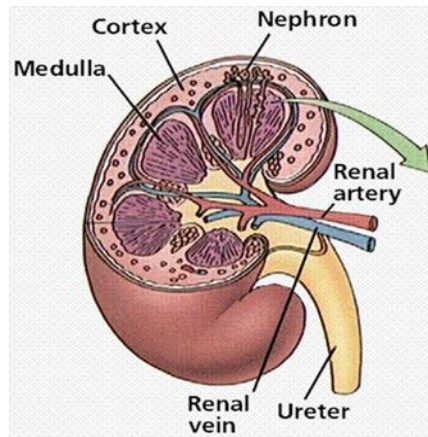
**Regulation of plasma volume.** Your kidneys are so important they even have an effect on your blood pressure. The kidneys control plasma volume by controlling how much water a person excretes. The plasma volume has a direct effect on the total blood volume, which has a direct effect on your blood pressure. Salt (NaCl) will cause osmosis to happen; the diffusion of water into the blood.

**Regulation of plasma hydrogen ion concentration (pH).** The kidneys partner up with the lungs and they together control the pH. The kidneys have a major role because they control the amount of bicarbonate excreted or held onto. The kidneys help maintain the blood pH mainly by excreting hydrogen ions and reabsorbing bicarbonate ions as needed.

**Removal of metabolic waste products and foreign substances from the plasma.** One of the most important things the kidneys excrete is nitrogenous waste. As the liver breaks down amino acids it also releases ammonia. The liver then quickly combines that ammonia with carbon dioxide, creating urea which is the primary nitrogenous end product of metabolism in humans. The liver turns the ammonia into urea because it is much less toxic. We can also excrete some ammonia, creatinine and uric acid. The creatinine comes from the metabolic breakdown of creatine phosphate (a high-energy phosphate in muscles). Uric acid comes from the breakdown of nucleotides. Uric acid is insoluble and too much uric acid in the blood will build up and form crystals that can collect in the joints and cause gout.

**Secretion of Hormones** The endocrine system has assistance from the kidney's when releasing hormones. Renin is released by the kidneys. Renin leads to the secretion of aldosterone which is released from the adrenal cortex. Aldosterone promotes the kidneys to reabsorb the sodium (Na<sup>+</sup>) ions. The kidneys also secrete erythropoietin when the blood doesn't have the capacity to carry oxygen. Erythropoietin stimulates red blood cell production. The Vitamin D from the skin is also activated with help from the kidneys. Calcium (Ca<sup>+</sup>) absorption from the digestive tract is promoted by vitamin D.

## Organs in the Urinary System Kidneys and Their Structure:



### **The kidneys**

The kidneys are a pair of bean shaped, reddish brown organs about the size of your fist. It measures 10-12 cm long. They are covered by the renal capsule, which is a tough capsule of fibrous connective tissue. Adhering to the surface of each kidney is two layers of fat to help cushion them. There is a concaved side of the kidney that has a depression where a renal artery enters, and a renal vein and a ureter exit the kidney. The kidneys are located at the rear wall of the abdominal cavity just above the waistline, and are protected by the ribcage. They are considered retroperitoneal, which means they lie behind the peritoneum. There are three major regions of the kidney, renal cortex, renal medulla and the renal pelvis. The outer, granulated layer is the renal cortex. The cortex stretches down in between a radially striated inner layer. The inner radially striated layer is the renal medulla. This contains pyramid shaped tissue called the renal pyramids, separated by renal columns. The ureters are continuous with the renal pelvis and is the very center of the kidney. 1. Renal pyramid 2. Interlobar artery 3. Renal artery 4. Renal vein 5. Renal hilum 6. Renal pelvis 7. Ureter 8. Minor calyx 9. Renal capsule 10. Inferior renal capsule 11. Superior renal capsule 12. Interlobar vein 13. Nephron 14. Minor calyx 15. Major calyx 16. Renal papilla 17. Renal column.

### **Renal Vein**

The renal veins are veins that drain the kidney. They connect the kidney to the inferior vena cava. Because the inferior vena cava is on the right half of the body, the left renal vein is generally the longer of the two. Unlike the right renal vein, the left renal vein often receives the left gonadal vein (left testicular vein in males, left ovarian vein in females). It frequently receives the left suprarenal vein as well.

### **Renal Artery**

The renal arteries normally arise off the abdominal aorta and supply the kidneys with blood. The arterial supply of the kidneys are variable and there may be one or more renal arteries supplying each kidney. Due to the position of the aorta, the inferior vena cava and the kidneys

in the body, the right renal artery is normally longer than the left renal artery. The right renal artery normally crosses posteriorly to the inferior vena cava. The renal arteries carry a large portion of the total blood flow to the kidneys. Up to a third of the total cardiac output can pass through the renal arteries to be filtered by the kidneys.

## **Ureters**

The ureters are two tubes that drain urine from the kidneys to the bladder. Each ureter is a muscular tube about 10 inches (25 cm) long. Muscles in the walls of the ureters send the urine in small spurts into the bladder, (a collapsible sac found on the forward part of the cavity of the bony pelvis that allows temporary storage of urine). After the urine enters the bladder from the ureters, small folds in the bladder mucosa act like valves preventing backward flow of the urine. The outlet of the bladder is controlled by a sphincter muscle. A full bladder stimulates sensory nerves in the bladder wall that relax the sphincter and allow release of the urine. However, relaxation of the sphincter is also in part a learned response under voluntary control. The released urine enters the urethra. Urinary Bladder The urinary bladder is a hollow, muscular and distensible or elastic organ that sits on the pelvic floor (superior to the prostate in males). On its anterior border lies the pubic symphysis and, on its posterior border, the vagina (in females) and rectum (in males).

## **The urinary bladder**

The urinary bladder can hold approximately 17 to 18 ounces (500 to 530 ml) of urine, however the desire to micturate is usually experienced when it contains about 150 to 200 ml. When the bladder fills with urine (about half full), stretch receptors send nerve impulses to the spinal cord, which then sends a reflex nerve impulse back to the sphincter (muscular valve) at the neck of the bladder, causing it to relax and allow the flow of urine into the urethra. The Internal urethral sphincter is involuntary. The ureters enter the bladder diagonally from its dorsolateral floor in an area called the trigone. The trigone is a triangular shaped area on the postero-inferior wall of the bladder. The urethra exits at the lowest point of the triangle of the trigone. The urine in the bladder also helps regulate body temperature. If the bladder becomes completely void of fluid, it causes the patient to chill.

## **The urethra**

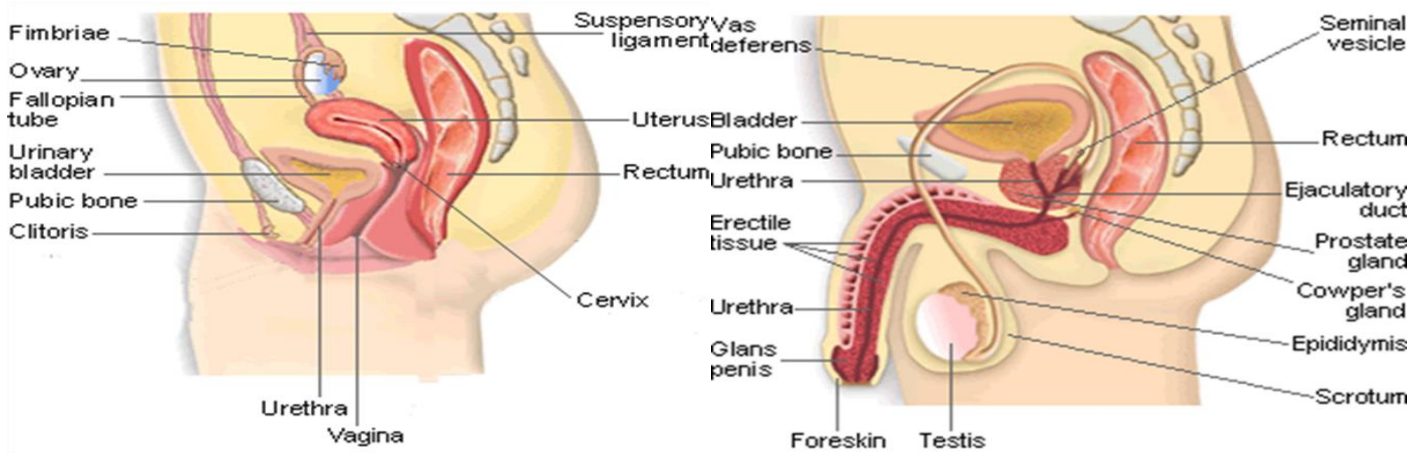
The urethra is a muscular tube that connects the bladder with the outside of the body. The function of the urethra is to remove urine from the body. It measures about 1.5 inches (3.8 cm) in a woman but up to 8 inches (20 cm) in a man. Because the urethra is so much shorter in a woman it makes it much easier for a woman to get harmful bacteria in her bladder this is commonly called a bladder infection or a UTI. The most common bacteria of a UTI is E-coli from the large intestines that have been excreted in fecal matter.

**Female urethra in the human female**, the urethra is about 1-2 inches long and opens in the vulva between the clitoris and the vaginal opening. Men have a longer urethra than women. This means that women tend to be more susceptible to infections of the bladder (cystitis) and the urinary tract.

**Male urethra in the human male**, the urethra is about 8 inches long and opens at the end of the head of the penis. The length of a male's urethra, and the fact it contains a number of bends, makes catheterisation more difficult.

The urethral sphincter is a collective name for the muscles used to control the flow of urine from the urinary bladder. These muscles surround the urethra, so that when they contract, the urethra is closed.

- There are two distinct areas of muscle: the internal sphincter, at the bladder neck and • the external, or distal, sphincter. Human males have much stronger sphincter muscles than females, meaning that they can retain a large amount of urine for twice as long, as much as 800mL, i.e . "hold it".



## Nephrons

The filtering units of the kidneys is the nephrons. There are approximately one million nephrons in each kidney. The nephrons are located within the cortex and medulla of each kidney.

The tubes of the nephron are surrounded by cells and a network of blood vessels spreads throughout the tissue. Therefore, material that leaves the nephron enters the surrounding cells and returns to the bloodstream by a network of vessels.

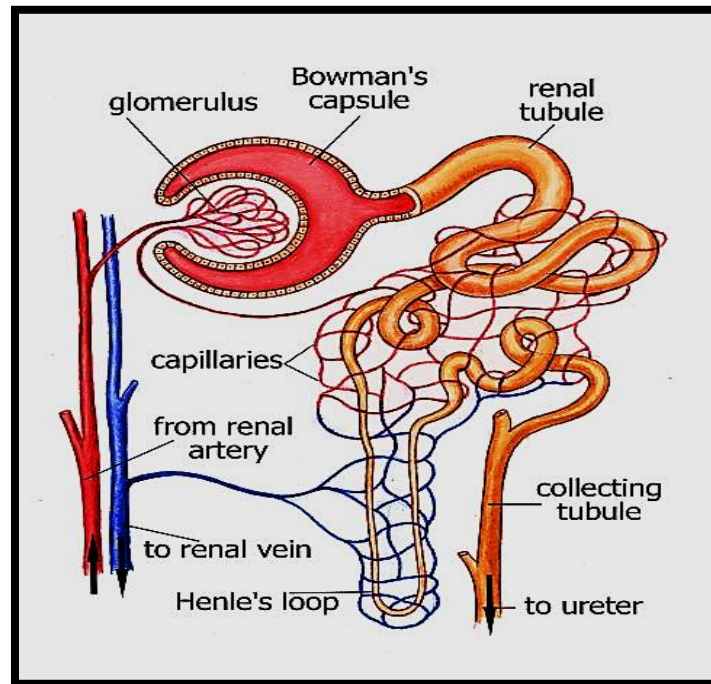
### Parts of the Nephron

Each nephron consists of the following parts:

- 1) glomerulus
- 2) Bowman's capsule
- 3) proximal tubule



- 4) loop of Henle
- 5) distal tubule
- 6) collecting duct



The **glomerulus** is a mass of thin-walled capillaries.

The **Bowman's capsule** is a double-walled, cup-shaped structure.

The **proximal tubule** leads from the Bowman's capsule to the Loop of Henle.

The **loop of Henle** is a long loop which extends into the medulla.

The **distal tubule** connects the loop of Henle to the collecting duct.

### Function of the Kidney

The principal function of the kidney is to filter blood to remove cellular waste products from the body. At any given time, 20 % of blood is in the kidneys. Humans can function with one kidney.

The kidney has other functions but it is usually associated with the excretion of cellular waste such as:

- 1) urea (a nitrogenous waste produced in the liver from the breakdown of protein. It is the main component of urine).
  - 2) uric acid (usually produced from breakdown of DNA or RNA).
  - 3) creatinine (waste product of muscle action).
- All these compounds have nitrogen as a major component.
  - The kidneys are more than excretory organs.
  - They are one of the major homeostatic organs of the body.
  - They control blood pH
  - Secrete erythropoietin (a hormone that stimulates red blood cell production)
  - Activate vitamin D production in the skin.

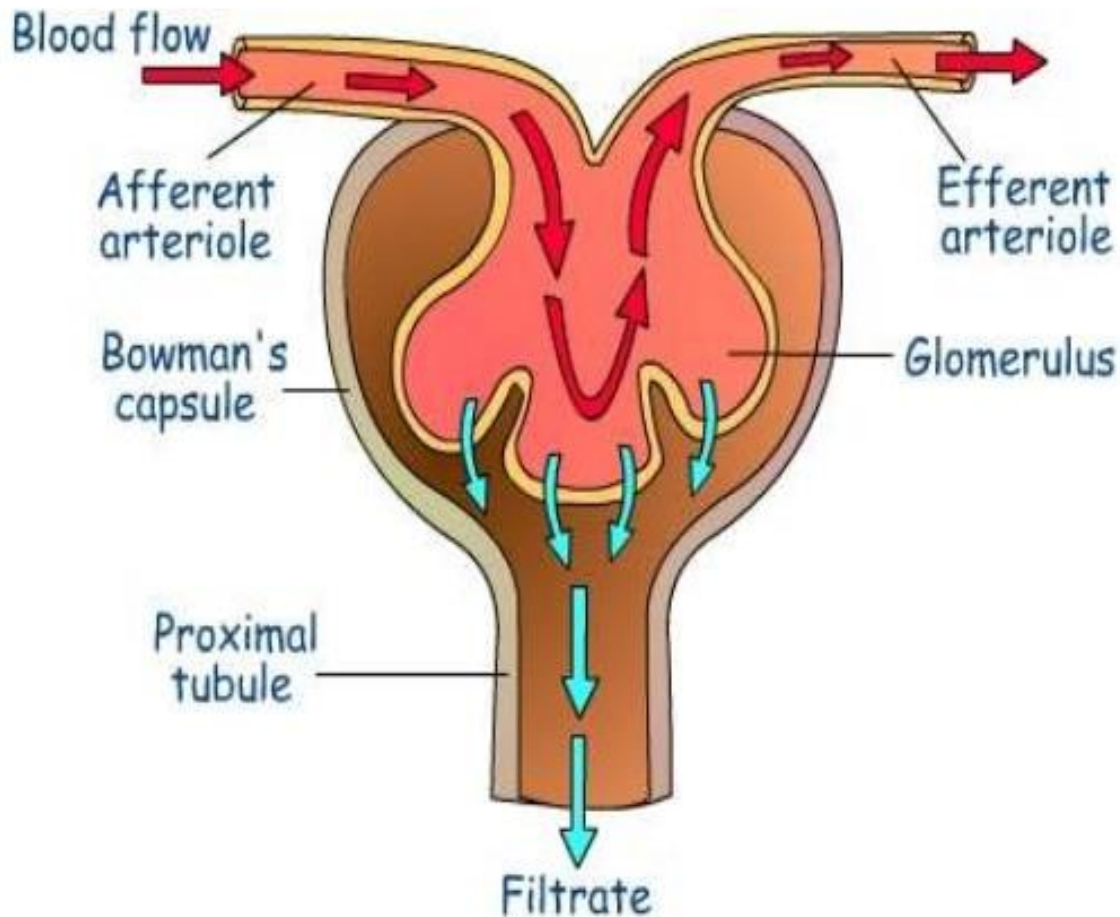
## **Formation of Urine**

Urine is formed in three steps: Filtration, Reabsorption, and Secretion.

**Filtration** Blood enters the afferent arteriole and flows into the glomerulus. Blood in the glomerulus has both filterable blood components and non-filterable blood components. Filterable blood components move toward the inside of the glomerulus while non-filterable blood components bypass the filtration process by exiting through the efferent arteriole. Filterable Blood components now take on plasma like form called glomerular filtrate. A few of the filterable blood components are water, nitrogenous waste, nutrients and salts (ions). Nonfilterable blood components include formed elements such as blood cells and platelets along with plasma proteins. The glomerular filtrate is not the same consistency as urine, as much of it is reabsorbed into the blood as the filtrate passes through the tubules of the nephron.

**Reabsorption** Within the peritubular capillary network, molecules and ions are reabsorbed back into the blood. Sodium Chloride reabsorbed into the system increases the osmolarity of blood in comparison to the glomerular filtrate. This reabsorption process allows water (H<sub>2</sub>O) to pass from the glomerular filtrate back into the circulatory system. Glucose and various amino acids also are reabsorbed into the circulatory system. These nutrients have carrier molecules that claim the glomerular molecule and release it back into the circulatory system. If all of the carrier molecules are used up, excess glucose or amino acids are set free into the urine. A complication of diabetes is the inability of the body to reabsorb glucose. If too much glucose appears in the glomerular filtrate, it increases the osmolarity of the filtrate, causing water to be released into the urine rather than reabsorbed by the circulatory system. Frequent urination and unexplained thirst are warning signs of diabetes, due to water not being reabsorbed. Glomerular filtrate has now been separated into two forms: Reabsorbed Filtrate and Non-reabsorbed Filtrate. Non-reabsorbed filtrate is now known as tubular fluid as it passes through the collecting duct to be processed into urine.

**Secretion** Some substances are removed from blood through the peritubular capillary network into the distal convoluted tubule or collecting duct. These substances are Hydrogen ions, creatinine, and drugs. Urine is a collection of substances that have not been reabsorbed during



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## Glomerular filtration

Glomerular filtration occurs when blood enters the glomerulus through the afferent arteriole. Due to glomerular blood pressure, water and small solutes (such as glucose, amino acids, nitrogenous wastes and ions) present in blood move from the glomerulus to the inside of the glomerulus capsule. This is a filtration process because large molecules (such as proteins) and formed elements (blood cells and platelets) are unable to pass through the capillary wall. The filtered fluid (called the glomerular filtrate) is essentially protein-free and devoid of cells. On an average, the daily volume of glomerular filtrate is about 180 L in adult males and 150 L in adult females. More than 99% of the glomerular filtrate returns to the bloodstream via tubular reabsorption, so only 1-1.5 L is excreted as urine. The volume of fluid filtered by the renal corpuscle is much larger than in other capillaries of the body for three reasons:

1. Glomerular capillaries present a large surface area for filtration because they are long.
2. The filtration membrane (the endothelial cells of glomerular capillaries and the podocytes, which completely encircle the capillaries, form a leaky barrier known as the filtration membrane) is thin and porous.

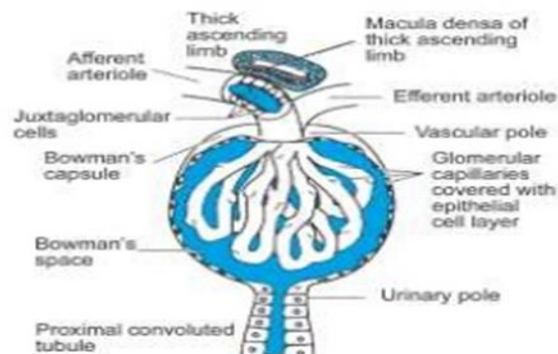
3. Glomerular capillary blood pressure is high. Because the efferent arteriole is smaller in diameter than the afferent arteriole.

### **Glomerular filtration rate (GFR)**

The amount of filtrate formed in all the renal corpuscles of both kidneys per minute is called GFR. The GFR in a healthy individual is approximately 125 ml/minute (7.5 L/hr or 180 L/day). The GFR is determined by the net filtration pressure and the glomerular filtration coefficient. The net filtration pressure represents the sum of the hydrostatic and colloid osmotic pressures that either favour or oppose filtration across the glomerular capillaries. Glomerular hydrostatic pressure is the blood pressure in glomerular capillaries. It is about 55 mm Hg. It promotes filtration by forcing water and solutes present in blood plasma to pass through the filtration membrane. Capsular hydrostatic pressure is the hydrostatic pressure exerted against the filtration membrane by fluid already present in the capsular space. It opposes filtration. Its value is about 15 mm Hg. Blood colloid osmotic pressure is the pressure develops due to the pressure of proteins such as albumin, globulins and fibrinogen in blood plasma. Because plasma proteins cannot be filtered, they are in the glomerular capillaries but not in Bowman's capsule. Blood colloid osmotic pressure also opposes filtration. Its value in glomerular capillaries is about 30 mm Hg. Net filtration pressure: The force favoring filtration is the glomerular hydrostatic pressure which is about 55 mm Hg. The total of the two forces opposing filtration is about 45 mm Hg. The net difference favoring filtration (about 10 mm Hg) is called net filtration pressure.

## **Introduction**

### **❖ Glomerular filtration rate (GFR):**



- Rate at which plasma is filtered from the glomerular capillaries into bowman's capsule per unit time.
- In average, GFR is about 125ml/min or 180 l/day and filtration fraction is about 0.2(20%)

## **Regulation of Glomerular filtration rate (GFR)**

The mechanisms that regulate glomerular filtration rate operate in two main ways: (1) By adjusting blood flow into and out of the glomerulus. GFR increases when blood flow into the glomerular capillaries increases. (2) By altering the glomerular capillary surface area available for filtration. Coordinated control of the diameter of both afferent and efferent arterioles regulates glomerular blood flow. Constriction of the afferent arteriole decreases blood flow into the glomerulus; dilation of the afferent arteriole increases it. Three mechanisms control GFR: renal autoregulation, neural regulation, and hormonal regulation. (A) Renal autoregulation The kidneys themselves help maintain a constant renal blood flow and GFR despite normal, everyday changes in blood pressure, like those that occur during exercise. This capability is called renal autoregulation and consists of two mechanisms—the myogenic mechanism and tubuloglomerular feedback. A1. Myogenic autoregulation: Myogenic constriction of the afferent arteriole occurs due to the ability of the smooth muscle to sense and respond to an increase in arterial pressure. As blood pressure rises, GFR also rises because renal blood flow increases. However, the elevated blood pressure stretches the walls of the afferent arterioles. In response, smooth muscle fibers in the wall of the afferent arteriole contract, which narrows the arteriole's lumen. As a result, renal blood flow decreases, thus reducing GFR to its previous level. Conversely, when arterial blood pressure drops, the smooth muscle cells are stretched less and thus relax. The afferent arterioles dilate, renal blood flow increases, and GFR increases. The myogenic mechanism normalizes renal blood flow and GFR within seconds after a change in blood pressure. A2. Tubuloglomerular feedback: It is so named because part of the renal tubules—the macula densa—provides feedback to the glomerulus. When GFR is above normal due to elevated systemic blood pressure, filtered fluid flows more rapidly along the renal tubules. As a result, the proximal convoluted tubule and loop of Henle have less time to reabsorb  $\text{Na}^+$ ,  $\text{Cl}^-$ , and water. Macula densa cells are thought to detect the increased delivery of  $\text{Na}^+$ ,  $\text{Cl}^-$ , and water and to inhibit release of nitric oxide (NO) from cells in the juxtaglomerular apparatus (JGA).<sup>5</sup> Because NO causes vasodilation, afferent arterioles constrict when the level of NO declines. As a result, less blood flows into the glomerular capillaries, and GFR decreases. When blood pressure falls, causing GFR to be lower than normal, the opposite sequence of events occurs, although to a lesser degree. Tubuloglomerular feedback operates more slowly than the myogenic mechanism. (B) Neural regulation All the blood vessels of the kidneys, including the afferent and the efferent arterioles, are richly innervated by sympathetic nerve fibres. Activation of renal sympathetic nerves releases norepinephrine. Norepinephrine causes vasoconstriction of afferent arterioles and thus decreases the GFR. (C) Hormonal regulation Two hormones contribute to regulation of GFR. Angiotensin II reduces GFR; atrial natriuretic peptide (ANP) increases GFR. Angiotensin II is a very potent vasoconstrictor that narrows both afferent and efferent arterioles and reduces renal blood flow, thereby decreasing GFR. Cells in the atria of the heart secrete atrial natriuretic peptide (ANP). Stretching of the atria, as occurs when blood volume increases, stimulates

secretion of ANP. By causing relaxation of the glomerular mesangial cells, ANP increases the capillary surface area available for filtration. Glomerular filtration rate rises as the surface area increases.

مقرر (علم الحيوان Zoo 101– Zoology I "جزء الخلية والأنسجة") لطلاب الفرقة  
الأولى شعبة العلوم البيولوجية والجيولوجية باللغة الإنجليزية (101 عل ح)

أستاذ المقرر

د/ نادية سمير (جزء الخلية والأنسجة)

## Introduction

The **cell** (from Latin *cellula* 'small room') is the basic structural, functional, and biological unit of all known organisms. A cell is the smallest unit of life. Therefore, cells are often described as the "building blocks of life".

Cells consist of cytoplasm enclosed within a membrane, which contains many biomolecules such as proteins and nucleic acids. Most plant and animal cells are only visible under a light microscope, with dimensions between 1 and 100 micrometres. Electron microscopy gives a much higher resolution showing greatly detailed cell structure. Organisms can be classified as unicellular (consisting of a single cell such as bacteria) or multicellular (including plants and animals). Most unicellular organisms are classed as microorganisms.

The number of cells in plants and animals varies from species to species; it has been approximated that the human body contains roughly 40 trillion ( $4 \times 10^{13}$ ) cells. The brain accounts for around 80 billion of these cells.

Cells were discovered by Robert Hooke in 1665, who named them for their resemblance to cells inhabited by Christian monks in a monastery. Cell theory, first 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. Cells emerged on Earth about 4 billion years ago.

## **Types of Cells**

Cells are of two types: eukaryotic, which contain a nucleus, and prokaryotic cells, which do not have a nucleus, but a nucleoid region is still present. Prokaryotes are single-celled organisms, while eukaryotes can be either single-celled or multicellular.

### **Prokaryotic cells**

A **prokaryote** is a typically unicellular organism that lacks a nuclear membrane-enclosed nucleus. The word *prokaryote* comes from the Greek *πρό* (*pro*, 'before') and *κάρυον* (*karyon*, 'nut' or 'kernel'). In the two-empire system arising from the work of Édouard Chatton, prokaryotes were classified within the empire **Prokaryota**. But in the three-domain system, based upon molecular analysis, prokaryotes are divided into two domains: *Bacteria* (formerly Eubacteria) and *Archaea* (formerly Archaeobacteria). Organisms with nuclei are placed in a third domain, Eukaryota. In the study of the origins of life, prokaryotes are thought to have arisen before eukaryotes.

Prokaryotes lack mitochondria, or any other eukaryotic membrane-bound organelles; and it was once thought that prokaryotes lacked cellular compartments, and therefore all cellular components within the cytoplasm were unenclosed, except for an outer cell membrane. But bacterial microcompartments, which are thought to be simple organelles enclosed in protein shells, have been discovered, along with other prokaryotic organelles. While typically being unicellular, some prokaryotes, such as cyanobacteria, may form large colonies. Others, such as myxobacteria, have multicellular stages in their life cycles. Prokaryotes are asexual, reproducing without fusion of gametes, although horizontal gene transfer also takes place.

Molecular studies have provided insight into the evolution and interrelationships of the three domains of life. The division between prokaryotes and eukaryotes reflects the existence of two very different levels of cellular organization; only eukaryotic cells have an enveloped nucleus that contains its chromosomal DNA, and other characteristic membrane-bound organelles including mitochondria. Distinctive types of prokaryotes include extremophiles and methanogens; these are common in some extreme environments.

## **History of Cells**

The distinction between prokaryotes and eukaryotes was firmly established by the microbiologists Roger Stanier and C. B. van Niel in their 1962 paper The concept of a bacterium (though spelled procaryote and eucaryote there). That paper cites Édouard Chatton's 1937 book *Titres et Travaux Scientifiques*<sup>1</sup> for using those terms and recognizing the distinction. One reason for this classification was so that what was then often called blue-green algae (now called cyanobacteria) would not be classified as plants but grouped with bacteria.

## **Cell Theory**

- The cell is the basic functional and structural unit of life. All the living organisms are composed of cells.
- All cells are formed by the division of the already existing cells which in terms of biology means reproduction. Every cell of our body comprises of genetic material which is passed down during the process.
- All the basic physiological and chemical functions i.e. the growth, repair, movement, communication, immunity and digestions are performed inside the cells.
- All the activities of the cell depend mainly on the activities of the subcellular structures that lie within the cell. These subcellular structures comprise of the plasma membrane, organelles and if present, the nucleus.

Here is the revised introduction to cell. In all the living beings, cells are the basic structural units. We can compare the presence of cells in our body to the bricks in a building. All the bricks are assembled to make a building. Similarly, all the cells are assembled to make the body of an organism.

Thus, it is the basic structural and functional unit of life and all the organisms are made up of cells. The subcellular structures of the cell comprise of the plasma membrane, organelles and in some cases a nucleus as well. As for the size of the cell, it is variable and maybe anything from 1 to 100 micrometre.

### **How Are Cells Produced?**

Every cell is produced by the division of a cell that is already existing in our body. This is possible because of the genetic material contained in the cell. The genetic material is passed down from one cell to another during the reproduction process. Unlike the non-living bricks, the cells of the living organisms are rather complex living structures. Therefore, the cells can divide the genetic material and form two new cells.

### **Types of Cell**

Broadly, there are two key types of cells i.e. the Prokaryotic Cell and the Eukaryotic Cell. The difference

between the two is defined mainly by the presence or the absence of the nuclear membrane. Let's know more about the two types of cells.

### **1) Prokaryotic Cell**

If a cell has a nuclear material without a nuclear membrane, then it is known as the prokaryotic cell. Those organisms which have these type of cells are commonly known as the prokaryotes where 'pro' stands for primitive and 'karyon' stands for the nucleus. Some of the organisms that have prokaryotic cells include bacteria and the blue-green algae.

### **2) Eukaryotic Cell**

If a cell has a nuclear material with a nuclear membrane, then it is known as the Eukaryotic Cell. Those organisms which have these type of cells are commonly known as eukaryotes where 'eu' stands for true and 'karyon' stands for the nucleus. All the living organisms except bacteria and blue-green algae have Eukaryotic Cells.

## **Functions of a Cell**

As you already know that a cell is a structural and functional unit of living. Let us study 6 of the most vital functions performed by a cell.

### **Structure and Support**

You know a house is made of bricks. Similarly, an organism is made up of cells. Though there are certain cells such as collenchyma and sclerenchyma are

present for offering structural support however in general too, all cells generally provide the structural basis of all organisms.

## **Growth**

In complex organisms such as humans, the tissues grow by simple multiplication of cells. Hence, cells are responsible for the growth of the organism. The entire thing takes place via a process of mitosis.

## **Transport**

Cells import the nutrients that are used in the different chemical process which take place inside them. As a result of these processes, a waste product is produced. Cells then work to get rid of this waste. In this manner, the small molecules like the such as oxygen, carbon dioxide, and ethanol pass through the cell membrane by diffusion. This method is known as passive transport. On the other hand, the larger molecules like the proteins and polysaccharides, go in and out of the cell via active transport.

## **Energy Production**

Organisms need energy to perform different chemical reactions. In plants, the energy comes from the process of photosynthesis while in the animals the energy comes via respiration.

## **Metabolism**

Cell is responsible for metabolism that includes all the chemical reactions that take place inside an organism to keep it alive.

## Reproduction

A cell helps in reproduction by the processes of mitosis (in more evolved organisms) and meiosis.

## Different Cell Organelles and their Functions

**Organelles** make up the subunits of a cell. There are numerous each with their own function.

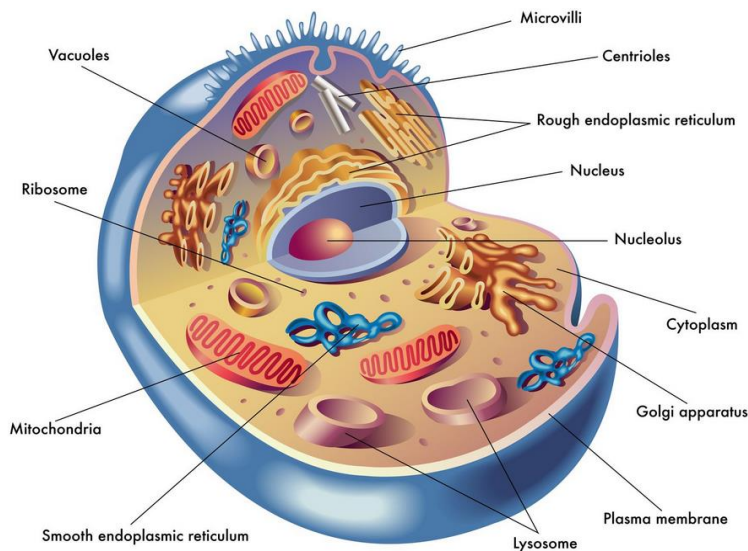


Fig. (1): Cell Organelles

## **Plasma Membrane**

The plasma membrane is the organelle that encapsulates the contents of the cell. Apart from encapsulating cell contents, the plasma membrane also plays a vital role in regulating the movement of substances in and out of the cell.

As such, it is actively involved in such both passive and active transportation to and from the cell. These processes also help maintain balance even when conditions outside the cell change.

The plasma membrane is made up of two layers of phospholipids (phospholipids bilayer).

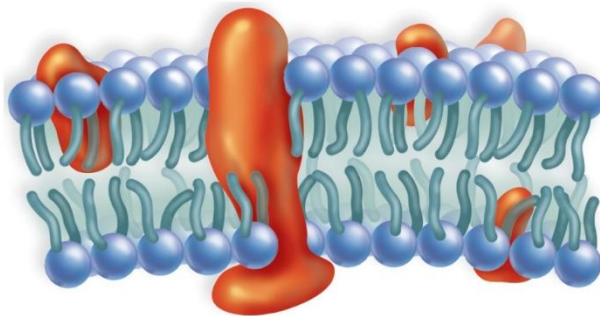


Fig. (2): The plasma membrane



## Nucleus/DNA

Some of the main components of the nucleus include the chromatin, nucleoplasm/nuclear sap and the nucleolus. The nucleus houses DNA (the hereditary material) as well as various proteins and the nucleolus. In eukaryotic cells, the nucleus is enclosed in a nuclear membrane. It is the organelle that controls the hereditary traits of an organism by directing such processes as protein synthesis and cell division among others. For prokaryotes, the DNA lacks a nuclear membrane. The genetic material is therefore bound in the nucleotide region. The nucleolus plays an important role in ribosome production.

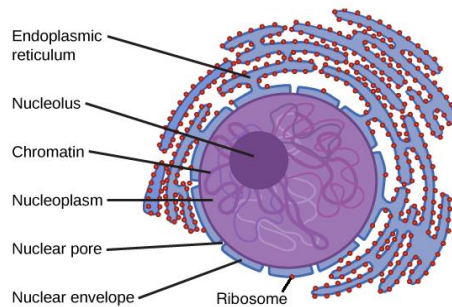


Fig. (3): The nucleolus Structural

## Ribosome

Ribosomes are tiny organelles that contain RNA and specific proteins within the cytoplasm. Within the cell, ribosomes are directly involved in the manufacture of proteins by using their RNA and amino acids.

This process involves decoding the information contained in the mRNA and using amino acids to produce the required proteins.

## **Mitochondria**

Mitochondria are some of the largest organelles within a cell. Compared to some of the other organelles, mitochondria contain DNA which makes them semiautonomous. Mitochondria also contain a double-membrane with the inner membrane folding to form cristae. Also known as the powerhouse, mitochondria play an important role in respiration where they generate ATP (adenosine triphosphate) from substrates in the presence of oxygen. Using their DNA, mitochondria are able to encode for some of the components they require to perform their functions.

ATP stores energy in the form of chemical bonds and is released whenever it is needed for various cell functions.

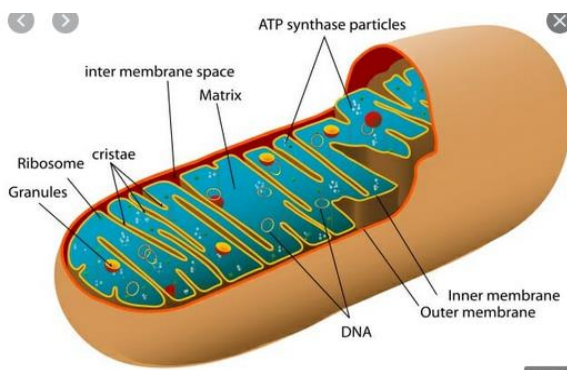


Fig. (4): The Mitochondria structural

## **Endoplasmic Reticulum**

Found in eukaryotic cells, Endoplasmic reticulum (ER) is the organelle that forms an interconnected network of flattened sacs (cisternae). Like some of the other organelles found in eukaryotes, ER is enclosed in a membrane. The ER is divided into two regions that vary in structure and function.

These include:

### **Smooth endoplasmic reticulum**

The smooth ER is named so because it lacks a ribosome on its surface. As a result, it is more smooth in appearance as compared to the rough ER. It is involved in the synthesis of lipids (e.g. phospholipids) and carbohydrates that are used to build the cell membrane.

**Some of the other functions of the smooth ER include:**

- Transportation of vesicles
- Enzyme production in the liver
- Contraction of muscle cells in the muscles
- Synthesis of hormones in the brain cells

### **Rough endoplasmic reticulum**

Unlike the smooth ER, rough ER has ribosome attached to its surface. It's involved in the manufacture of various proteins in the cell. On the other hand, the rough ER is involved in the production of antibodies, insulin as well as transportation of proteins into the smooth ER.

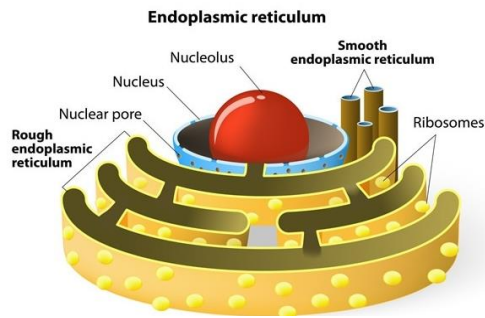


Fig. (5): Rough endoplasmic reticulum

## Golgi Apparatus

Golgi apparatus are found in eukaryotic and are highly folded into cisternae (flattened sacs). They are enclosed in a membrane that varies in thickness from different regions.

In the cell, Golgi apparatus are actively involved in the manufacturing, storage as well as transportation of products from the ER.

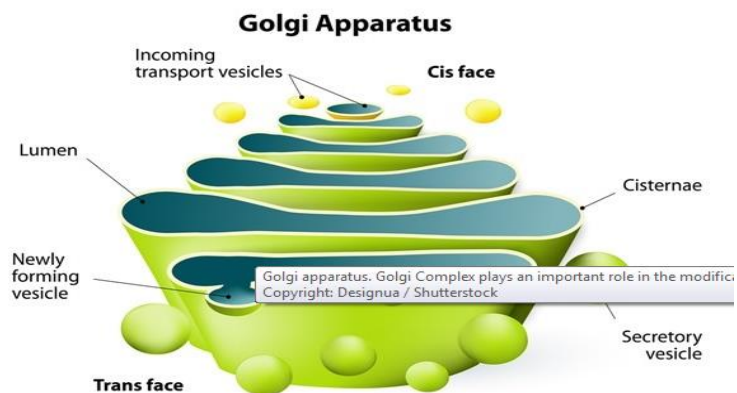


Fig. (6): Golgi Apparatus

## **Vacuoles**

A vacuole may be described as a space inside the cell that does not contain cytoplasm. It is surrounded by a membrane and filled with a fluid. Vacuoles store various molecules including enzymes, waste products of the cell, water, and even food material depending on the type of cell.

In cases where vacuoles contain waste products of the cell, they are also involved in the exportation of waste from the cell thus protecting the cell from toxicity.

\* Some vacuoles also play a role in maintaining the internal hydrostatic pressure of the cell as well as regulating pH.

## **Cytoskeleton**

The cytoskeleton is made up of microtubules and microfilaments. By spreading throughout the cell (in the cytoplasm), the cytoskeleton helps maintain the shape of the cell while also ensuring its elasticity.

\* The cytoskeleton is also involved in anchoring the nucleus and supporting cell contents.

## **Centriole**

Centrioles are cylindrical organelles found in most eukaryotic cells. They contain tube-shaped molecules known as microtubules that help separate chromosomes and move them during cell division.

## **Lysosome**

A lysosome is commonly referred to as sacs of enzymes. They are membranous organelles that contain acidic enzymes (hydrolase enzymes) that serve to digest various macromolecules (e.g. lipids and nucleic acids) in the cell.

Conditions inside lysosomes have been shown to be acidic. These conditions are maintained by the lysosome membrane thus providing favorable conditions for the enzymes to perform their functions.

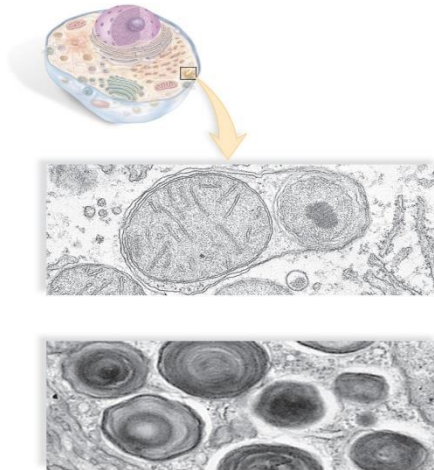


Fig. (7): lysosome

Other features of a cell include:

## **Cell Wall**

- Some books do not consider the cell wall to be an organelle. However, it's one of the most important components of plant cells. The cell wall surrounds the

cell membrane and serves to strengthen and protect the cell.

For instance, in the cells of plant roots, the cell wall protects the cell as they grow deeper in the soil. The cell wall also serves as a filter that controls the movement of molecules in and out of the cell.

## **Cytoplasm**

is also not considered as an organelle in some books. However, it is an important component of the cell. Cell cytoplasm is composed of protoplasm in which all the other cell organelles are suspended.

Many of the cell processes (protein synthesis, respiration etc) take place in the cytoplasm. The cytoplasm also plays an important role in the movement of various materials around the cell.

## **Chemical Composition of the Cell**

Chemical compounds in the cell can be divided into two major groups: Organic and Inorganic compounds

Organic compounds are chemical compounds that contain the element carbon. Organic compounds in the cell include carbohydrates, protein, lipids and nucleic acids. Some of these compounds are synthesized by the cell itself.

Water is an inorganic compound which is composed of hydrogen and oxygen. It is an important compound in the cell.

**Table 1: inorganic chemical compounds in the cell.**

<b>Percentage of Body Weight</b>	<b>Element</b>	<b>Usage</b>
65%	Oxygen	This element is obviously the most important element in the human body. Oxygen atoms are present in water, which is the compound most common in the body, and other compounds that make up tissues. It is also found in the blood and lungs due to respiration.
18.6%	Carbon	Carbon is found in every organic molecule in the body, as well as the waste product of respiration (carbon dioxide). It is typically ingested in food that is eaten.
9.7%	Hydrogen	Hydrogen is found in all water molecules in the body as well as many other compounds making up the various tissues.
3.2%	Nitrogen	Nitrogen is very common in proteins and organic compounds. It is also present in the lungs due to its abundance in the atmosphere.
1.8%	Calcium	Calcium is a primary component of the skeletal system, including the teeth. It is also found in the nervous system, muscles, and the blood.
1.0%	Phosphorus	This element is common in the bones and teeth, as well as nucleic acids.
0.4%	Potassium	Potassium is found in the muscles, nerves, and certain tissues.
0.2%	Sodium	Sodium is excreted in sweat, but is also found in muscles and nerves.
0.2%	Chlorine	Chlorine is present in the skin and facilitates water absorption by the cells.
0.06%	Magnesium	Magnesium serves as a cofactor for various enzymes in the body.
0.04%	Sulfur	Sulfur is present in many amino acids and proteins.
0.007%	Iron	Iron is found mostly in the blood since it facilitates the transportation of oxygen.
0.0002%	Iodine	Iodine is found in certain hormones in the thyroid gland.



## **The Importance of Organic Compounds in the Cell**

### **1. Carbohydrates**

- Supply energy for cell processes
- A means of storing energy
- Give structural support to cell walls

### **2. Lipids**

- Store large amounts of energy over long periods of time
- Act as an energy source
- Play a major role in the structure of the cell membranes
- Act as a source of metabolic water
- Reduce the loss of water by evaporation

### **3. Proteins**

- Act as building blocks of many structural components of the cell ; required for growth
- Form enzymes which catalyze chemical reactions
- Form hormones which control growth and metabolism

### **4. Nucleic acids**

- Contain the genetic information of cells
- Play a vital role in protein synthesis

### **The importance of water in the cell**

- Water is important for life because its chemical and physical properties allow it to sustain life.
- Water is a polar molecule which consists of 2 hydrogen atoms and 1 oxygen atom. A polar molecule is a molecule with an unequal distribution of charges. Each molecule has a positively charged and a negatively charged end. Polar molecules attract one another as well as ions. Because of this property, water is considered the solvent of life.

- It is the transport medium in the blood
- It acts as a medium for biochemical reactions.
- Water helps in the maintenance of a stable internal environment within a living organism. The concentration of water and inorganic salts that dissolve in water is important in maintaining the osmotic balance between the blood and interstitial fluid.
- It helps in lubrication.
- Water molecules have very high cohesion. Water molecules tend to stick to each other and move in long unbroken columns through the vascular tissues in plants.

## **Cell Division Definition**

Cell division is the process cells go through to divide. There are several types of cell division, depending upon what type of organism is dividing. Organisms have evolved over time to have different and more complex forms of cell division. Most prokaryotes, or bacteria, use binary fission to divide the cell. Eukaryotes of all sizes use *mitosis* to divide. Sexually-reproducing eukaryotes use a special form of cell division called *meiosis* to reduce the genetic content in the cell. This is necessary in sexual reproduction because each parent must give only half of the required genetic material, otherwise the offspring would have too much DNA, which can be a problem. These different types of cell division are discussed below.

## **Types of Cell Division**

### **Prokaryotic Cell Division**

Prokaryotes replicate through a type of cell division known as *binary fission*. Prokaryotes are simple organism, with only one membrane and no division internally. Thus, when aprokaryote divides, it simply replicates the DNA and splits in half. The process is a

little more complicated than this, as DNA must first be unwound by special proteins. Although the DNA in prokaryotes usually exists in a ring, it can get quite tangled when it is being used by the cell. To copy the DNA efficiently, it must be stretched out. This also allows the two new rings of DNA created to be separated after they are produced. The two strands of DNA separate into two different sides of the prokaryote cell. The cell then gets longer, and divides in the middle.

The DNA is the tangled line. The other components are labeled. Plasmids are small rings of DNA that also get copied during *binary fission* and can be picked up in the environment, from dead cells that break apart. These plasmids can then be further replicated. If a plasmid is beneficial, it will increase in a population. This is in part how antibiotic resistance in bacteria happens. The ribosomes are small protein structures that help produce proteins. They are also replicated so each cell can have enough to function.

### **Eukaryotic Cell Division: Mitosis**

Eukaryotic organisms have membrane bound organelles and DNA that exists on chromosomes, which makes cell division harder. Eukaryotes must replicate their DNA, organelles, and cell mechanisms before dividing. Many of the organelles divide using a process that is essentially *binary fission*, leading scientist to believe that eukaryotes were formed by prokaryotes living inside of other prokaryotes.

After the DNA and organelles are replicated during *interphase* of the cell cycle, the eukaryote can begin the process of mitosis. The process begins during prophase, when the chromosomes condense. If mitosis proceeded without the chromosomes condensing, the DNA would become tangled and break. Eukaryotic DNA is associated with many proteins which can fold it into complex structures. As mitosis proceeds to *metaphase* the chromosomes are lined up in the middle of the cell. Each half of a chromosome, known *assister chromatids* because they are replicated copies of each other, gets separated into each half of the cell as mitosis proceeds. At the end of mitosis, another process called *cytokinesis* divides the cell into two new daughter cells.

All eukaryotic organisms use mitosis to divide their cells. However, only single-celled organisms use mitosis as a form of reproduction. Most multicellular organisms are sexually reproducing and combine their DNA with that of another organism to reproduce. In these cases, organisms need a different method of cell division. Mitosis yields identical cells, but meiosis produces cells with half the genetic information of a regular cell, allowing two cells from different organisms of the same species to combine.

### **Eukaryotic Cell Division: Meiosis**

In sexually reproducing animals, it is usually necessary to reduce the genetic information before fertilization. Some plants can exist with too many copies of the genetic code, but in most organisms it is highly detrimental to have too many copies. Humans with even one extra copy of one chromosome can experience detrimental changes to their body. To counteract this, sexually reproducing organisms undergo a type of cell division known as meiosis. As before mitosis, the DNA and organelles are replicated. The process of meiosis contains two different cell divisions, which happen back-

to-back. The first meiosis, *meiosis I*, separates homologous chromosomes. The homologous chromosomes present in a cell represent the two alleles of each gene an organism has. These alleles are recombined and separated, so the resulting daughter cells have only one allele for each gene, and no homologous pairs of chromosomes. The second division, *meiosis II*, separated the two copies of DNA, much like in mitosis. The end result of meiosis in one cell is 4 cells, each with only one copy of the genome, which is half the normal number.

Organisms typically package these cells into *gametes*, which can travel into the environment to find other gametes. When two gametes of the right type meet, one will fertilize the other and produce a *zygote*. The zygote is a single cell that will undergo mitosis to produce the millions of cells necessary for a large organism. Thus, most eukaryotes use both mitosis and meiosis, but at different stages of their lifecycle.

## **Cell Division Stages**

Depending upon which type of cell division an organism uses, the stages can be slightly different.

### **Mitosis Stages**

Mitosis starts with *prophase* in which the chromosome is condensed. The cell proceeds to *metaphase* where the chromosomes are aligned on the metaphase plate. Then the chromosomes are separated in *anaphase* and the cell's cytoplasm is pinched apart during *telophase*. *Cytokinesis* is the final process that breaks the cell membrane and divides the cell into two.

### **Meiosis Stages**

The stages of meiosis are similar to mitosis, but the chromosomes act differently. Meiosis has two phases, which include two separate cell divisions without the DNA replicating between them. *Meiosis I* and *meiosis II* have the same 4 stages as mitosis: prophase, metaphase, anaphase, and telophase. Cytokinesis concludes both rounds of meiosis.

In prophase I, the chromosomes are condensed. In metaphase I, the chromosomes line up across from



their homologous pairs. When they are separated in anaphase I and telophase I, there is only one form of each gene in each cell, known as a reduction division. Meiosis II proceeds in the same manner as mitosis, which sister chromatids dividing on the metaphase plate. By telophase II, there are 4 cells, each with half of the alleles as the parent cell and only a single copy of the genome. The cells can now become gametes and fuse together to create new organisms.

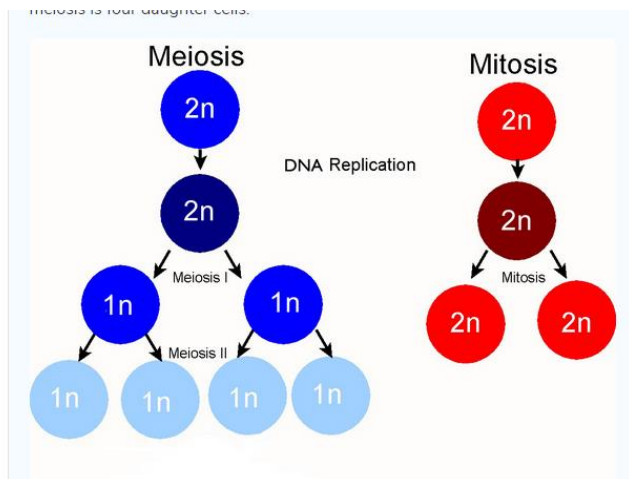


Fig.(8): Types of Cell Division

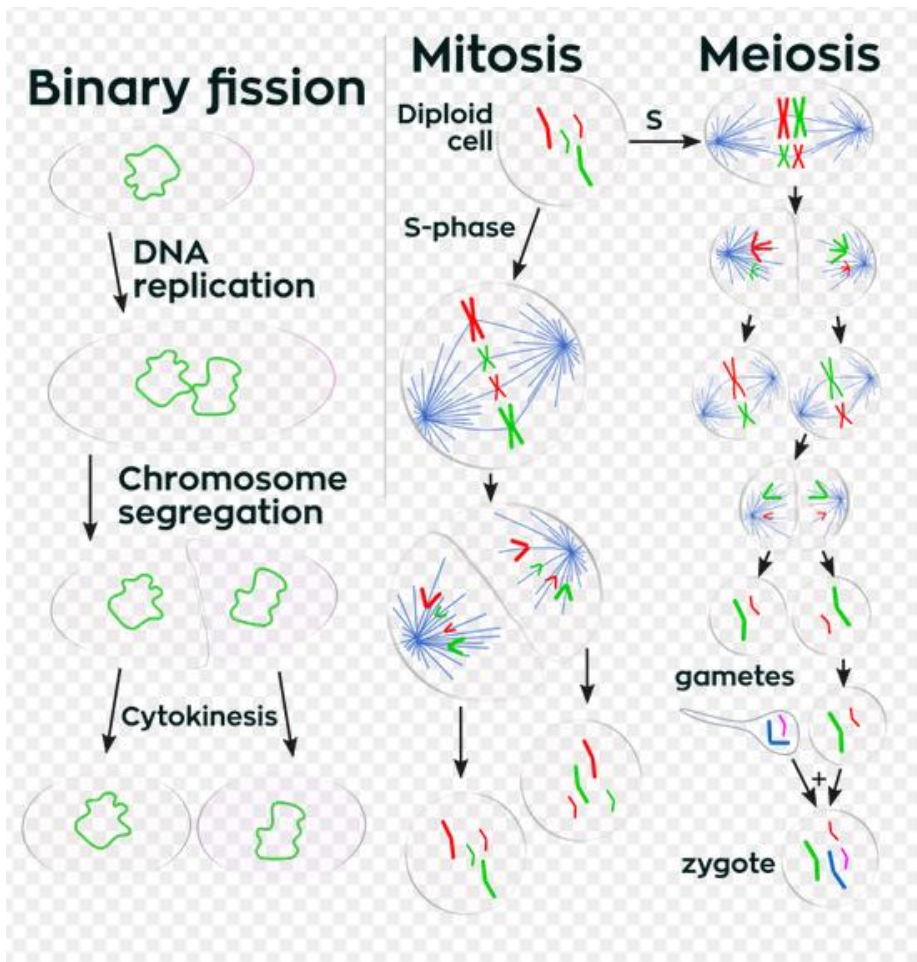


Fig.(9): Types of Cell Division

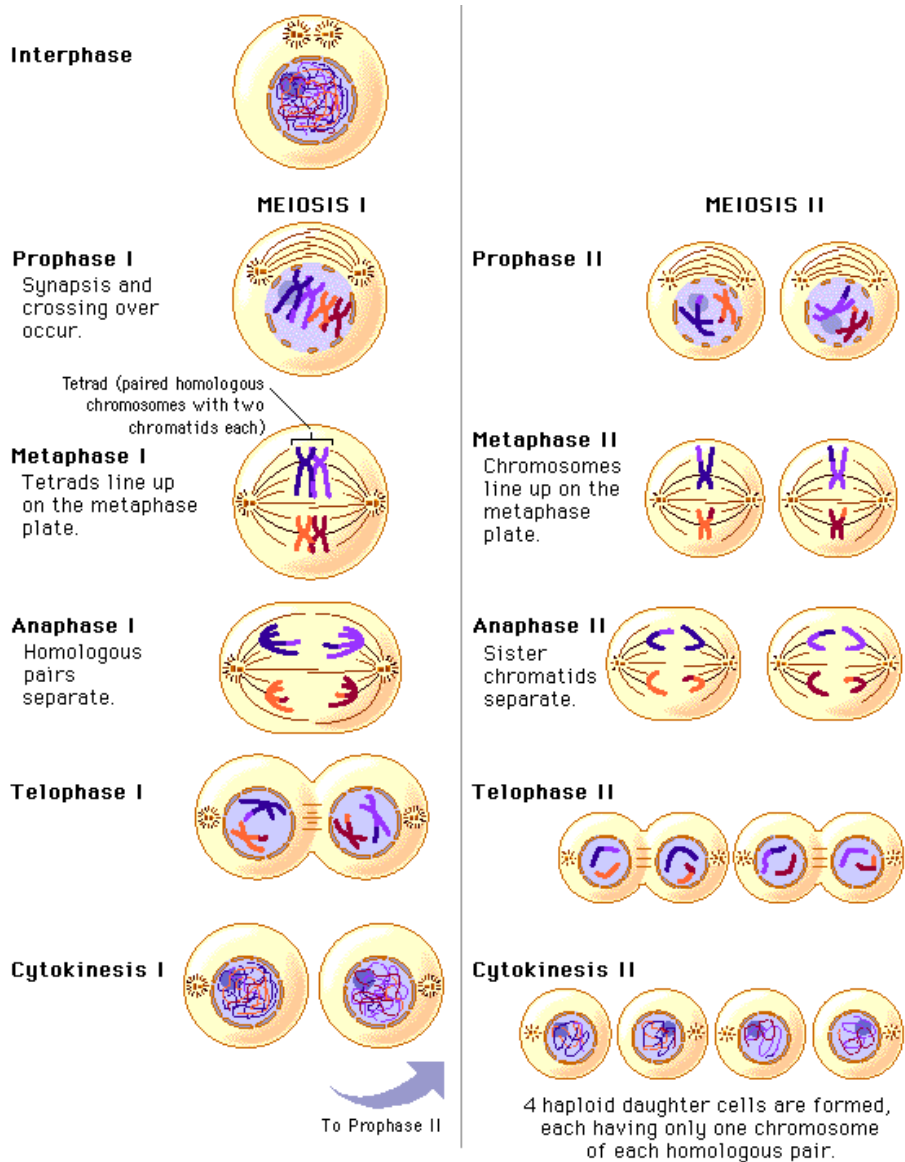


Fig.(10): Cell Division

# **Part 2**

## **Tissues**

**Tissues**

Cells work together in functionally related groups called tissues

### **Types of tissues:**

- 1- Epithelial – lining and covering
- 2-Connective – support
- 3-Muscle – movement
- 4-Nervous – control

## **Epithelial Tissue**

### **General Characteristics & Functions**

Covers a body surface or lines a body cavity  
Forms most glands

### **Functions of epithelium**

- Protection
- Absorption, secretion, and diffusion
- Filtration
- Forms slippery surfaces (mucus secretion)

## **Classifications of Epithelia**

**First name of tissue indicates number of layers**

**Simple – one layer of cells**

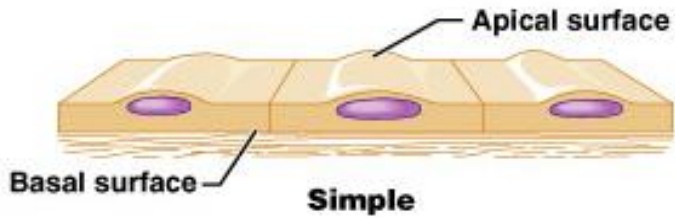


Fig. (11): Simple Epithelia cells

**\*Stratified – more than one layer of cells**

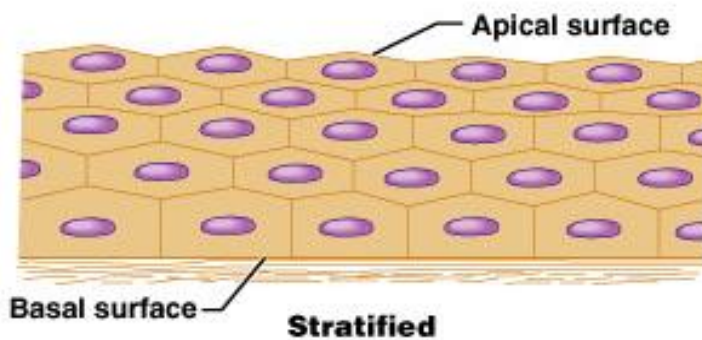


Fig. (12): Stratified – Epithelia cells

**\*Last name of tissue describes shape of cells**  
**Squamous – cells wider than tall (plate or “scale” like)**



Fig. (13): Squamous cells

**\*Cuboidal – cells are as wide as tall, as in cubes**

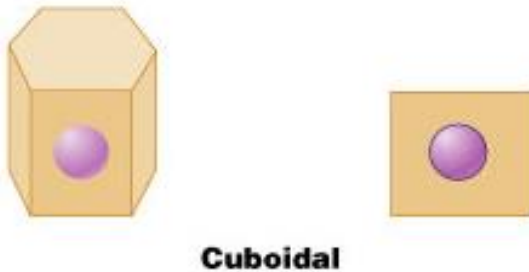
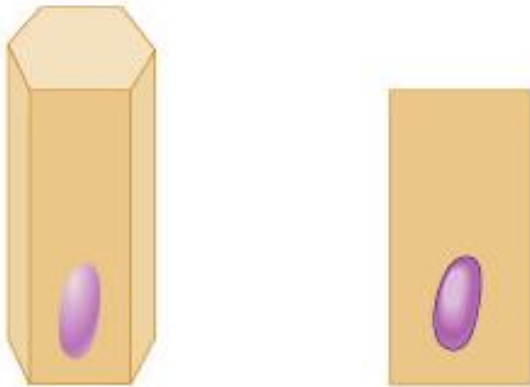


Fig. (14): Cuboidal cells

**\*Columnar – cells are taller than they are wide, like columns.**



**Columnar**

Fig. (15): Columnar – cells

### **Naming Epithelia**

Naming the epithelia includes both the layers (first) and the shape of the cells (second)

i.e. stratified cuboidal epithelium

The name may also include any accessory

### **structures**

Goblet cells

Cilia

Keratin

### **Simple Squamous Epithelium**

#### **Description**



single layer of flat cells with disc-shaped nuclei, Special types  
Endothelium (inner covering) slick lining of hollow organs

Mesothelium (middle covering)  
Lines peritoneal, pleural, and pericardial cavities  
Covers visceral organs of those cavities

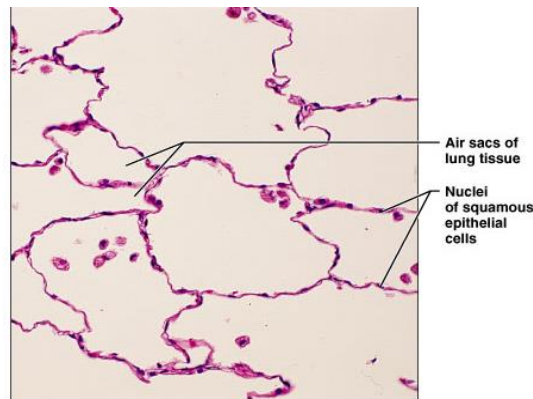
### **Function**

Passage of materials by passive diffusion and filtration

Secretes lubricating substances in serous membranes

### **Location**

- Renal corpuscles (kidneys)
- Alveoli of lungs
- Lining of heart, blood and lymphatic vessels
- Lining of ventral body cavity (serosae/serous memb.)



**Photomicrograph:** Simple squamous epithelium forming part of the alveolar (air sac) walls (400x).

## **Fig. (16): Simple Squamous Epithelium**

### **Simple Cuboidal Epithelium**

### **Description**

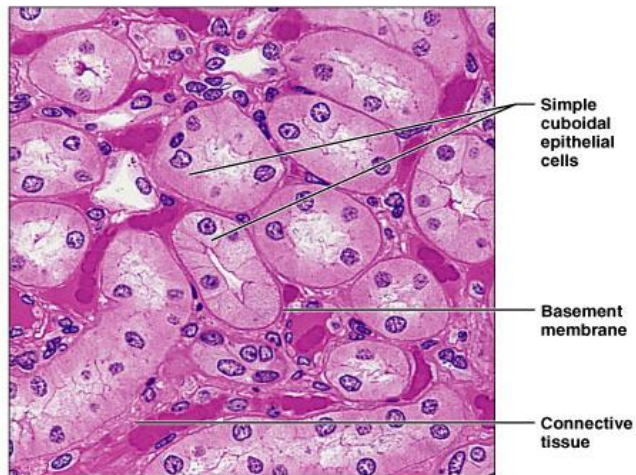
single layer of cube-like cells with large, spherical central nuclei

**Function**

secretion and absorption

**Location**

kidney tubules, secretory portions of small glands, ovary surface



**Photomicrograph:** Simple cuboidal epithelium in kidney tubules (400x).

Fig. (17): Simple Cuboidal Epithelium

**Simple Columnar Epithelium**

**Description**

single layer of column-shaped  
(rectangular) cells with oval nuclei  
Some bear cilia at their apical surface  
May contain goblet cells

### **Function**

Absorption; secretion of mucus, enzymes, and  
other substances

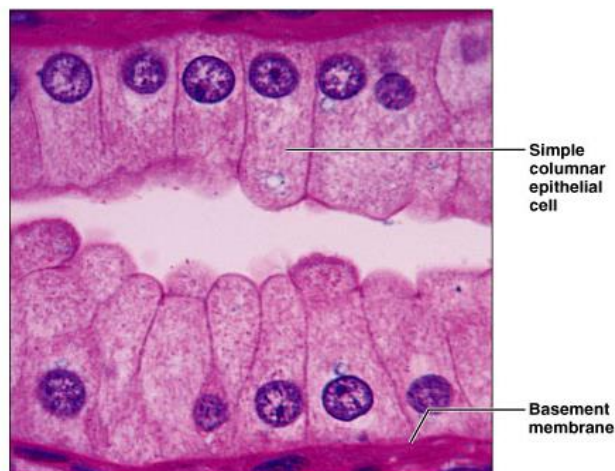
Ciliated type propels mucus or reproductive  
cells by ciliary action

### **Location**

Non-ciliated form

-Lines digestive tract, gallbladder, ducts of -  
some glands

-



**Photomicrograph:** Simple columnar epithelium  
of the stomach mucosa (1300 $\times$ ).

**Fig. (18): Simple Columnar Epithelium**

## **Ciliated Columnar Epithelium**

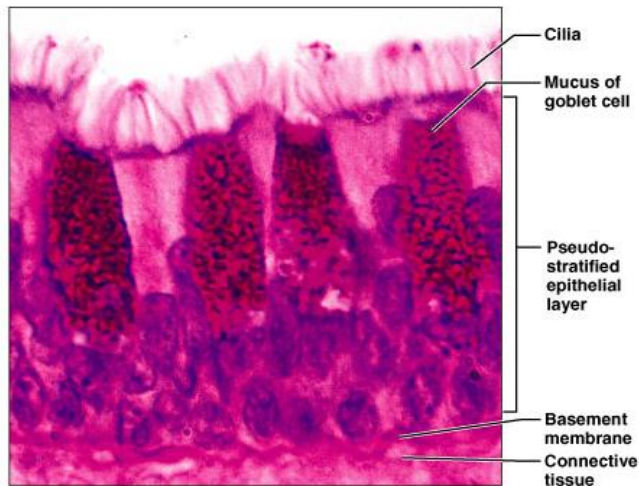
**Lines small bronchi,**

## **Description**

- All cells originate at basement membrane
  - Only tall cells reach the apical surface
  - May contain goblet cells and bear cilia
  - Nuclei lie at varying heights within cells
- Gives false impression of stratification

## **Function**

Secretion of mucus; propulsion of mucus by cilia



**Photomicrograph:** Pseudostratified ciliated columnar epithelium lining the human trachea (400 $\times$ ).

**Fig. (19): ciliated Simple Columnar Epithelium**

## **Stratified Epithelial cell**

Contain two or more layers of cells

Regenerate from below  
Major role is protection  
Are named according to the shape of cells at apical layer

## **Stratified Squamous Epithelium**

### **Description**

Many layers of cells – squamous in shape  
Deeper layers of cells appear cuboidal or columnar

Thickest epithelial tissue – adapted for protection

### **Function**

Protects underlying tissues in areas subject to abrasion

### **Location**

Keratinized – forms epidermis

Non-keratinized – forms lining of esophagus, mouth, and vagina

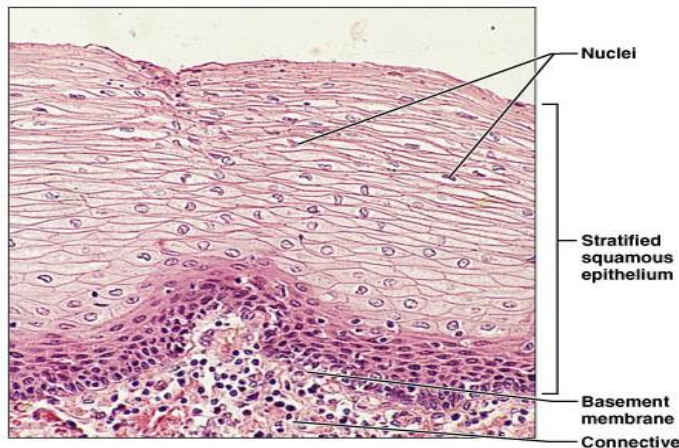


Fig. (20): Stratified Squamous Epithelium

## **Transitional Epithelium**

## Description

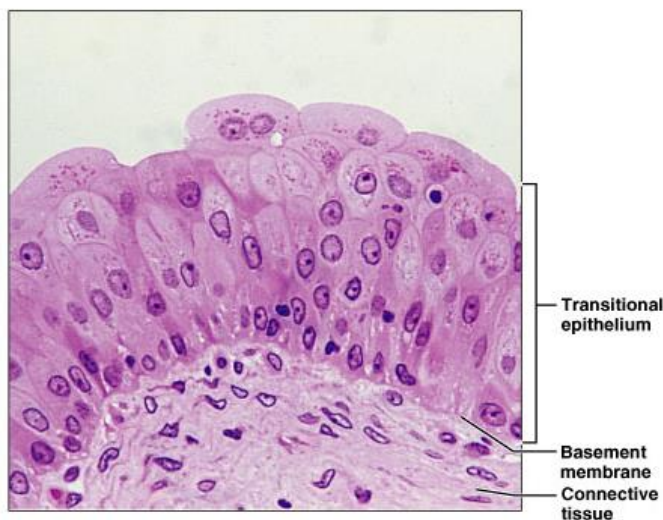
Basal cells usually cuboidal or columnar  
Superficial cells dome-shaped or squamous

## Function

stretches and permits distension of urinary bladder

## Location

Lines ureters, urinary bladder and part of urethra



**Photomicrograph:** Transitional epithelium lining of the bladder, relaxed state (500 $\times$ ); note the bulbous, or rounded, appearance of the cells at the surface; these cells flatten and become elongated when the bladder is filled with urine.

Fig. (21): Transitional Epithelium

## Connective Tissue

Most diverse and abundant tissue

### **Main classes**

- Connective tissue proper
- Cartilage
- Bone tissue
- Blood

### **Components of connective tissue:**

- Cells (varies according to tissue)
- Matrix
- Fibers (varies according to tissue)
- Ground substance (varies according to tissue) dermatin sulfate, hyaluronic acid, keratin sulfate, chondroitin sulfate...

### **Common embryonic origin**

- mesenchyme

### **Connective Tissue Proper**

- Loose Connective Tissue
- Areolar
- Reticular
- Adipose
- Dense Connective Tissue
- Regular
- Irregular
- Elastic

### **Areolar Connective Tissue**

#### **Description**

-Gel-like matrix with:

all three fiber types (collagen, reticular, elastic) for support

-Ground substance is made up by glycoproteins also made and secreted by the fibroblasts.

Cells – fibroblasts, macrophages, mast cells, white blood cells

### **Function**

-Wraps and cushions organs

-Holds and conveys tissue fluid

-Important role in inflammation Main battlefield in fight against infection

-Defenders gather at infection sites

Macrophages

Plasma cells

Mast cells

Neutrophils, lymphocytes, and eosinophils

### **Location**

-Widely distributed under epithelia

-Packages organs

Surrounds capillaries

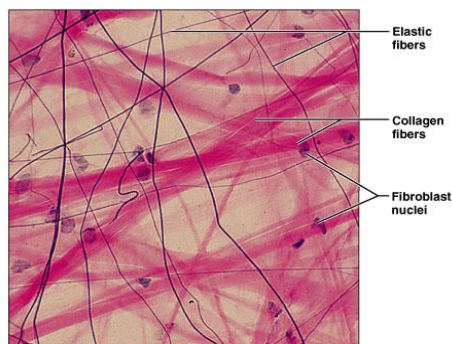


Fig. (22): Areolar Connective Tissue



## **Adipose Tissue**

### **Description**

Closely packed adipocytes  
Have nucleus pushed to one side by fat droplet

### **Function**

Provides reserve food fuel  
Insulates against heat loss  
Supports and protects organs

### **Location**

Under skin  
Around kidneys  
Behind eyeballs, within abdomen and in breasts

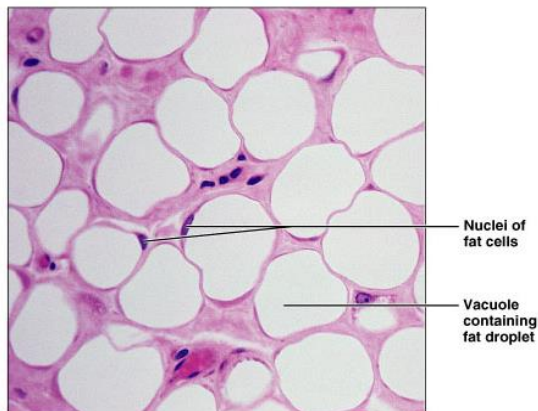


Fig. (23): Adipose Tissue

## Reticular Connective Tissue

### **Description**

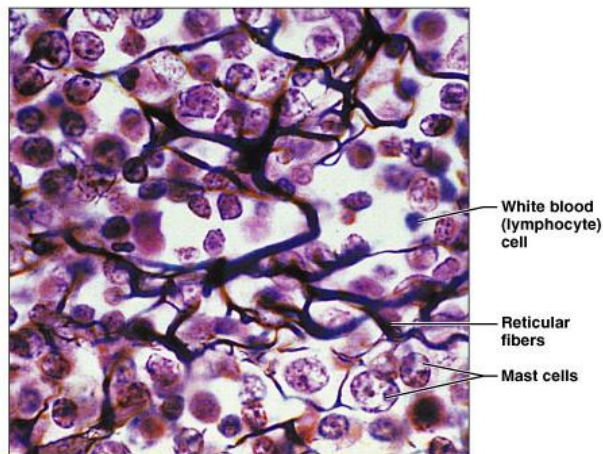
– network of reticular fibers in loose ground substance

### **Function**

– form a soft, internal skeleton (stroma) – supports other cell types

### **Location**

– lymphoid organs  
- Lymph nodes, bone marrow, and spleen



**Photomicrograph:** Dark-staining network of reticular connective tissue fibers forming the internal skeleton of the spleen (350 $\times$ ).

Fig. (24): Reticular Connective Tissue

## Dense Regular Connective Tissue

### **Description**

Primarily *parallel* collagen fibers  
Fibroblasts and some elastic fibers  
Poorly vascularized

### **Function**

Attaches muscle to bone  
Attaches bone to bone  
Withstands great stress in  
one direction

### **Location**

Tendons and ligaments  
Aponeuroses  
Fascia around muscle

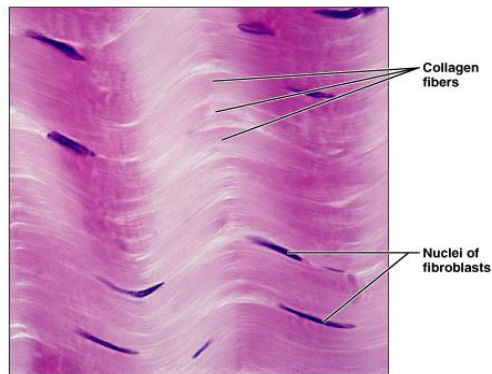


Fig. (25): Dense Regular Connective Tissue

## **Cartilage**

### **Characteristics:**

Firm, flexible tissue

Contains no blood vessels or nerves

Matrix contains up to 80% water

Cell type – chondrocyte

### **Types:**

Hyaline

Elastic

Fibrocartilage

## Hyaline Cartilage

### **Description**

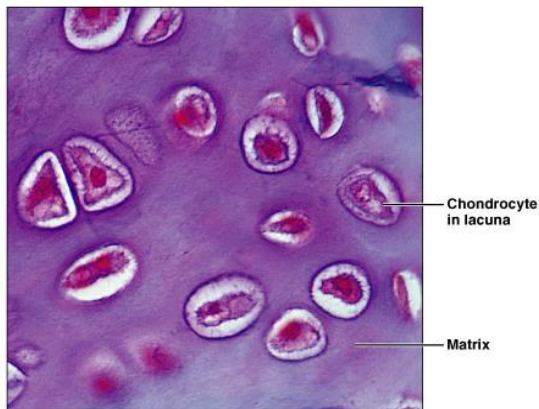
Imperceptible collagen fibers (hyaline = glassy)  
Chondroblasts produce matrix  
Chondrocytes lie in lacunae

### **Function**

- Supports and reinforces
- Resilient cushion
- Resists repetitive stress

### **Location**

Fetal skeleton  
Ends of long bones  
Costal cartilage of ribs  
Cartilages of nose,  
trachea, and larynx



**Photomicrograph:** Hyaline cartilage from the trachea (300×).

Fig. (26): Hyaline Cartilage

## Elastic Cartilage

### **Description**

Similar to hyaline cartilage

More elastic fibers in matrix

### **Function**

-Maintains shape of structure

-Allows great flexibility

### **Location**

-Supports external ear

-Epiglottis

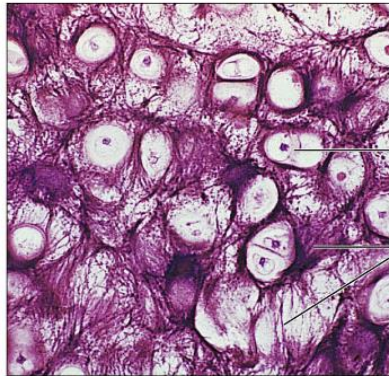


Fig. (27): Elastic Cartilage

## Fibrocartilage

## **Description**

Matrix similar, but less firm than hyaline cartilage

Thick collagen fibers predominate

## **Function**

Tensile strength and ability to absorb compressive shock

## **Location**

Intervertebral discs

Pubic symphysis

Discs of knee joint

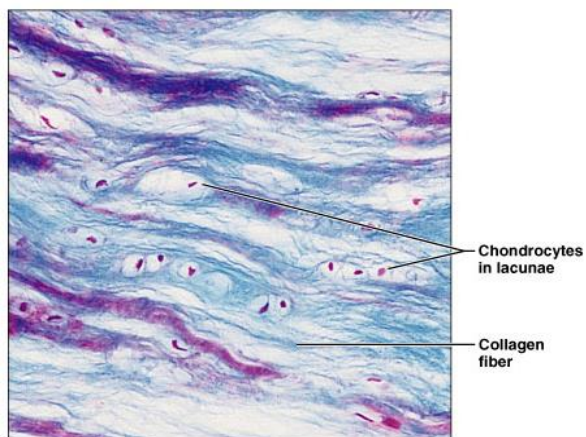


Fig. (28): Fibrocartilage

## **Bone Tissue**

## **Function**

- Supports and protects organs
- Provides levers and attachment site for muscles
- Stores calcium and other minerals
- Stores fat
- Marrow is site for blood cell formation

## **Location**

Bones

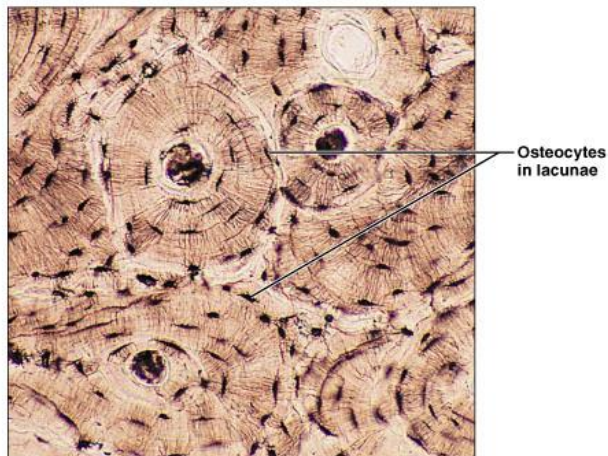


Fig. (29) Bone Tissue

:

## **Blood Tissue**



## Description

red and white blood cells  
in a fluid matrix

## Function

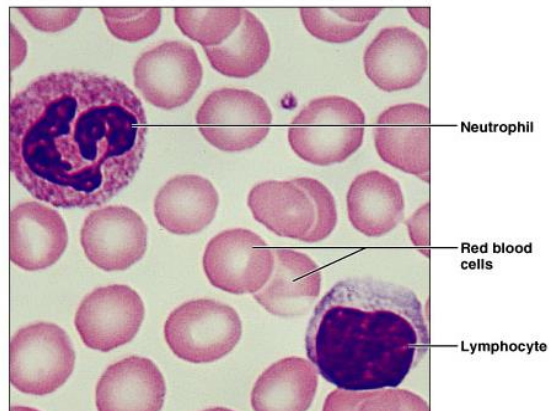
-transport of respiratory gases, nutrients, and  
wastes

## Location

within blood vessels

## Characteristics

An atypical connective tissue  
Develops from mesenchyme  
Consists of cells surrounded by nonliving  
matrix



**Photomicrograph:** Smear of human blood (1500 $\times$ ); two white blood cells (neutrophil in upper left and lymphocyte in lower right) are seen surrounded by red blood cells.

Fig. (30): Blood Tissue

## Muscle Tissue

## Types

- Skeletal muscle tissue
- Cardiac muscle tissue
- Smooth muscle tissue

## Skeletal Muscle Tissue

### Characteristics

Long, cylindrical cells, Multinucleate Obvious striations

### Function

Voluntary movement  
Manipulation environment  
Facial expression

### Location

Skeletal muscles attached to bones  
(occasionally to skin)

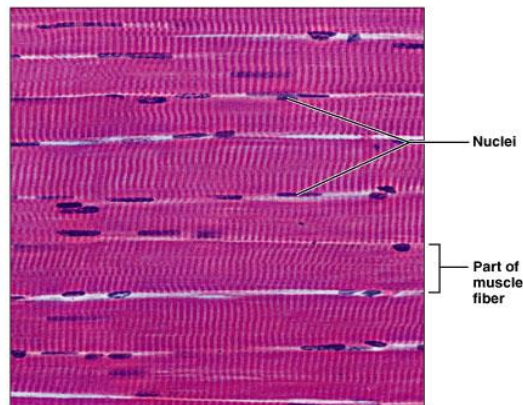


Fig. (31): Skeletal Muscle Tissue  
Cardiac Muscle Tissue

### Function

Contracts to propel blood into circulatory system

### **Characteristics**

Branching cells, Uninucleate, Striations  
Intercalated discs

### **Location**

Occurs in walls of heart

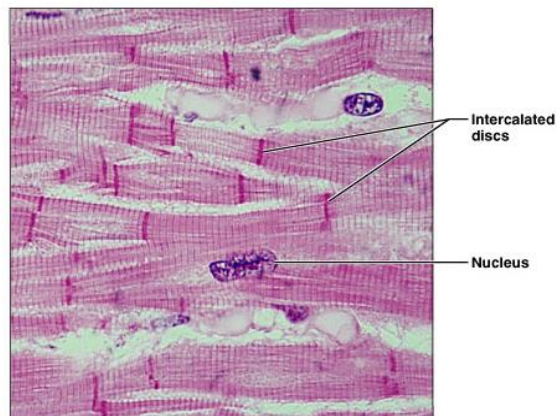


Fig. (32): Cardiac Muscle Tissue

## **Smooth Muscle Tissue**

### **Characteristics**

Spindle-shaped cells with  
central nuclei  
Arranged closely to form  
sheets  
No striations

**Function**

Propels substances along  
internal passageways  
Involuntary control

**Location**

Mostly walls of hollow organs

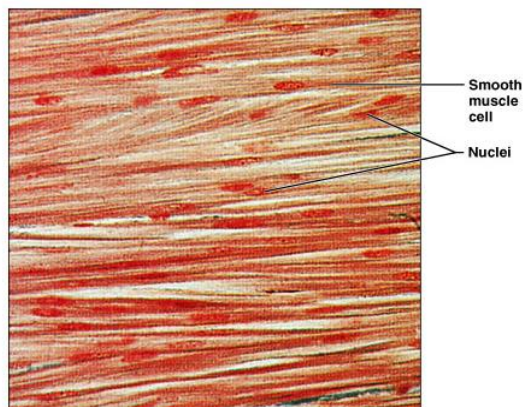


Fig. (33): Smooth Muscle Tissue

**Nervous Tissue**

## Function

Transmit electrical signals from sensory receptors to effectors.

## Location

Brain, spinal cord, and nerves

## Description

Main components are brain, spinal cord, and nerves

Contains two types of cells

Neurons – excitatory cells

### Supporting cells (neuroglial cells)

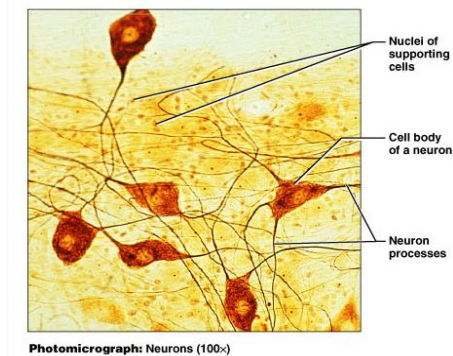


Fig. (34): Nervous Tissue



مقرر

# علم الأجنة Embryology

الفرقة الأولى شعبة العلوم البيولوجية والجيولوجية باللغة الإنجليزية (101 عل ح)

أستاذ المقرر

د/ سهام علي مبارك محمد







قسم علم الحيوان - كلية العلوم بقنا

العام الجامعي  
2023 / 2022م

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## Introduction


-  Human development is a continuous process that begins when an oocyte (ovum) from a female is fertilized by a sperm (spermatozoon) from a male.
-  Cell division, cell migration, programmed cell death (apoptosis), differentiation, growth, and cell rearrangement transform the fertilized oocyte, a highly specialized, totipotent cell, a zygote, into a multicellular human being.
-  Most changes occur during the embryonic and fetal periods; however, important changes also occur during later periods of development.
-  The field of study which includes investigations of the molecular, cellular, and structural factors contributing to the formation of an organism is called embryology.
-  It is a branch of science that is related to the formation, growth, and development of an embryo.
-  It mostly deals with the prenatal stage of development beginning from the formation of gametes, fertilization, the formation of a zygote, development of embryo and fetus to the birth of a new individual.




## Definitions of Embryology

-  Embryology is a branch of science that is related to the fertilization, formation, growth, and development of embryo. In mammals, it deals with the prenatal stage of development beginning from formation of gametes, fertilization, formation of zygote, development of embryo and fetus to the birth of a new individual.



 Embryology is the study of the early development of living organisms till it reaches to adult form.

 All animals and insects can reproduce new individuals to ensure the survival of their kind.



## A Brief History of Embryology

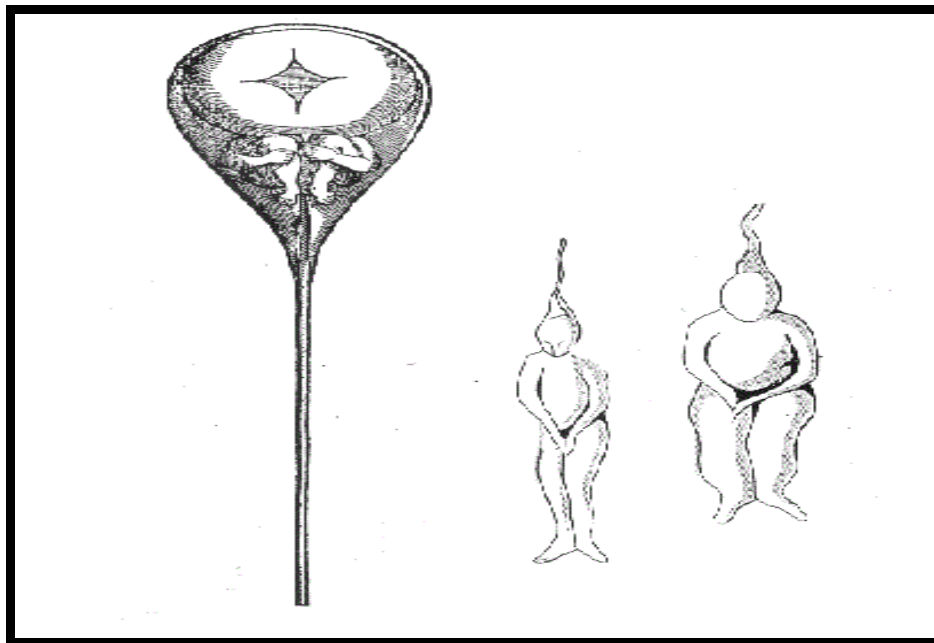
### The theory of preformationism

- ✓ Early embryology was proposed by Marcello Malpighi, and known as **preformationism**, a theory held that the generation of offspring occurs as a result of an unfolding and growth of preformed parts. There were two competing models of preformationism: **the ovism model**, in which the location of these preformed parts prior to gestation was the maternal egg, and the **spermism model**, in which a preformed individual or homunculus was thought to exist in the head of each sperm.
- ✓ Preformationism was the first theory of generation and development that applied to all organisms in the plant and animal kingdoms.
- ✓ The theory of preformation gained much traction before the invention of microscopes and more advanced imaging techniques.
- ✓ The theory also suggested women were simply vessels to carry the growing child, and that girls came from the left testicle, while boys came from the right.

### The theory of epigenesis

- ✓ Aristotle first proposed the correct mechanism for the development of an embryo, without having a microscope to observe his theory.

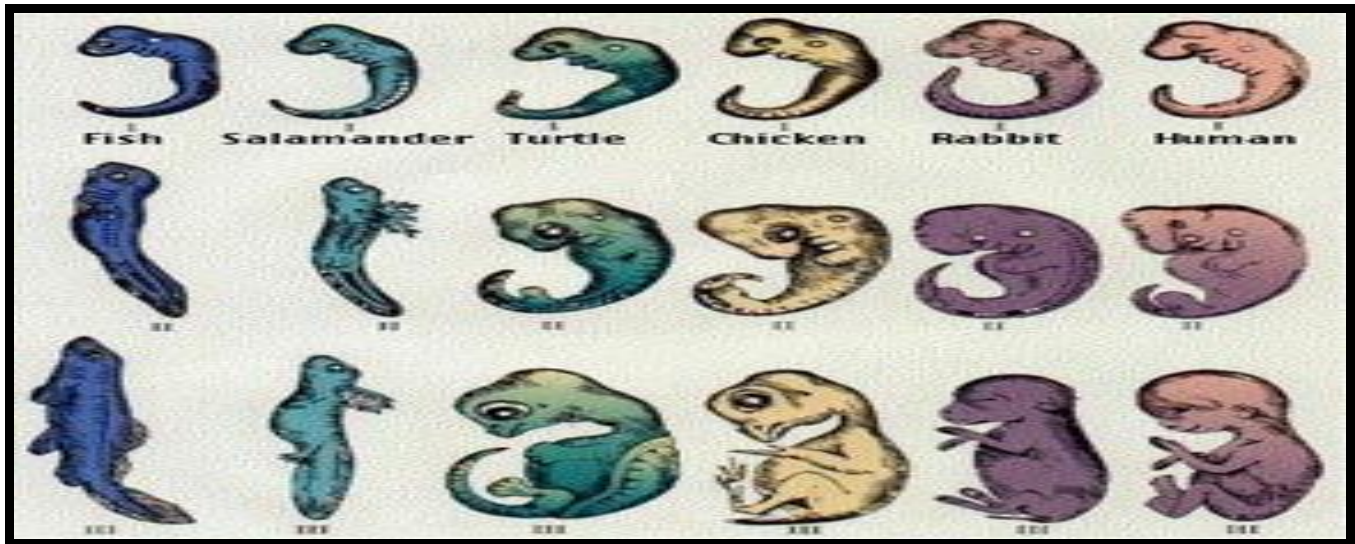
- ✓ He suggested that animals form through the process of epigenesis, in which a single cell divides and differentiates into the many tissues and organs of an animal.
- ✓ It wasn't until 1827 that clear evidence was obtained that female mammals also produce a sex cell, the ovum.
- ✓ The discovery of a female sex cell directly contradicted many aspects of the preformation theory and led to wider acceptance of the epigenesis theory.



A tiny person (a *homunculus*) inside a sperm, as drawn by Nicolaas Hartsoecker in 1695

### The theory of recapitulation

- ✓ The embryonic forms of an animal resembled the adult organisms in its evolutionary ancestry.
- ✓ A historical hypothesis that the development of the embryo of an animal, from fertilization to gestation or hatching (ontogeny), goes through stages resembling or representing successive adult stages in the evolution of the animal's remote ancestors (phylogeny).



[https://thebrain.mcgill.ca/flash/capsules/outil\\_bleu12.html](https://thebrain.mcgill.ca/flash/capsules/outil_bleu12.html)

## The Cell Theory



The cell theory (proposed independently in 1838 and 1839) is a cornerstone of biology.



All organisms are composed of one or more cells.



Cells are the smallest living things.



Cells arise only by division of previously existing cells.



Ovum was discovered as a single cell and that fertilization is the union of the ovum and spermatozoon to form the zygote.



## Importance of embryology

For example:



Embryology is the basis for understanding the intimate relation between structures in different organ systems, such as the nervous system and muscle, and is primordial for understanding disorders of development that in the human may present as one of the congenital

myopathies.



Provide knowledge essential for creating health care strategies for better reproductive outcomes



A better understanding of embryology results in new techniques for prenatal diagnoses and treatments, therapeutic procedures to circumvent problems with infertility, and mechanisms to prevent birth defects, the leading cause of infant mortality.



Supports the research and application of stem cells for the treatment of certain chronic diseases.



## **Subspecialties (Fields) of embryology**



**Descriptive embryology:** This field of embryology associated with the morphological description of different embryonic stages in the ontogenetic development of individuals of different species.



**Comparative embryology:** the study of how anatomy changes during the development of different organisms.



**Evolutionary embryology:** the study of how changes in development may cause evolutionary changes and of how an organism's ancestry may constrain the types of changes that are possible.



**Experimental embryology:** It involves all those studies that attempt to understand the various fundamental mechanism in the development of different animals, like fertilization, cleavage, gastrulation, embryonic induction, determination, and differentiation.



**Behavioral embryology:** the study of the early development of the

nervous system and behavior with a view toward understanding how the formative periods of neural and behavioral development affect later stages of neurobehavioral ontogeny.



**Chemical embryology:** The branch of embryology includes all those studies which employ various biochemical, biophysical and physiological techniques for understanding embryological events at a molecular level.



**Teratology:** Teratology is the division of embryology and pathology that deals with abnormal development (birth defects). This branch of embryology is concerned with various genetic and/or environmental factors that disturb normal development and produce birth defects.



## Reproduction

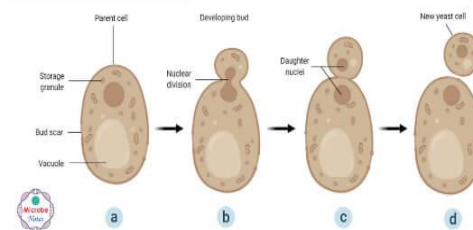
Reproduction may be defined as the biological process by which organisms give rise to their own kind. Reproduction may occur in two ways: **Asexual and Sexual reproduction.**

- Bacterial and protozoan offspring may be produced by single individuals. This is termed asexual reproduction. Lower animals and many plants reproduce asexually. When two individuals are involved in reproduction, it is termed sexual reproduction. Some methods of asexual reproduction are - fission, budding, fragmentation and spore formation.

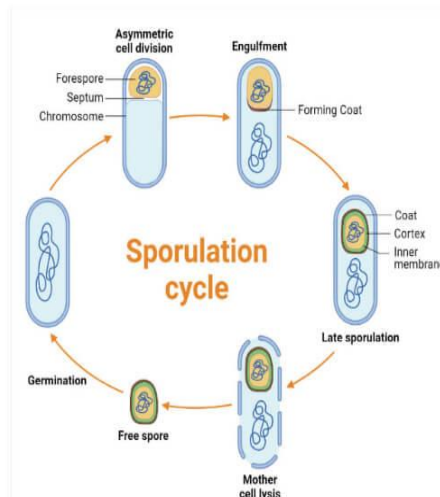
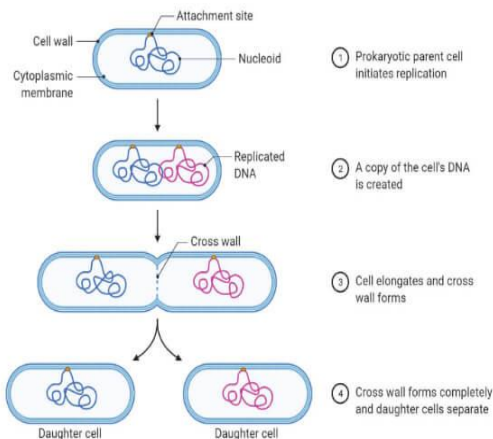
# Asexual Reproduction

## Definition, Features, Types, Examples, Advantages

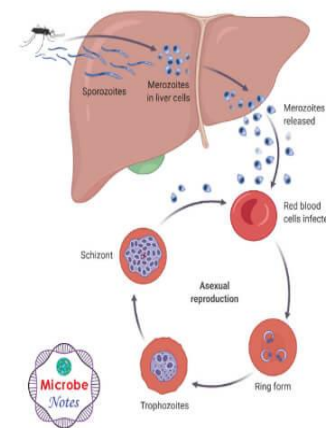
### BUDDING IN YEAST



### Prokaryotic Cell Division by Binary Fission



### Malaria Asexual Reproduction

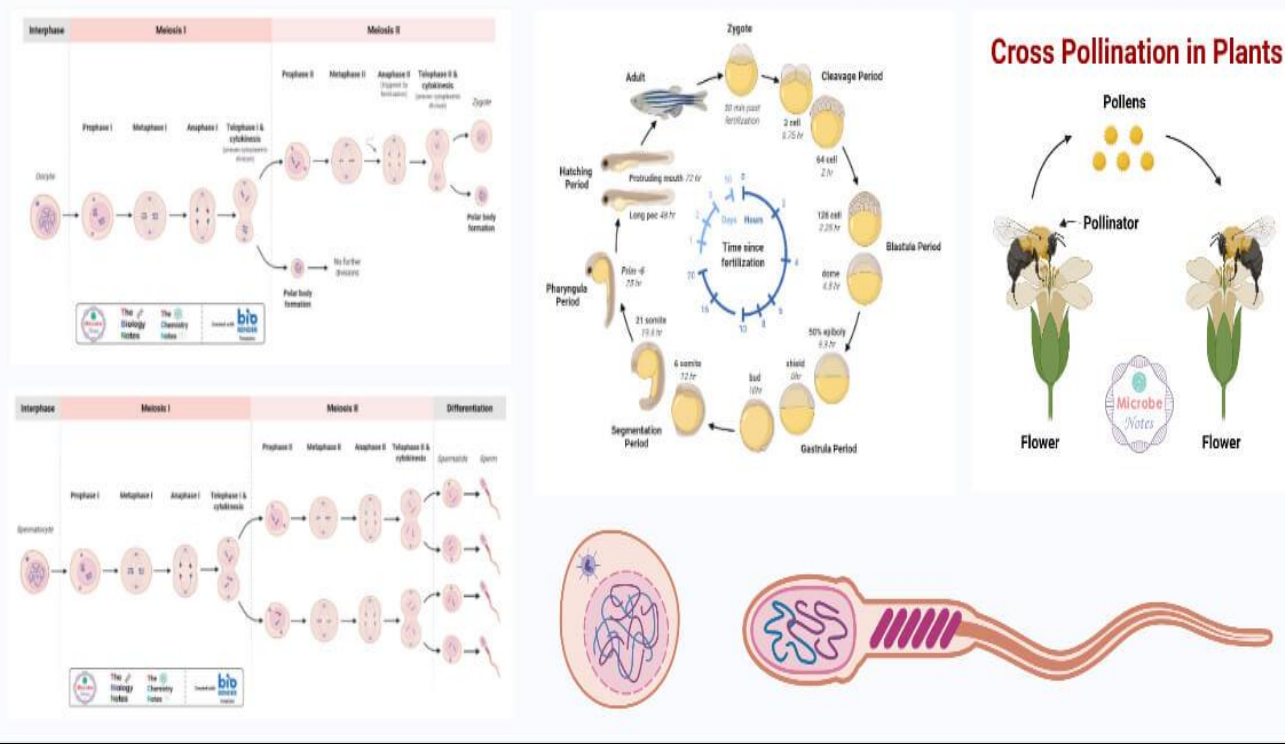


Asexual Reproduction. Created with biorender.com

- In sexual reproduction, is a type of reproduction that involves a complex life cycle in which a gamete (such as a sperm or egg cell) with a single set of chromosomes (haploid) combines with another to produce a zygote that develops into an organism composed of cells with two sets of chromosomes (diploid). Sexual reproduction is the most common life cycle in multicellular eukaryotes, such as animals, fungi and plants. This is a more common mode of reproduction in plants and animals.

# Sexual Reproduction

## Definition, Features, Stages, Types, Examples



<https://thebiologynotes.com/sexual-reproduction/>



**Basic Concepts of embryonic development include:**

1. Gametogenesis
2. Fertilization
3. Cleavage
4. Blastulation
5. Gastrulation
6. Organization (Organogenesis)



## Gametogenesis

Gametogenesis is the process by which male and female sex cells or gametes, i.e., sperms and ova are formed respectively in the male and female gonads (testes and ovaries). The gametes differ from all other cells (= somatic cells) of the body in that their nuclei contain only half the number of chromosomes found in the nuclei of somatic cells. Meiosis forms the most significant part of process of gametogenesis.

Gametogenesis for the formation of sperms is termed **spermatogenesis**, while that of ova is called **oogenesis**.

- ❖ Both spermatogenesis and oogenesis comprise similar phases of sequential changes as: **multiplication phase, growth, and maturation phases**.



### Spermatogenesis:

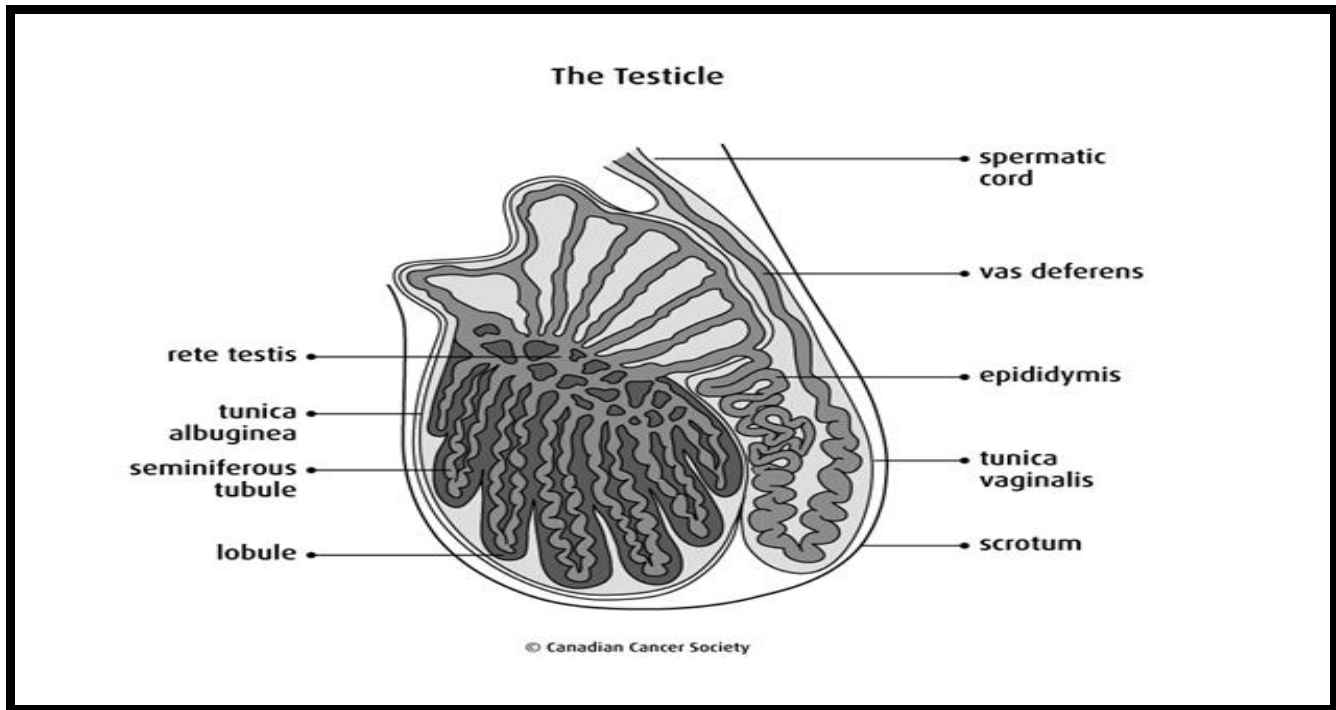
The process of formation of sperms is called spermatogenesis. It occurs in the seminiferous tubules of the testes. The seminiferous tubules are lined by germinal epithelium. The germinal epithelium consists largely of cuboidal primary or primordial germ cells (PGCs) and contains certain tall somatic cells called Sertoli cells (= nurse cells). Spermatogenesis includes formation of spermatids and formation of spermatozoa.



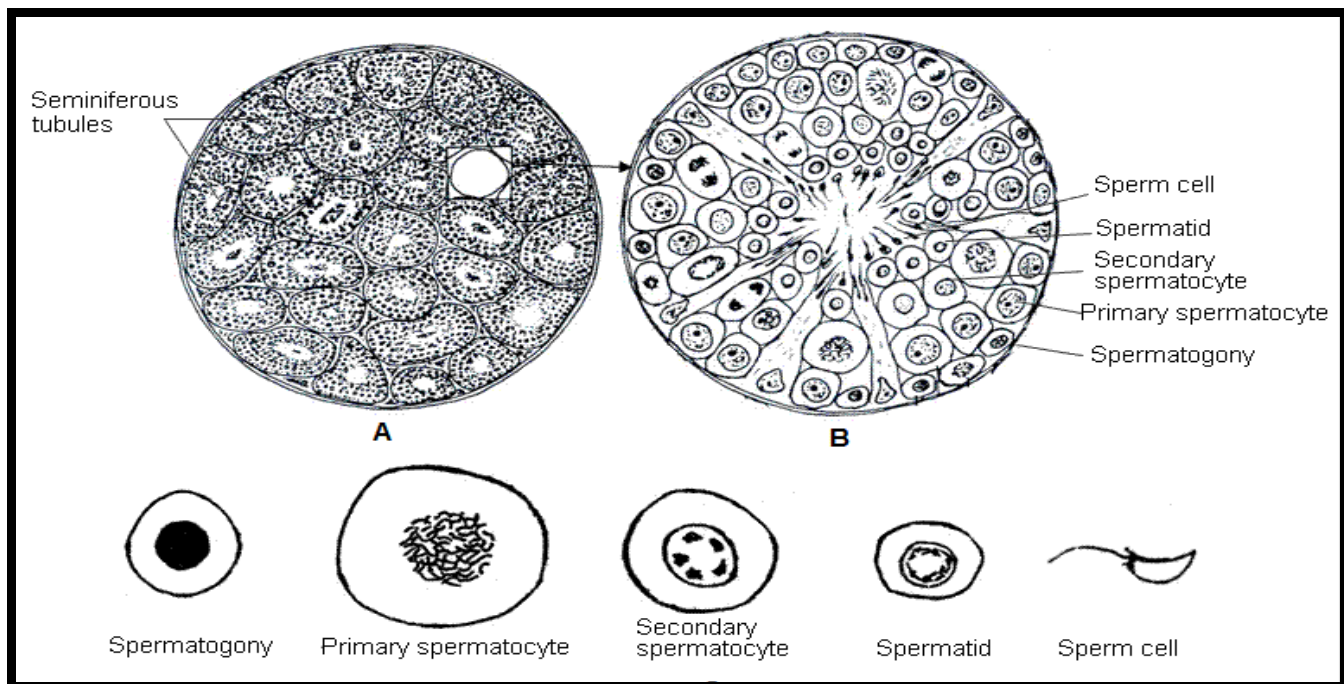
### The testes structure

The testicles (testes) are part of a man's reproductive system. A man has 2 testicles. Each testicle is egg-shaped and about 5 cm long. The testicles are covered by a sac of skin called the scrotum. The scrotum hangs below the penis, between the legs. The testicles make sperm. They also make testosterone, which is a male sex hormone.





T.s. of rat testis



[https://cit.vfu.cz/frvs2011/?title=ukoly-rozmnozovani\\_a\\_vyvoj-meioza&lang=en](https://cit.vfu.cz/frvs2011/?title=ukoly-rozmnozovani_a_vyvoj-meioza&lang=en)

- ✓ Each testicle is covered by tough, fibrous layers of tissue called the tunica. The outer layer is called the tunica vaginalis and the inner layer is called the tunica albuginea.

- ✓ The testicle is divided into parts called lobules. Each lobule contains tiny U-shaped tubes called seminiferous tubules. There are about 800 seminiferous tubules tightly coiled within each testicle.
- ✓ The seminiferous tubules open into a series of uncoiled, interconnected channels called the rete testis. Ducts, or tubes, connect the rete testis to a tightly coiled tube called the epididymis. The epididymis joins to a long, large duct called the vas deferens.
- ✓ Each testicle is held in the scrotum by a spermatic cord. Each spermatic cord is made of tough connective tissue and muscle. It contains the vas deferens, blood vessels, lymph vessels and nerves.
- ✓ Lymph fluid
- ✓ travels through vessels in the spermatic cord and drains from the testicles into several groups of lymph nodes at the back of the abdomen. These lymph nodes are called the retroperitoneal lymph nodes.
- ✓ Sertoli cells are the cells that are randomly scattered throughout the seminiferous tubules and provide nutrients to the developing spermatogonia.

 **Spermatogenesis:** includes the following phases:

#### **Multiplication Phase:**

At sexual maturity, the undifferentiated primordial germ cells divide several times by mitosis to produce a large number of spermatogonia (Gr. sperma = seeds, gonos- generation). Spermatogonia (2N) are of two types: type A spermatogonia and type B spermatogonia. Type A spermatogonia serve as the stem cells which divide to form additional spermatogonia. Type B spermatogonia are the precursors of sperms.

#### **Maturation Phase:**

Each primary spermatocyte undergoes two successive divisions, called maturation divisions. The first maturation division is reductional or meiotic.

Hence, the primary spermatocyte divides into two haploid daughter cells called secondary spermatocytes. Both secondary spermatocytes now undergo second maturation division which is an ordinary mitotic division to form, four haploid spermatids, by each primary spermatocyte.

### Growth Phase:

Each type B spermatogonium actively grows to a larger primary spermatocyte by obtaining nourishment from the nursing cells.

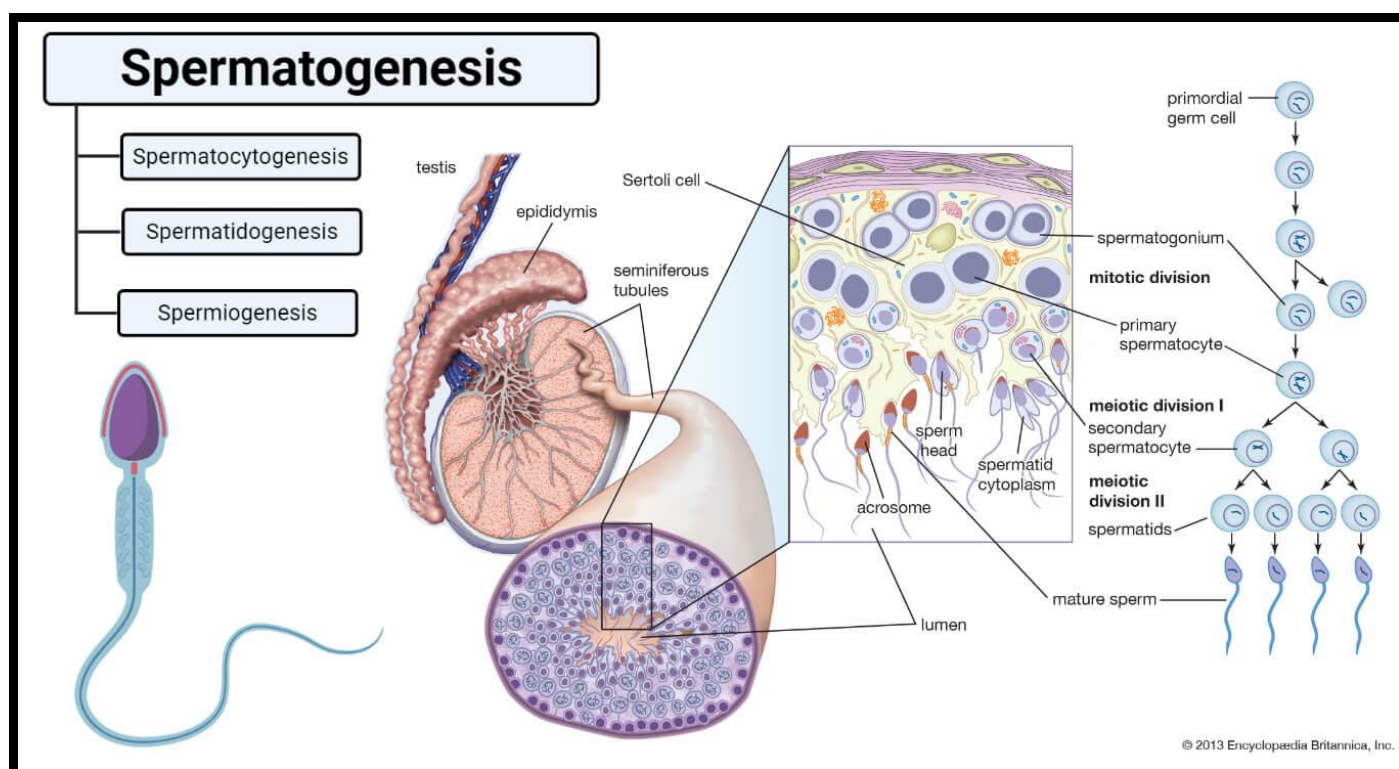
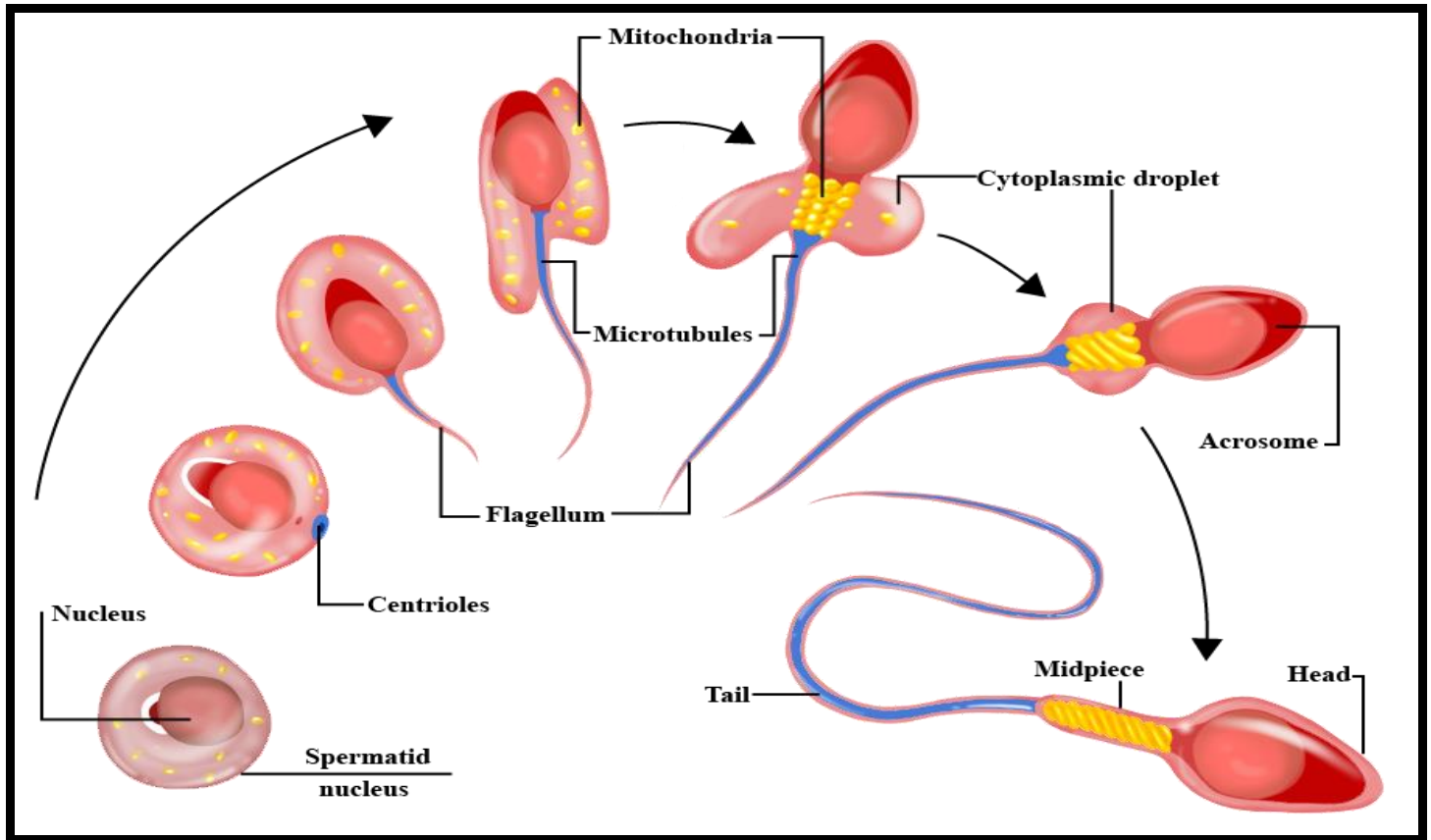


Image Source: [Britannica](#), Created with BioRender.com.

### Formation of Sperms from Spermatids (Spermiogenesis):

The transformation of spermatids into spermatozoa is called spermiogenesis. The spermatozoa are later on known as sperms. Thus four sperms are formed from one spermatogonium. After spermiogenesis sperm heads become embedded in the Sertoli cells and are finally released from the seminiferous tubules by the process called spermiation.



<https://byjus.com/question-answer/spermiogenesis-spermateliosis-is-formation-of-spermatozoa-from/>

## Spermatozoon (Sperm):

The sperms are microscopic and motile cells. Sperms remain alive and retain their ability to fertilize an ovum (egg) from 24 to 48 hours after having been released in the female genital tract. A typical mammalian sperm consists of a head, neck, middle piece and tail.

### Head:

It contains anterior small acrosome and posterior large nucleus. Acrosome is formed from Golgi body of the spermatid. Acrosome contains hyaluronidase proteolytic enzymes which are popularly known as sperm lysins that are used to contact and penetrate the egg (ovum) at the time of fertilization.

### Neck:

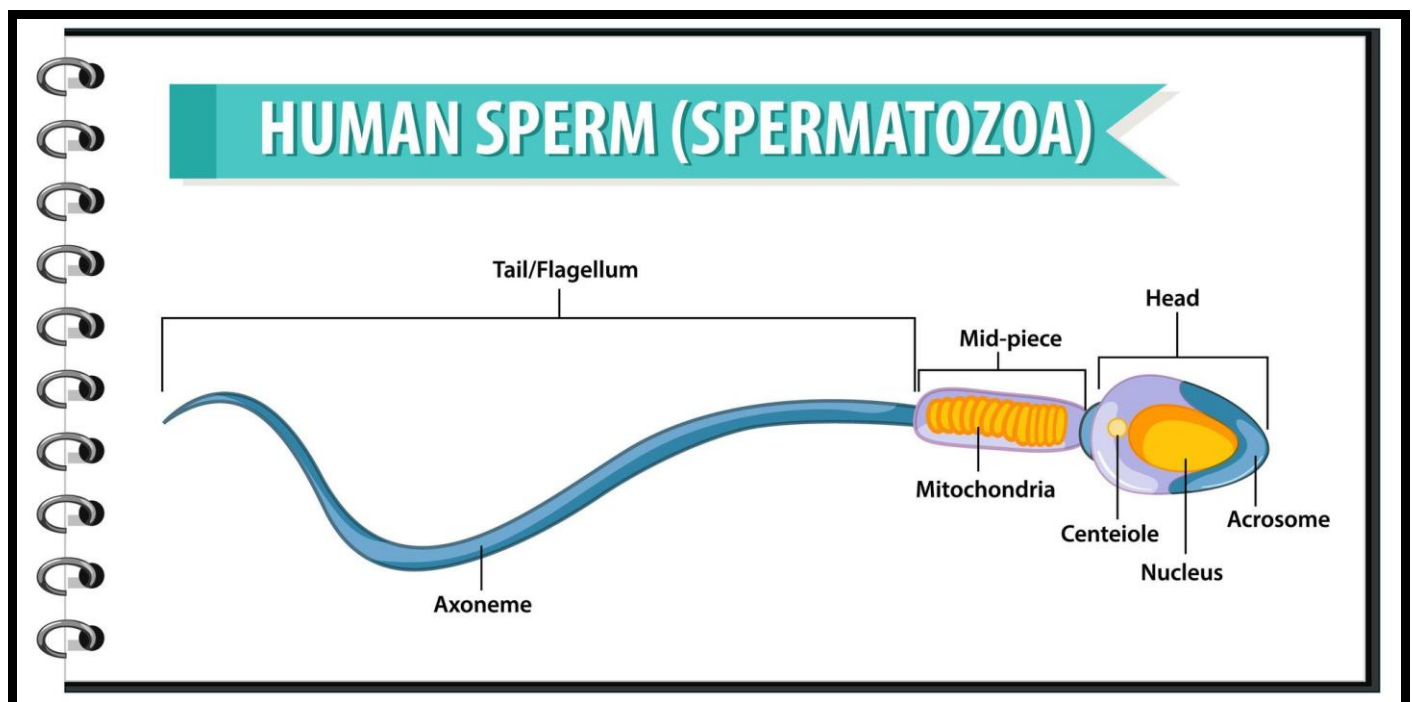
It is very short and is present between the head and middle piece. It contains the proximal centriole towards the nucleus which plays a role in the first cleavage of the zygote and the distal centriole which gives rise to the axial filament of the sperm.

**Middle piece:**

The middle piece of human sperm contains the mitochondria coiled around the axial filament called mitochondrial spiral. They provide energy for the movement of the sperm. So, it is the “power house of the sperm”. At the end of the middle piece there is a ring centriole (annulus) with unknown function. Posterior half of nucleus, neck and middle piece of sperm are covered by a sheath called manchette.

**Tail:**

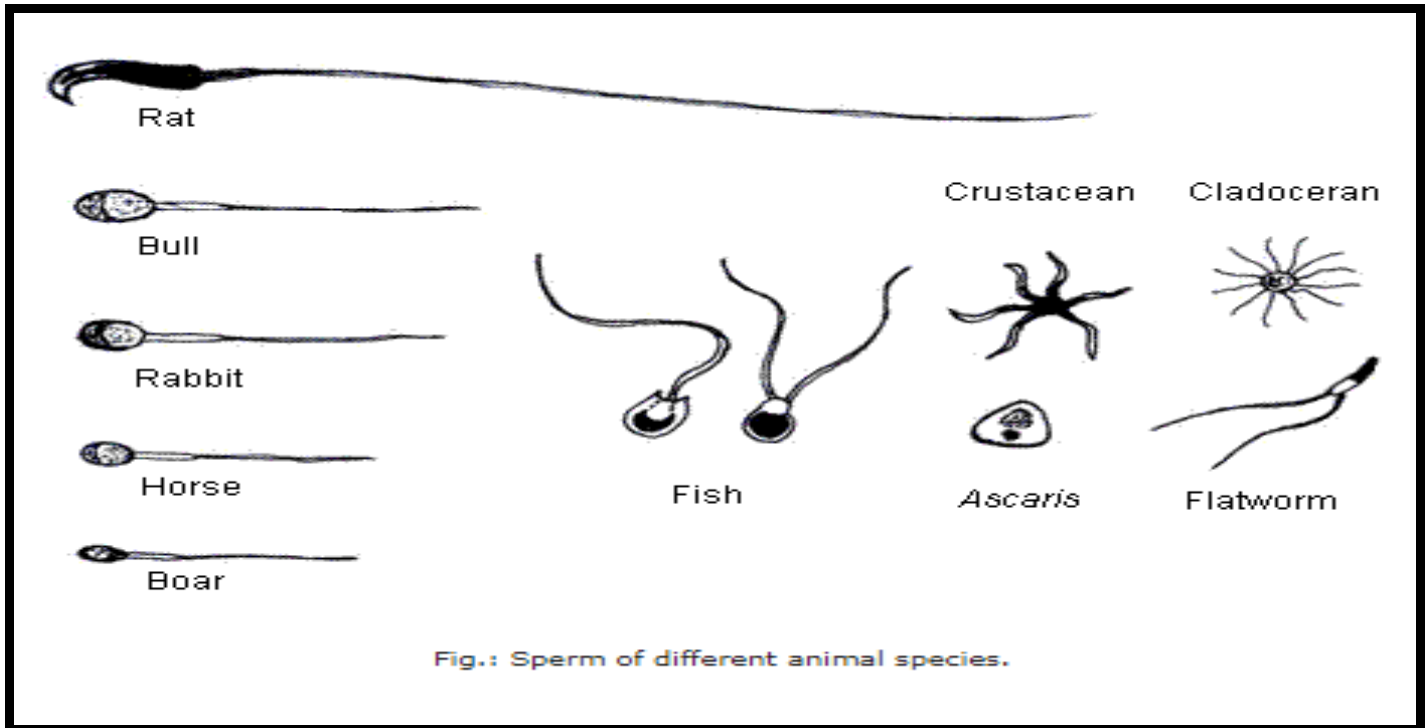
The tail is several times longer than the head. In its most part called main piece, the axial filament is surrounded by a thin layer of cytoplasm. The part behind the main piece is called end piece which consists of naked filament alone. The sperm swims about by its tail in a fluid medium.



<https://www.vecteezy.com/vector-art/1434164-human-sperm-or-spermatozoa-cell-structure>

**Different shape and size of sperm**

Observe sperm of different animal species and compare the shape of sperm head, the size of acrosome (structure on the top of head containing enzymes important for penetrating the egg) and size and number of sperm tails.



[https://cit.vfu.cz/frvs2011/?title=ukoly-rozmnzovani\\_a\\_vyvoj-meioza&lang=en](https://cit.vfu.cz/frvs2011/?title=ukoly-rozmnzovani_a_vyvoj-meioza&lang=en)

## Purpose of Spermatogenesis

The process of Spermatogenesis occurs to create mature male gametes, which then fertilize female gametes to create a zygote, a single-celled organism. This results in cell division and multiplication to create a fetus. For a healthy offspring, the number of chromosomes must be maintained properly across the body as failure can lead to some abnormalities.



## Oogenesis

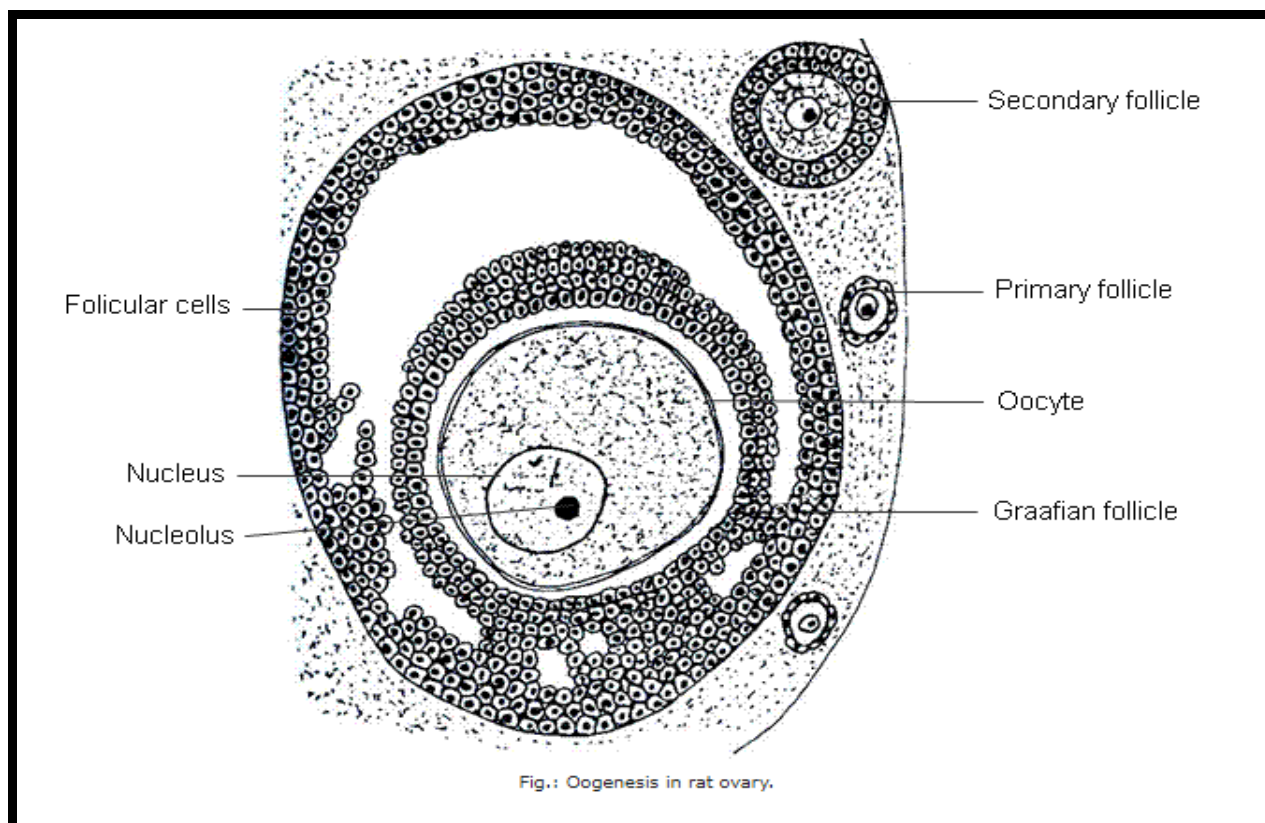
The process of formation of a mature female gamete (ovum) is called oogenesis. It occurs in the ovaries (female gonads).

### Ovum structure:

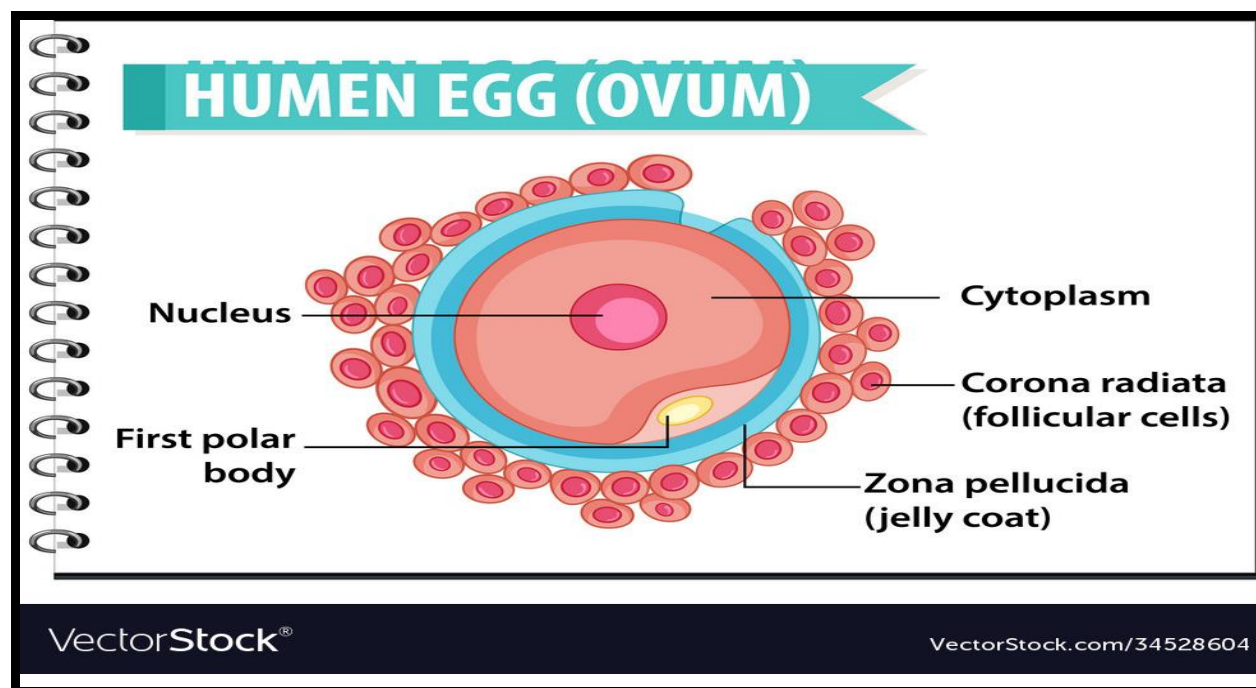
The ovum is one of the largest cells that measures approximately 120  $\mu\text{m}$  in diameter. The ovum has a large, centrally located nucleus which is covered by cytoplasm. This oocyte nucleus and nucleolus are termed **germinal vesicle** and **germinal disc** respectively. Likewise, the cytoplasm (yolk) of an ovum is termed **ooplasm**. It has less amount of yolk (in humans) and hence it is alecithal. This ooplasm is enclosed by a peripheral layer called the cortex which has many microvilli. These microvilli are tubular projections of the plasmalemma that aids in the transportation of substances in and out of the cytoplasm.

### The human ovum is typically covered by 3 layers:

1. Inner thin vitelline membrane
  2. Middle zona pellucida
  3. Outer corona radiata
- ✓ The narrow space present between the vitelline membrane and zona pellucida is called the perivitelline space. The zona pellucida is often seen as a thick girdle that is surrounded by the corona radiata.
  - ✓ For fertilization to occur, the hyaluronidase in the acrosome of the sperm has to scatter the outer corona radiata from the middle zona pellucida of an ovulated oocyte. This invariably allows contact between the oocyte's core and the sperm.



[https://cit.vfu.cz/frvs2011/?title=ukoly-rozmnozovani\\_a\\_vyvoj-meioza&lang=en](https://cit.vfu.cz/frvs2011/?title=ukoly-rozmnozovani_a_vyvoj-meioza&lang=en)



### Types of Egg:

According to the proportion of the yolk to the cytoplasm of the ovum there are three types of egg:

#### [I] Microlecithal egg:

In microlecithal eggs the amount of yolk is much less than the amount of



cytoplasm. These eggs are very small in size. Some embryologists described microlecithal eggs as alecithal eggs or oligolecithal eggs or miolecithal (i.e. little yolk) eggs.

- ❖ The eggs of Amphioxus and mammals are of this type.
- ❖ The mammalian eggs contain so little yolk that they are sometimes called alecithal (without yolk) eggs.

### [II] Mesolecithal eggs:

Here yolk is moderate in amount and such eggs are called mesolecithal or medialecithal (i.e. median yolk). The distribution of yolk is distinctly unequal.

- ❖ The eggs of sharks, fishes and many amphibians are of this type.

### [III] Macrolecithal or polylecithal eggs:

Enormous amount of yolk is present in macrolecithal eggs and here yolk is several times greater than cytoplasm. These eggs may be small or big.

- ❖ The eggs of teleost fishes, reptiles, birds and monotremates (egg laying mammals) are of this type.

According to distribution of yolk granules or platelets in the cytoplasm of the ova or egg, the eggs are classified as follows:

#### 1. Homolecithal/Isolecithal:

The yolk in these eggs is uniformly distributed through the cytoplasm.

- ❖ Examples are of Amphioxus, many invertebrates and mammals including man.

#### 2. Centrolecithal:

Yolk is concentrated in the interior of the egg and the cytoplasm is distributed as a thin layer on the outside of the yolk.

- ❖ As in insects and many other arthropodes.

#### 3. Teleolecithal:

Yolk becomes more abundant and tends to concentrate in one hemisphere of the egg. Because of the uneven distribution of yolk, such an egg is said to have

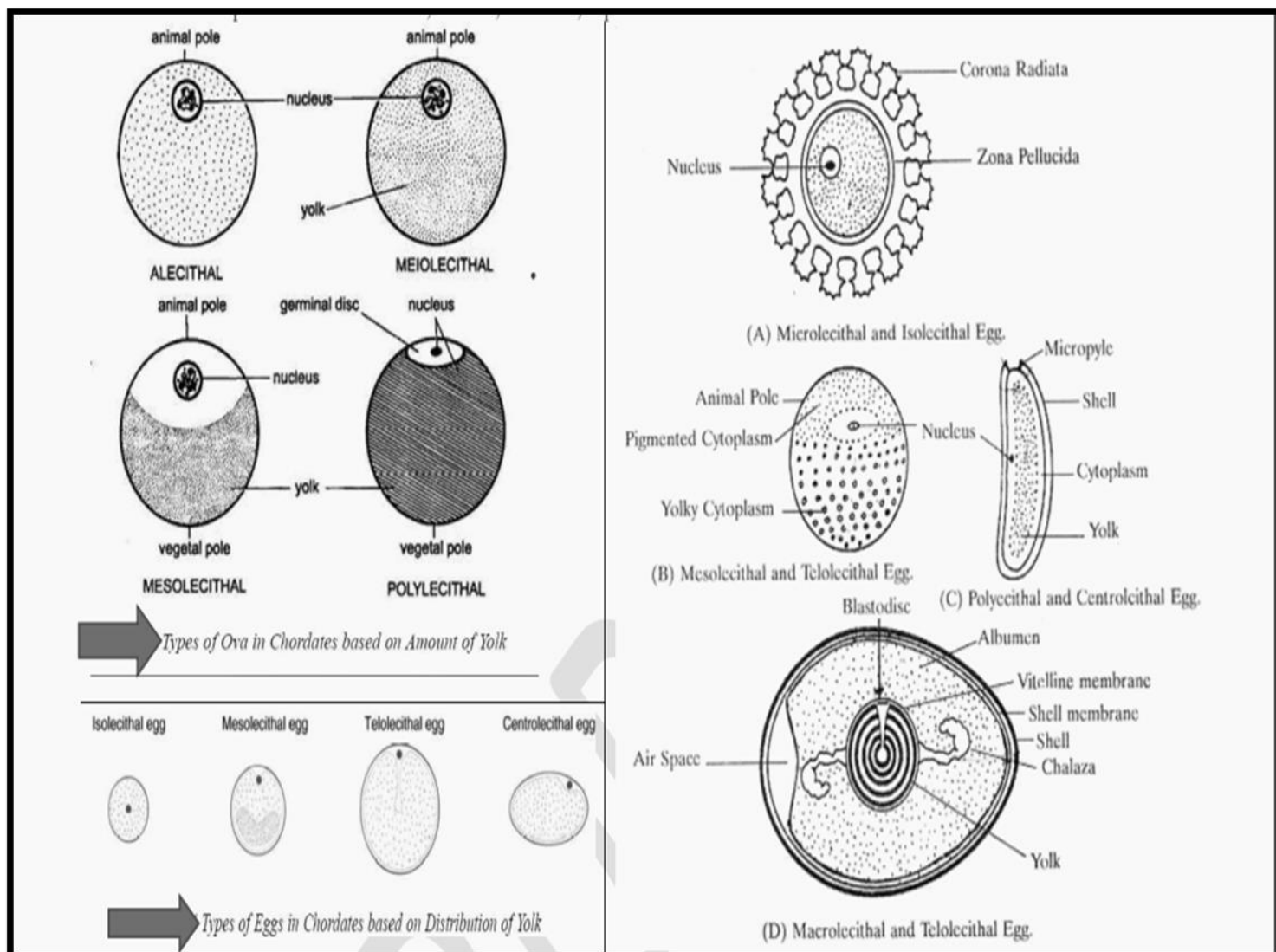
a vegetal pole, where the concentration of yolk is greatest and an animal pole, where the concentration of yolk is smallest.

In fact, in macrolecithal eggs, the amount of yolk is so massive that it occupies almost all the vegetal pole, and the active cytoplasm and germinal vesicles (nucleus) remain confined to a small cap at the animal pole.

- ❖ Examples are of fishes, amphibians, and reptiles, birds and monotremes eggs.

### **Functions of Yolk:**

Although yolk of egg is used for two purposes—supply of energy and synthesis of the products required for elaboration of the embryonic body. The yolk also has important influence on—(i) the size of the ovum, (ii) differentiation of ooplasm, (iii) patterns of cleavage, (iv) morphogenetic movements of blastomeres during gastrulation, and (v) the type of development whether direct or indirect.



<https://www.bajkulcollegeonlinestudy.in>

**Oogenesis:** consists of three phases: multiplication, growth and maturation.

### Multiplication phase:

In the foetal development, certain cells in the germinal epithelium of the ovary of the foetus are larger than others. These cells divide by mitosis, producing a couple of million egg mother cells or oogonia in each ovary of the foetus. No more oogonia are formed or added after birth. The oogonia multiply by mitotic divisions forming the primary oocytes.

**Growth phase:**

This phase of the primary oocyte is very long. It may extend over many years. The oogonium grows into a large primary oocyte. Each primary oocyte then gets surrounded by a layer of granulosa cells to form primary follicle. A large number of these follicles degenerate during the period from birth to puberty. So at puberty only 60,000- 80,000 primary follicles are left in each ovary.

**Maturation phase:**

Like a primary spermatocyte, each primary oocyte undergoes two maturation divisions, first meiotic and the second meiotic. The results of maturation divisions in oogenesis are, however, very different from those in spermatogenesis. In the first, meiotic division, the primary oocyte divides into two very unequal haploid daughter cells—a large secondary oocyte and a very small first polar body or polocyte. In the second maturation division, the first polar body may divide to form two second polar bodies. The secondary oocyte again divides into unequal daughter cells, a large ootid and a very small second polar body. The ootid grows into a functional haploid ovum. Thus, from one oogonium, one ovum and three polar bodies are formed. The ovum is the actual female gamete. The polar bodies take no part in reproduction and, hence, soon degenerate.

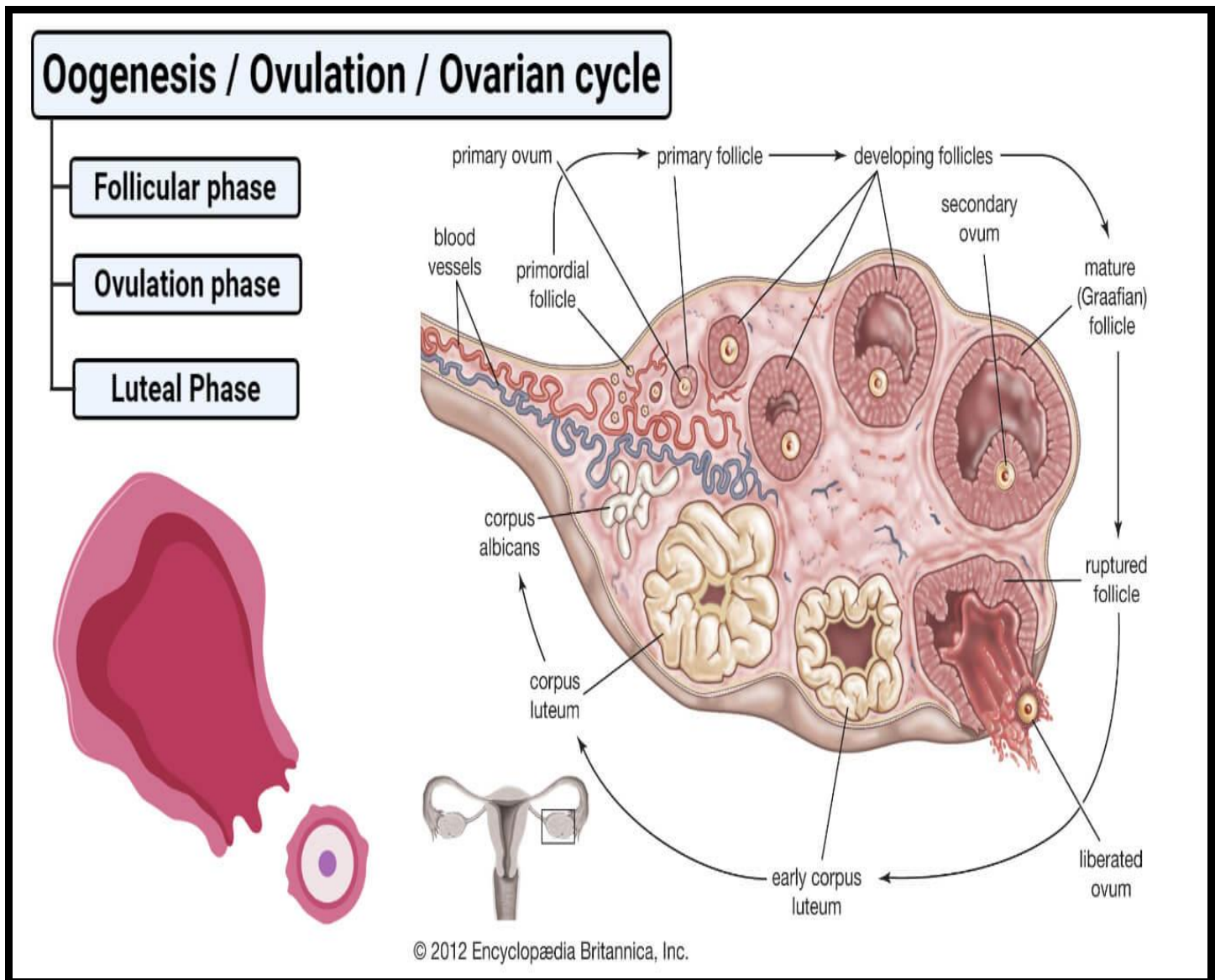


Image Source: [Britannica](#), Created with BioRender.com.



## The ovarian cycle

Of the roughly 500'000 follicles that are present in the two ovaries at the beginning of sexual maturity, only around 480 reach the graafian follicle stage and are thus able to release oocytes (ovulation). This number is simply derived by multiplying the number of cycles per year and the number of years in which a woman is fertile.

**Ovulation represents an exceptional fate of a follicle.**

- It is a series of cyclic changes occurring in the reproductive tract of female with the periodicity of 28 days
- It is also known as menstrual cycle,
- also known as endometrial cycle

- It occurs from puberty to menopause
- It is characterized by loss of vaginal blood (breaking of endometrium wall of uterus)
- The cycle is under influenced of hormones secreted by pituitary gland (FSH and LH), and ovary (progesterone and oestrogen).
- Menstrual phase
- Proliferative phase or follicular or ovulatory phase
- Luteal or secretory phase

### Menstrual phase

- This phase is characterized by discharge of blood, connective tissues and mucus due to cast off of epithelial lining of endometrium wall
- It lasts for **3-5 days**
- Ovum remain unfertilized,
- At this time level of oestrogen and progesterone is very low in blood resulting in breaking of endometrium wall of uterus
- About 50-100 ml blood with mucus are discharged as menstrual flow.

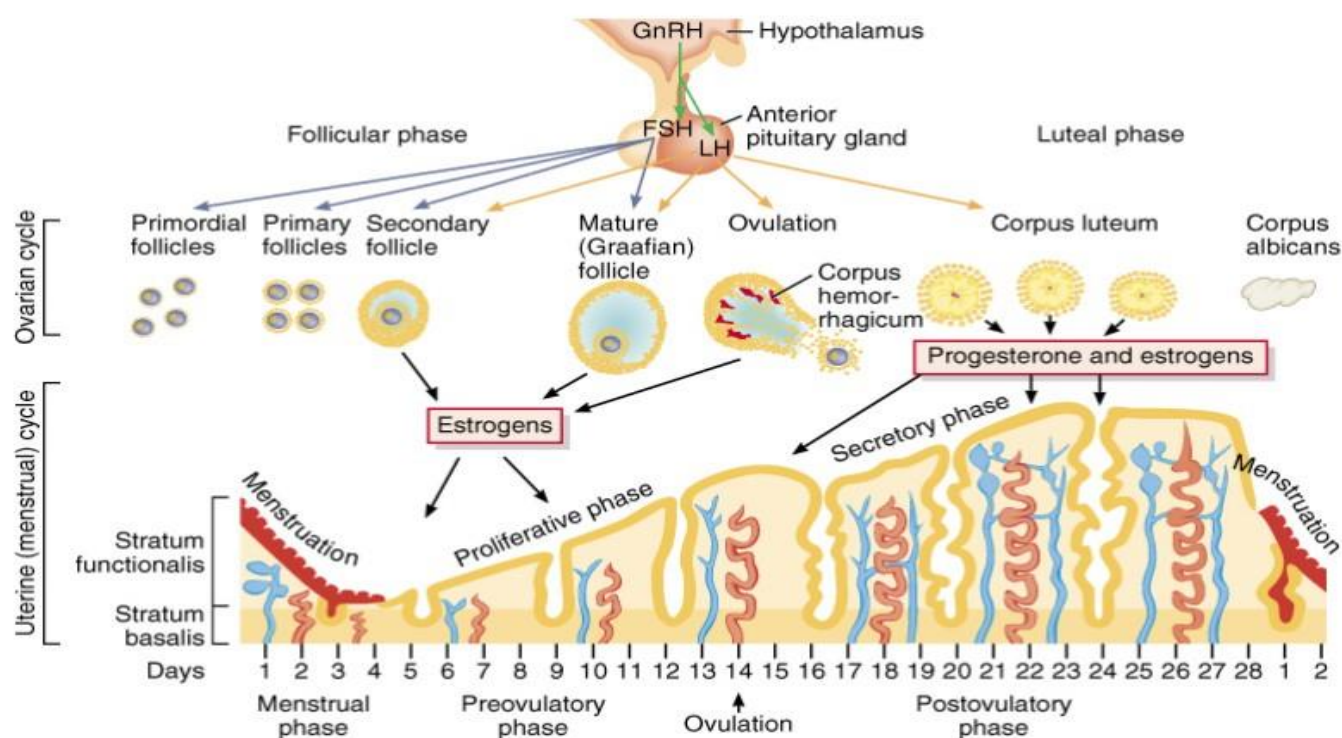
### Proliferative phase

- This phase is characterized by rapid proliferation and repair of damaged endometrium wall
- It lasts for **9-10 days** (5th – 14th days)
- Anterior pituitary gland releases **Follicular stimulating Hormone (FSH)** which stimulates development and maturation of graafian follicle. So, it is also known as Follicular Phase.
- Mature graafian follicle secrete **oestrogen**. Its level gradually increases and maximizes at 12th day
- Oestrogen stimulates endometrium repair and proliferation. It also stimulates ovulation

- Endometrium become 2-3 mm thick and highly vascular

## Luteal phase

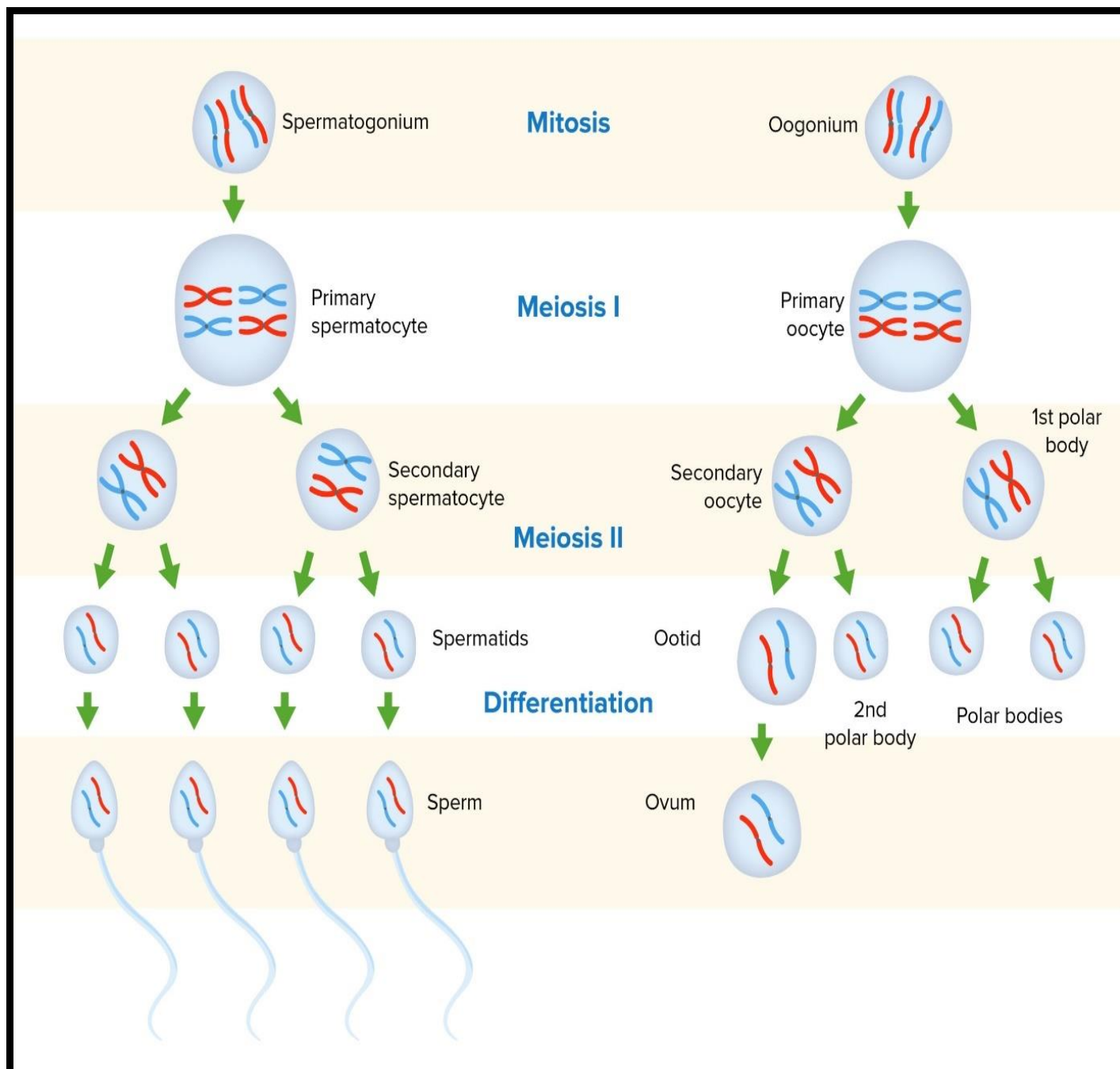
- This phase is characterized by release of Ovum from mature graffian follicle which is stimulated by the secretion of **luteinizing hormone** (LH) by pituitary gland.
- It Lasts for **12-14 days** (14th-28th day)
- LH along with FSH stimulate ovulation.
- Mature graffian follicle release ovum and the rapture follicular cell form **corpus luteum**
- Corpus luteum secrete progesterone, high level of progesterone inhibits maturation of any other follicles
- Progesterone also stimulates thickening of endometrium wall
- When ovum remain unfertilized, corpus luteum degenerate; level of both hormone (progesterone and oestrogen) decreases, causing breaking of endometrium wall continuing the menstrual phase.





## Purpose of Oogenesis

The goal of oogenesis is to produce one egg with massive amounts of cytoplasm.



<https://cdn.lecturio.com/assets/Gametogenesis-1200x956.jpg>



	Spermatogenesis	Oogenesis
<b>Process</b>		
<i>Location</i>	Occurs <i>entirely</i> in testes	Occurs <i>mostly</i> in ovaries
<i>Meiotic divisions</i>	Equal division of cells	Unequal division of cytoplasm
<i>Germ line epithelium</i>	Is involved in gamete production	Is not involved in gamete production
<b>Gametes</b>		
<i>Number produced</i>	Four	One (plus 2 – 3 polar bodies)
<i>Size of gametes</i>	Sperm smaller than spermatocytes	Ova larger than oocytes
<b>Timing</b>		
<i>Duration</i>	Uninterrupted process	In arrested stages
<i>Onset</i>	Begins at puberty	Begins in foetus (pre-natal)
<i>Release</i>	Continuous	Monthly from puberty (menstrual cycle)
<i>End</i>	Lifelong (but reduces with age)	Terminates with menopause



<http://ib.bioninja.com.au/higher-level/topic-11-animal-physiology/114-sexual-reproduction/gametogenesis.html>



## The timing of meiosis differs in females and males



### In males

- 1- The spermatogonia enter meiosis and produce sperm from puberty until death.
- 2-The process of sperm production takes only a few weeks.
- 3- Four sperms result from spermatogenesis.



### In females

- 1-This process is more complex. The first meiotic division starts before birth but fails to proceed. It is eventually completed about one month before ovulation, the second meiotic division occurs just before the actual process of fertilization occurs. Thus, in females, the completion of meiosis can be delayed for over 50 years.
- 2- All meiosis is ended in females at menopause.  
The result is the egg cell and three polar bodies.
- 3- Only 1 egg produced A polar body consists of the chromosomes and nucleus resulting from meiotic division but it receives almost no cytoplasm. All the cytoplasm is found in the secondary oocyte.  
The cytoplasm will provide energy to the developing embryo until it implants.

In the mammalian ovary, the oocytes are closely associated with somatic cells called granulosa cells which aid oocyte maturation and ovulation.



## Fertilization

Fertilization, the process by which male and female gametes nuclei fuses together to produce diploid zygote.

### Types of Fertilization:

#### 1. External:

Eggs are librated in water.

- Occurs outside the female genital system.
- Female laid a large number of eggs, them the male pour its sperms in the same region in water
- e.g. in fish and amphibian.

#### 2. Internal:

- land-dwellers
- specialized structures for housing gametes.
- embryo more protected during development.
- Occurs in animals that have a well developed reproductive system, animals may be:

a) **Oviparous:** zygote develops in a shell e.g. birds.

b) **Viviparous:** zygote develops inside uterus e.g. mammals.

The intrauterine life is about 21 days in the rat, 70 days in the in the Guinea pig while its about 280 days in human.

c) **Ovoviviparous:** - e.g. dog fish

It has 4 major steps:

1. Contact and recognition between sperm and egg. (Same species)
2. Regulation of sperm entry into the egg. (Only one and inhibiting the others)
3. Fusion of the genetic material of sperm and egg.
4. Activation of egg metabolism to start development.

### **Egg Maturation at Sperm Entry**

Most eggs are not fully mature at the time of fertilization. Sperm entry activates metabolism and relieves meiotic arrest.

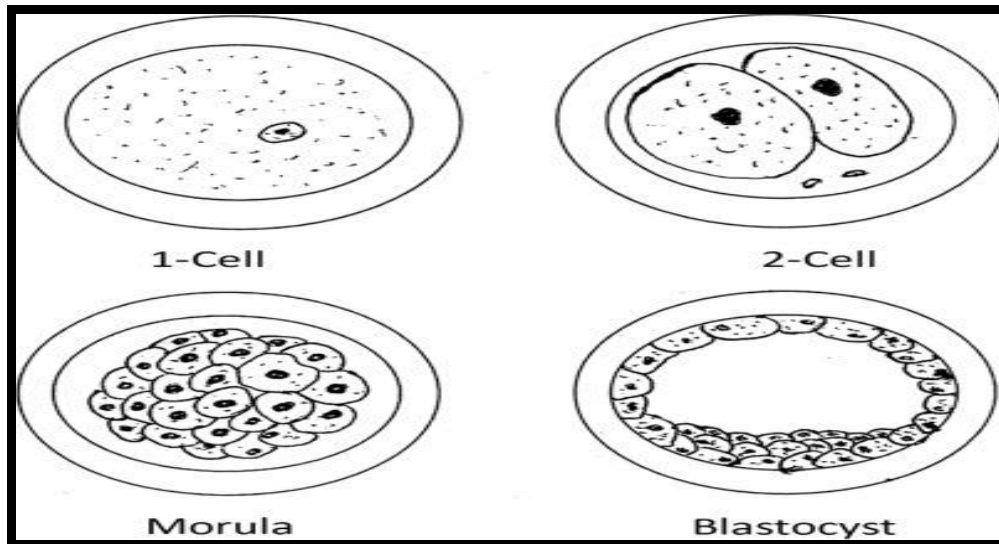
Polar body formation results in egg maturation to be ready for actual fertilization (Pronucleus formation).



### **Cleavage and Blastula Formation**

#### **Meaning of Cleavage:**

- Is the process of repeated rapid mitotic cell divisions of the zygote (unicellular structure) to form the Blastula (multicellular structure).
- The produced cells named Blastomeres.
- During this stage the size of the embryo does not change, the blastomeres become smaller with each division.
- The type & pattern of cleavage differ from species to species.
- continues divisions to form a ball of 32 cells called the morula.
- The morula continues divisions to form the hollow blastula with up to several hundred cells.
- The cavity of the blastula is the blastocoel.



<https://www.sciencedirect.com/topics/agricultural-and-biological-sciences/ova>



### Planes of Cleavage:

Considerable amount of reorganization occurs during the period of cleavage and the types of cleavage depend largely upon the cytoplasmic contents.

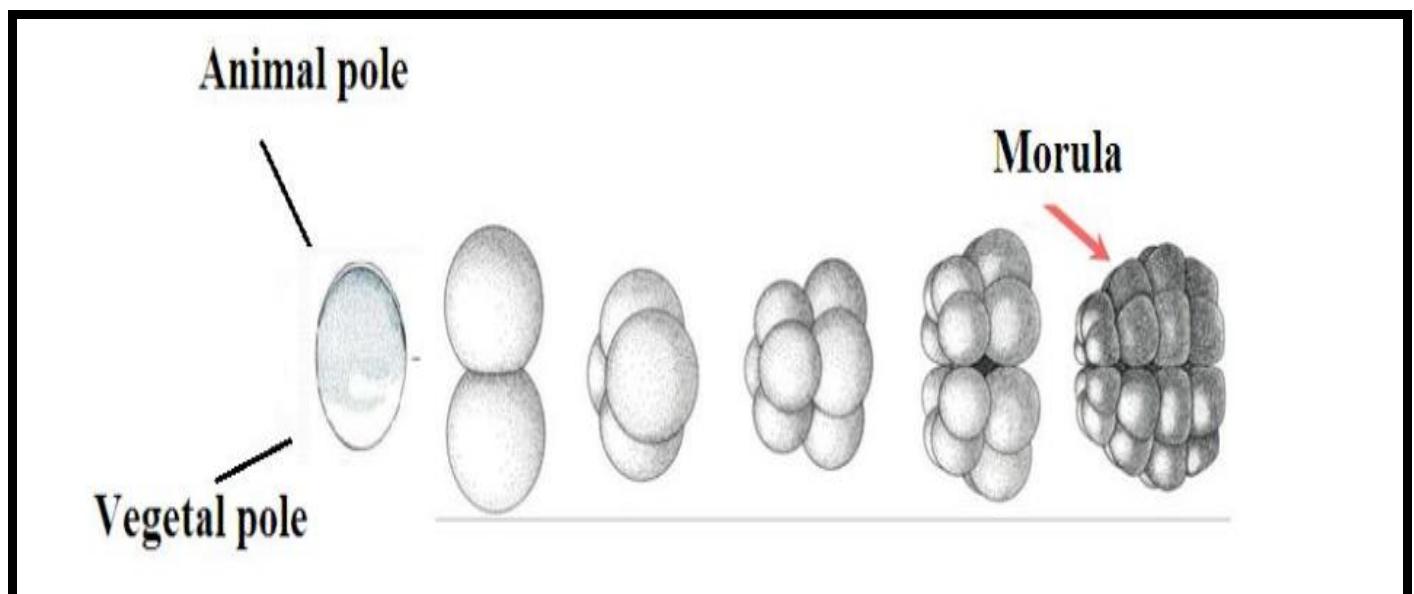
#### A: Holoblastic or total cleavage:

When the cleavage furrows divide the entire egg.

It may be:

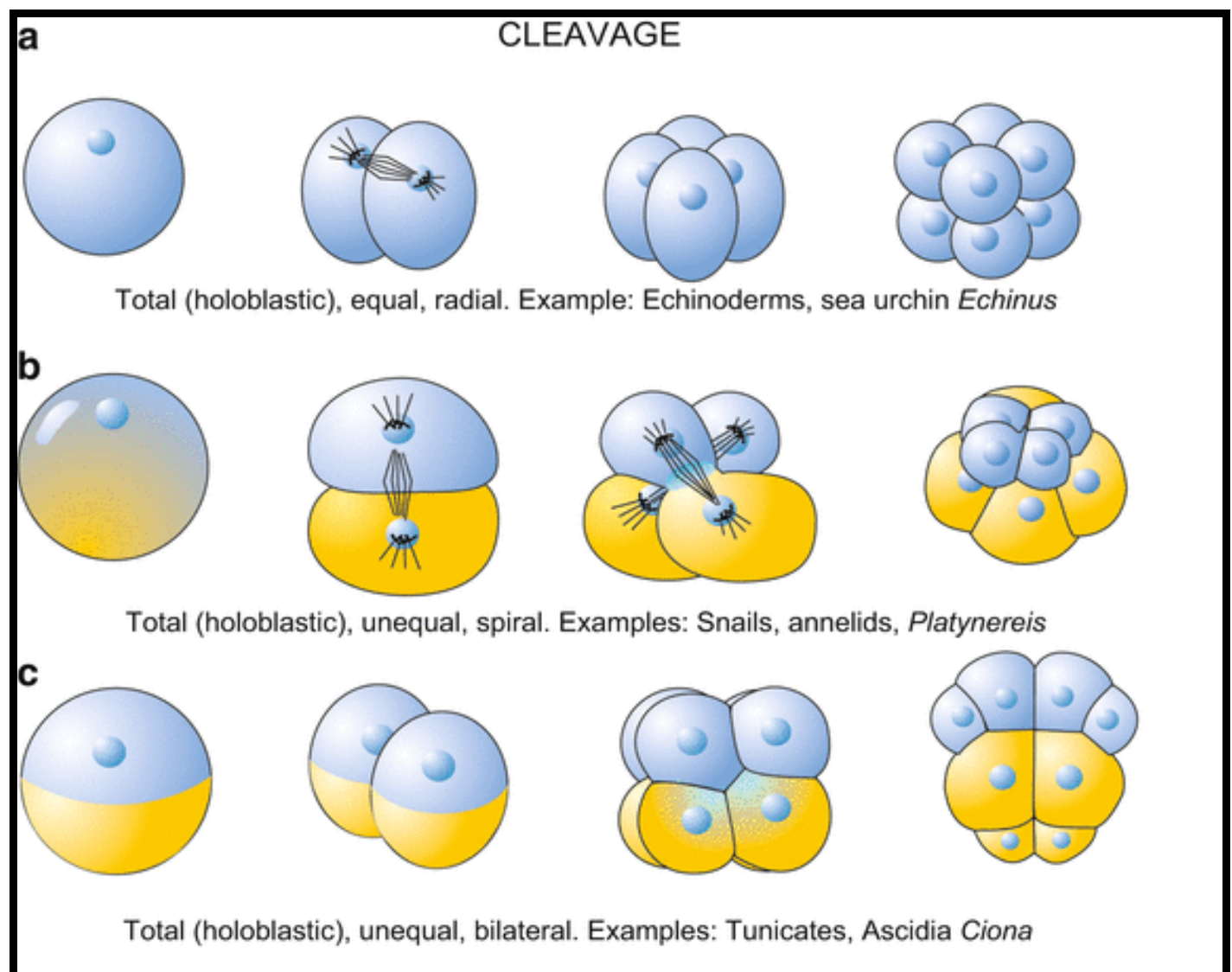
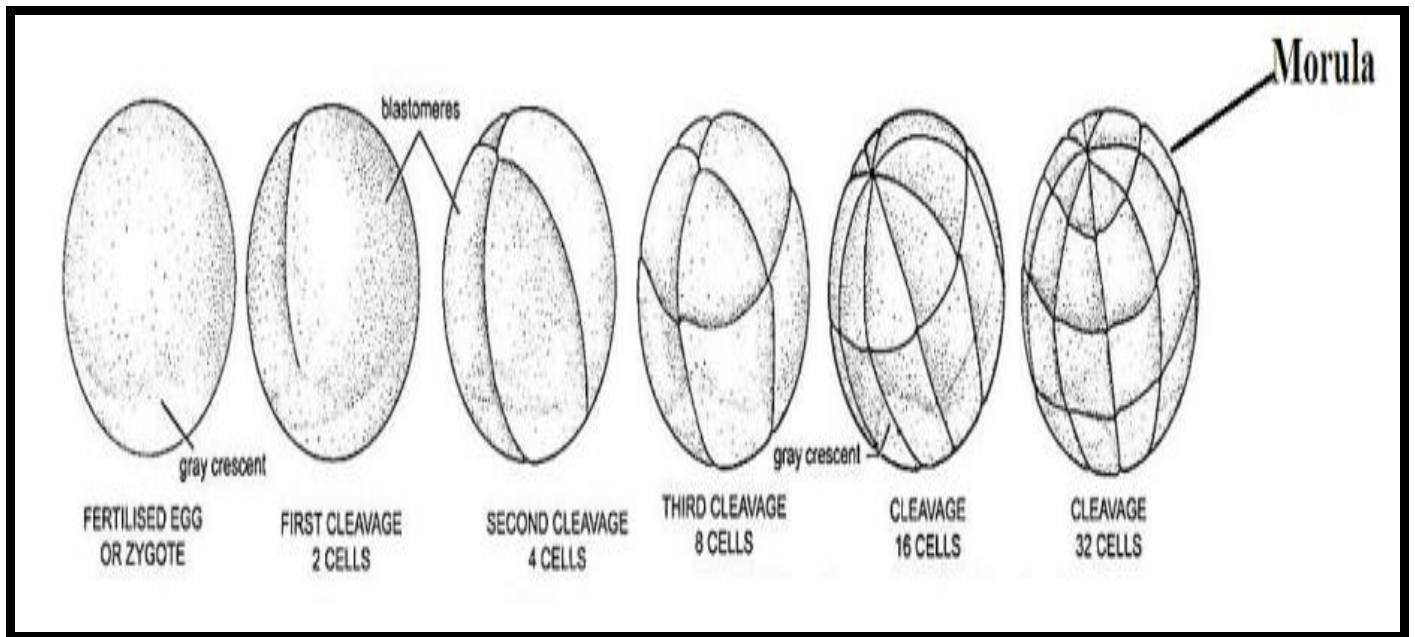
#### Equal:

When the cleavage furrow cuts the egg into two equal cells. It may be radially symmetrical, bilaterally symmetrical, spirally symmetrical or irregular.



**Unequal:**

When the resultant blastomeres become unequal in size.



## B. Meroblastic cleavage:

When segmentation takes place only in a small portion of the egg resulting in the formation of blastoderm, it is called meroblastic cleavage. Usually the blastoderm is present in the animal pole and the vegetal pole becomes laden with yolk which remains in an uncleaved state, i.e., the plane of division does not reach the periphery of blastoderm or blastodisc.

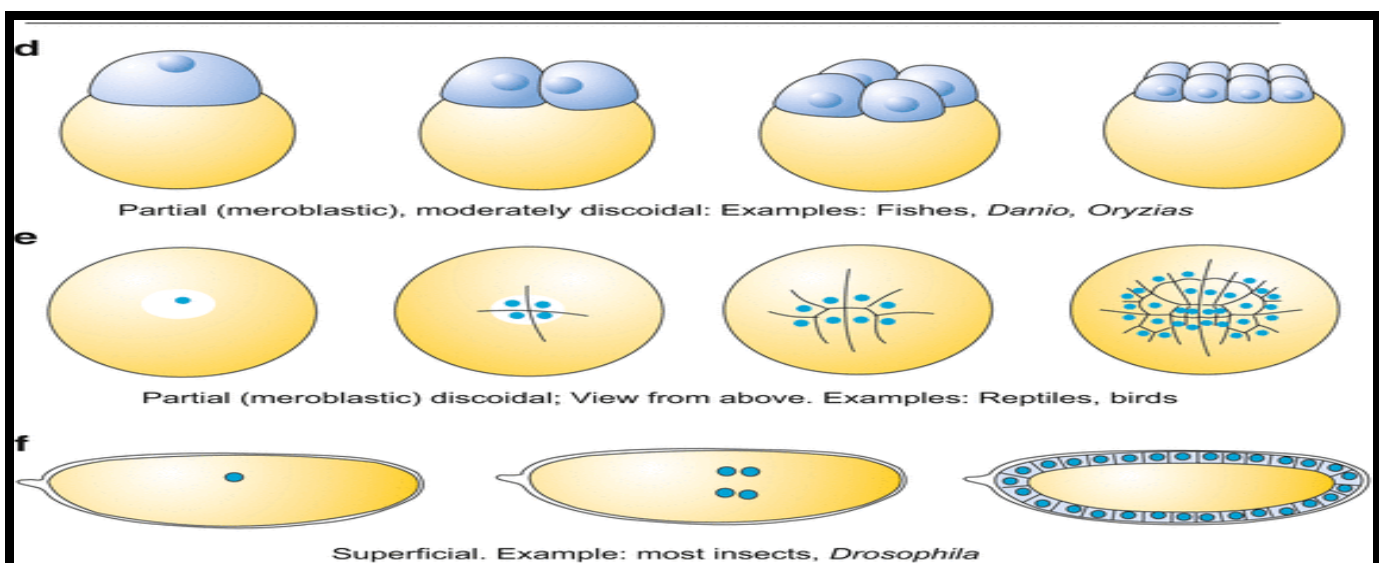
Two major types of meroblastic cleavage are discoidal and superficial:

### 1- Discoidal

In discoidal cleavage, the cleavage furrows do not penetrate the yolk. The embryo forms a disc of cells, called a blastodisc, on top of the yolk. Discoidal cleavage is commonly found in monotremes, birds, reptiles, and fish that have telolecithal egg cells (egg cells with the yolk concentrated at one end).

### 2-Superficial

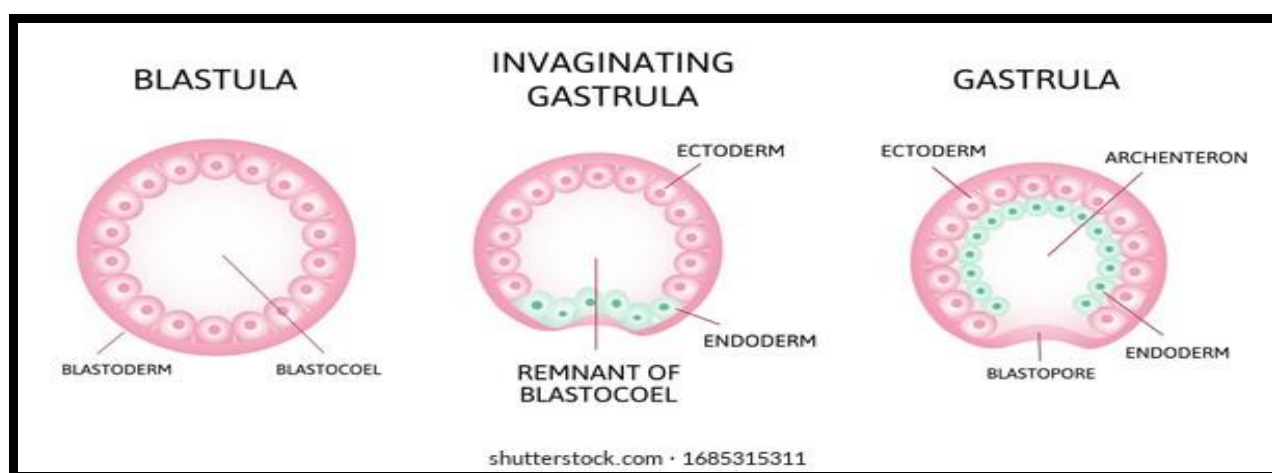
In superficial cleavage, mitosis occurs but not cytokinesis, resulting in a polynuclear cell. With the yolk positioned in the center of the egg cell, the nuclei migrate to the periphery of the egg, and the plasma membrane grows inward, partitioning the cytoplasm into individual cells. Superficial cleavage occurs in arthropods that have centrolecithal eggs.





## Gastrulation

The morphogenetic process called gastrulation rearranges the cells of a blastula into a three-layered (triploblastic) embryo, called a gastrula, that has a primitive gut. It means rearrangement of blastula cells that transforms the blastula into a gastrula. The blastula develops a hole in one end and cells start to migrate into the hole; this forms the gastrula which is characterized by cell movement. Blastocoel is gradually disappeared and a new cavity is formed archenteron.



### Germ Layer Patterns

**Diploblastic gastrula = 2 germ layers**

Endoderm (inner)

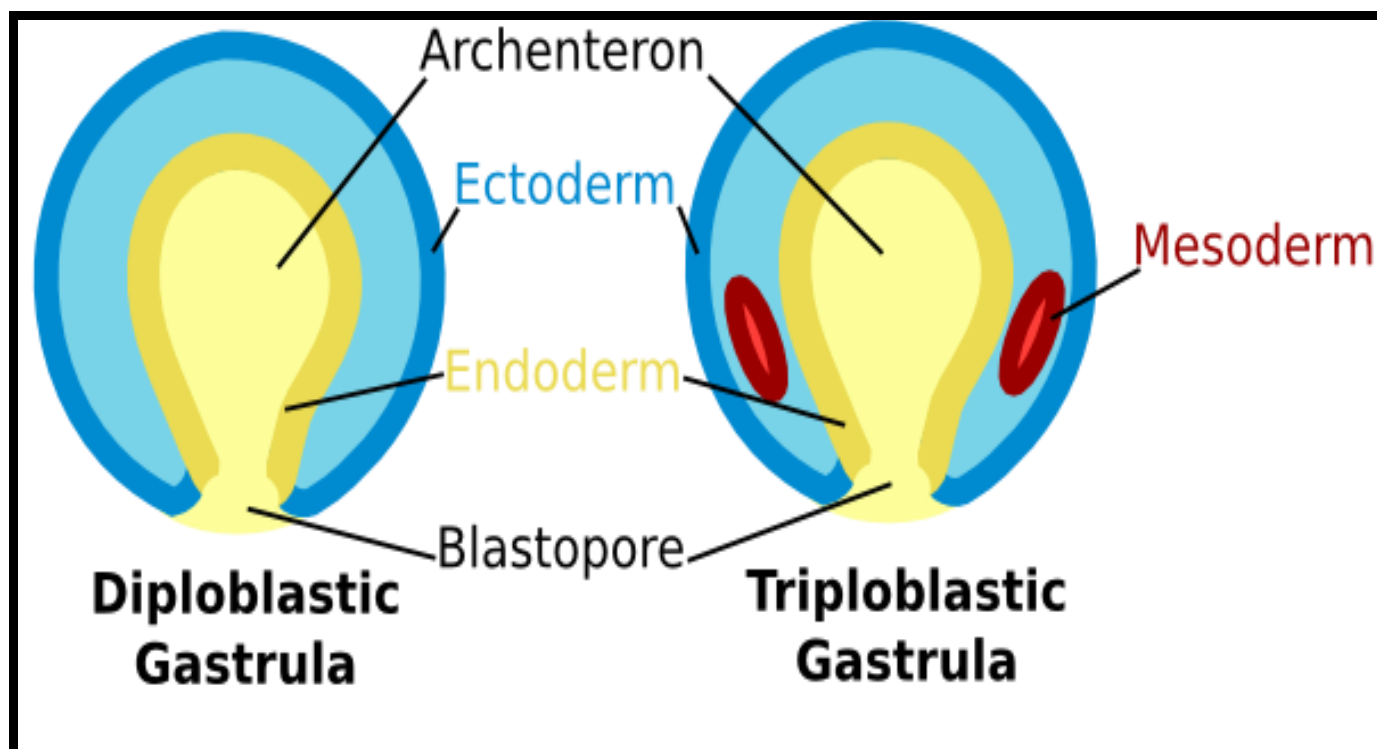
Ectoderm (outer)

**Triploblastic = 3 germ layers**

Endoderm (inner)

Mesoderm (middle)

Ectoderm (outer)



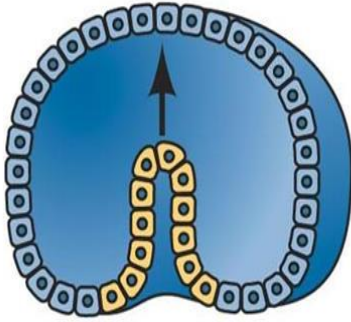
The pattern of gastrulation is affected by the amount of yolk. The cells at the vegetal pole invaginate, initiating gastrulation.

### **Gastrulation patterns**

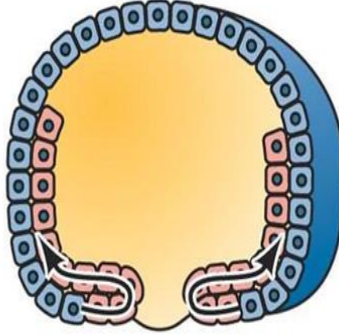
gastrulation patterns exhibit enormous variation throughout the animal kingdom, they are unified by the five basic types of cell movements that occur during gastrulation:

- Invagination.
- Involution.
- Ingression.
- Delamination.
- Epiboly

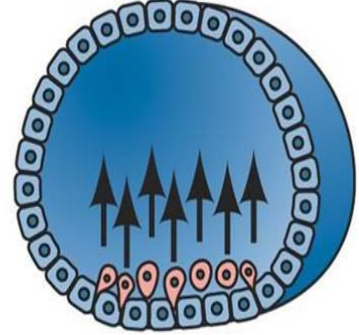


**Invagination**

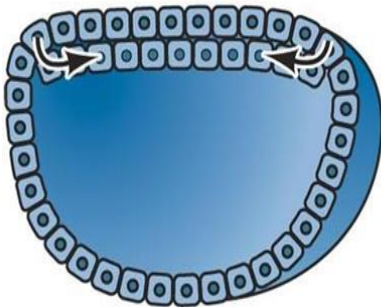
Local inward buckling  
of an epithelium

**Involution**

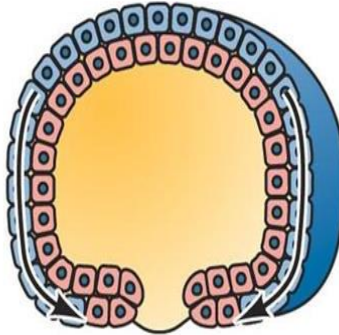
Inward movement of a cell  
layer around a point or edge

**Ingression**

Movement of individual cells  
or small groups from an  
epithelium into a cavity

**Delamination**

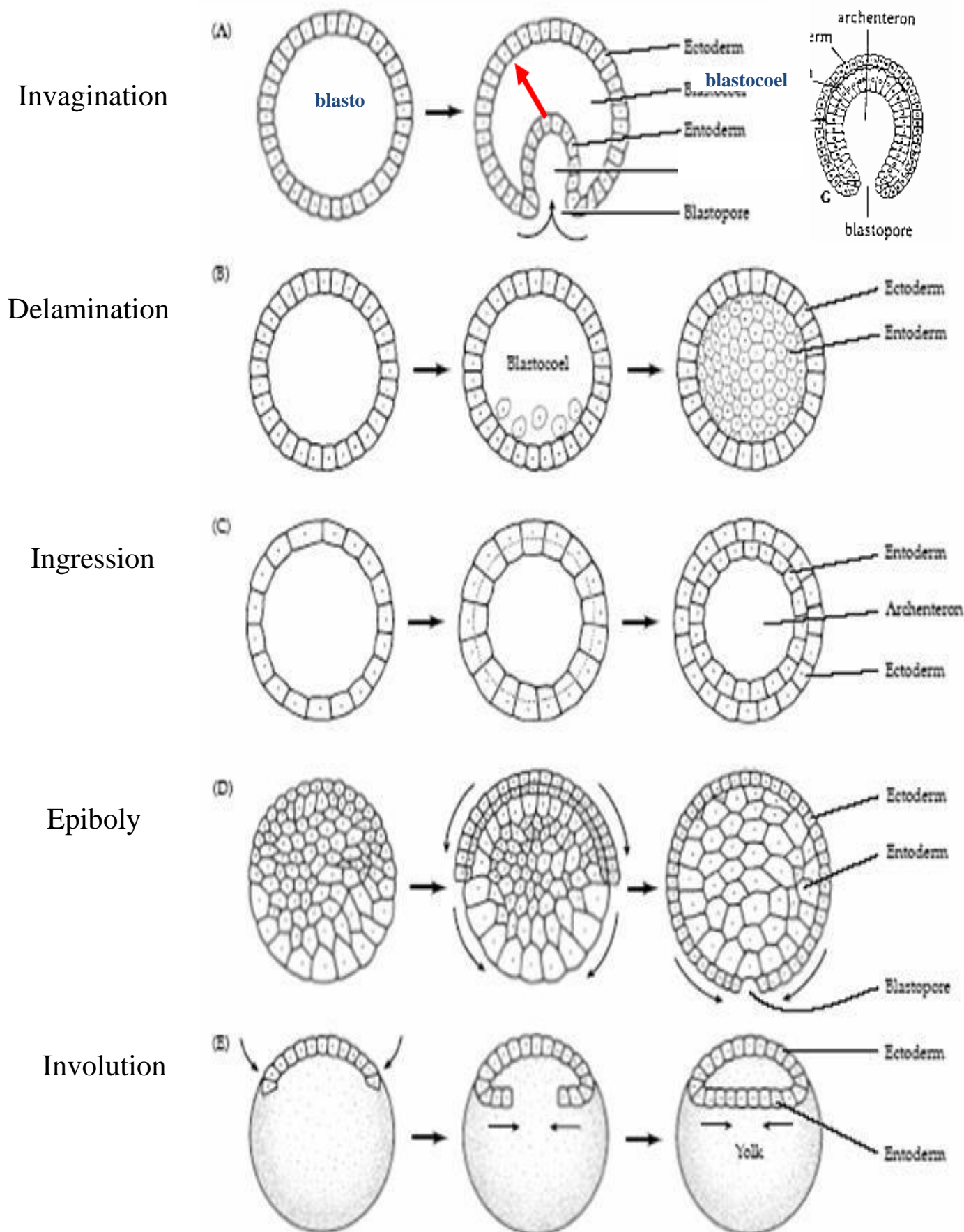
Splitting layers of cells  
(sometimes used to  
describe coordinated  
ingression)

**Epiboly**

Spread of an outside cell layer  
(as a unit) to envelop a  
yolk mass or deeper layer

**Migration**

Movement of individual cells  
over other cells or matrix





## Blastopore

Blastopore means a mouth-like opening of the archenteron on the surface of the embryo during the invagination of the archenteron. In many animals, the Blastopore becomes the anus.

The Blastopore is responsible for organizing and defining the germ layers. Through this layer, there is communication which takes place between the embryo and the outside environment in the womb. There is also a transfer of the necessary fluids until the gastrulation period lasts.

- ✓ In Coelenterate it becomes the mouth.
- ✓ In Protostomia (including Annelida, Mollusca, Arthropoda ous), it becomes subdivided into two opening, one of which becomes the mouth and other the anus.
- ✓ In Deuterostomia (including Echinodermata and Chordata), only the anus is formed.



## Organogenesis

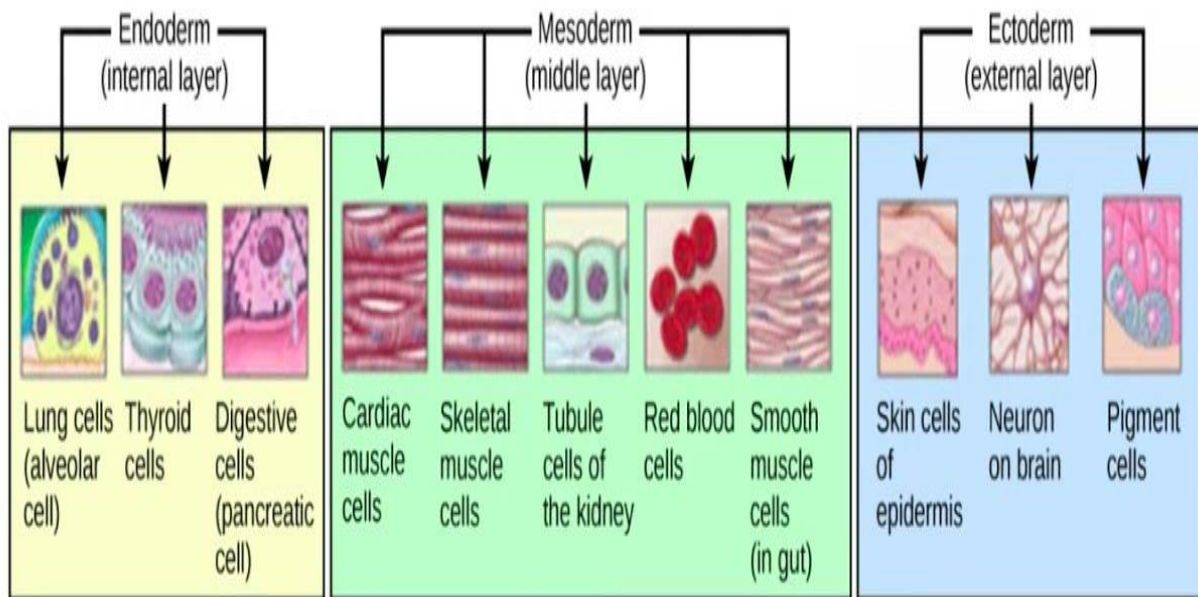
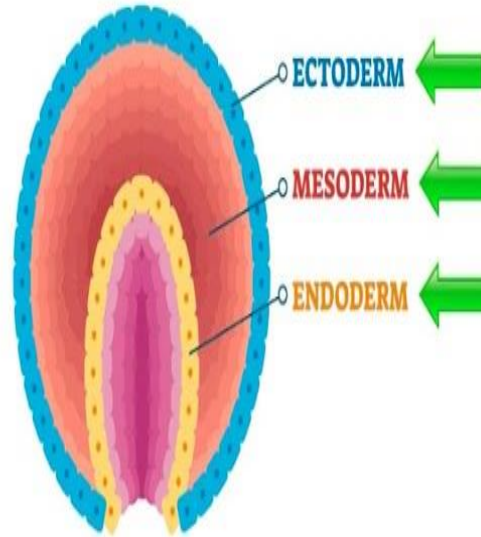
- ✓ Organ formation it is basically cell differentiation. The embryo is called “fetus”
- ✓ During the fifth phase of development, the *organogenesis* (organs formation), the continuous masses of cells of the three germinal layers split up into smaller groups of cells, called the *primary organ rudiments*, each of which is destined to produce a certain organ or part of the adult animal body.

### Germ layers

<b>Embryonic Germ Layer</b>	<b>Vertebrate Adult Structures</b>
<b>Ectoderm (outer layer)</b>	<b>Epidermis of skin; epithelial lining of oral cavity and rectum; nervous system</b>
<b>Mesoderm (middle layer)</b>	<b>Skeleton; muscular system; dermis of skin; cardiovascular system; excretory system; reproductive system—including most epithelial linings; outer layers of respiratory and digestive systems</b>
<b>Endoderm (inner layer)</b>	<b>Epithelial lining of digestive tract and respiratory tract; associated glands of these systems; epithelial lining of urinary bladder</b>

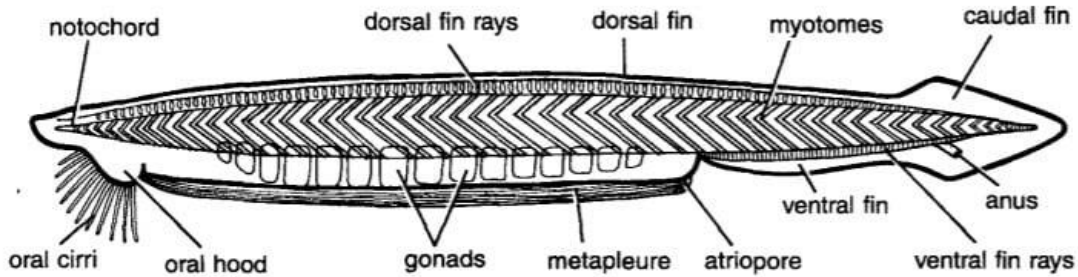
Differentiation of the Three Germ Layers

# Organogenesis





## Early embryonic development of Amphioxus

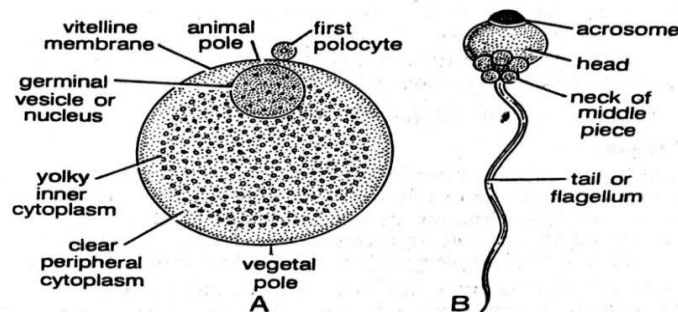


### Egg:

- 1 : The egg of Amphioxus is microlecithal and isolecithal type.
- 2 : The nucleus is almost centric because the yolk content is very less and does not affect the nucleus of the egg considerably.
- 3 : It can be differentiated into upper animal hemisphere and lower vegetal hemisphere containing animal pole and vegetable pole respectively.

### Sperm:

1. The sperm of Amphioxus is extremely minute about  $4\mu$  in length and consist of a beak or acrosome, a head with a large compact nucleus, a neck or middle piece and a very long vibratile tail.



*Amphioxus: A. Unfertilized egg. B. Sperm*

## **FERTILIZATION:**

Only one sperm can fuse with the egg. It is not yet known whether the entire sperm enters the egg or only the head enters. After the entry of sperm, the membrane becomes fibrous and is called fertilization membrane. A fluid filled space then appears between the fertilization membrane and the cell membrane. The fertilization membrane prevents the entry of more sperm. The chromosome of the egg and sperm come very close, develop a nuclear membrane around them and form a single nucleus- zygote nucleus. The egg is now called the zygote.

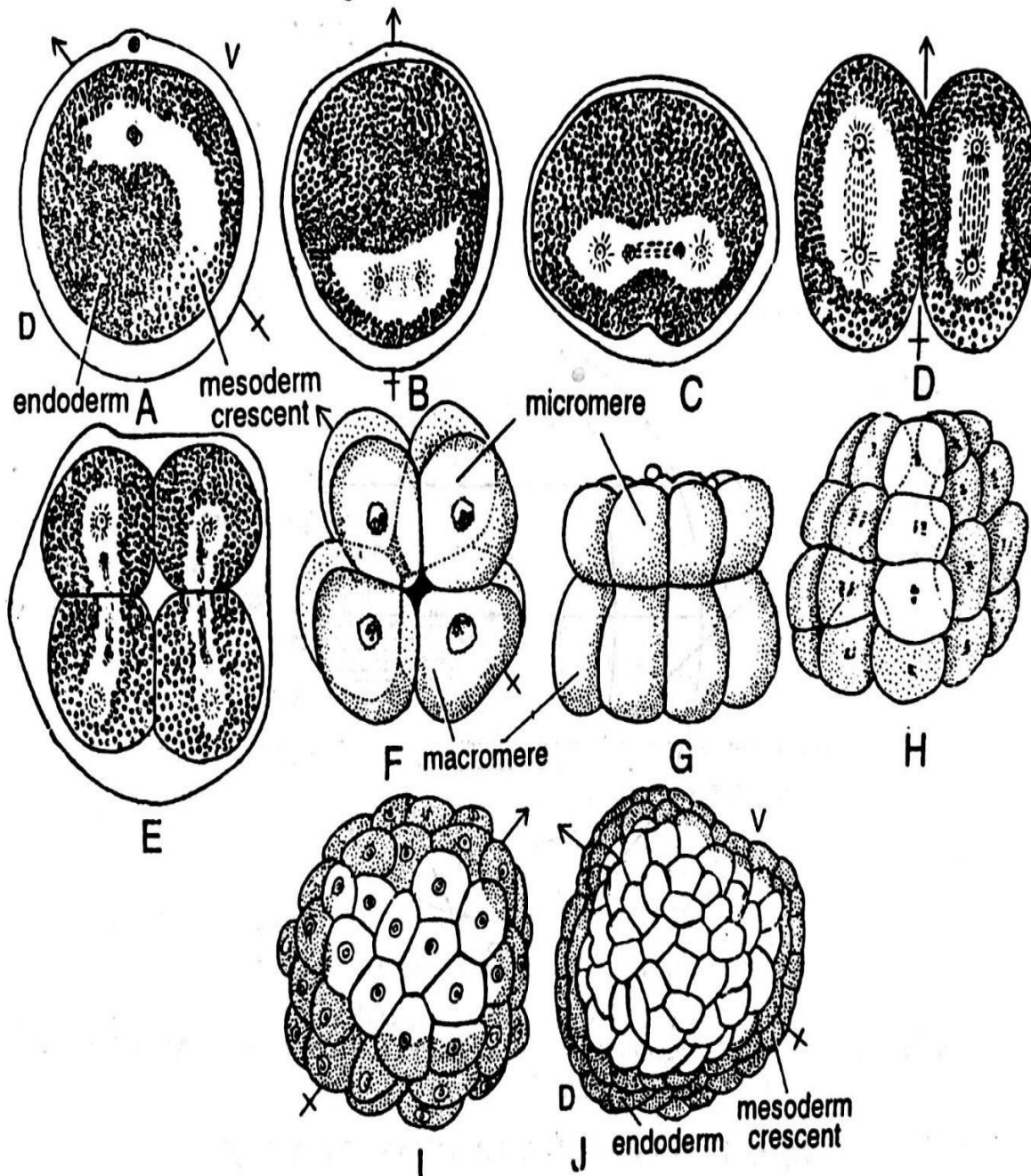
## **CLEAVAGE:**

It is complete i.e., holoblastic which divides the egg completely into blastomeres.

1. First cleavage plane is meridional that is passing through the animal pole to vegetal pole axis forming two equal blastomeres.
2. Second plane of cleavage is also meridional but at right angle to the first one forming four equals sized blastomeres.
3. Third plane of cleavage is latitudinal which is slightly above the equatorial plane, The product is the 8-cell stage of which four upper are smaller cells called micro mere and four lower larger are called megameres.
4. Fourth set of cleavage is meridional forming 16 cell stage.
5. Fifth set of cleavage is latitudinal forming 32 cells in four tiers.
6. Sixth set of cleavage is meridional forming 64 cell stage.

7. The cleavage till now is synchronous i.e., all cells at a particular cleavage divide at a time.
8. The cleavage plane on seventh cleavage onwards is asynchronous i.e., all cells at particular cleavage do not divide at a time.
9. As the division advances, the embryo is converted into a solid ball of cell called as morula.
10. Soon a small cavity appears in the interior of the embryo. it became fluid filled and expand gradually pushing the cells on periphery and as a result a hollow ball of cells is formed having a spacious fluid filled cavity called blastocoel surrounded by a single layer of cells. This is called Blastula.

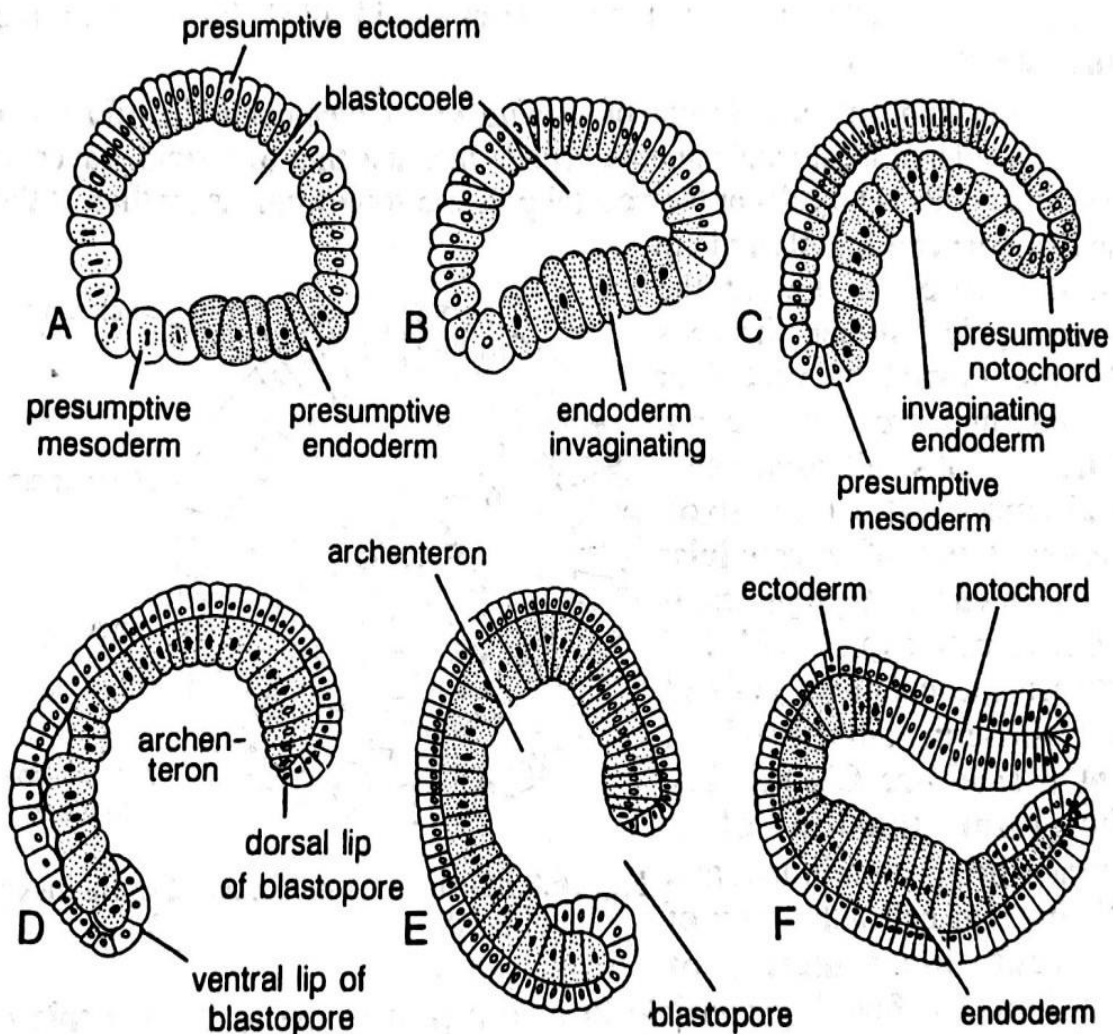




Cleavage and blastulation in Amphioxus- A-Fertilized egg, B.Mitosis of 1st cleavage .C- nuclear division. D-Two cell stage.E-Four cell stage.F-Eight cell stage. G-sixteen cell stage. H-Thirty cell stage. I-Morula stage J-Blastula stage

## GASTRULATION:

Gastrulation is a process by which the monoblastic blastula is converted into a structure containing well-defined three germinal layer from which different organs can be formed.



*Gastrulation of Amphioxus: A series of consecutive stages.*

---

## Summary

- ✓ Phylum: Chordata – Class: Cephalochordata
- ✓ Sexes are separate
- ✓ The gonads which are in the form of hollow sacs enclosed in coelomic pouches- twenty six in number on each side
  - genital ducts are lacking
- ✓ On maturity of gonads the sperms and ova are liberated into the atrium and from where they are discharged outside through the atriopore in breeding season
- ✓ The spermatozoa contain spherical head, very short mid-piece and tail
- ✓ The ovum of is 0.10 mm to 0.12 mm in diameter
- ✓ Type of Egg: According to amount of yolk oligolecithal or microlecithal

According to distribution of yolk isolecithal

- ✓ Fertilization: External
- ✓ **Type of cleavage:** holoblastic cleavage

**The first** is holoblastic and meridional passes through the egg axis from pole to pole. Result in two identical blastomeres.

**The second** in a vertical plane, at right angles to the first plane, forming four cells.

**The third** is horizontal (latitudinal) nearer the animal pole. Each of the four blastomeres dividing into a smaller micromere at the animal pole and a larger macromere at the vegetal pole. Eight blastomeres are produced.

**The Fourth** cleavage is double plane each one oriented from animal to vegetal pole. Results in eight animal micromeres and eight vegetal macromeres.

**The Fifth** is latitudinal and double parallel to plane three – one occurring in the animal, the other in the vegetal hemisphere. They result in 32-cells, arranged in four tiers.

**The sixth cleavage** are approximately meridional, producing 64-blastomeres

The cleavages pattern beyond this is irregular and difficult to follow.

The blastomeres remain loosely packed and form the embryonic stage, called morula.

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## Early embryonic development of Frog



*Xenopus laevis*

There are three groups of present-day amphibians:

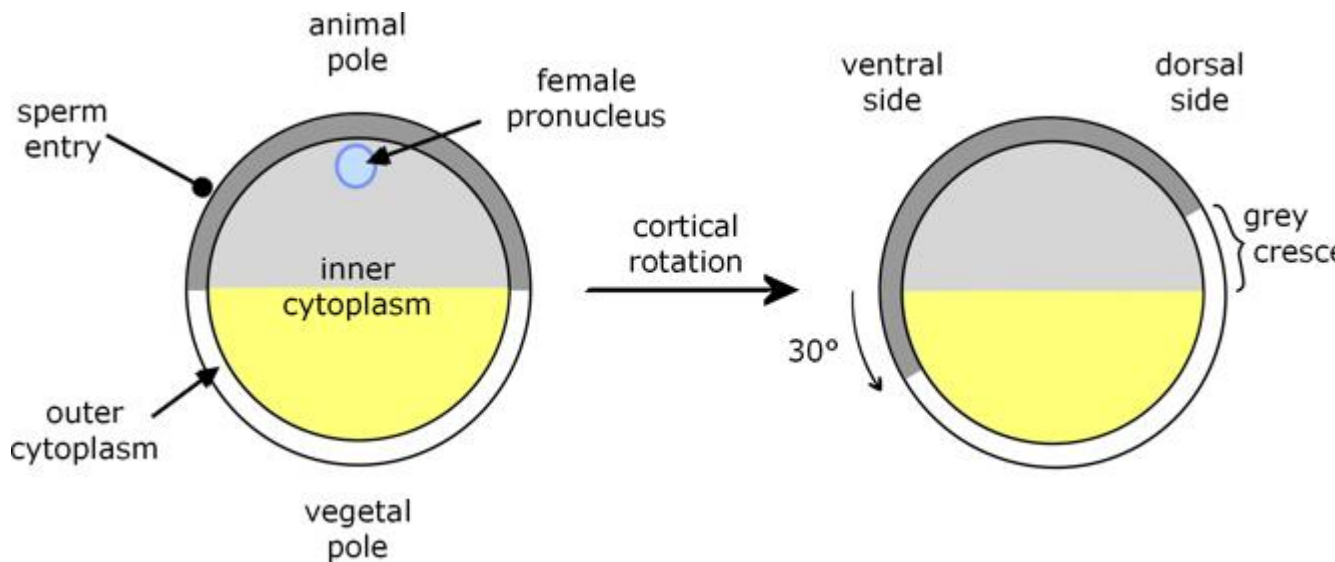
- urodeles (having a tail) e.g. salamanders;
- anurans (without a tail) e.g. frogs and toads; and
- caecilians (without limbs) so resemble worms or snakes.

Most embryological studies have been on frogs because their development can be observed easily, especially the African clawed frog (*Xenopus laevis*) because it is easy to stimulate egg laying, and the following description is based mainly on this species.

### **Egg and fertilisation**

Amphibian eggs are generally large, with a substantial quantity of yolk. In *Xenopus* the cytoplasm is in two distinct parts, inner and outer (or cortical). The inner cytoplasm of the animal hemisphere

is darker than that of the vegetal hemisphere; and, corresponding with this, the animal cortical cytoplasm is pigmented, whereas the vegetal cortical cytoplasm is not. In the haploid egg meiosis is halted at metaphase II, and on fertilisation meiosis completes to yield the diploid zygote.

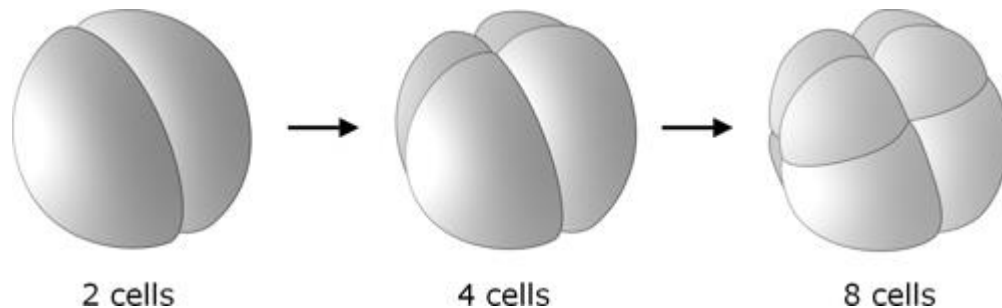


*Xenopus* fertilisation

Sperm entry and completion of meiosis prompts the cortical cytoplasm to rotate by about  $30^\circ$  relative to the deep cytoplasm, such that part of the underlying animal hemisphere becomes visible as a 'grey crescent'. Due to this rotation, sperm entry (which always occurs in the animal hemisphere) specifies the dorsal-ventral (back-front) axis of the embryo, because gastrulation begins opposite sperm entry, on the same side as the grey crescent.

## Cleavage and Blastula

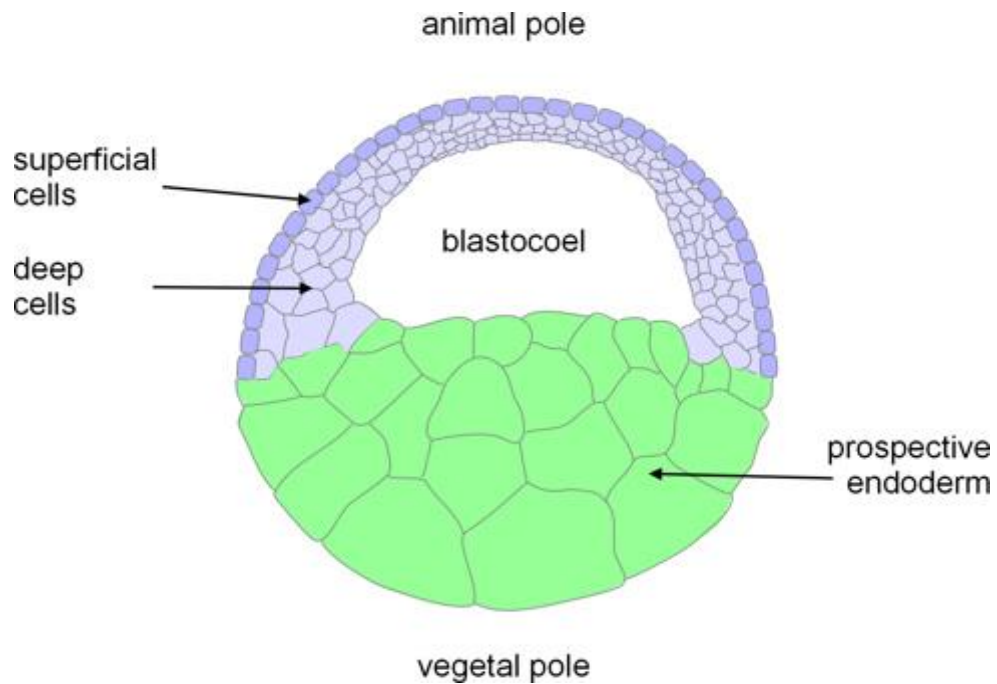
Cleavage is holoblastic: the first divisions extend right through the zygote. The first cell division goes through the poles of the zygote (meridional), as does the second but at right angles to the first, to produce four approximately symmetrical cells. The third division is at right angles to the first two (latitudinal), and approximately equatorial except that it is somewhat towards the animal pole. This occurs because, although cleavage is holoblastic, division is slower through the yolky vegetal hemisphere; and this pattern continues with subsequent divisions being more frequent in the animal hemisphere, which leads to cells there being smaller than in the vegetal hemisphere.



*Xenopus* cleavage: the first 3 cell divisions.

A blastocoel begins to develop from the first cell division and is evident from the 8-cell stage. Eventually it occupies a large part of the animal half of the blastula. Its dome-like roof is formed by numerous small, pigmented micromeres, whilst the vegetal half is composed of large yolk-laden, pale macromeres. Also, the cells of the animal hemisphere and upper part of the vegetal

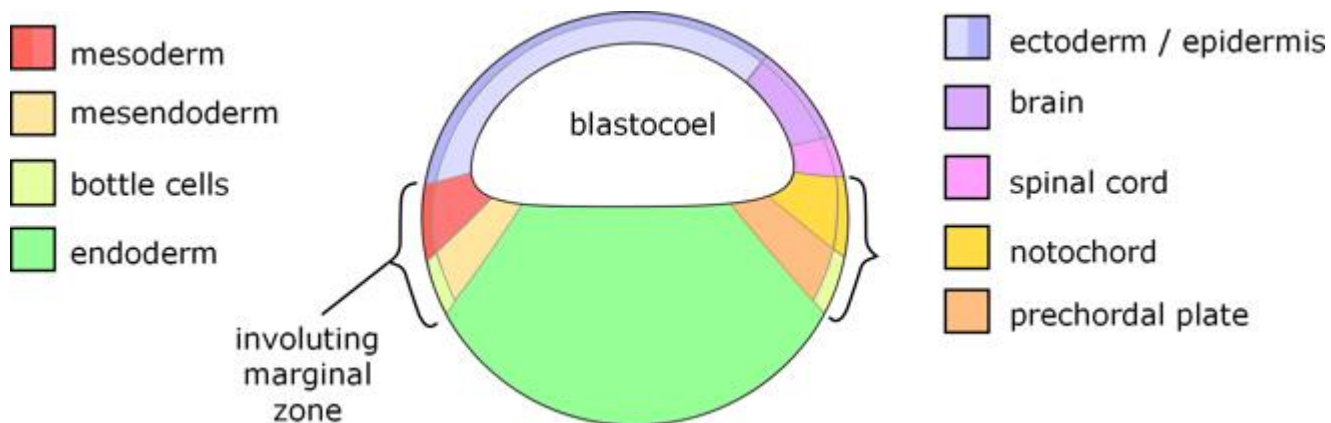
hemisphere form an outer epithelial layer and an inner or deep layer whose cells are more mesenchymal in nature.



Section through *Xenopus* blastula.

## Gastrulation

Before describing the cell movements that occur during gastrulation it's probably helpful to summarise the fates of different parts of the blastula.





Section through *Xenopus* blastula section, showing prospective fates following gastrulation.

The cells of the animal hemisphere spread (epiboly) to cover the whole of the embryo, to form the ectoderm. Cells along the dorsal centreline form the neural plate, which produces the neural tube and then reforms the epidermis above the neural tube.

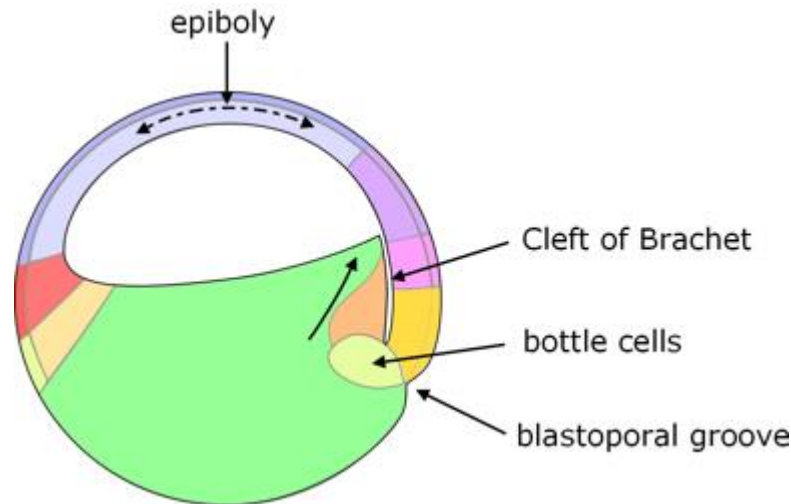
- A band of cells (the involuting marginal zone, IMZ) around the equator of the blastula moves inwards and then upwards. Mainly, these form mesoderm, the most dorsal part of which is the notochord.
- Most of the cells of the vegetal hemisphere are enveloped by the expanding ectoderm, are incorporated within the body of the embryo, and develop into endoderm.

The first external sign of gastrulation is when cells just below the equator, called bottle cells, on the dorsal side invaginate to form a crescent-shaped groove, which is the beginning of the blastopore.

However, before this, some deep cells (presumptive prechordal plate) of the dorsal side of the vegetal hemisphere begin to move toward the animal pole. They move close to the inside of the animal dome, but separated from it by a small gap known as the Cleft of Brachet.

Bottle cells. Bottle cells occur on the outer surface of the embryo where invagination occurs. Their outer surface

constricts, and this constriction creates a local depression which develops into the invagination through which cells involute. They are called 'bottle cells' because the constriction makes them somewhat bottle-shaped.

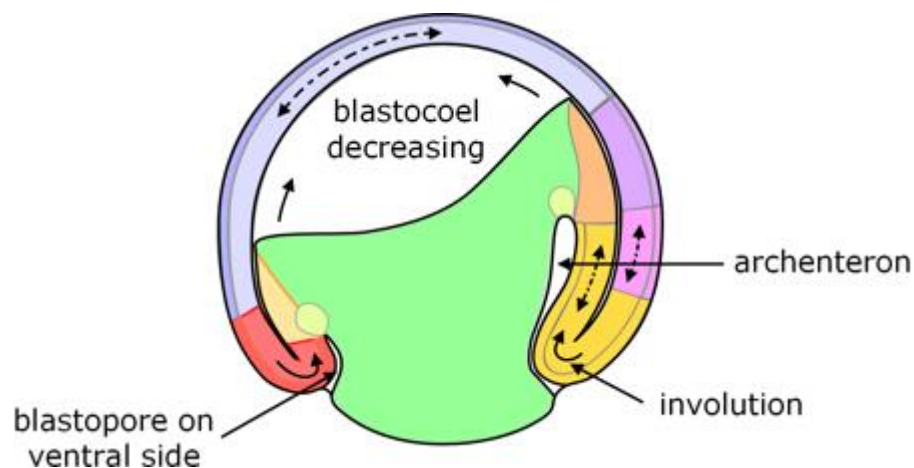


Start of gastrulation

At the same time, the animal hemisphere begins epiboly (partly by flattening of its cells, and partly by intercalation of cells from different layers) and, as it does so, its cells along the upper edge (dorsal lip) of the blastopore start to roll over the lip (involute) into the blastopore. These cells, which are predominantly presumptive notochord, move towards the animal pole, close to the overlying outer layer of cells, behind the presumptive prechordal plate.

The dorsal part of the blastopore deepens to form a pouch known as the archenteron which gets progressively larger, gradually displacing the blastocoel which in due course almost disappears.

The blastoporal groove progressively extends laterally on both sides until the two ends of the groove meet on the ventral side of the blastula, and the resulting blastoporal groove completely encircles the vegetal pole of the blastula. At the ventral side of the blastopore, cells of the expanding animal hemisphere involute and move towards the animal pole. What were part of the outer layer of cells becomes endoderm, and the deep cells become predominantly mesoderm.



Mid-gastrulation: the archenteron has formed, the blastopore encircles the vegetal pole, and involution is taking place all around the lip of the blastopore.

It will be apparent from the foregoing that involution begins at the dorsal side before the ventral side. Consistent with this, the involuted cells on the dorsal side move further than those on the ventral side, and there is an overall rotation of the inner cells with respect to the outer cell layers.

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## Summary

✓ Phylum: Chordata – Class: Amphibia

✓ Fertilization: External.

✓ Type of Egg: According to amount of yolk mesolecithal.  
According to distribution of yolk telolecithal.

Spawning:

The mesolecithal eggs of frog enclosed in a protective gelatinous albumen are laid in water. The cluster or masses of eggs which remain stick together is called spawn. A spawn of *Rana tigrina* may have 3000 to 4000 ova. The spawn is laid during pseudocopulation or amplexus.

✓ Type of cleavage: Unequal holoblastic cleavage.

✓ Adult toads live on land most of the time and rely on water for hydration, breeding, and temperature regulation, Mating.

✓ Cleavage and blastulation the period of cleavage and blastula formation completed within 24 hours.

✓ The cleavage furrow elongates at a rate of about 1mm/minute in the animal hemisphere but slows to 0.03mm/minute in the vegetal pole.

**The first** is meridional. It cuts the egg through its median animals-vegetal polar axis and result in two equals sized blastomeres.

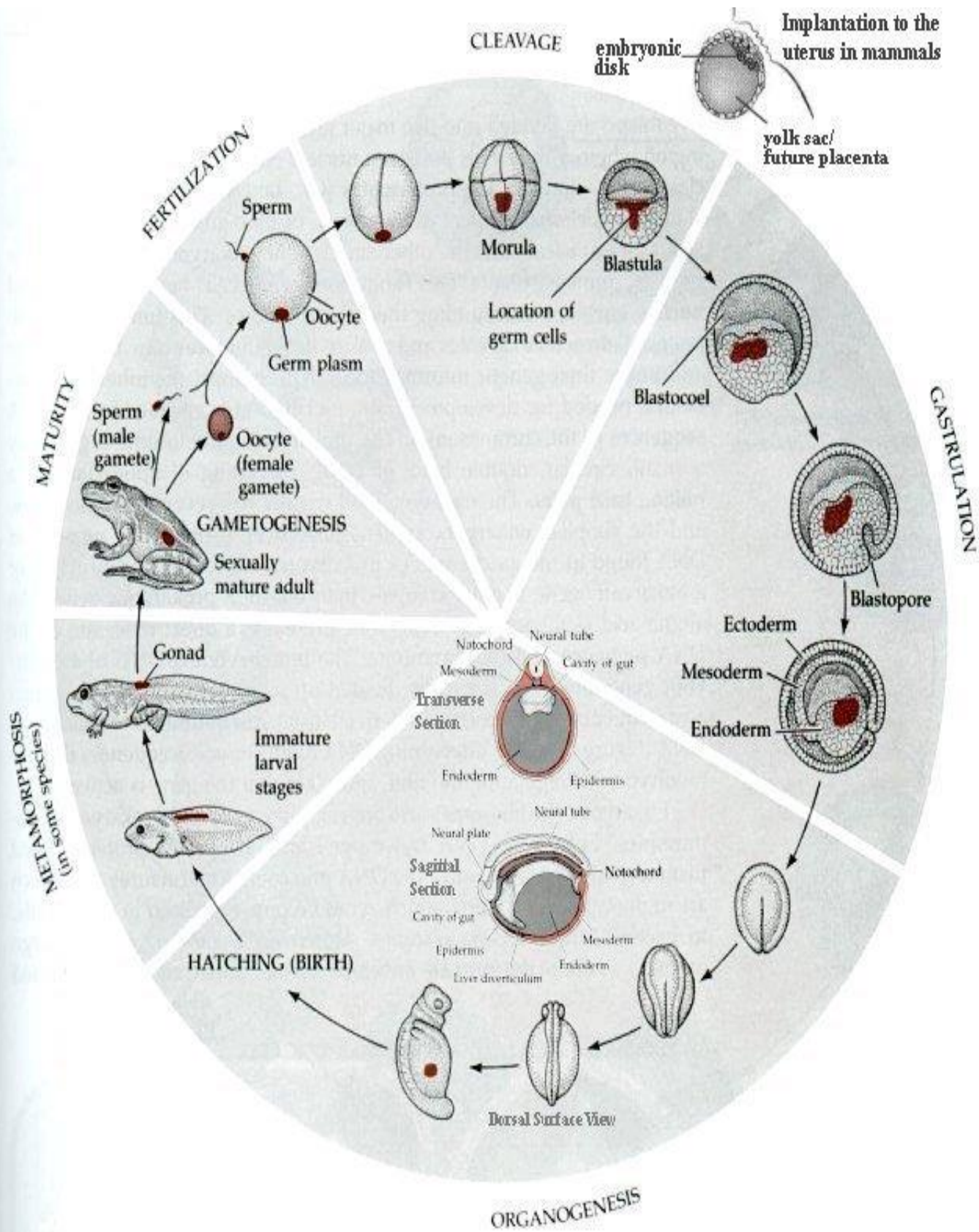
**The second** at the right angles to the first plane.

**The third** orient parallel to the polar axis and displaced near the animal pole. The eight blastomeres stage consists of four large-sized, yolk rich, vegetal cells (macromeres) and four small-sized, yolk poor, animal cells (micromeres).

**The fourth** cleavage is double plane each one oriented from animal to vegetal pole result in 16-cell stage.

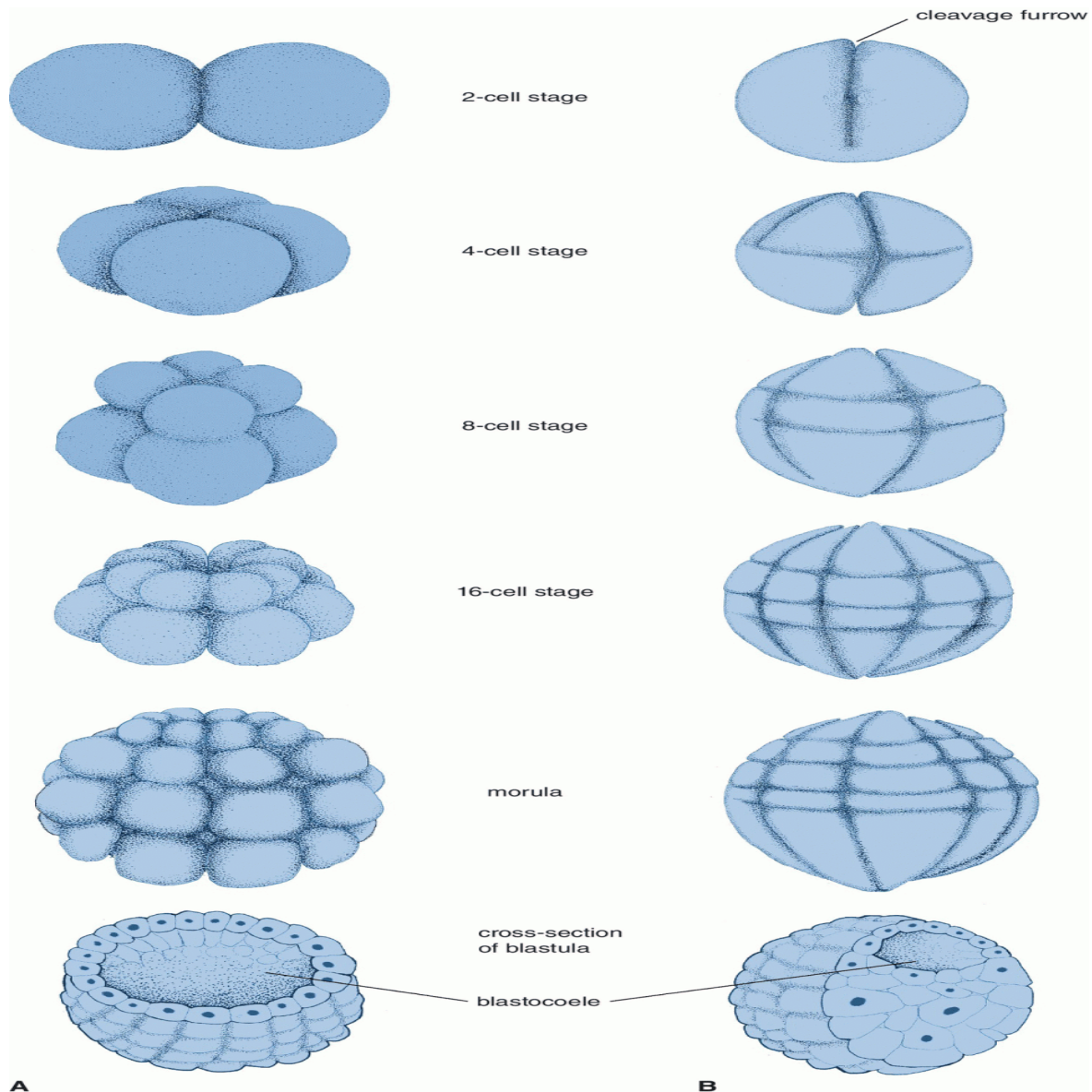
In amphibians, an embryo between the 16-and 64-cell stages is commonly called a morula. A cavity (blastocoel) appears in the animal's hemisphere above the mass of yolk. The blastula Blastula is a hollow spherical embryonic stage. The blastoderm remains two-cell thick towards the animal pole of the egg, the sides and floor of the blastocoel are multilayered blastoderm of large yolky blastomeres. -The blastoderm encloses ecentric (peripheral) blastocoel.

-The blastocoel becomes infiltrated by water and albuminous fluid secreted by the surrounding blastomeres.



## Quick comparative review between Amphioxus and Frog early development

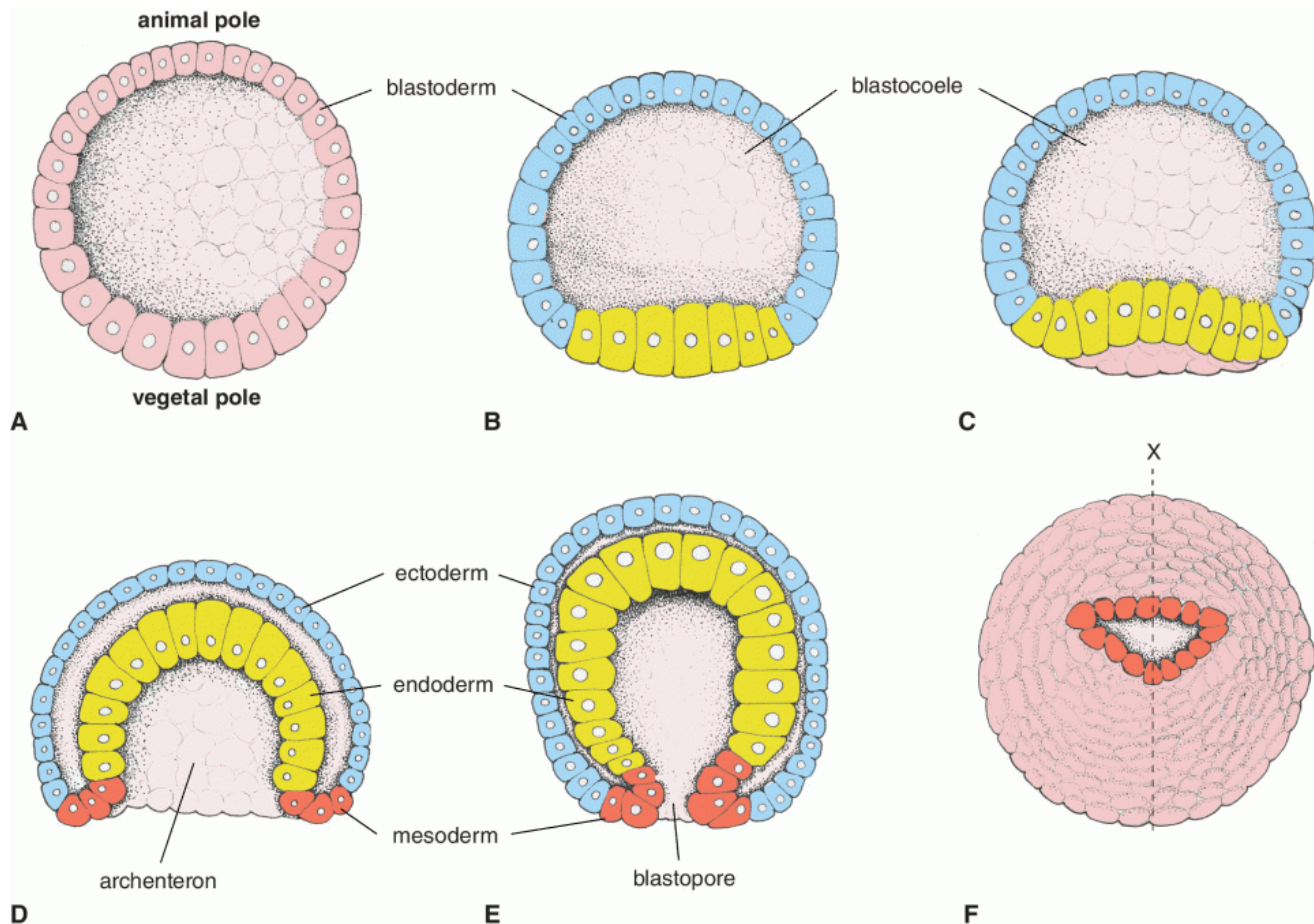
1- Stages of cleavage from the 2-cell stage to the early blastula stage in Amphioxus, A, and amphibians, B.



2- Sections showing sequential stages of gastrulation in Amphioxus from the blastula stage A to the gastrula stage E. The section shown in E is at the level indicated in the embryo at the gastrula stage in F.

The pattern of gastrulation in *Amphioxus* represents a comparatively simple model for illustrating the major cellular events in germ layer formation observed in more evolutionarily advanced species. Gastrulation in *Amphioxus* begins when the blastoderm at the vegetal pole flattens and invaginates. The embryo then undergoes a series of morphological changes. As cells at the vegetal pole invaginate, the spherical shape of the embryo changes with the sequential formation of a cavity referred to as the archenteron or primitive gut. The opening of the archenteron to the exterior is known as the blastopore. The outer layer of cells form the ectoderm, and the inner layer the endoderm. Cells responsible for the formation of the notochord and other mesodermal structures originally occupy a position at the edge of the blastopore. Later, these cells migrate to a position between the ectoderm and endoderm. Thus, the endodermal and mesodermal structures relocate from the surface of the embryo to its interior, forming a trilaminar embryo referred to as a gastrula.

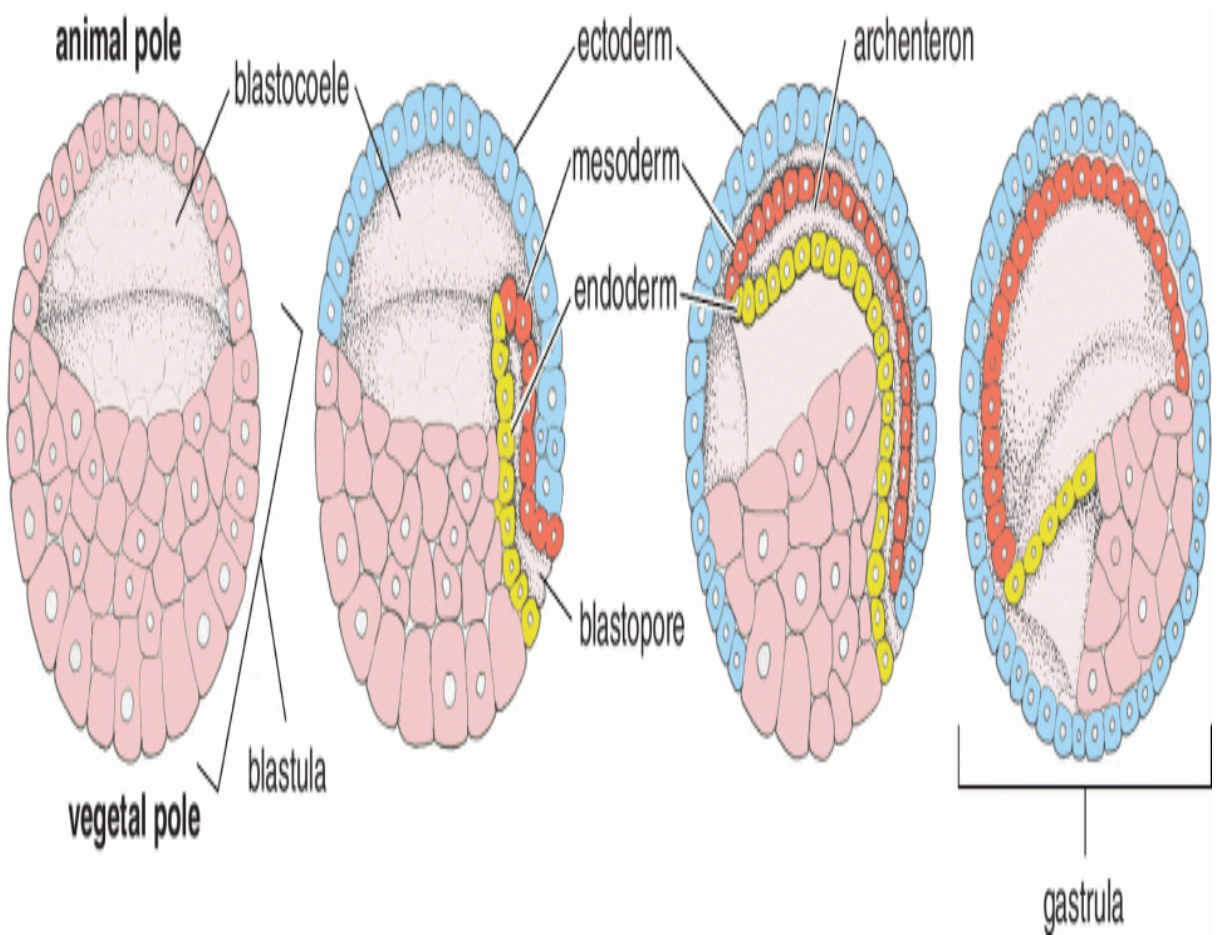




### 3- Sequential stages of gastrulation in amphibians from the blastula stage to the gastrula stage.

Because of the presence of yolk filled cells in the vegetal hemisphere of the amphibian blastula, invagination, as observed in *Amphioxus*, cannot occur. At the junction of the animal and vegetal hemispheres, cells from the surface move to the interior forming a cleft, the forerunner of the primitive gut. Following an influx of endodermal cells from below the cleft and mesodermal cells from above, the cleft deepens. With the constant movement of cells from the surface to the interior, a circular blastopore is formed. The blastocoele becomes obliterated and the yolk-laden cells at the vegetal pole move to the interior. Finally, a trilaminar

embryo, similar to that observed in *Amphioxus*, is formed.





## Early embryonic development of Birds

### Egg and fertilization

#### Testes

The male bird or rooster, possess a pair of testes, each testis is an oval body, cream white in colour and. From its inner border, a vas deferens emerges to run backwards, lateral to the ureter to open into the cloaca. Just before its opening, it dilates forming a small vesicular seminalis.

#### Ovaries

In the adult of most flying birds only one ovary of the left side, the right one degenerates.

#### The structure of ovum:

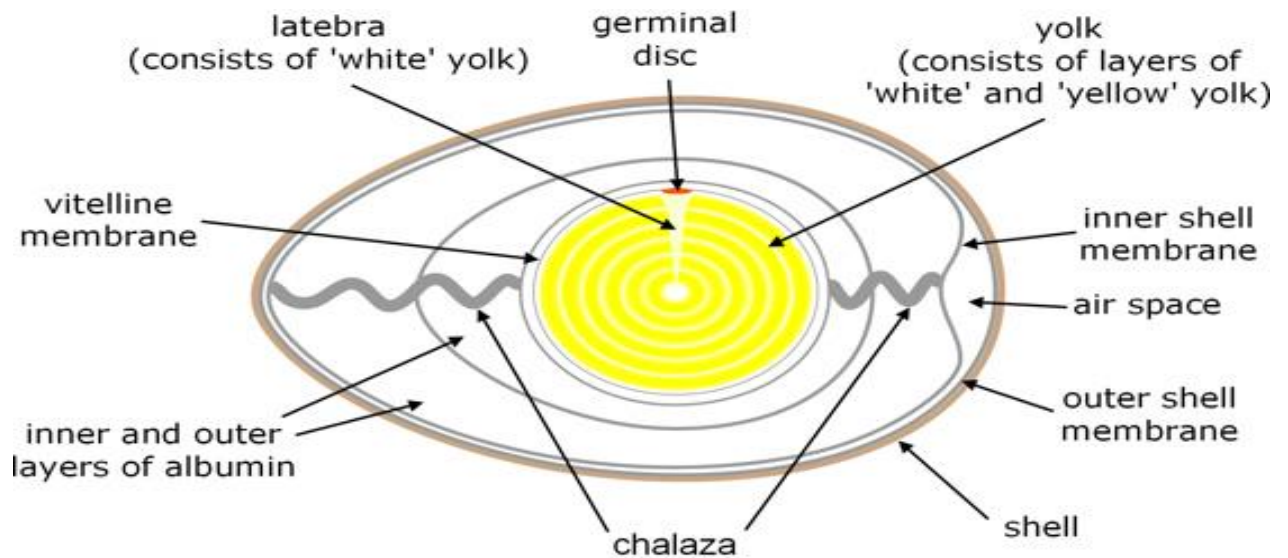
The oocyte grows due to the accumulation of yolk in it, the cytoplasm localized at the animal pole in which the nucleus embedded.

The mature oocyte migrates towards the periphery and finally bulged out from the ovary, connected with the ovary only by means of stalk. The fully formed egg contains a large amount of yolk. The cytoplasm is very little and is in the form of a small disc (the blastodisc or germinal disc).

After fertilization the ovum is surround by various envelopes added to the delicate vitelline membrane.

The cleavage starts immediately in the germinal disc. The fully formed and laid egg is surrounded on the outer side by a

calcareous shell. The shell consisting chiefly of calcium carbonate.



Section through domestic hen's egg.

Fertilisation of the egg occurs in the oviduct, before the albumen and shell are added to it. The egg is laid about 24 hours after fertilisation, by which time the development has reached the blastula stage.

### **Cleavage and blastula**

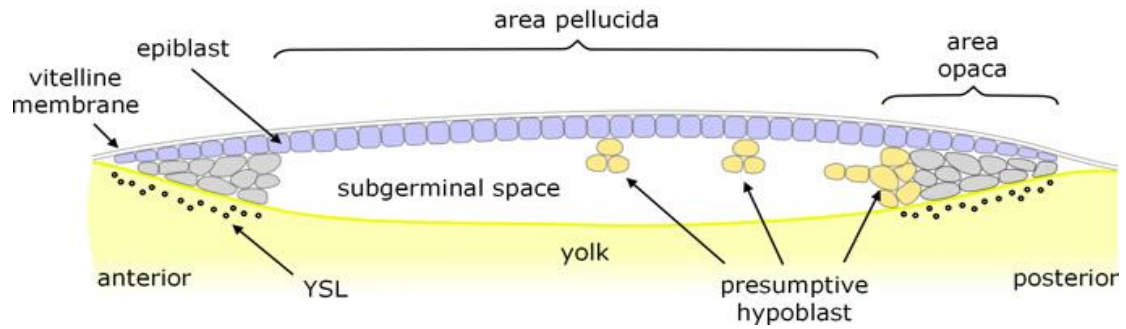
Consistent with having a large yolk, cleavage is meroblastic and is restricted to the germinal disc, and cell divisions do not extend into the yolk at all. The first division passes near the centre of the germinal disc and the next few divisions are at right angles to the preceding one, but then divisions become more irregular and asymmetric.



View from above of the germinal disc after the first few cleavages.

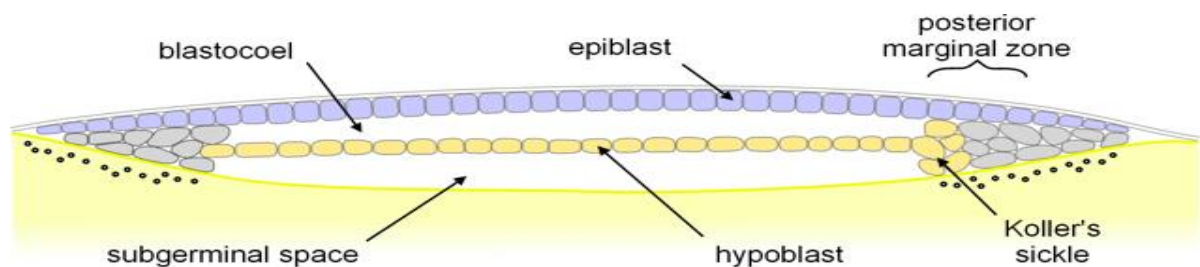
Hence, the early divisions give rise to a disc of cells (blastoderm), several layers thick, sitting on top of the yolk, with a subgerminal cavity in-between. Cells on the periphery of the disc are open to the cytoplasm of the germinal disc, i.e. they form a yolk syncytial layer (YSL).

Across most of the disc, the cells thin to a substantially single layer (but see below) of epithelial cells called the epiblast. This thinning appears to be mainly through cells of the deeper layers being shed, and possibly through some of the lower layer cells being intercalated with the uppermost layer. The central area is relatively translucent and is called the area pellucida, and beneath this is a substantially cell-free subgerminal space. Around the margins of the germinal disc, the cells remain several layers thick, and this relatively opaque peripheral ring is called the area opaca.



Longitudinal section through the blastoderm.

The hypoblast (sometimes called the primary hypoblast) then forms. Some of its cells arise as separate ‘islands’ beneath the epiblast; it is not known to what extent these are left-overs from the preceding thinning process or arise by delamination (ingression) from the overlying epiblast. Most of the hypoblast cells arise from the posterior marginal zone (especially the area known as Koller’s sickle, named because it is crescent shaped when viewed from above) by some of its underlying cells spreading anteriorly and incorporating the ‘islands’. The space between the epiblast and hypoblast is the blastocoel.

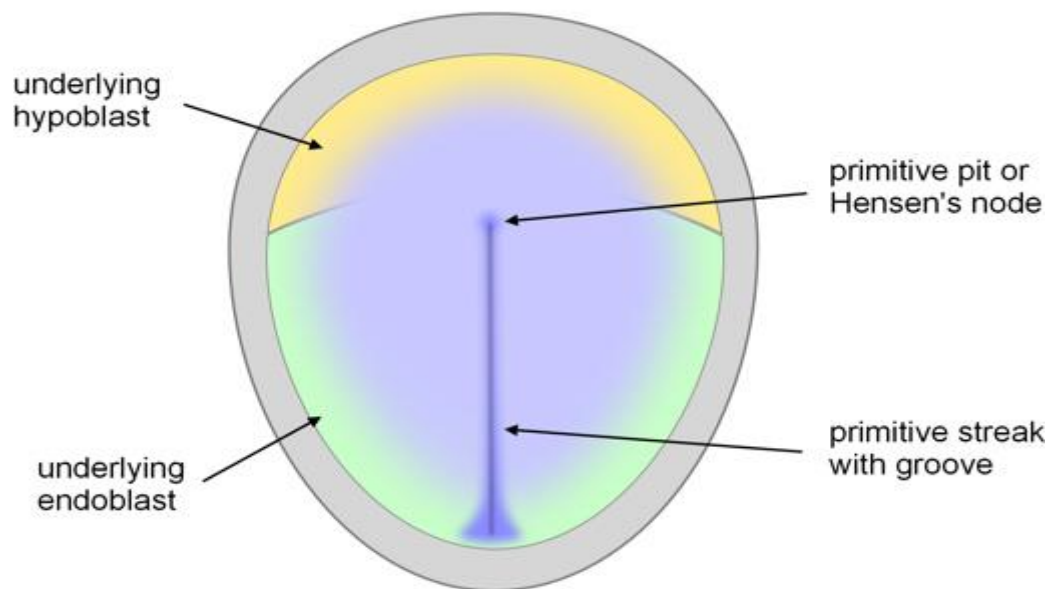


Longitudinal section through the blastula.

## Gastrulation

The key feature of gastrulation in birds is the primitive streak.

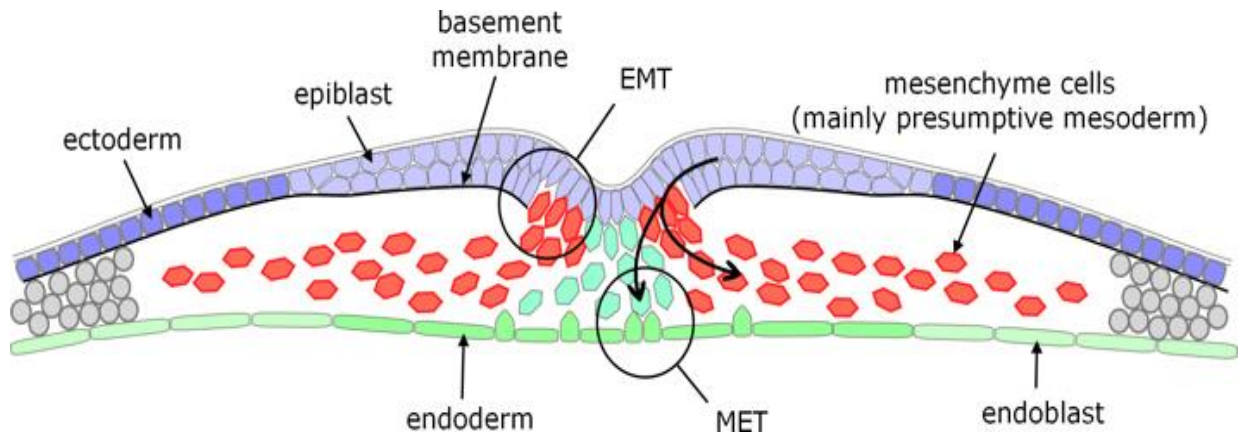
Initially this is a thickening of the epiblast along its midline, originating close to its posterior end (just forward of Koller's sickle) and then extending anteriorly until it reaches a maximum extent of about 2/3 across the area pellucida. As these thickening progresses along the epiblast, in tandem with it, a lower layer of cells spreads from the posterior margin. This endoblast (or secondary hypoblast) displaces the (primary) hypoblast anteriorly (figure 5). When the primitive streak reaches its maximum length, a groove develops on its dorsal (upper) surface along its length, culminating in a funnel-shaped depression at its anterior end, known as the primitive pit or Hensen's node.



Early primitive streak, at its maximum extent, viewed from above, with the epiblast faded at the margin to view the underlying hypoblast and endoblast.

The epiblast spreads (epiboly); and it is through the primitive groove and pit that epiblast cells now ingress between the

epiblast and endoblast. This translocation involves a change from an epithelial nature of the epiblast cells to a mesenchymal nature (epithelial-to-mesenchymal transition, EMT) such that the cells can migrate.



Transverse section through the primitive streak.

Some cells entering the primitive groove move across the intervening space and enter the endoblast to become endoderm (progressively displacing the endoblast to the sides of the embryo). Because the endoderm is an epithelial tissue, these migrating cells must revert to an epithelial nature i.e. undergo a mesenchymal-to-endothelial transition (MET).

Other cells spread out to form mesoderm between the overlying epiblast (ectoderm) and underlying endoderm.

In general, early cells entering the primitive streak become endoderm and later ones become mesoderm; but at any particular time some cells entering the streak are presumptive endoderm and some presumptive mesoderm.



Epiblast cells that do not enter the primitive streak remain as the ectoderm.

Early cells entering the primitive pit (Hensen's node) move anteriorly, enter the endoblast and become the endoderm of the presumptive foregut. Cells following these become head mesenchyme, prechordal plate mesoderm, and then chordamesoderm which becomes the most anterior part of the notochord, i.e. starting at the most anterior, progressively more posterior parts of the mesoderm are laid down. When the posterior-ward growth of the chordamesoderm reaches the primitive pit, the pit itself starts to retreat posteriorly; and, as it moves progressively further posteriorly, cells entering the node become progressively more posterior parts of the chordamesoderm (and subsequently notochord).

While gastrulation is proceeding, the cells of the marginal zone start to spread outwards, beginning the extraembryonic membranes.



## Embryonic development of mammals (humans)

Humans are classified within the group of mammals called eutherians which (with a few exceptions) use a placenta to nourish the developing embryo within the mother.

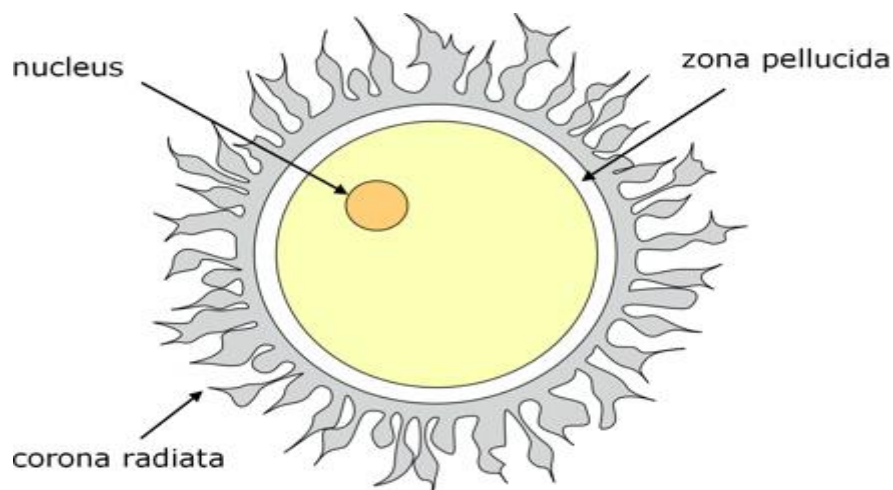
The other main groups of present-day mammals are:

marsupials (pouched), and

monotremes (egg-laying).

### Egg and fertilisation

The human ovum comprises a single cell, about 0.1 mm in size, containing the haploid nucleus. It is enveloped by a clear jelly-like coat called the zona pellucida; and the whole is surrounded by a population of follicular cells (originating from the ovary) which form the corona radiata.

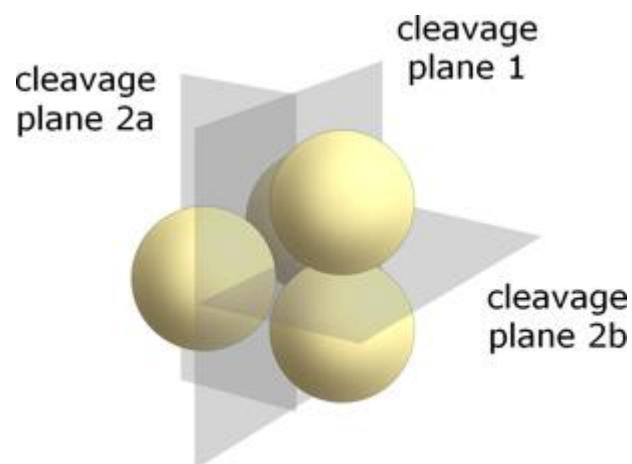


Section through human ovum.

Fertilisation takes place in the upper region of the oviduct (Fallopian tube) and the early stages of development unfold as the embryo travels along the oviduct. During its passage along the oviduct, the embryo loses some of the cells of the corona radiata, but the zona pelludica remains intact. Implantation in the uterus occurs when development has reached the blastocyst (blastula) stage.

## Cleavage to blastocyst

Consistent with the absence of a significant yolk, cleavage in mammals is holoblastic, meaning that the first division extends right through the egg cell. The first division is typical of other organisms having holoblastic cleavage, with the division being meridional – extending from one pole to the other. However, in mammals the second cleavage is unusual: one cell divides meridionally, but the other divides equatorially; which is called rotational cleavage.



Rotational cleavage: cleavage 2a is meridional, 2b is equatorial.

In most other organisms (having holoblastic cleavage) both of the second divisions are meridional, and the third or subsequent divisions are equatorial.

Also, whereas in other vertebrates the early cell divisions tend to be synchronous, this is often not the case with mammals, such that there is an odd number of cells instead of the usual geometric increase 2, 4, 8 ... .

Early cell divisions also tend to be slower than for other animals.

The cell divisions up to just before implantation take place within the zona pellucida, so there is no overall increase in size of the embryo, but its cells become smaller as they proliferate.

## Compaction and morula

Up to 8 – 16 cells, they form a loose association within the zona pellucida, but then they compact, with tight junctions forming between the outer cells. This is called the morula stage.

It is about now that the embryonic genome is activated, and this is followed by the first clear differentiation of cells, with further cell divisions resulting in inner cells being distinct from the peripheral ones, and the beginning of a fluid-filled cavity, the blastocoel.

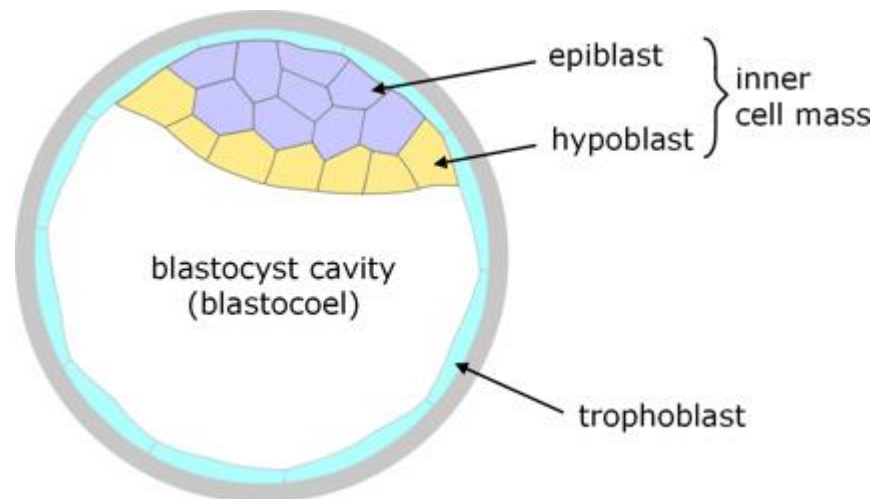
## Blastocyst

The blastocyst is the mammalian equivalent of the blastula in other vertebrates. It comprises three populations of cells:

an outer cell layer (trophoblast) which develops into the placenta; and

an inner cell mass:

most of which is the epiblast, which is the source of embryonic tissues, and the amnion, although the layer of cells in contact with the blastocoel is the hypoblast, most of which forms extraembryonic tissues.



Human blastocyst.

## Implantation and embryonic disc

About 7 days after fertilisation the embryo loses the zona pellucida, and implants into the lining of the uterus, by which time it comprises about 200 cells.

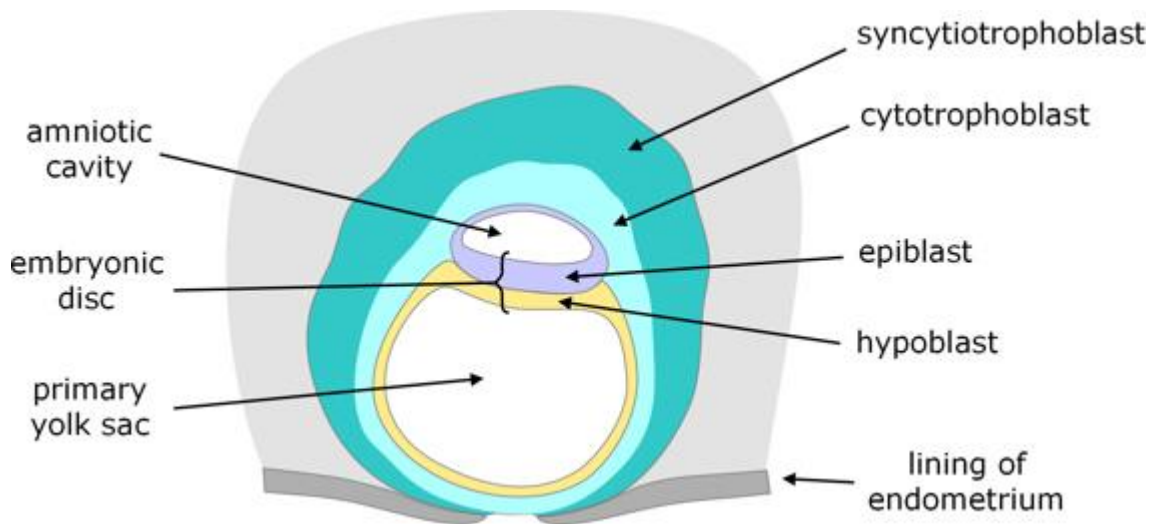
Around the time of implantation:

- the amniotic cavity arises within the epiblast,
- the blastocyst cavity becomes the primary yolk sac, which is lined by cells that spread from the hypoblast.

In addition, cells from the epiblast and hypoblast organise into a two-layered structure known as the embryonic disc, positioned between the primary yolk sac and amniotic cavity. Previously these two layers

had been thought to correspond with the first two germ layers, the ectoderm and endoderm, but see below.

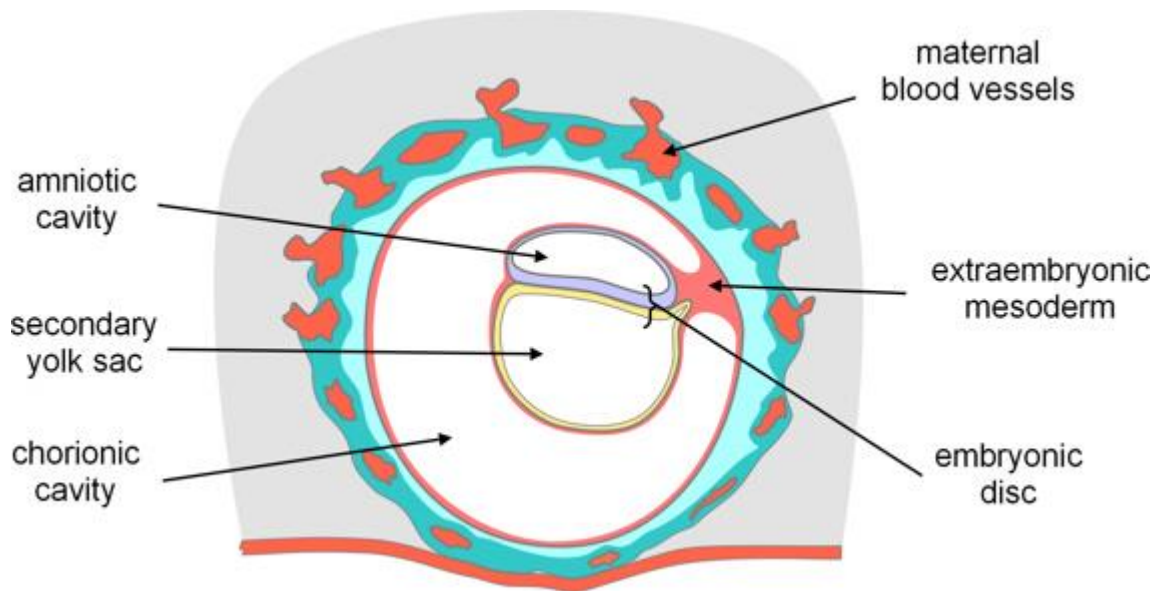
The trophoblast penetrates the uterine wall and begins to form the placenta; at an early stage it differentiates into two distinct layers, known as the cytotrophoblast and the syncytio trophoblast.



Embryo shortly after implantation, with embryonic disc.

## Extraembryonic mesoderm

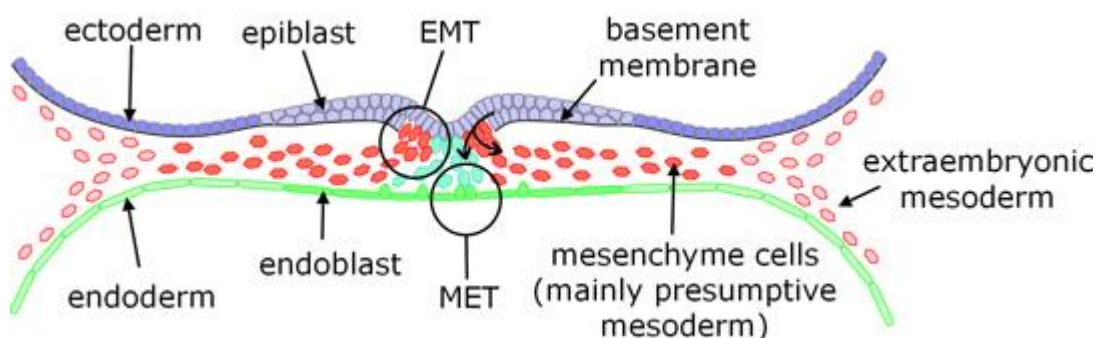
Extraembryonic mesoderm is a layer of tissue that arises between the lining of the primary yolk sac and the cytotrophoblast, and spreads to cover the amniotic cavity as well. As this tissue thickens, cavities form within it, and coalesce to form the chorionic cavity which is lined with extraembryonic mesoderm (figure 5). (Note that this is different from the germ-layer mesoderm which forms later, see below.) In this process some of the primary yolk sac is lost, and what remains is called the secondary yolk sac. The embryo remains attached to the internal lining of the chorion by a connecting stalk of extraembryonic mesoderm which becomes the umbilical cord.



Longitudinal section pre-gastrulation (approx. day 14).

## Gastrulation

From about day 14 after fertilisation, a groove appears near the caudal end of the epiblast, it extends about two-thirds of the way along the midline towards the cranial end, terminating in a widening with a depression at its centre. This primitive groove and primitive pit are where gastrulation occurs. As gastrulation proceeds, the primitive node recedes caudally, with a corresponding shortening of the primitive streak.



Transverse section (perpendicular to figure 5) through the embryonic disc, showing gastrulation.

In the course of gastrulation, cells of the epiblast proliferate and move towards the groove where they transition from epithelial to mesenchymal in character, and ingress below the surface. In the early phase, these ingressing cells enter the hypoblast, reverting to epithelial cells (mesenchymal to epithelial transition) to become the definitive endoderm; at the same time displacing the hypoblast cells from the embryonic disc to line the yolk sac. As this stage proceeds, further cells ingressing from the epiblast move into the space between the epiblast and endoderm to form a middle layer of cells called mesoderm. (At the edges of the embryonic disc this embryonic mesoderm merges with the previously formed extraembryonic mesoderm.) Once the mesoderm is formed, the remaining epiblast is called ectoderm, and the three germ layers are complete.



### **Four extraembryonic membranes (or embryonic membranes or foetal membranes):**

Formed in amniotes (reptiles, birds and mammals) outside the body of embryo by the cells of presumptive ectoderm, mesoderm and endoderm. These are:

- Chorion
- Amnion
- Allantois
- Yolk sac



**1- Chorion (serosa):** The outermost covering, formed by ectoderm and mesoderm as a protective layer head fold and tail fold of ectoderm and mesoderm emerge from respective parts of embryo, start growing and folding upon the dorsal side where both fuse (= sero-amniotic connection) to form outer chorion and inner amnion.

✓ The site of exchange between the embryo and the environment around it.

✓ **In reptiles and birds**, the principal function of the chorion is the respiratory exchange of gases.

✓ In mammals, the chorion serves a much more all-embracing function which includes not only respiration but also nutrition, excretion, filtration, and synthesis-with hormone production begin an important example of the last function.

**2- Amnion:** It forms private (closest) chamber of embryo filled with amniotic fluid, isotonic to the body fluid. The aquatic medium for embryo to float and grow, denotes the aquatic origin of life. This fluid having cells of embryo is used (amniocentesis) to test its sex and genetic disorders.

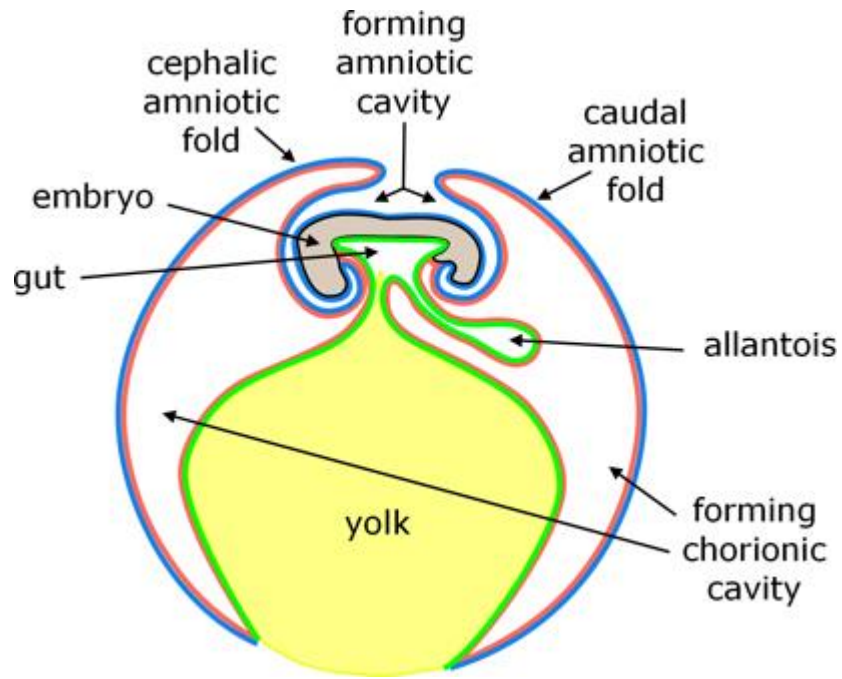
✓ A thin ectodermally derived membrane which eventually encloses the entire embryo in a fluid-filled sac. The amniotic membrane is functionally specialized for the secretion and absorption of the amniotic fluid that bathes the embryo. So characteristic is this structure that the reptiles, birds, and mammals as a group are often called amniotes. The fishes and amphibians, lacking an amnion, are collectively called an amniotes.

**3- Allantois:** It develops from the inner endoderm and outer mesoderm. Its principal functions are to act as a reservoir for storing or removing urinary wastes and to mediate gas exchange between the embryo and its surroundings.

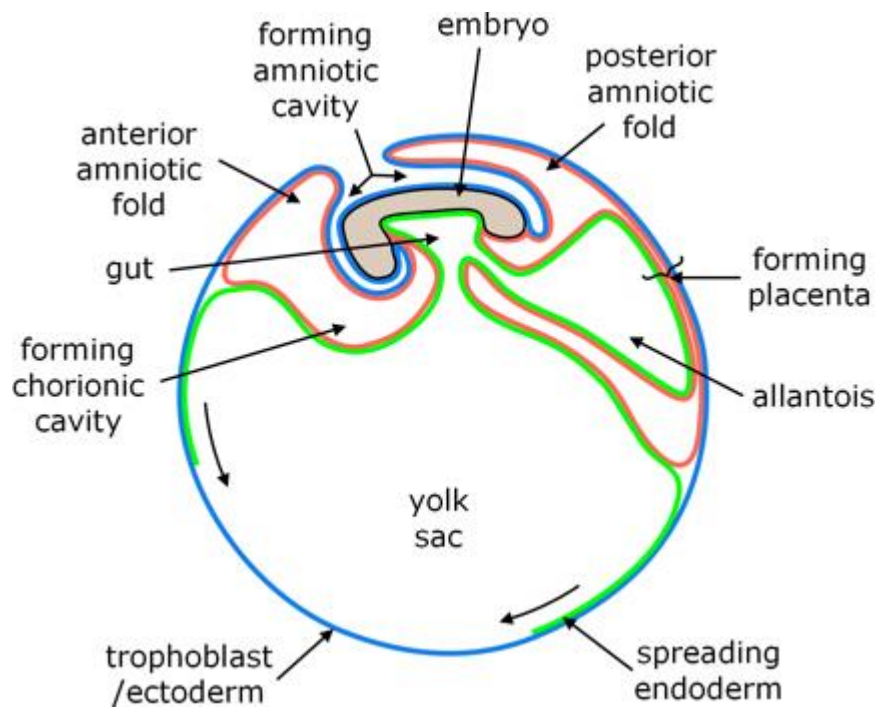
✓ **In reptiles and birds,** the allantois is a large sac, and because the egg is a closed system with respect to urinary wastes, the allantois must sequester nitrogenous by-product so that they do not subject the embryo to osmotic stress or toxic effects.

✓ **In mammals** the role and prominence of the allantois vary with the efficiency of the interchange that takes place at the fetal-maternal interface. The allantois of the pig embryo rivals that of the bird in both size and functional importance, whereas the human allantois has been reduced to a mere vestige that contributes only a well-developed vascular network to the highly efficient placenta.

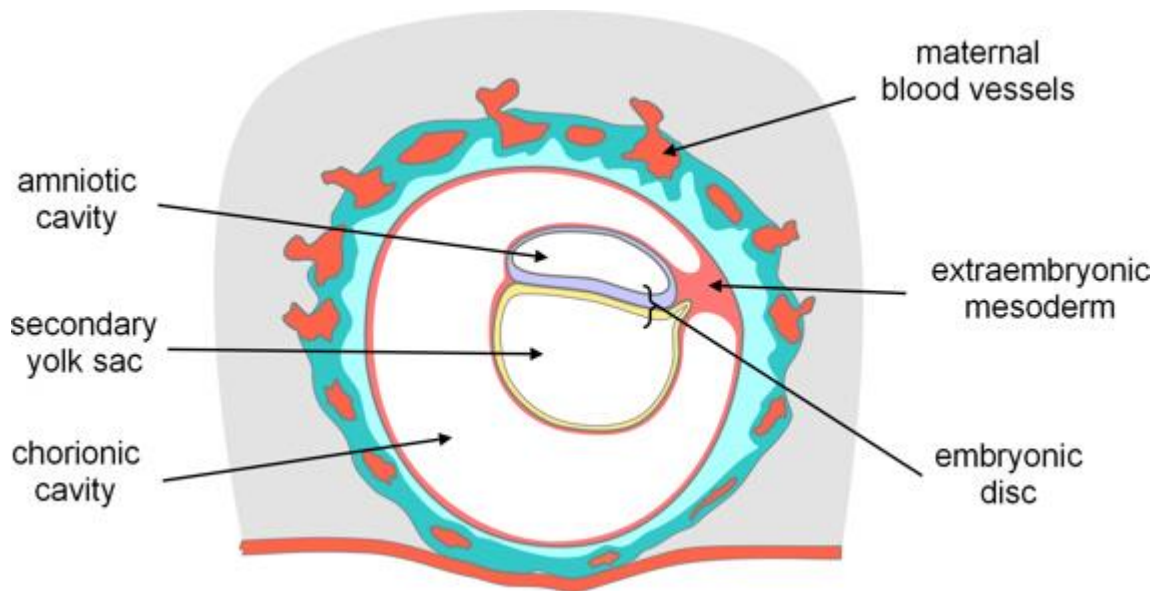
**4- The yolk sac:** The endodermal yolk sac is intimately involved with nutrition of the embryo in large-yolked forms such as reptiles and birds. Despite the lack of stored in mammalian eggs, the yolk sac has been preserved, possibly because other important secondary functions are associated with it. For example, the yolk sac endoderm induces the surrounding extraembryonic mesoderm to form the first blood cells and blood vessels.



Chick extraembryonic membranes



Rabbit extraembryonic membranes



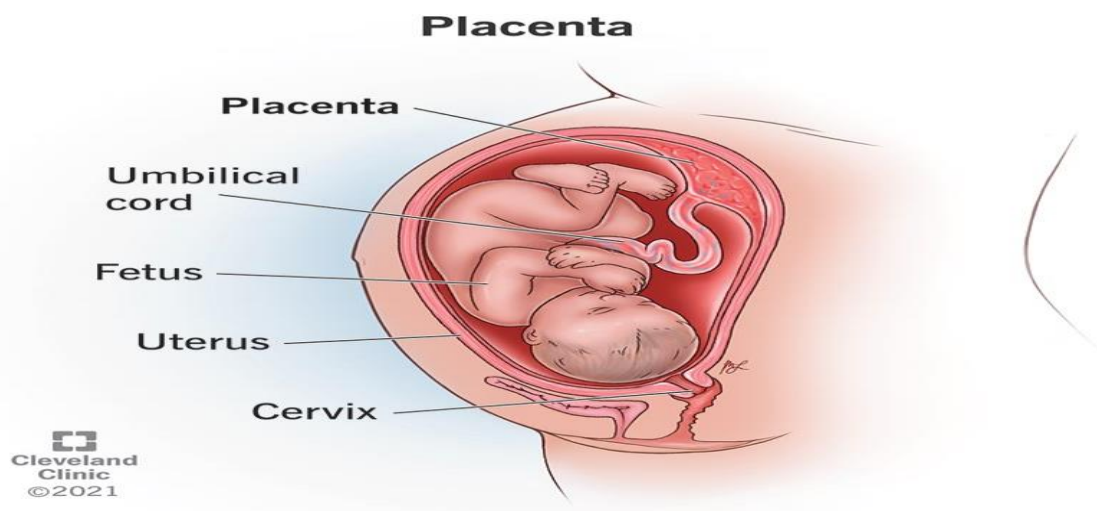
Human extraembryonic membranes



## Placenta

### What is the placenta?

The placenta is a temporary organ that forms in your uterus during pregnancy. It attaches to your uterine wall and provides nutrients and oxygen to your baby through the umbilical cord. Certain conditions of the placenta can cause pregnancy complications.



The placenta is a temporary organ that connects your baby to your uterus during pregnancy. The placenta develops shortly after conception and attaches to the wall of your uterus. Your baby is connected to the placenta by the umbilical cord. Together, the placenta and umbilical cord act as your baby's lifeline while in the uterus. Functions of the placenta include:

- Provides your baby with oxygen and nutrients.
- Removes harmful waste and carbon dioxide from your baby.
- Produces hormones that help your baby grow.
- Passes immunity from you to your baby.
- Helps protect your baby.

### **When does the placenta form?**

- The placenta begins to form after a fertilized egg implants in your uterus around seven to 10 days after conception. It continues to grow throughout your pregnancy to support your baby. The placenta starts as a few cells and grows to be several inches long.

### **When does the placenta take over?**

- The placenta takes over hormone production by the end of the first trimester (12 weeks of pregnancy). Up until this time, the corpus luteum handles most of the hormone production. Many people's first-trimester symptoms of nausea and fatigue go away once the placenta takes over in the second trimester.

### **What does the placenta do?**

- The placenta helps to keep your baby alive and healthy during pregnancy. Your blood passes through the placenta and provides oxygen, glucose and nutrients to your baby through the umbilical cord. The placenta can also filter out harmful waste and carbon dioxide from your baby's blood. The placenta enables the exchange of oxygen and nutrients between the bloodstreams of you and your baby without ever mixing them. It acts as your baby's lungs, kidneys and liver until birth.
- As you get closer to delivery, the placenta passes antibodies to your baby to jumpstart its immunity. This immunity sticks with your baby for the first several months of life.
- The placenta produces several important hormones like lactogen, estrogen and progesterone during pregnancy. These pregnancy hormones are beneficial to both you and your baby. For example, the placenta produces a hormone that suppresses milk production during pregnancy.

### **Does the placenta move?**

- The placenta appears to move only because the uterus expands as the pregnancy and fetus grow. Your healthcare provider will look at the location of your placenta during your 20-week anatomy ultrasound and determine if its position may cause complications. Most placentas move to the top or side of the uterus by 32 weeks of pregnancy.

### **Where does the placenta form?**

The placenta can form anywhere in your uterus. It develops wherever the fertilized egg implants into your uterine wall. Some of the positions of the placenta are:

- **Posterior placenta:** The placenta grows on the back wall of your uterus.
- **Anterior placenta:** The placenta grows on the front wall of your uterus closest to your abdomen.
- **Fundal placenta:** The placenta grows at the top of your uterus.
- **Lateral placenta:** The placenta grows on the right or left wall of your uterus.

The placenta can move up until about 32 weeks of pregnancy. It's common to have a placenta that moves upwards and away from your cervix as your baby gets bigger.

### **What does the placenta look like?**

The placenta looks like a disc of bumpy tissue rich in blood vessels, making it appear dark red at term. Most of the mature placental tissue is made up of blood vessels. They connect with the baby through the umbilical cord and branch throughout the placenta disc like the limbs of a tree.

### **What color is the placenta?**

The placenta has two sides: the side attached to your uterus and the side closest to your baby. The side attached to your uterine wall is a deep reddish blue color, while the side facing your baby is gray.

### **How big is a normal placenta?**

The placenta is about 10 inches long and 1 inch thick at its center. It weighs around 16 ounces (1 pound) by the time your baby is born.

### **What is the placenta made of?**

The placenta begins to develop when the fertilized egg implants into your uterine wall. The placenta contains mostly blood vessels contained within structures called “villi.” The blood vessels connect with the baby’s bloodstream through the umbilical cord. The rest of the placental tissues mainly connect the villi to the umbilical cord and allow your blood to bathe the villi, supplying the baby with oxygen and nutrients.

### **What types of substances are bad for the placenta?**

Medicine, drugs, alcohol, and nicotine can all transfer from your bloodstream to your baby through the placenta. Talk to your healthcare provider before taking any prescription or over-the-counter medications (including vitamins and supplements) during pregnancy. Drinking alcohol or smoking cigarettes is not recommended during pregnancy.

### **How is the placenta delivered?**

The placenta is delivered shortly after your baby is born (usually between five and 30 minutes after). This is called the afterbirth or the third stage of labor. If you've delivered your baby vaginally, your uterus will continue to contract to expel the placenta. Your healthcare provider may push on your belly or ask you for one final push. If your baby was born via C-section, your healthcare provider removes the



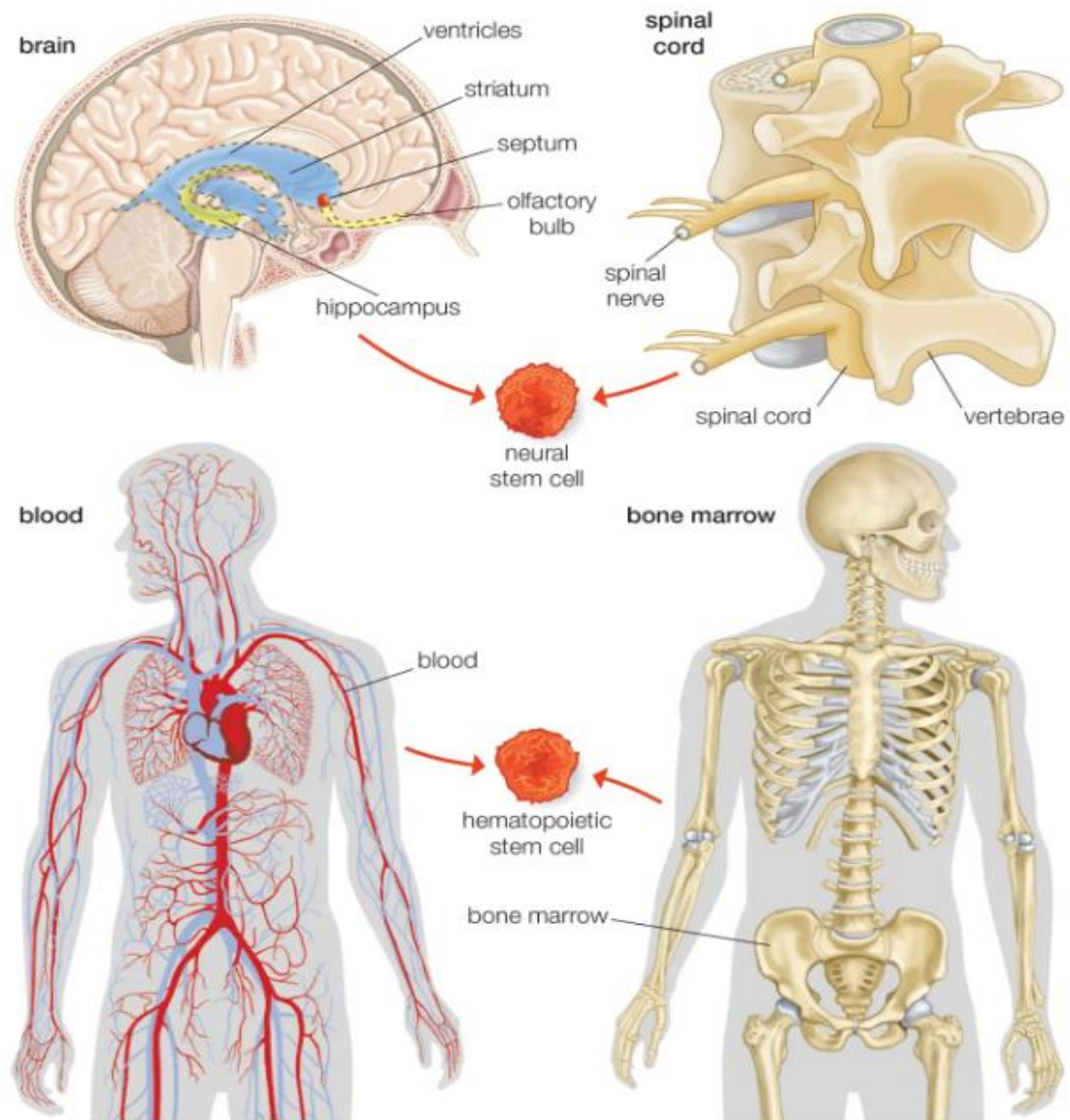
placenta through the incision used to deliver your baby. In rare cases, parts of the placenta stay in your uterus after delivery. This can cause bleeding, pain and infection.



## Stem cells

*Stem cell*, an undifferentiated cell that can divide to produce some offspring cells that continue as stem cells and some cells that are destined to differentiate (become specialized). Stem cells are an ongoing source of the differentiated cells that make up the tissues and organs of animals and plants. There is great interest in stem cells because they have potential in the development of therapies for replacing defective or damaged cells resulting from a variety of disorders and injuries, such as Parkinson disease, heart disease, and diabetes. There are two major types of stem cells: embryonic stem cells and adult stem cells, which are also called tissue stem cells.

### Anatomical sources of neural and hematopoietic stem cells



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## Types of stem cells

There are several types of stem cells that can be used for different purposes.

## **Embryonic stem cells**

Embryonic stem cells come from human embryos that are three to five days old. They are harvested during a process called in-vitro fertilization. This involves fertilizing an embryo in a laboratory instead of inside the female body. Embryonic stem cells are known as pluripotent stem cells. These cells can give rise to virtually any other type of cell in the body.

## **Non-embryonic (adult) stem cells**

Adult stem cells have a misleading name, because they are also found in infants and children. These stem cells come from developed organs and tissues in the body. They're used by the body to repair and replace damaged tissue in the same area in which they are found.

For example, hematopoietic stem cells are a type of adult stem cell found in bone marrow. They make new red blood cells, white blood cells, and other types of blood cells. Doctors have been performing stem cell transplants, also known as bone marrow transplants, for decades using hematopoietic stem cells in order to treat certain types of cancer.

Adult stem cells can't differentiate into as many other types of cells as embryonic stem cells can.

### **Induced pluripotent stem cells (iPSCs)**

Scientists have recently discovered how to turn adult stem cells into pluripotent stem cells. These new types of cells are called induced pluripotent stem cells (iPSCs). They can differentiate into all types of specialized cells in the body. This means they can potentially produce new cells for any organ or tissue. To create iPSCs, scientists genetically reprogram the adult stem cells, so they behave like embryonic stem cells.

The breakthrough has created a way to “de-differentiate” the stem cells. This may make them more useful in understanding how diseases develop. Scientists are hoping that the cells can be made from someone’s own skin to treat a disease. This will help prevent the immune system from rejecting an organ transplant. Research is underway to find ways to produce iPSCs safely.

### **Cord blood stem cells and amniotic fluid stem cells**

Cord blood stem cells are harvested from the umbilical cord after childbirth. They can be frozen in cell banks for use in the future. These cells have been successfully used to treat children with blood cancers, such as leukemia, and certain genetic blood disorders.

Stem cells have also been found in amniotic fluid. This is the fluid that surrounds a developing baby inside the mother’s womb. However, more research is needed to help understand the potential uses of amniotic fluid stem cells.



## **In vitro fertilization (IVF)**

In vitro fertilization (IVF) is a complex series of procedures used to help with fertility or prevent genetic problems and assist with the conception of a child.

During IVF, mature eggs are collected (retrieved) from ovaries and fertilized by sperm in a lab. Then the fertilized egg (embryo) or eggs (embryos) are transferred to a uterus. One full cycle of IVF takes about three weeks. Sometimes these steps are split into different parts and the process can take longer.

IVF is the most effective form of assisted reproductive technology. The procedure can be done using a couple's own eggs and sperm. Or IVF may involve eggs, sperm or embryos from a known or anonymous donor. In some cases, a gestational carrier — someone who has an embryo implanted in the uterus — might be used.

Your chances of having a healthy baby using IVF depend on many factors, such as your age and the cause of infertility. In addition, IVF can be time-consuming, expensive and invasive. If more than one embryo is transferred to the uterus, IVF can result in a pregnancy with more than one fetus (multiple pregnancy).

## Why it's done

In vitro fertilization (IVF) is a treatment for infertility or genetic problems. If IVF is performed to treat infertility, you and your partner might be able to try less-invasive treatment options before attempting IVF, including fertility drugs to increase production of eggs or intrauterine insemination — a procedure in which sperm are placed directly in the uterus near the time of ovulation.

Sometimes, IVF is offered as a primary treatment for infertility in women over age 40. IVF can also be done if you have certain health conditions. For example, IVF may be an option if you or your partner has:

- **Fallopian tube damage or blockage.** Fallopian tube damage or blockage makes it difficult for an egg to be fertilized or for an embryo to travel to the uterus.
- **Ovulation disorders.** If ovulation is infrequent or absent, fewer eggs are available for fertilization.
- **Endometriosis.** Endometriosis occurs when tissue similar to the lining of the uterus implants and grows outside of the uterus — often affecting the function of the ovaries, uterus and fallopian tubes.
- **Uterine fibroids.** Fibroids are benign tumors in the uterus. They are common in women in their 30s and 40s. Fibroids can interfere with implantation of the fertilized egg.
- **Previous tubal sterilization or removal.** Tubal ligation is a type of sterilization in which the fallopian tubes are cut or blocked to

permanently prevent pregnancy. If you wish to conceive after tubal ligation, IVF may be an alternative to tubal ligation reversal surgery.

- **Impaired sperm production or function.** Below-average sperm concentration, weak movement of sperm (poor mobility), or abnormalities in sperm size and shape can make it difficult for sperm to fertilize an egg. If semen abnormalities are found, a visit to an infertility specialist might be needed to see if there are correctable problems or underlying health concerns.
- **Unexplained infertility.** Unexplained infertility means no cause of infertility has been found despite evaluation for common causes.
- **A genetic disorder.** If you or your partner is at risk of passing on a genetic disorder to your child, you may be candidates for preimplantation genetic testing — a procedure that involves IVF. After the eggs are harvested and fertilized, they're screened for certain genetic problems, although not all genetic problems can be found. Embryos that don't contain identified problems can be transferred to the uterus.
- **Fertility preservation for cancer or other health conditions.** If you're about to start cancer treatment — such as radiation or chemotherapy — that could harm your fertility, IVF for fertility preservation may be an option. Women can have eggs harvested from their ovaries and frozen in an unfertilized state for later use. Or the eggs can be fertilized and frozen as embryos for future use.

Women who don't have a functional uterus or for whom pregnancy poses a serious health risk might choose IVF using another person to carry the pregnancy (gestational carrier). In this case, the woman's eggs are fertilized with sperm, but the resulting embryos are placed in the gestational carrier's uterus.

**Risks of IVF include:**

- **Multiple births.**
- **Premature delivery and low birth weight.**
- **Ovarian hyperstimulation syndrome.**
- **Miscarriage.**
- **Egg-retrieval procedure complications**
- **Ectopic pregnancy.**
- **Birth defects.**
- **Cancer.**
- **Stress.**

**The End**

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## Glossary of embryological terms

### Gametes

#### **gamete**

Reproductive cell: e.g. ovum or sperm.

#### **meiosis**

The process where a single cell divides twice to give four cells containing half of the original genetic material; typically produces sperm or ovum.

#### **haploid**

Sperm and egg cells result from meiosis and are haploid: they have only half of the number of chromosomes of somatic (body) cells, e.g. in humans, haploid cells have 23 chromosomes, whereas somatic cells have 46 chromosomes.

#### **polar body**

Production of an ovum involves 2 rounds of cell division, giving rise to 4 (sometimes only 3) nuclei, only one of which is incorporated within an ovum. The resulting 3 (or 2) nuclei are small polar bodies which eventually degrade. One polar body may remain associated with the ovum for a while. [check]

#### **animal pole and vegetal pole**

In most ova, the nucleus is not in the centre but displaced to one end - this is called the animal pole, and the opposite end is the vegetal pole. In general there is more yolk towards the vegetal pole.

#### **germinal disc**

The embryo-forming part of the egg, e.g. excluding the body of the yolk.

**zona pellucida**

A clear layer surrounding the cell of the mammalian ovum (c.f. area pellucida of bird and reptile blastula).

**corona radiata**

The outermost layer of the mammalian ovum.

**Zygote****zygote**

Fertilised egg.

**pronucleus**

A pronucleus is the male or female polar body or 'nucleus' within the zygote, at the start of fertilisation.

**Cleavage****cleavage**

The first few cell divisions of the zygote, during which there is an increase in number of cells, but not in overall size or mass, so the cells get smaller.

**holoblastic cleavage**

The cell divisions pass right through the zygote.

**meroblastic cleavage**

The cell divisions do not extend right through the zygote (generally there is a large amount of yolk)

**discoidal cleavage**

A form of meroblastic cleavage where the cell divisions are restricted to the germinal disc.

**rotational cleavage**

A form of holoblastic cleavage where the two second cell divisions are in different planes, one meridional and one equatorial.

**morula**

An early stage of embryonic development, especially in mammals, when the cells are in the form of a loose clump.

**compaction**

A stage of embryonic development when the loose clump of cells becomes more compact, usually involving the formation of tight junctions between peripheral cells and the formation of a central blastocoel.

**blastomere**

One of the cells resulting from cleavage; or one of the cells of the blastoderm.

**blastodisc**

The germinal disc during and after cleavage, up to the stage of the blastula.

**Blastula****blastula**

The early embryo, after cleavage and the first differentiation of cells, just before gastrulation.

**blastocyst**

The name for the blastula in mammals.

**blastoderm**

A surface layer of cells of the blastula.

**blastocoel**

A space within the body of cells of the blastula, generally arising during cleavage.

**subgerminal space / cavity**

A space between the cells of the blastula and the underlying yolk.

**yolk syncytial layer (YSL)**

A layer at the surface of the yolk containing nuclei, but the cytoplasm of the cells is continuous with the yolk, i.e. the cell membranes are incomplete.

**mid blastula transition (MBT)**

Generally refers to the stage of the blastula when the embryonic genome becomes active and/or the cells begin to differentiate.

**epiblast**

In amniotes: the upper, epithelial layer of the blastula, i.e. before gastrulation, which in most cases is the source of all of the germ layers.

In anamniotes: sometimes used to refer to the upper, epithelial layer of the gastrula, i.e. after gastrulation, which usually becomes the ectoderm.

**hypoblast**

In amniotes: a layer of cells below the epiblast, which is substantially displaced in the course of gastrulation and does not become part of the embryo.

In anamniotes: sometimes used to refer to a lower layer of cells that have involuted in the course of gastrulation, and usually becomes mesoderm and endoderm (except amphibians).

**trophoblast**

The outer layer of cells of the mammalian blastocyst.

**inner cell mass**

Generally refers to mammals where the blastocyst comprises an outer layer of cells (the trophoblast), an inner cell mass (generally containing epiblast and hypoblast) and a blastocoel.

**area pellucida**

Central part of the reptile or bird blastula, that is over the blastocoel and is relatively translucent (c.f. zona pellucida of the mammalian ovum).

**area opaca**

Area of the reptile or bird blastula, that is around the area pellucida and above the marginal cells, and is relatively opaque.

**enveloping layer (EVL)**

A thin outer layer of cells of the teleost blastula, which persists through early embryonic development, but is shed at hatching.

**Gastrulation****gastrulation**

The overall term for the various processes through which the blastula develops into the gastrula in which all three germ layers - ectoderm, mesoderm and endoderm - are formed.

**presumptive and definitive**

Presumptive refers to cells which are still undifferentiated, or partly differentiated, but destined to become the final or definitive tissue.

### **epithelial**

Surface cells are generally epithelial in nature, and move as a sheet of cells.

### **mesenchymal**

Mesenchymal cells are usually below the surface, and migrate individually through tissues.

### **EMT and MET**

When cells transition between epithelial and mesenchymal in nature by 'epithelial-to-mesenchymal transition' or 'mesenchymal-to-epithelial transition'.

### **epiboly**

Spreading of a layer of cells in the course of gastrulation. This can be achieved by thinning of individual cells, and/or intercalation of cells from within the same or adjacent cell layer.

### **invagination**

The process whereby a layer of cells indents and forms a cavity or pouch.

### **involution**

Movement of cells from the outside to the inside, the cells remaining epithelial in nature and moving as a layer of cells.

### **ingression**

Movement of cells, typically from the outside to the inside, the cells being (or transitioning to) mesenchymal in nature, and moving individually.

**blastopore**

A sac-like cavity in the outer surface of the blastula through which cells are internalised, usually by involution.

**primitive streak**

A depression in the epiblast of birds and mammals through which cells are internalised by ingression.

**ectoderm**

The outer germ layer; generally produces the outer layer of the embryo, and central nervous system via the neural tube.

**mesoderm**

The middle germ layer; from it forms the notochord, somites, some of the skeleton and musculature.

**endoderm**

The inner germ layer; it forms most of the digestive tract and associated organs.

**extraembryonic membranes**

These are membranes that develop along with the embryo but do not form part of the final embryo.

Usually these are the amniote amnion, chorion, and membranes of the yolk sac and allantois.

The yolk sac of teleosts is part of the embryo, and the yolk sac of chondrichthyans is reabsorbed, so these are not strictly extraembryonic.

Although the teleost enveloping layer (EVL) does not become part of the embryo, it is not usually regarded as an extraembryonic membrane.

## References

1. EMBRYOLOGY . Mathur,Ramesh. RAMESH MATHUR, Meenakshi Mehta. INDIA : ANMOL PUBLICATIONS P.V.T LTD. 2005 .
2. Müller, W.A., Hassel, M. and Greal, M., 2015. Development and reproduction in humans and animal model species. Springer.
3. [en.wikipedia.org/wiki/Embryology](http://en.wikipedia.org/wiki/Embryology).
4. [www.embryology.ch/indexen.html](http://www.embryology.ch/indexen.html)
5. <http://courses.biology.utah.edu/bastiani/3230/DB%20Lecture/Lectures/a6Cleav.html>
6. <http://www.yourarticlelibrary.com/biology/the-pattern-of-cleavage-due-to-organization-of-egg-may-be-of-the-following-types-biology/5129>.
7. <http://www.vcbio.science.ru.nl/en/virtuallessons/embryology/seaurchinslides/>
8. [https://faculty.cascadia.edu/ccollin/frog\\_development.htm](https://faculty.cascadia.edu/ccollin/frog_development.htm)
9. <http://www.notesonzoology.com/vertebrates/chick/development-of-chick-with-diagram-vertebrates-chordata-zoology/8645>
10. <http://www.notesonzoology.com/embryology/gastrulationembryology/gastrulation-in-amphioxus-and-amphibianembryology/13392>
11. <http://www.yourarticlelibrary.com/biology/5-steps-involved-in-the-development-of-chick-explained/23153>.
12. <http://www.notesonzoology.com/phylumchordata/branchiostoma/development-of-branchiostomacephalochordata-chordata-zoology/8606>.
13. [https://embryology.med.unsw.edu.au/embryology/index.php/Book\\_-\\_Text-Book\\_of\\_Embryology\\_5](https://embryology.med.unsw.edu.au/embryology/index.php/Book_-_Text-Book_of_Embryology_5)
14. <https://thebiologynotes.com>



15. Britannica
16. <https://byjus.com>
17. <https://www.bajkulcollegeonlinestudy.in>
18. <https://cdn.lecturio.com>
19. <https://www.sciencedirect.com>
20. <https://veteriankey.com>