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PRACTICAL HANDBOOK OF ZOOLOGY MINOR (B. SC. II, SEM IV)

AS PER NEP-2020 (2.0) SYLLABUS OF SHIVAJI UNIVERSITY, KOLHAPUR

K. J. Adate

S. A. Vhanalakar

V. V. Ajagekar



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PREFACE

Dear Students,

It is with great pleasure that we present this practical manual, carefully prepared in accordance with the B.Sc. Part II, Semester IV (NEP 2.0) Minor Zoology Practical – IV syllabus prescribed by Shivaji University, Kolhapur. This book has been designed as a comprehensive and reliable companion to support students in exploring the diverse and fascinating areas of Physiology, Endocrinology, Histology, Economic Zoology, and Parasitology through systematic practical work.

In keeping with the vision of the National Education Policy (NEP 2.0), the syllabus places strong emphasis on experiential and skill-based learning. Accordingly, this manual aims not only to reinforce theoretical concepts but also to nurture essential scientific skills such as observation, analysis, interpretation, and application of knowledge to real-life biological problems.

The content of this practical book is organized in a student-friendly manner, with each exercise presented through clear objectives, step-by-step procedures, suitable diagrams, and expected observations. Students are encouraged to actively participate in practical sessions, maintain accuracy in experimentation, and critically evaluate their results. Practical learning plays a vital role in developing scientific temperament, logical thinking, and problem-solving abilities.

We are confident that this manual will prove to be an invaluable resource for understanding and mastering the fundamentals of Reproductive Biology and Applied Entomology. We wish students a productive, insightful, and enriching learning journey.

Best regards,

- Authors

B. Sc. PART – II SEMESTER – IV (NEP 2.0)

MINOR ZOOLOGY PRACTICAL - IV

(Based on Physiology, Endocrinology, Histology, Economic Zoology, and Parasitology)

PRACTICAL: 60 Hrs. MARKS-50 (CREDITS: 02)

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Unit I: Physiology and Endocrinology

- A. Estimation of Haemoglobin percentage using Sahli's Hemoglobinometer.
- B. Determination of Bleeding time (own or provided blood sample)
- C. Determination of Coagulation time (own or provided blood sample)
- D. Microscopic Examination of Blood Smear
- E. Peak expiratory flow rate
- F. Study of vitamins: Water and fat soluble
- G. Study of Endocrine Glands (Slides/Charts/Models)
 - Pituitary gland
 - Thyroid gland
 - Adrenal gland
 - Pancreas (Islets of Langerhans)

Unit II: Histology

A. Study of following Permanent Slide

- Oesophagus
- Stomach
- Small intestine
- Large intestine
- Liver
- Pancreas

Unit III: Fisheries

A. Study of Economically Important Fishes

(Using preserved specimens/charts/models)

- Rohu (*Labeo rohita*)
- Catla (*Catla catla*)
- Mrigal (*Cirrhinus mrigala*)
- Tilapia (*Oreochromis mossambicus*)

B. Study of Economic importance of followings

(using preserved specimens/charts/models)

- Prawn
- Lobster
- Crab
- Sepia
- Mussel

C. Study of Fish Farm Design (Chart/Model)

- Structure and layout of fish pond:
- Hatchery, nursery, grow-out pond
- Water inlet/outlet, bunds, aerators

D. Fishing Crafts

- Gill Net, Cast Net, Trawl Net, Long Lines, Seine Net, Drift Net, Purse Seines,

Unit IV: Goat Farming and Dairy Science

A. Study of Goat Breeds

Sirohi, Jamunapari, Osmanabadi, Boer, Beetal, Saanen

B. Study dairy products

Milk, Curd, Ghee, Paneer, Cheese, Khoa

Unit V: Parasitology

A. Ascaris

- a. Morphology (Male, Female, Sexual dimorphism)
- b. Life cycle
- c. Parasitic adaptations

Study Tour: visit to any one Sea Shore or National Park, Sanctuary or Zoo to study animal diversity. Submission of report during the practical examination. Duration for study tour may be of 2 to 7 days.

ESTIMATION OF HAEMOGLOBIN PERCENTAGE USING SAHLI'S HEMOGLOBINOMETER

Aim:

To estimate the haemoglobin concentration (percentage) in the given blood sample using Sahli's Hemoglobinometer.

Principle

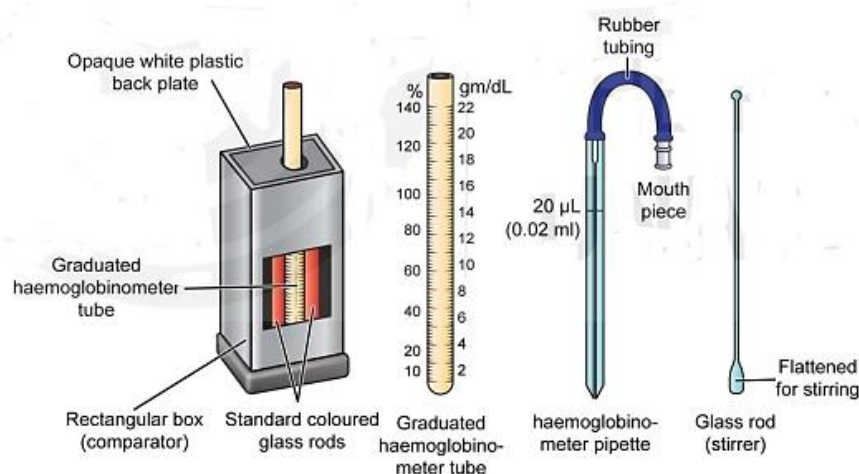
Sahli's method is based on the conversion of haemoglobin into acid haematin by the action of 0.1 N hydrochloric acid (HCl). When blood is added to dilute HCl, haemoglobin reacts to form brown-coloured acid haematin. This colour is then diluted with distilled water until it matches the standard brown glass comparator. The haemoglobin concentration is directly read from the calibrated graduated tube in g/dL (or %).

Requirements

- Sahli's Hemoglobinometer (comparator, graduated tube, pipette)
- Sahli's pipette (20 μ L or 0.02 mL)
- 0.1 N Hydrochloric acid (N/10 HCl)
- Distilled water
- Sterile lancet
- Cotton swab
- Spirit / antiseptic solution
- Dropper
- Tissue paper

Procedure

1. Clean the fingertip with spirit and allow it to dry.
2. Prick the fingertip using a sterile lancet.
3. Draw blood up to the 0.02 mL mark in Sahli's pipette, ensuring no air bubbles.
4. Immediately transfer the blood into the Sahli's graduated tube containing 0.1 N HCl up to the 2 g% mark.
5. Mix gently and allow it to stand for 5–10 minutes for complete formation of acid haematin.
6. Add distilled water drop by drop, mixing after each addition, until the colour matches the standard comparator.
7. Read the haemoglobin value directly from the graduated scale on the tube.
8. Record the observation.



Observation Table

Sr. No.	Volume of Blood Taken (mL)	Colour Matching Level (g/dL or %)
1		
2		
3		

Calculation

The haemoglobin concentration is directly read from the calibrated Sahli's tube.

(No mathematical calculation required)

Result

The haemoglobin percentage of the given blood sample was found to be _____ g/dL (or %).

Precautions

- Ensure accurate filling of the Sahli's pipette without air bubbles.
- Colour matching should be done in **natural daylight**.
- Clean the pipette immediately after use.
- Avoid hemolysis of blood sample.

Normal Values

- Adult Male: **13–18 g/dL**
- Adult Female: **12–16 g/dL**

DETERMINATION OF BLEEDING TIME (OWN OR PROVIDED BLOOD SAMPLE)

Aim

To determine the bleeding time of the given subject by Duke's method.

Principle

Bleeding time is the time interval between the puncture of a blood vessel and the cessation of bleeding. It mainly reflects the platelet function, capillary integrity, and the process of primary haemostasis. In Duke's method, a standardized skin puncture is made and the time taken for bleeding to stop is measured using a stopwatch.

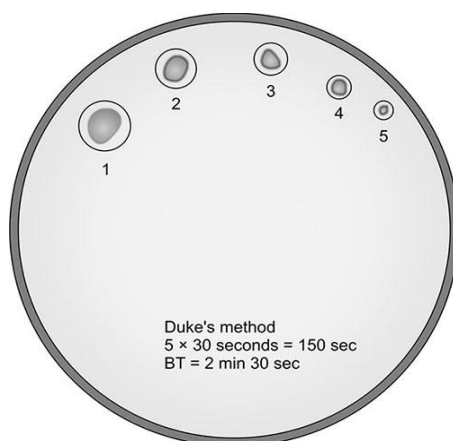
Requirements

- Sterile lancet
- Stopwatch
- Filter paper / blotting paper
- Cotton swab
- Spirit / antiseptic solution
- Gloves

Procedure

1. Clean the fingertip or ear lobe with spirit and allow it to dry.
2. Prick the fingertip or ear lobe with a sterile lancet.
3. Start the stopwatch immediately after puncture.
4. Gently blot the blood at 30-second intervals using filter paper, without squeezing the site.
5. Continue blotting until no blood stain appears on the filter paper.
6. Stop the stopwatch and note the time taken for bleeding to stop.
7. Record the observation.

Observation



Observation Table

Sr. No.	Site of Prick	Time of Puncture	Time of Cessation of Bleeding	Bleeding Time (minutes)
1	Finger / Ear lobe			
2				

Calculation

$$\text{Bleeding Time} = \text{Time of cessation} - \text{Time of puncture}$$

Result

The bleeding time of the given subject was found to be _____ minutes.

Precautions

- Do not squeeze or massage the puncture site.
- Blot gently without touching the wound directly.
- Use a sterile lancet for each subject.
- Maintain proper aseptic conditions.

Normal Values

- Duke's Method: 2–5 minutes

DETERMINATION OF COAGULATION TIME **(OWN OR PROVIDED BLOOD SAMPLE)**

Aim

To determine the coagulation (clotting) time of the given blood sample by the Capillary tube method.

Principle

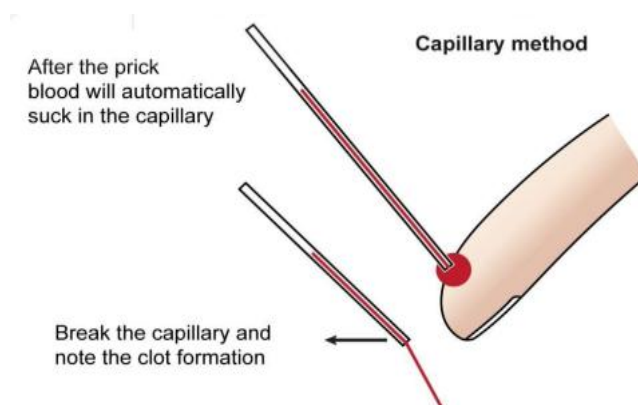
Coagulation time is the time required for blood to clot after it is shed from the body. It reflects the efficiency of the intrinsic pathway of blood coagulation and the availability of clotting factors. In the capillary tube method, freshly drawn blood is collected in a capillary tube and the time taken for the appearance of a fibrin thread is noted as the clotting time.

Requirements

- Sterile lancet
- Capillary tubes
- Stopwatch
- Cotton swab
- Spirit / antiseptic solution
- Gloves

Procedure

1. Clean the fingertip with spirit and allow it to dry.
2. Prick the fingertip using a sterile lancet.
3. Start the stopwatch immediately.
4. Fill **2–3 capillary tubes** with blood.
5. After **1 minute**, break a small piece of the capillary tube every **30 seconds**.
6. Observe for the formation of a **fibrin thread** between the broken ends.
7. Note the time at which the fibrin thread first appears.
8. Record the observation.



Observation Table

Sr. No.	Time of Prick	Time of Fibrin Thread Appearance	Coagulation Time (minutes)
1			
2			
3			

Calculation

Coagulation Time = Time of fibrin thread appearance – Time of prick

Result

The coagulation (clotting) time of the given blood sample was found to be _____ minutes.

Precautions

- Use clean and dry capillary tubes.
- Avoid squeezing the finger excessively.
- Perform the experiment at room temperature.
- Start the stopwatch immediately after pricking.

Normal Values

- Capillary Tube Method: 3–6 minutes

MICROSCOPIC EXAMINATION OF BLOOD SMEAR

Microscopic examination of a blood smear is a fundamental technique in zoology and veterinary science. It allows students to visually identify and differentiate various blood cells, assess their morphology (shape and appearance), and detect abnormalities or parasitic infections. This hands-on experience provides a deeper understanding of hematology beyond numerical counts.

I. Introduction & Principle

- **Blood Smear (Blood Film):** A thin layer of blood spread on a glass microscope slide, which is then dried and stained to make the different blood components visible under a microscope.
- **Purpose:** The primary goals of examining a blood smear are:
 - **Differential White Blood Cell (WBC) Count:** To determine the relative percentages of the different types of leukocytes (neutrophils, lymphocytes, monocytes, eosinophils, basophils). This "manual differential" is crucial as automated analyzers may not always accurately distinguish abnormal cells or provide complete morphological information.
 - **Red Blood Cell (RBC) Morphology:** To evaluate the size, shape, color (hemoglobin content), and inclusions of red blood cells. Abnormalities can indicate various types of anemia or other red blood cell disorders.
 - **Platelet Estimation and Morphology:** To estimate platelet numbers and observe their size and aggregation.
 - **Detection of Blood Parasites:** To identify and quantify blood-borne parasites (e.g., *Plasmodium* species causing malaria, *Trypanosoma*, *Babesia*, microfilariae). This is particularly important in zoology.
 - **Identification of Abnormal Cells:** To detect immature or abnormal blood cells (e.g., blast cells in leukemia, atypical lymphocytes) that automated counters might not flag.
- **Staining Principle (Romanowsky Stains):** Most blood smears are stained using Romanowsky-type stains (e.g., Leishman's, Giemsa, Wright's stain). These stains contain a mixture of methylene blue (a basic dye that stains acidic components, such as DNA/RNA blue/purple) and eosin (an acidic dye that stains basic components, such as hemoglobin and eosinophilic granules red/pink/orange). This differential staining allows for clear visualization and distinction of cellular components.
 - **Fixation:** Before staining, the blood smear is fixed (typically with methanol). This rapidly denatures proteins, "fixes" the cells to the slide, and preserves their morphology, preventing them from washing off during subsequent staining steps.

II. Materials Required

1. Clean, grease-free glass microscope slides: Essential for a good smear.
2. Spreader slide: A clean glass slide with a smooth, unbroken edge.
3. Blood sample
 - Fresh capillary blood: Obtained from a finger prick (human) or suitable site in an animal (e.g., ear vein in rabbit, tail vein in rat, wing vein in bird).
 - Anticoagulated venous blood: Collected in an EDTA (purple top) tube. EDTA is the preferred anticoagulant as it preserves cell morphology well.
4. Sterile lancet or needle: For pricking.
5. Cotton swabs and 70% ethanol/spirit: For antiseptic precautions.
6. Romanowsky stain (e.g., Leishman's stain, Giemsa stain, or Wright's stain):
 - Stain solution.
 - Buffer solution (usually distilled water adjusted to a specific pH, e.g., pH 6.8).
7. Distilled water.
8. Microscope with oil immersion objective (100x).
9. Immersion oil.
10. Waste containers (biohazard, sharps).
11. Gloves.

III. Procedure: Preparation of a Blood Smear

1. **Cleaning Slides:** Ensure the slides are perfectly clean and grease-free. You can wash them with soap and water, rinse thoroughly with distilled water, and dry. Handle slides by their edges to avoid fingerprints.
2. **Obtaining Blood Sample**
 - **Capillary Blood:** Clean the puncture site with alcohol and let it dry. Prick the area with a sterile lancet. Wipe away the first drop of blood.
 - **Venous Blood:** If using EDTA blood, gently invert the tube several times to ensure thorough mixing with the anticoagulant.
3. **Placing the Blood Drop:** Place a **small drop of blood** (about 2-3 mm in diameter) approximately 1-2 cm from one end of a clean slide (the "specimen slide"). *The size of the drop is crucial – too large will result in a thick smear.*
4. **Spreading the Smear (Wedge Method)**
 - Take the spreader slide and place its edge on the specimen slide just in front of the blood drop, holding it at an angle of 30-45 degrees (steeper angle for thicker blood, shallower for thinner blood).
 - Pull the spreader slide backwards until its edge just touches the blood drop. The blood will spread along the entire edge of the spreader slide by capillary action.

- Once the blood has spread, push the spreader slide forward smoothly and rapidly in one continuous motion, maintaining the angle and light pressure, until the blood runs out and the spreader leaves the slide.
 - **Aim for an ideal smear:** It should be about 2/3 to 3/4 the length of the slide, have a smooth, even thickness, and end in a thin, "feathered edge." It should not touch the side edges of the slide.
- 5. Drying the Smear:** Allow the smear to air dry completely by waving it gently in the air. This is crucial for proper staining. Do not blow on it, as this can introduce contaminants or red blood cell artifacts. Protect from dust.
- 6. Labeling:** Once dry, label the frosted end of the slide (if available) with relevant information (e.g., animal ID, date, species).

IV. Procedure: Staining the Blood Smear (Using Leishman's Stain as an example)

1. **Fixation:** Place the dried blood smear on a level staining rack. Flood the slide completely with undiluted Leishman's stain.
 - Let it stand for **1-2 minutes**. This step fixes the cells.
2. **Dilution & Staining:** After the fixation time, add double the volume of distilled water (or buffer solution) to the stain on the slide. Gently mix the stain and water by tilting the slide or blowing gently on the surface until a metallic green scum (sheen) appears.
 - Let this mixture stand for 8-10 minutes. This is the differential staining phase. (Adjust time based on stain batch and desired intensity).
3. **Washing:** Gently pour off the stain mixture. Wash the slide thoroughly but gently with a continuous stream of distilled water (or buffer) until the film appears pinkish-purple and the metallic scum is washed away. Do not let water directly hit the smear as it can dislodge cells.
4. **Drying:** Stand the slide upright in a drying rack or blot it very gently with filter paper (do not blot over the smear itself). Allow it to air dry completely.

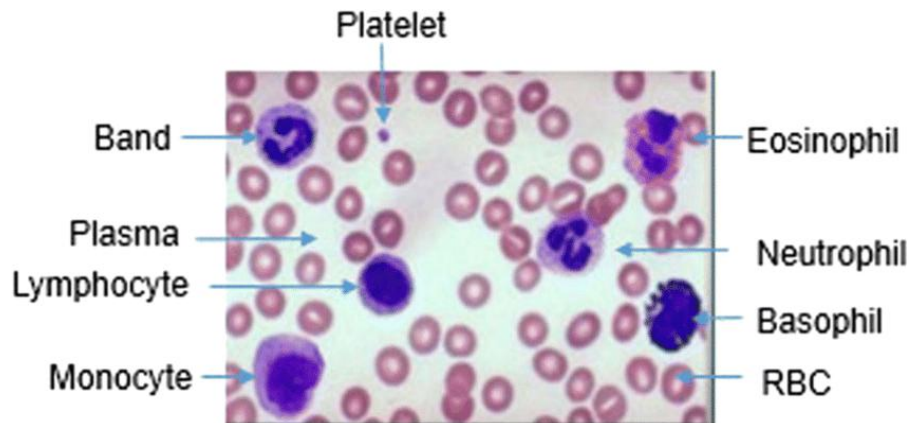
V. Procedure: Microscopic Examination

1. **Initial Scan (Low Power - 10x or 40x)**
 - Place a drop of immersion oil on the feathered edge (thin area) of the dried, stained smear.
 - Begin by scanning the smear under lower magnification (10x or 40x objective) to get an overall impression of the quality of the smear, distribution of cells, and to locate the "monolayer" or optimal viewing area. This is the area where red blood cells are just touching or slightly overlapping, not clumped or widely dispersed.
 - Note any obvious abnormalities, such as large cell clumps, rouleaux formation (RBCs stacking like coins), or agglutination (irregular clumping of RBCs).

2. Oil Immersion Examination (100x Objective)

- Switch to the 100x oil immersion objective. Place a drop of immersion oil directly on the smear over the area you wish to examine. Lower the objective into the oil until it makes contact with the oil.
- Focus carefully. You will now be able to see individual blood cells in detail.
- Systematic Scan: Move the slide systematically across the monolayer area (e.g., in a serpentine or battlement pattern) to examine different fields and ensure a representative sample of cells is observed.

3. Cell Identification and Morphology



- **Red Blood Cells (Erythrocytes)**
 - **Size:** Normal (normocytic), smaller (microcytic), larger (macrocytic).
 - **Color:** Normal (normochromic, with central pallor), paler (hypochromic), darker (hyperchromic, rare).
 - **Shape (Poikilocytosis):** Normal (biconcave disc), or abnormal shapes like sickle cells, target cells, spherocytes, schistocytes (fragments), acanthocytes, echinocytes, etc. (Relevant for diagnosing anemias).
 - **Inclusions:** Howell-Jolly bodies, Heinz bodies (require special stains), basophilic stippling, blood parasites.
 - **Arrangement:** Rouleaux formation, agglutination.
- **White Blood Cells (Leukocytes) - Differential Count**
 - Identify and count at least 100-200 WBCs in the optimal area to determine their relative percentages.
 - **Neutrophils:** Segmented nucleus (2-5 lobes), pale pink/lilac granules. Primary phagocytes.
 - **Lymphocytes:** Large, round nucleus, scanty blue cytoplasm. Involved in adaptive immunity.

- **Monocytes:** Large, kidney-bean-shaped or lobulated nucleus, abundant grayish-blue cytoplasm, often vacuolated. Phagocytic, differentiate into macrophages.
- **Eosinophils:** Bilobed nucleus, prominent red/orange granules. Involved in allergic reactions and parasitic infections.
- **Basophils:** Lobulated nucleus often obscured by large, dark purple/black granules. Involved in allergic reactions, releasing histamine.
- **Note:** Also look for immature forms (bands, myelocytes, blasts) or abnormal cells.
- **Platelets (Thrombocytes)**
 - **Number:** Estimate the approximate number per high-power field (HPF). (e.g., 8-20 platelets/HPF generally indicates adequate numbers, but this varies by species). Look for clumping.
 - **Morphology:** Normal size, large platelets, giant platelets, mega thrombocytes.
- 4. **Reporting Findings:** Record your observations systematically, including percentages of different WBCs, any morphological abnormalities of RBCs, WBCs, or platelets, and the presence of any parasites or other unusual findings.

Microscopic examination of a blood smear is a cornerstone of practical in hematology and provides invaluable visual data that complements quantitative blood tests. It develops keen observational skills crucial for students.

PEAK EXPIRATORY FLOW RATE

Aim

To determine the Peak Expiratory Flow Rate (PEFR) of the given subject using a Peak Flow Meter.

Principle

Peak Expiratory Flow Rate is the maximum rate of air flow achieved during a forceful expiration following a deep inspiration. It reflects the patency of airways and the functional status of the respiratory system, particularly the large airways. PEFR is reduced in obstructive airway disorders such as asthma and chronic bronchitis.

Requirements

- Peak flow meter
- Disposable mouthpiece
- Stopwatch (optional)
- Nose clip (if required)
- Recording sheet

Procedure

1. Instruct the subject to stand upright.
2. Set the pointer of the peak flow meter to zero.
3. Ask the subject to take a deep inspiration.
4. The subject should seal lips tightly around the mouthpiece.
5. Instruct the subject to blow out as hard and as fast as possible in a single breath.
6. Note the reading shown on the peak flow meter.
7. Allow adequate rest and repeat the procedure three times.
8. Record all readings and select the highest value as PEFR.

Observation Table

Sr. No.	Age (Years)	Sex	Height (cm)	Trial 1 (L/min)	Trial 2 (L/min)	Trial 3 (L/min)	Best PEFR (L/min)
1							
2							
3							

Calculation

The highest value obtained among the three trials is taken as the Peak Expiratory Flow Rate.

$$\text{PEFR} = \text{Highest of three readings (L/min)}$$

Result

The Peak Expiratory Flow Rate of the given subject was found to be _____ L/min.

Precautions

- Ensure a tight seal of lips around the mouthpiece.
- The subject should not bend forward while blowing.
- Reset the meter to zero before each trial.
- Use a clean mouthpiece for each subject.

Normal Values (Approximate)

- Adult Male: **400–600 L/min**
- Adult Female: **300–500 L/min**

STUDY OF VITAMINS: WATER AND FAT SOLUBLE

The study of vitamins is a fundamental aspect of nutrition and biochemistry, crucial for students to understand how different animals obtain and utilize these essential organic compounds for growth, development, metabolism, and overall health. Vitamins are broadly classified based on their solubility: water-soluble and fat-soluble. This classification dictates their absorption, transport, storage, excretion, and the likelihood of toxicity.

Study of Vitamins: Water and Fat-Soluble

I. What are Vitamins?

Vitamins are organic compounds required in small quantities in the diet for normal metabolic function and good health. They are generally not synthesized by the body (or synthesized in insufficient amounts) and must be obtained from external sources.

II. Classification of Vitamins

The primary classification is based on their solubility

A. Water-Soluble Vitamins

These vitamins dissolve in water and are generally not stored in the body in significant amounts (with the exception of Vitamin B12). Because they are easily excreted in urine, a regular dietary intake is required to prevent deficiency. The risk of toxicity from excessive intake of water-soluble vitamins is generally low, but some can still cause adverse effects at very high doses.

Members

- Vitamin C (Ascorbic Acid)
- B-Complex Vitamins
 - B1 (Thiamine)
 - B2 (Riboflavin)
 - B3 (Niacin - Nicotinic Acid & Nicotinamide)
 - B5 (Pantothenic Acid)
 - B6 (Pyridoxine, Pyridoxal, Pyridoxamine)
 - B7 (Biotin)
 - B9 (Folate / Folic Acid)
 - B12 (Cobalamin)

General Characteristics

- **Absorption:** Directly absorbed into the bloodstream.
- **Storage:** Minimal storage (except B12 in the liver).
- **Excretion:** Excess amounts are readily excreted in urine.
- **Frequency of Intake:** Needed regularly in the diet.
- **Toxicity:** Generally low risk, but megadoses can cause side effects.

Detailed Study of Water-Soluble Vitamins

Vitamin	Primary Functions	Deficiency Symptoms	Food Sources	Potential Toxicity (at very high doses)
Vitamin C	Antioxidant, collagen synthesis (connective tissue, wound healing), iron absorption, immune function.	Scurvy: Bleeding gums, petechiae (small red spots on skin), poor wound healing, joint pain, fatigue, tooth loss.	Citrus fruits, berries (strawberries), kiwi, bell peppers, broccoli, tomatoes, leafy greens.	Gastrointestinal upset (nausea, diarrhea, cramps), kidney stones (rare), migraines (at 6g/day).
B1 (Thiamine)	Coenzyme in carbohydrate metabolism (energy production), nerve function.	Beriberi: - Wet Beriberi: High-output heart failure, edema. - Dry Beriberi: Polyneuritis, symmetrical muscle wasting, neurological symptoms. Wernicke-Korsakoff Syndrome (in alcoholics): Confusion, ophthalmoplegia, ataxia, memory loss, confabulation.	Whole grains, pork, legumes, nuts, fortified cereals.	None clearly documented from oral intake, readily excreted.
B2 (Riboflavin)	Coenzyme in redox reactions (FAD, FMN) for energy metabolism.	Ariboflavinosis: Cheilosis (cracks at corners of mouth), glossitis (inflamed tongue), sore throat, seborrheic dermatitis, corneal vascularization.	Milk, dairy products, eggs, leafy green vegetables, fortified cereals, lean meats.	None clearly documented from oral intake.

B3 (Niacin)	Coenzyme in redox reactions (NAD ⁺ , NADP ⁺) for energy metabolism; DNA repair.	Pellagra: "3 Ds" - Dermatitis (photosensitive rash), Diarrhea , Dementia . Can also include glossitis, apathy.	Meat (poultry, beef), fish, peanuts, mushrooms, fortified grains. Can also be synthesized from tryptophan.	Niacin Flush: Redness, itching, burning sensation (especially nicotinic acid form). Liver damage, high blood pressure, impaired vision at very high doses (1-3g/day).
B5 (Pantothenic Acid)	Component of Coenzyme A (CoA), essential for fatty acid synthesis and breakdown, energy metabolism, hormone synthesis.	Extremely rare; "Burning feet syndrome" (paresthesia), fatigue, GI distress, restlessness, sleep disturbances, adrenal insufficiency.	Found in virtually all foods (hence "panto" meaning "from everywhere"). Meat, eggs, legumes, whole grains, vegetables.	Gastrointestinal distress (diarrhea).
B6 (Pyridoxine)	Coenzyme (PLP) in amino acid metabolism, neurotransmitter synthesis, red blood cell formation, glycogenolysis.	Microcytic anemia, dermatitis, cheilosis, glossitis, neurological symptoms (depression, confusion, convulsions, peripheral neuropathy).	Meat, fish, poultry, potatoes, bananas, whole grains, fortified cereals.	Severe neurological symptoms (sensory neuropathy), skin lesions, sensitivity to light at high doses (e.g., >1g/day chronically).
B7 (Biotin)	Coenzyme in fatty acid synthesis, gluconeogenesis, amino acid metabolism.	Hair loss (alopecia), scaly skin rash, conjunctivitis, neurological symptoms (depression, lethargy, hallucinations), muscle pain. (Rare, often associated with excessive raw egg white consumption which contains avidin, a biotin binder).	Egg yolks, liver, nuts, seeds, sweet potatoes, some intestinal bacteria.	None clearly documented.

B9 (Folate)	Coenzyme in DNA synthesis and repair, cell division, amino acid metabolism. Crucial for red blood cell maturation.	Megaloblastic anemia: Large, immature red blood cells. Neural tube defects in newborns (if deficient during pregnancy). Fatigue, weakness, irritability, headaches, shortness of breath.	Leafy green vegetables, legumes, citrus fruits, fortified grains, liver.	May mask B12 deficiency (by correcting anemia but not neurological damage); can affect mental function at very high supplemental doses.
B12 (Cobalamin)	Coenzyme in DNA synthesis, red blood cell formation, neurological function (myelin sheath integrity).	Megaloblastic (Pernicious) anemia: (often due to intrinsic factor deficiency). Neurological symptoms (paresthesia, gait disturbance, cognitive impairment, memory loss). Fatigue, weakness, glossitis.	Animal products only: Meat, fish, poultry, eggs, dairy. (Vegans need supplementation or fortified foods).	None clearly documented.

B. Fat-Soluble Vitamins

These vitamins dissolve in fats and oils. They are absorbed along with dietary fats, transported via lymphatic system in chylomicrons, and can be stored in the body's fatty tissues and liver. Because they can accumulate, excessive intake of fat-soluble vitamins carries a higher risk of toxicity (hypervitaminosis) than water-soluble vitamins.

Members

- Vitamin A (Retinol, Retinal, Retinoic Acid)
- Vitamin D (Cholecalciferol - D3, Ergocalciferol - D2)
- Vitamin E (Tocopherols, Tocotrienols)
- Vitamin K (Phylloquinone - K1, Menaquinone - K2)

General Characteristics

- Absorption: Absorbed with dietary fats; requires bile acids.
- Storage: Stored in liver and adipose tissue.
- Excretion: Not easily excreted; can accumulate to toxic levels.
- Frequency of Intake: Not needed daily due to storage.
- Toxicity: Higher risk, especially with supplements.

Detailed Study of Fat-Soluble Vitamins

Vitamin	Primary Functions	Deficiency Symptoms	Food Sources	Potential Toxicity (Hypervitaminosis)
Vitamin A	Vision (component of rhodopsin), cell differentiation, immune function, bone growth, reproduction.	Night blindness (earliest symptom), Xerophthalmia (dry eyes), Bitot's spots (foamy spots on conjunctiva), eventual blindness, impaired immune function, stunted growth.	Preformed A (Retinol): Liver, fish oil, dairy products, eggs. Provitamin A Carotenoids (Beta-carotene): Carrots, sweet potatoes, spinach, kale, mangoes, apricots.	Acute: Nausea, vomiting, headache, blurred vision, dizziness, muscle pain, skin peeling, irritability, increased intracranial pressure. Chronic: Dry skin, hair loss, brittle nails, fatigue, headache, liver damage, bone abnormalities, birth defects (teratogenic in pregnancy). Toxicity more common from supplements than food.
Vitamin D	Calcium and phosphate homeostasis (bone health), immune function, cell growth. Can be synthesized in skin upon UV exposure.	Children: Rickets (soft, deformed bones, bowed legs). Adults: Osteomalacia (softening of bones, bone pain, muscle weakness), Osteoporosis. Impaired immune function.	Fatty fish (salmon, mackerel, tuna), fish liver oils, fortified milk/cereals, egg yolks. Sunlight exposure.	Hypercalcemia (high blood calcium) leading to calcification of soft tissues (kidneys, blood vessels), nausea, vomiting, constipation, polyuria (excessive urination), weakness, kidney damage, heart arrhythmias. More common from excessive supplementation.

Vitamin E	Potent antioxidant, protects cell membranes from oxidative damage.	Very rare in healthy individuals; usually associated with malabsorption or genetic disorders. Symptoms include neurological problems (neuropathy, ataxia), muscle weakness, hemolytic anemia.	Vegetable oils (wheat germ, sunflower, corn, soybean), nuts (almonds, peanuts), seeds, leafy green vegetables, fortified cereals.	Relatively low toxicity, but very high doses can interfere with Vitamin K activity, increasing bleeding risk (especially in individuals on anticoagulants), and can cause GI upset, fatigue, and muscle weakness.
Vitamin K	Essential for blood coagulation (synthesis of clotting factors II, VII, IX, X) and bone metabolism (osteocalcin activation). Can be synthesized by gut bacteria.	Prolonged bleeding/clotting time, easy bruising, hemorrhages (internal or external). (Rare in adults, more common in newborns due to underdeveloped gut flora and low placental transfer, thus prophylactic injection is often given).	K1 (Phylloquinone): Green leafy vegetables (spinach, kale, broccoli), vegetable oils. K2 (Menaquinones): Fermented foods, some animal products, synthesized by gut bacteria.	Generally low toxicity for K1. Synthetic K3 (menadione) can be toxic (hemolytic anemia, liver damage), but it is not used in supplements. No known toxicity for K1 and K2 at high doses in healthy individuals.

III. Practical Application for Students

The study of vitamins involves understanding

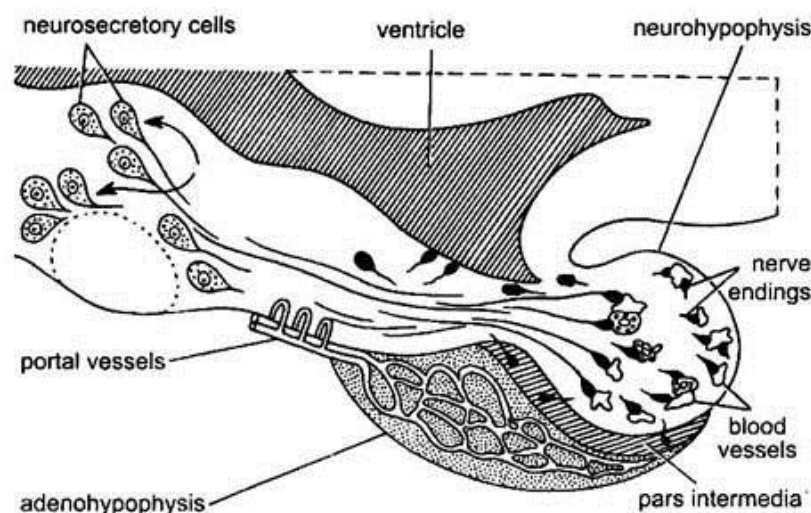
1. **Species-Specific Requirements:** Vitamin needs vary widely across different animal species. For example, most mammals can synthesize Vitamin C, but humans, other primates, guinea pigs, and some birds/fish cannot and thus require it in their diet.
2. **Dietary Sources in Animal Feed:** Recognizing what natural food sources provide specific vitamins for different animals (e.g., pasture for ruminants, insects for insectivores, fortified diets for captive animals).
3. **Deficiency Diseases in Animals:** Identifying clinical signs of vitamin deficiencies in various animal models (e.g., rickets in young animals due to Vitamin D deficiency, neurological issues in poultry with B-vitamin deficiencies, poor vision in zoo animals lacking Vitamin A).
4. **Vitamin Toxicity in Animals:** Understanding that excessive supplementation can lead to hypervitaminosis, which can be particularly problematic for fat-soluble vitamins (e.g., Vitamin A toxicity in polar bears from eating seal liver, Vitamin D toxicity from over-supplementation in companion animals).
5. **Role in Comparative Physiology:** How vitamins act as coenzymes or have specific roles in metabolic pathways that are conserved or vary across different animal groups.
6. **Nutritional Management:** The importance of balanced diets and appropriate supplementation in animal husbandry and conservation efforts to prevent deficiency and toxicity.

This study helps students appreciate the intricate role of micronutrients in animal health, disease, and adaptation.

STUDY OF ENDOCRINE GLANDS (SLIDES/CHARTS/MODELS)

I. PITUITARY GLAND

Often referred to as the "master gland," the pituitary gland is a small, pea-sized endocrine gland located at the base of the brain, housed within a bony depression called the sella turcica of the sphenoid bone. Despite its small size, it plays a critical role in regulating the function of many other endocrine glands and various bodily processes. It is intimately connected to the hypothalamus, which controls its activity.



T. S. OF PITUITARY GLAND

Structure of the Pituitary Gland

The pituitary gland is anatomically and functionally divided into two main lobes:

1. Anterior Pituitary (Adenohypophysis)

- This is the larger, glandular part of the pituitary.
- It develops from an out-pouching of the roof of the embryonic oral cavity (Rathke's pouch).
- It is connected to the hypothalamus by a specialized system of blood vessels called the **hypothalamic-hypophyseal portal system**. Releasing and inhibiting hormones produced by the hypothalamus are transported through this portal system to the anterior pituitary, where they regulate the synthesis and release of anterior pituitary hormones.
- The anterior pituitary contains various types of hormone-secreting cells, typically classified by their staining properties:
 - **Somatotrophs:** Secrete Growth Hormone (GH).
 - **Lactotrophs (Mammotrophs):** Secrete Prolactin (PRL).
 - **Corticotrophs:** Secrete Adrenocorticotrophic Hormone (ACTH).
 - **Thyrotrophs:** Secrete Thyroid-Stimulating Hormone (TSH).

- **Gonadotrophs:** Secrete Follicle-Stimulating Hormone (FSH) and Luteinizing Hormone (LH).

2. Posterior Pituitary (Neurohypophysis)

- This part is essentially an extension of the hypothalamus.
- It is composed mainly of nerve fibers (axons) and specialized glial cells called pituicytes.
- It does **not** synthesize its own hormones. Instead, it stores and releases hormones that are produced by neurosecretory cells in the hypothalamus. These hormones travel down the axons from the hypothalamus to the posterior pituitary.
- It is connected to the hypothalamus by the hypothalamic-hypophyseal tract, a bundle of nerve fibers.

There is also an Intermediate Lobe (Pars Intermedia), which is rudimentary in humans and largely integrated into the anterior lobe. It produces Melanocyte-Stimulating Hormone (MSH) in some species, though its role in humans is not fully understood.

Functions of the Pituitary Gland

The pituitary gland's hormones regulate a vast array of bodily functions:

A. Hormones of the Anterior Pituitary

1. Growth Hormone (GH) or Somatotropin

- **Function:** Promotes growth of bones, muscles, and other tissues, especially during childhood and adolescence. It influences metabolism by promoting protein synthesis, increasing fat breakdown, and raising blood glucose levels (anti-insulin effect).
- **Regulation:** Stimulated by Growth Hormone-Releasing Hormone (GHRH) and inhibited by Somatostatin (Growth Hormone-Inhibiting Hormone, GHIH) from the hypothalamus.

2. Thyroid-Stimulating Hormone (TSH) or Thyrotropin

- **Function:** Stimulates the thyroid gland to produce and secrete thyroid hormones (T3 and T4).
- **Regulation:** Stimulated by Thyrotropin-Releasing Hormone (TRH) from the hypothalamus.

3. Adrenocorticotrophic Hormone (ACTH) or Corticotropin

- **Function:** Stimulates the adrenal cortex to produce and secrete glucocorticoids (primarily cortisol).
- **Regulation:** Stimulated by Corticotropin-Releasing Hormone (CRH) from the hypothalamus.

4. Follicle-Stimulating Hormone (FSH)

- **Function**
 - **Females:** Stimulates the growth and development of ovarian follicles and estrogen production.
 - **Males:** Stimulates sperm production (spermatogenesis) in the testes.
- **Regulation:** Stimulated by Gonadotropin-Releasing Hormone (GnRH) from the hypothalamus.

5. Luteinizing Hormone (LH)

- **Function:**
 - **Females:** Triggers ovulation and stimulates the formation of the corpus luteum (which produces progesterone and estrogen).
 - **Males:** Stimulates testosterone production by the Leydig cells in the testes.
- **Regulation:** Stimulated by Gonadotropin-Releasing Hormone (GnRH) from the hypothalamus.

6. Prolactin (PRL)

- **Function:** Primarily stimulates milk production (lactation) in the mammary glands after childbirth. It also plays a role in reproductive function in both sexes.
- **Regulation:** Primarily inhibited by Dopamine (Prolactin-Inhibiting Hormone, PIH) from the hypothalamus. Stimulated by Prolactin-Releasing Hormone (PRH) (though PRH is not as well-defined as other releasing hormones).

B. Hormones of the Posterior Pituitary (Stored and Released)

1. Antidiuretic Hormone (ADH) or Vasopressin:

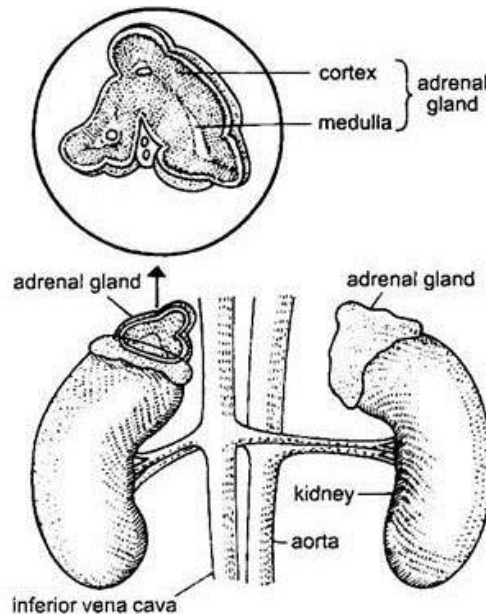
- **Function:** Increases water reabsorption by the kidneys, thereby decreasing urine output and helping to regulate blood volume and blood pressure. It also has vasoconstrictor effects at higher concentrations.
- **Production Site:** Synthesized in the supraoptic nucleus of the hypothalamus.

2. Oxytocin

- **Function**
 - **Females:** Stimulates uterine contractions during childbirth and milk ejection (let-down) during breastfeeding.
 - **Males and Females:** Plays a role in social bonding, trust, and sexual behavior.
- **Production Site:** Synthesized in the paraventricular nucleus of the hypothalamus.

II. ADRENAL GLAND (SUPRARENAL GLAND)

The adrenal glands are a pair of small, triangular-shaped endocrine glands located on top of each kidney. Each adrenal gland is composed of two distinct parts, the outer cortex and the inner medulla, which function as separate endocrine glands, producing different sets of hormones.



ADRENAL GLAND AND ITS LOCATION

Structure of the Adrenal Gland

Each adrenal gland is encapsulated and consists of

1. Adrenal Cortex

- This is the outer, yellowish layer, comprising about 80-90% of the gland's volume.
- It is responsible for synthesizing and secreting steroid hormones (corticosteroids) from cholesterol.
- The cortex is further divided into three distinct zones, each producing specific types of steroid hormones:
 - Zona Glomerulosa (Outer layer): Produces mineralocorticoids, primarily aldosterone.
 - Zona Fasciculata (Middle and largest layer): Produces glucocorticoids, primarily cortisol.
 - Zona Reticularis (Inner layer): Produces weak androgens (sex hormones), such as dehydroepiandrosterone (DHEA).

2. Adrenal Medulla

- This is the inner, reddish-brown core of the adrenal gland.
- It is essentially a modified sympathetic ganglion, derived from neural crest cells.
- It contains chromaffin cells, which are specialized neurosecretory cells.

Functions of the Adrenal Gland

The adrenal glands are vital for stress response, metabolism, and maintaining electrolyte balance.

A. Hormones of the Adrenal Cortex

1. Mineralocorticoids (e.g., Aldosterone - produced by Zona Glomerulosa)

- **Function:** Primarily regulate electrolyte balance (sodium and potassium) and water balance in the body. Aldosterone acts on the kidneys to:
 - Increase sodium reabsorption (leading to water retention).
 - Increase potassium excretion.
 - Increase hydrogen ion excretion (contributing to acid-base balance).
- **Overall Effect:** Helps maintain blood volume and blood pressure.
- **Regulation:** Primarily regulated by the Renin-Angiotensin-Aldosterone System (RAAS), and to a lesser extent, by potassium levels in the blood. ACTH has a minor role.

2. Glucocorticoids (e.g., Cortisol - produced by Zona Fasciculata)

- **Function:** Play a crucial role in the body's response to stress, metabolism, and inflammation.
 - **Metabolic Effects:** Increase blood glucose levels (gluconeogenesis), promote protein breakdown, and mobilize fats for energy.
 - **Anti-inflammatory and Immunosuppressive Effects:** Suppress the immune system and reduce inflammation.
 - **Cardiovascular Effects:** Maintain blood pressure and cardiovascular function.
 - **Stress Response:** Essential for adapting to stress.
- **Regulation:** Regulated by the Hypothalamic-Pituitary-Adrenal (HPA) axis. CRH from the hypothalamus stimulates ACTH release from the pituitary, which then stimulates cortisol secretion from the adrenal cortex. This is a classic negative feedback loop.

3. Adrenal Androgens (e.g., DHEA - produced by Zona Reticularis)

- **Function:** These are weak male sex hormones. In males, their contribution to overall androgen levels is minor compared to testicular androgens. In females, they are a significant source of androgens, contributing to pubic and axillary hair growth, and libido. They can be converted into more potent androgens (like testosterone) or estrogens in peripheral tissues.
- **Regulation:** Primarily regulated by ACTH.

B. Hormones of the Adrenal Medulla

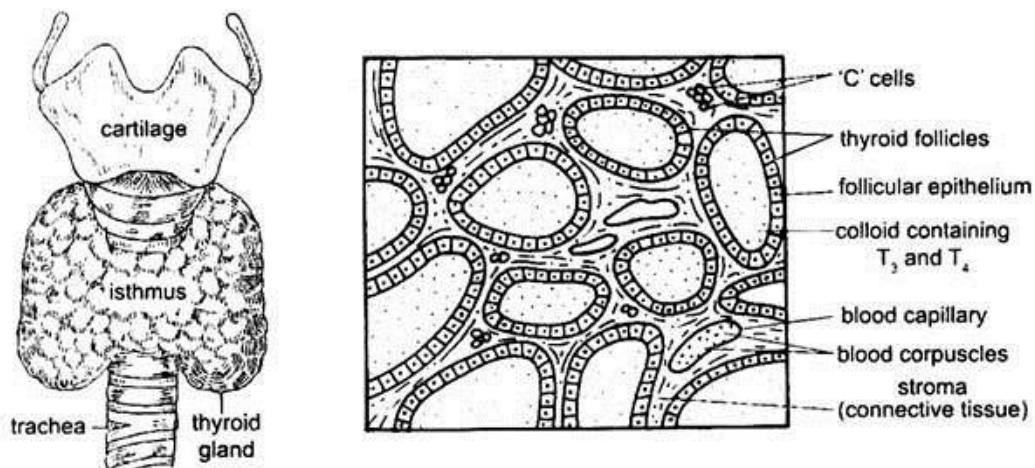
The adrenal medulla secretes **catecholamines**, which are part of the "fight-or-flight" response.

1. Epinephrine (Adrenaline) and Norepinephrine (Noradrenaline):

- **Function:** These hormones prepare the body for immediate action in stressful situations. Their effects include:
 - Increased heart rate and force of contraction.
 - Increased blood pressure.
 - Dilation of bronchioles (improving airflow).
 - Increased blood glucose levels (by promoting glycogenolysis and gluconeogenesis).
 - Redistribution of blood flow (away from digestion and towards muscles and brain).
 - Increased metabolic rate.
- **Regulation:** Secretion is directly stimulated by the sympathetic nervous system in response to stress, fear, excitement, and exercise.

III. THYROID GLAND

The thyroid gland is a butterfly-shaped endocrine gland located in the lower front of the neck, just below the Adam's apple, anterior to the trachea. It is a highly vascular organ and plays a central role in regulating metabolism throughout the body.



HUMAN THYROID GLAND AND T. S. OF THYROID GLAND

Structure of the Thyroid Gland

The thyroid gland consists of

1. **Two Lobes:** Right and left lobes, located on either side of the trachea.
2. **Isthmus:** A narrow band of tissue connecting the two lobes across the front of the trachea, usually at the level of the 2nd and 3rd tracheal rings.

3. **Pyramidal Lobe (Optional):** A small, often present extension of tissue superiorly from the isthmus or one of the lobes.

Microscopic Structure: The thyroid gland's functional units are spherical structures called **thyroid follicles**.

- **Follicular Cells:** These cuboidal epithelial cells form the walls of the follicles. They are responsible for synthesizing and secreting the thyroid hormones, T3 (triiodothyronine) and T4 (thyroxine). They actively transport iodine from the blood into the follicle.
- **Colloid:** The lumen (center) of each follicle is filled with a protein-rich fluid called colloid. The colloid primarily consists of thyroglobulin, a large glycoprotein that acts as a precursor and storage form for thyroid hormones.
- **Parafollicular Cells (C Cells):** These cells are located in the connective tissue between the thyroid follicles. They are distinct from follicular cells and produce the hormone **calcitonin**.

Functions of the Thyroid Gland

The thyroid gland produces hormones that influence nearly every cell in the body.

A. Thyroid Hormones (T3 and T4 - produced by Follicular Cells)

Thyroid hormones are unique because they contain iodine, which is essential for their synthesis.

- **Thyroxine (T4):** The main hormone secreted by the thyroid gland (about 90%). It is considered a prohormone as it is converted to the more active T3 in target tissues.
- **Triiodothyronine (T3):** The more potent and active form of thyroid hormone. About 10% is secreted directly, and the rest is formed from T4 deiodination.

The functions of T3 and T4 are broad and crucial for

1. **Regulation of Basal Metabolic Rate (BMR):** Thyroid hormones increase the BMR, influencing oxygen consumption and heat production in most body tissues. This impacts energy levels and body temperature.
2. **Growth and Development:** Essential for normal growth and maturation, especially of the nervous system and skeletal system, particularly during fetal development and childhood. Deficiency during these periods can lead to irreversible developmental problems (cretinism).
3. **Metabolism of Macronutrients**
 - **Carbohydrate Metabolism:** Increase glucose absorption from the GI tract, enhance glucose utilization, and increase gluconeogenesis and glycogenolysis.
 - **Fat Metabolism:** Increase lipolysis (fat breakdown) and reduce blood cholesterol and triglyceride levels.

- **Protein Metabolism:** Promote protein synthesis (anabolic effect at physiological levels), but can cause protein breakdown at high levels.
- 4. **Nervous System Development and Function:** Crucial for normal brain development and function, affecting alertness, reflexes, and mood.
- 5. **Cardiovascular System:** Increase heart rate, contractility, and cardiac output, making the heart more sensitive to catecholamine.
- 6. **Gastrointestinal System:** Increase gut motility.
- 7. **Reproductive System:** Play a role in maintaining normal reproductive function.

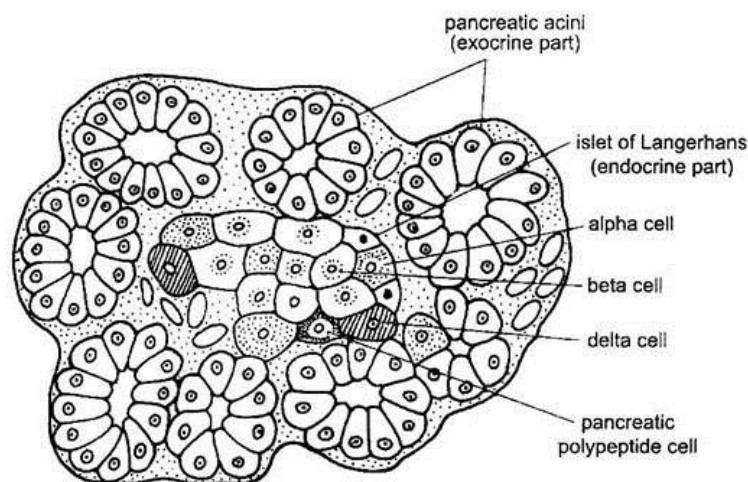
Regulation of Thyroid Hormones: The secretion of thyroid hormones is tightly regulated by the **Hypothalamic-Pituitary-Thyroid (HPT) axis** via a negative feedback mechanism:

- Hypothalamus releases Thyrotropin-Releasing Hormone (TRH).
- TRH stimulates the anterior pituitary to release Thyroid-Stimulating Hormone (TSH).
- TSH stimulates the thyroid gland to synthesize and release T3 and T4.
- High levels of T3 and T4 then inhibit the release of TRH from the hypothalamus and TSH from the anterior pituitary, completing the negative feedback loop.

B. Calcitonin (produced by Parafollicular Cells or C Cells)

1. **Function:** Plays a role in calcium homeostasis.
 - It lowers blood calcium levels by inhibiting osteoclast activity (bone breakdown) and promoting calcium deposition into bones.
 - It also inhibits calcium reabsorption by the kidneys.
2. **Regulation:** Secretion is stimulated by high blood calcium levels. Its role in adult calcium regulation is generally considered less significant than that of parathyroid hormone (produced by the parathyroid glands).

IV. PANCREATIC GLAND



T. S. OF PANCREAS

The pancreas is a unique and vital gland that serves as both an exocrine gland (producing digestive enzymes) and an endocrine gland (producing hormones that regulate blood glucose). It is an elongated, flattened organ located in the abdomen, nestled behind the stomach, with its head in the curve of the duodenum (the first part of the small intestine) and its tail extending towards the spleen.

Structure of the Pancreatic Gland

The pancreas is composed of two main types of tissue, reflecting its dual functions:

1. Exocrine Pancreas (Acinar Cells)

- This constitutes the vast majority (about 99%) of the pancreas.
- It is made up of clusters of cells called **acini** (singular: acinus).
- Acinar cells synthesize and secrete digestive enzymes (like amylase, lipase, proteases) into small ducts, which eventually merge to form the main pancreatic duct. This duct then carries the enzymes into the duodenum to aid in food digestion.

2. Endocrine Pancreas (Islets of Langerhans)

- These are small, spherical clusters of hormone-producing cells scattered throughout the exocrine tissue, accounting for only about 1-2% of the total pancreatic mass. There are typically 1 to 2 million islets in a healthy adult pancreas.
- Although small, the islets are highly vascularized, allowing their hormones to be directly secreted into the bloodstream.
- The islets contain at least four main types of hormone-secreting cells, each with a specific function:
 - **Alpha (α) cells:** Produce Glucagon. (Approximately 15-20% of islet cells)
 - **Beta (β) cells:** Produce Insulin and amylin. (Approximately 65-80% of islet cells, the most abundant)
 - **Delta (δ) cells:** Produce Somatostatin. (Approximately 3-10% of islet cells)
 - **PP (or Gamma, γ) cells:** Produce Pancreatic Polypeptide. (Less than 5% of islet cells)

Functions of the Pancreatic Gland (Endocrine Functions)

The endocrine pancreas primarily focuses on the regulation of blood glucose levels, which is critical for maintaining the body's energy supply.

A. Hormones of the Islets of Langerhans

1. Insulin (produced by Beta cells)

- **Nature:** A peptide hormone.
- **Function:** Insulin is the primary hormone responsible for lowering blood glucose levels when they are high (e.g., after a meal). It achieves this by:

- **Promoting glucose uptake:** Facilitating the entry of glucose from the blood into most body cells (especially muscle and adipose tissue) by increasing the number of glucose transporters on cell membranes.
 - **Promoting glycogen synthesis (glycogenesis):** Stimulating the liver and muscles to convert excess glucose into glycogen for storage.
 - **Promoting fat synthesis (lipogenesis):** Encouraging adipose tissue to convert glucose into triglycerides for storage.
 - **Promoting protein synthesis:** Increasing amino acid uptake and protein synthesis.
 - **Inhibiting gluconeogenesis:** Reducing the liver's production of glucose from non-carbohydrate sources.
 - **Inhibiting glycogenolysis:** Preventing the breakdown of stored glycogen into glucose.
 - **Overall Effect:** Anabolic hormone, promoting the storage of nutrients.
 - **Regulation:** Primarily stimulated by high blood glucose levels, but also by certain amino acids, fatty acids, and parasympathetic nervous system activity. Inhibited by sympathetic nervous system activity and somatostatin.
2. **Glucagon (produced by Alpha cells)**
- **Nature:** A peptide hormone.
 - **Function:** Glucagon is the main hormone responsible for raising blood glucose levels when they are low (e.g., during fasting or intense exercise). It achieves this by:
 - **Promoting glycogenolysis:** Stimulating the liver to break down stored glycogen into glucose and release it into the blood.
 - **Promoting gluconeogenesis:** Stimulating the liver to produce new glucose from non-carbohydrate sources (amino acids, glycerol).
 - **Promoting lipolysis:** Breaking down fats into fatty acids, which can be used as an alternative energy source.
 - **Overall Effect:** Catabolic hormone, mobilizing stored nutrients.
 - **Regulation:** Primarily stimulated by low blood glucose levels, high amino acid levels (e.g., after a high-protein meal, to prevent hypoglycemia from insulin's effect), and sympathetic nervous system activity. Inhibited by high blood glucose levels and insulin.
3. **Somatostatin (GHIH - Growth Hormone-Inhibiting Hormone) (produced by Delta cells)**
- **Nature:** A peptide hormone.

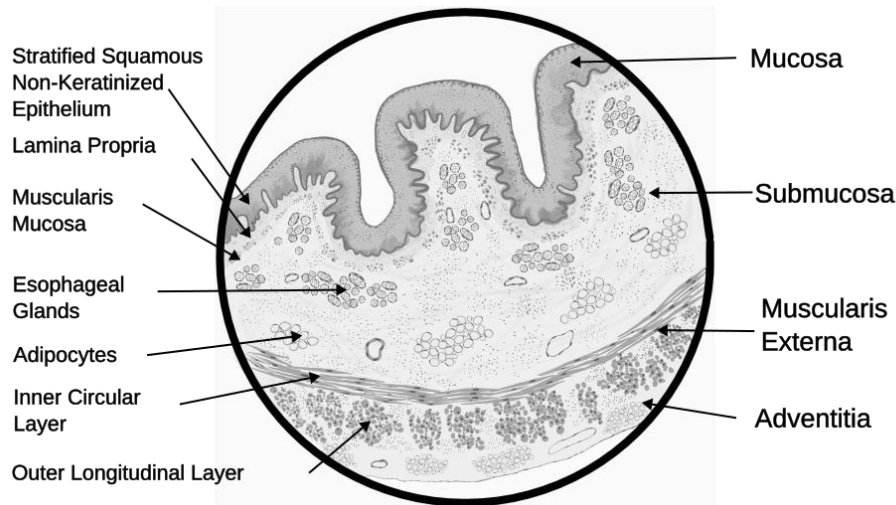
- **Function:** In the pancreas, somatostatin acts as a local paracrine regulator. It helps to modulate the secretion of both insulin and glucagon, primarily by inhibiting their release. This helps to slow down nutrient absorption and prevent rapid fluctuations in blood glucose after a meal. It also inhibits the secretion of pancreatic exocrine enzymes.
 - **Regulation:** Stimulated by high levels of glucose, amino acids, and fatty acids.
4. **Pancreatic Polypeptide (PP) (produced by PP cells)**
- **Nature:** A peptide hormone.
 - **Function:** The exact physiological role of pancreatic polypeptide is not fully understood, but it is believed to be involved in regulating pancreatic exocrine and endocrine secretion, as well as gastrointestinal motility and appetite. Its release is often stimulated by protein-rich meals, fasting, exercise, and acute hypoglycemia.

Importance of Pancreatic Hormones

The antagonistic actions of insulin and glucagon are crucial for maintaining glucose homeostasis. This precise balance ensures that cells have a continuous supply of glucose for energy, while preventing dangerously high (hyperglycemia) or dangerously low (hypoglycemia) blood sugar levels. Disruptions in insulin production or action are the hallmarks of diabetes mellitus, a widespread metabolic disorder.

HISTOLOGY

OESOPHAGUS



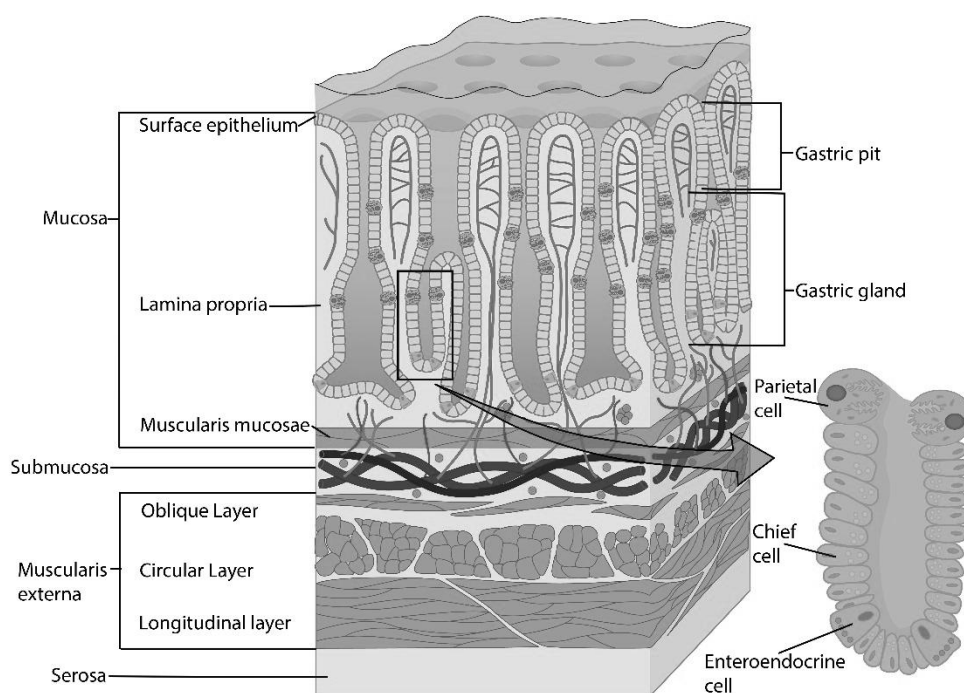
The histology of the esophagus follows the general structural plan of the gastrointestinal tract and consists of four concentric layers: mucosa, submucosa, muscularis externa (propria), and adventitia.

The Four Layers of the Esophageal Wall

- **Mucosa:** This is the innermost lining, which itself has three components:
 1. **Epithelium:** Nonkeratinized, stratified squamous epithelium lines the lumen, providing protection against the abrasive effects of food. This is a key distinguishing feature from the simple columnar epithelium of the stomach, with an abrupt transition occurring at the gastroesophageal junction.
 2. **Lamina propria:** A thin layer of loose connective tissue beneath the epithelium, containing blood vessels and occasional lymphoid aggregates.
 3. **Muscularis mucosae:** A thin layer of smooth muscle, which becomes thicker in the lower esophagus.
- **Submucosa:** This layer of dense, irregular connective tissue contains blood vessels, lymphatic vessels, nerves (Meissner plexus), and mucus-secreting submucosal glands (esophageal glands proper). These glands help lubricate the passage of food.
- **Muscularis externa (propria):** The main muscle layer responsible for peristalsis, consisting of an inner circular layer and an outer longitudinal layer. The composition of this layer changes along the length of the esophagus:
 1. **Upper third:** Primarily voluntary striated (skeletal) muscle.
 2. **Middle third:** A mixture of both striated and smooth muscle.
 3. **Lower third:** Entirely smooth muscle.

4. The myenteric (Auerbach) plexus, a network of nerve fibers involved in regulating muscle contraction, is located between the inner and outer muscle layers.
- **Adventitia:** For most of its length in the thorax, the esophagus is covered by an outer layer of loose connective tissue called the adventitia, which anchors it to surrounding structures. The short portion within the abdominal cavity is covered by a serosa (visceral peritoneum).

STOMACH



The histology of the stomach follows the typical four-layered organization of the gastrointestinal tract, comprising the mucosa, submucosa, muscularis externa, and serosa. The mucosa is lined by simple columnar epithelium, which forms invaginations known as gastric pits that open into specialized gastric glands.

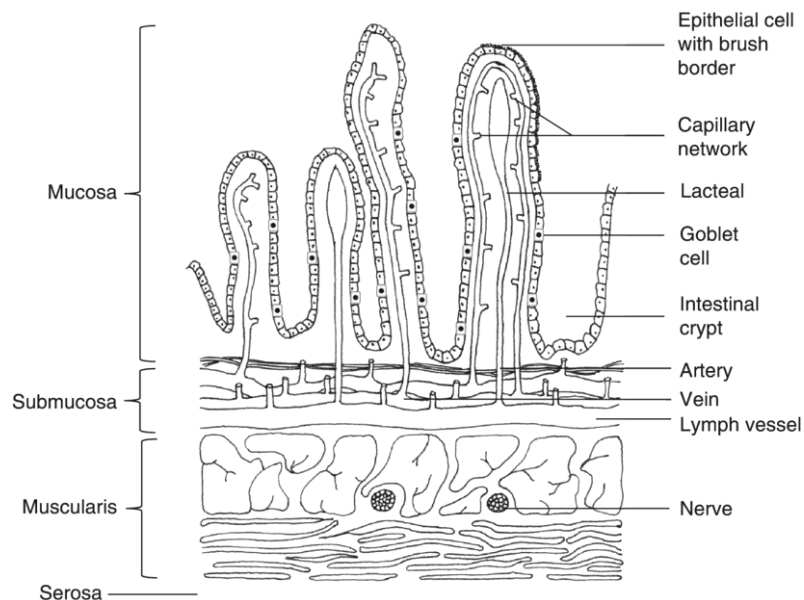
Layers of the Stomach Wall

The stomach wall features a unique adaptation in its muscular layer to facilitate mechanical digestion and churning.

- **Mucosa:** The innermost layer, consisting of a surface epithelium, an underlying connective tissue layer (lamina propria), and a thin smooth muscle layer (muscularis mucosae).
- **Submucosa:** A layer of dense connective tissue containing blood vessels, lymphatic vessels, and the submucosal (Meissner's) nerve plexus.

- **Muscularis Externa (Muscularis Propria):** This layer is unique to the stomach as it has three layers of smooth muscle instead of the typical two found in the rest of the GI tract.
 1. **Inner oblique layer** (unique to the stomach; responsible for churning)
 2. **Middle circular layer** (thickens to form the pyloric sphincter)
 3. **Outer longitudinal layer** -The myenteric (Auerbach's) nerve plexus is located between the circular and longitudinal muscle layers and controls muscle contractions.
- **Serosa:** The outermost layer, a serous membrane (visceral peritoneum) that provides protection and lubrication.

SMALL INTESTINE



The histology of the small intestine includes a layered structure (mucosa, submucosa, muscularis, serosa) and features like circular folds, villi, and microvilli to maximize surface area for absorption. The mucosa's epithelium contains enterocytes (absorptive cells), goblet cells, enteroendocrine cells, and Paneth cells, which reside in the crypts of Lieberkühn alongside stem cells. The three sections—duodenum, jejunum, and ileum—have specific characteristics, such as Brunner's glands in the duodenum and Peyer's patches in the ileum.

Layers

Mucosa: The innermost layer, featuring the epithelium, lamina propria, and muscularis mucosae.

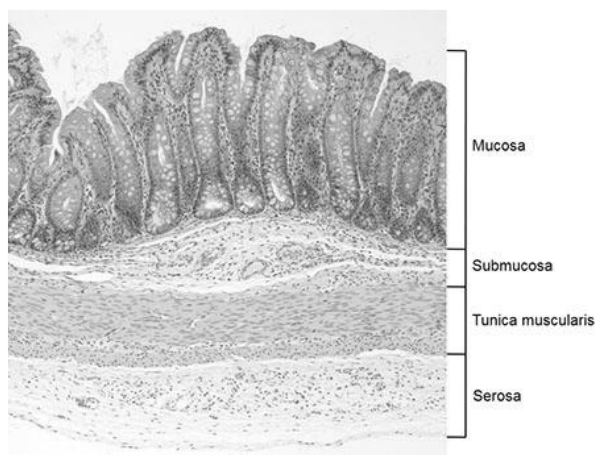
1. **Epithelium:** Lined with simple columnar epithelium that includes absorptive enterocytes, goblet cells, enteroendocrine cells, and Paneth cells.
2. **Lamina Propria:** The core of the villi, containing blood capillaries and lacteals (lymphatic vessels).

Submucosa: A layer of connective tissue containing blood vessels, nerves, and lymphatic vessels.

Muscularis Externa: Composed of an inner circular and an outer longitudinal layer of smooth muscle responsible for peristalsis.

Serosa: The outermost layer of connective tissue that allows the intestine to move freely.

LARGE INTESTINE



The histology of the large intestine follows the typical structural organization of the gastrointestinal tract and consists of four layers: mucosa, submucosa, muscularis externa, and serosa or adventitia. Distinctive features include the absence of villi, the presence of deep intestinal glands (crypts of Lieberkühn) rich in goblet cells, and the specialization of the outer longitudinal muscle layer into three distinct bands known as taeniae coli.

Layers of the Large Intestine

Mucosa: The innermost layer, responsible for absorption of water and electrolytes, and mucus secretion.

1. **Epithelium:** Lined by simple columnar epithelium, containing enterocytes (absorptive cells with short microvilli) and a high density of mucus-secreting goblet cells.
2. **Lamina Propria:** Loose connective tissue that supports the epithelium and contains numerous immune cells (lymphocytes, plasma cells, macrophages) and lymphoid nodules (especially abundant in the appendix).
3. **Muscularis Mucosae:** A thin layer of smooth muscle that separates the mucosa from the submucosa.

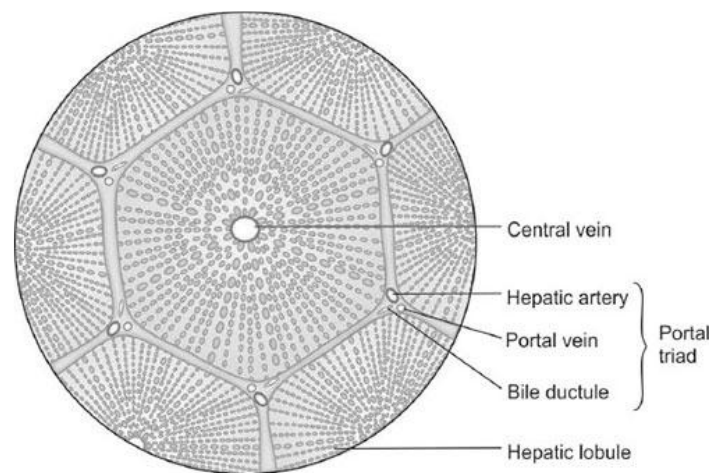
Submucosa: Consists of dense irregular connective tissue, blood and lymphatic vessels, and the submucosal nerve plexus (Meissner's plexus).

Muscularis Externa: Composed of two smooth muscle layers, important for motility (peristalsis and haustration).

1. **Inner Circular Layer:** A continuous layer of smooth muscle.
2. **Outer Longitudinal Layer:** This layer is concentrated into the three narrow, longitudinal bands known as the taeniae coli. These bands are shorter than the large intestine itself, causing it to form characteristic pouches called haustra.
3. The myenteric nerve plexus (Auerbach's plexus) is located between these two muscle layers and regulates muscle contraction.

Serosa/Adventitia: The outermost layer. Intraperitoneal parts (cecum, transverse colon, sigmoid colon) are covered by serosa, while retroperitoneal parts (ascending and descending colon, rectum) are covered by adventitia, which anchors them to surrounding structures. Small, fat-filled pouches called appendices epiploicae project from the serosal surface.

LIVER



The liver's histology is centered around a repeating unit known as the hepatic lobules, a roughly hexagonal structure organized around a central vein. Hepatocytes (liver cells) are arranged in plates that radiate from the central vein, separated by vascular channels called sinusoids. At the corners of the hexagon are portal tracts (or portal triads), which contain three key structures: a branch of the hepatic artery, a branch of the portal vein, and a bile ductule.

Structural Components

- **Hepatocytes:** These are the primary parenchymal cells of the liver, large and polygonal with abundant eosinophilic (pink) cytoplasm and round, central nuclei. They perform most of the liver's metabolic functions, including bile production, protein synthesis, and detoxification.
- **Hepatic Lobule:** The classic structural unit. Blood flows from the portal triad (periphery) through the sinusoids to the central vein.

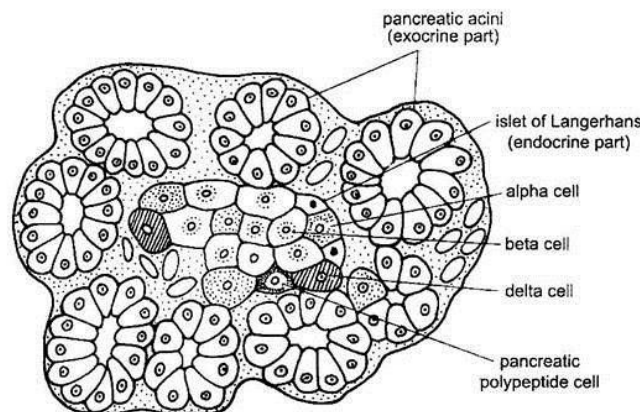
- **Hepatic Acinus:** Considered the functional unit, it emphasizes the gradient of blood flow and oxygenation, divided into three zones:
 1. **Zone 1 (Periportal):** Closest to the arterial and portal blood supply, receiving the most oxygen and nutrients; involved in oxidative metabolism and gluconeogenesis.
 2. **Zone 3 (Pericentral/Centrilobular):** Furthest from the blood supply, most susceptible to ischemic injury and involved in detoxification and lipogenesis.
 3. **Zone 2 (Midzonal):** The area between zones 1 and 3.
- **Portal Triad:** Located at the "corners" of the lobule, containing:

Portal Venule: A large lumen with a thin wall, carrying nutrient-rich, deoxygenated blood from the digestive tract.

Hepatic Arteriole: A smaller lumen with a thick muscular wall, carrying oxygenated blood.

Bile Ductule: Lined by a single layer of cuboidal epithelium; collects bile from the bile canaliculi.

PANCREAS



Structure of the Pancreatic Gland

The pancreas is composed of two main types of tissue, reflecting its dual functions:

1. Exocrine Pancreas (Acinar Cells)

- This constitutes the vast majority (about 99%) of the pancreas.
- It is made up of clusters of cells called **acini** (singular: acinus).
- Acinar cells synthesize and secrete digestive enzymes (like amylase, lipase, proteases) into small ducts, which eventually merge to form the main pancreatic duct. This duct then carries the enzymes into the duodenum to aid in food digestion.

2. Endocrine Pancreas (Islets of Langerhans)

- These are small, spherical clusters of hormone-producing cells scattered throughout the exocrine tissue, accounting for only about 1-2% of the total pancreatic mass. There are typically 1 to 2 million islets in a healthy adult pancreas.

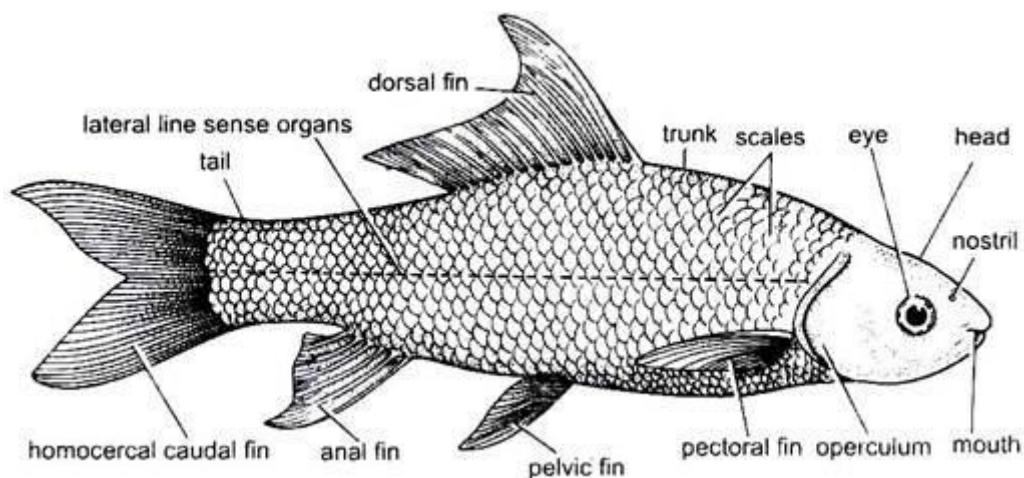
- Although small, the islets are highly vascularized, allowing their hormones to be directly secreted into the bloodstream.
- The islets contain at least four main types of hormone-secreting cells, each with a specific function:
- **Alpha (α) cells:** Produce **Glucagon**. (Approximately 15-20% of islet cells)
- **Beta (β) cells:** Produce **Insulin** and amylin. (Approximately 65-80% of islet cells, the most abundant)
- **Delta (δ) cells:** Produce **Somatostatin**. (Approximately 3-10% of islet cells)
- **PP (or Gamma, γ) cells:** Produce **Pancreatic Polypeptide**. (Less than 5% of islet cells)

Functions of the Pancreatic Gland (Endocrine Functions)

The endocrine pancreas primarily focuses on the regulation of blood glucose levels, which is critical for maintaining the body's energy supply.

STUDY OF ECONOMICALLY IMPORTANT FISHES

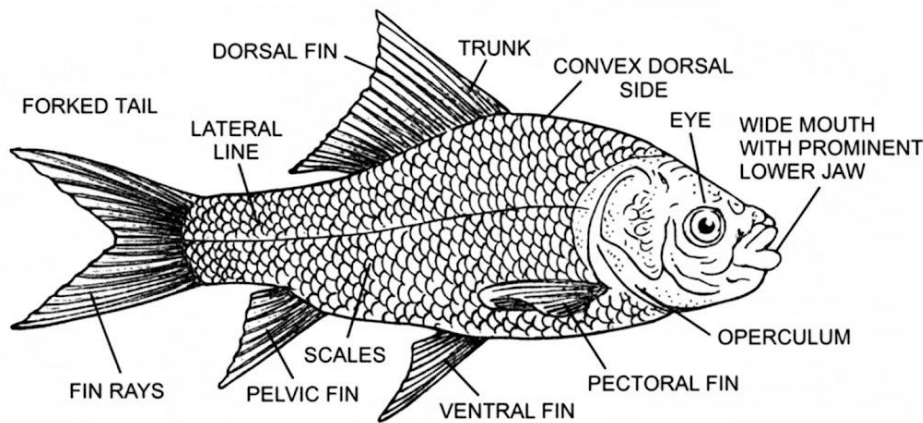
1. ROHU (*Labeo rohita*)



LABEO

- **Body Shape:** Typically, fusiform and elongated, moderately compressed laterally. It has a streamlined body suitable for swimming in currents.
- **Coloration:** Generally silvery-grey on the dorsal side, fading to silvery-white on the flanks and belly. Fins are often dusky.
- **Head:** Relatively small head.
 - **Snout:** Depressed and blunt, often projecting slightly beyond the jaws.
 - **Mouth:** Sub-terminal, crescentic slit, with thick, fleshy, and often fringed lips. This indicates its bottom-feeding (illithophagous) habit, grazing on detritus and algae.
 - **Barbels:** Usually one pair of small, delicate maxillary barbels (at the corner of the mouth). Rostral barbels are typically absent or very minute.
- **Scales:** Large, distinct, overlapping cycloid scales covering the trunk and tail. A complete lateral line with 40-43 scales.
- **Fins:**
 - **Dorsal fin:** Large, originating roughly over the middle of the body, with 13-15 soft rays.
 - **Caudal fin:** Deeply forked with two equal lobes.
 - Pectoral, pelvic, and anal fins are also present, with specific ray counts.
- **Distinguishing feature:** The combination of a sub-terminal mouth with fringed lips and usually one pair of small maxillary barbels is characteristic.

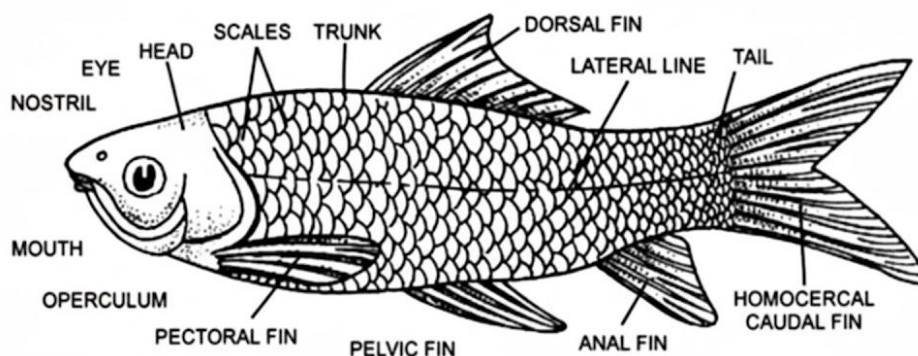
2. CATLA (*Catla catla*)



CATLA

- **Body Shape:** Short and very deep-bodied, significantly compressed laterally. Its depth is greater than its head length. This body shape is adapted for surface and mid-water feeding.
- **Coloration:** Greyish on the back and flanks, transitioning to silvery-white below. Fins are often dusky.
- **Head:** Very large head, conspicuously large, often making up more than one-third of the total length. The head is scaleless.
 - **Snout:** Broad and bluntly rounded.
 - **Mouth:** Wide and distinctly upturned (superior mouth). The lower jaw protrudes. This morphology is perfectly adapted for surface feeding on zooplankton and floating vegetation.
 - **Barbels:** Absent.
- **Scales:** Conspicuously large cycloid scales on the body. A complete lateral line with 40-43 scales.
- **Fins:**
 - **Dorsal fin:** High, inserted slightly in advance of the pelvic fins, with 14-16 branched rays.
 - **Caudal fin:** Deeply forked.
 - Pectoral fins are long, often extending to the pelvic fins.
- **Distinguishing feature:** Its very large head, deep body, and characteristic upturned, wide mouth with no barbels make it easily identifiable.

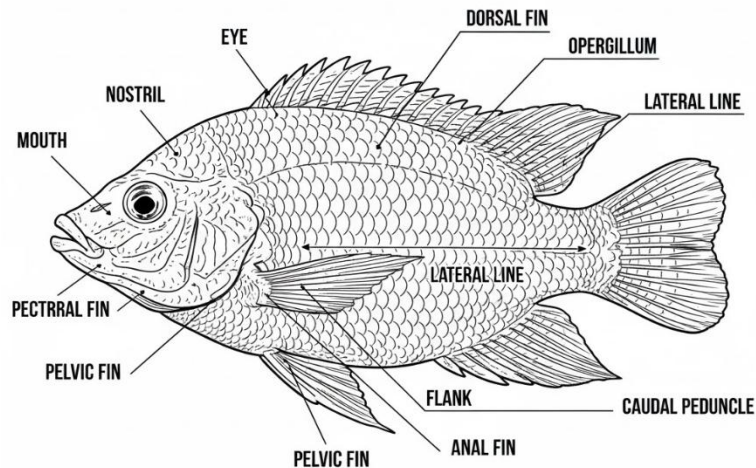
3. MRIGAL (*Cirrhinus mrigala*)



MRIGAL

- **Body Shape:** Streamlined and elongated, less deep-bodied than Catla, but more slender than Rohu. Its body is somewhat cylindrical, well-suited for bottom dwelling and feeding.
- **Coloration:** Coppery-silvery body, often with a golden tinge on the flanks. Eyes can be golden.
- **Head:** Relatively small head.
 - **Snout:** Blunt, often with pores.
 - **Mouth:** Sub-terminal, broad, and transverse. The upper lip is entire, and the lower lip is often indistinct or less prominent than Rohu. It's adapted for bottom feeding on detritus.
 - **Barbels:** Usually a single pair of short rostral barbels (at the tip of the snout, near the nostrils). These are often more prominent than Rohu's maxillary barbels.
- **Scales:** Cycloid scales, though not as large as Catla. A complete lateral line with 40-45 scales.
- **Fins:**
 - **Dorsal fin:** Originates nearer to the end of the snout than the caudal fin base, with 12 or 13 branched rays.
 - **Caudal fin:** Deeply forked.
- **Distinguishing feature:** The streamlined body, sub-terminal mouth, and presence of only one pair of distinct rostral barbels help differentiate it from Rohu and Catla.

4. TILAPIA (*Oreochromis mossambicus*)



TILAPIA

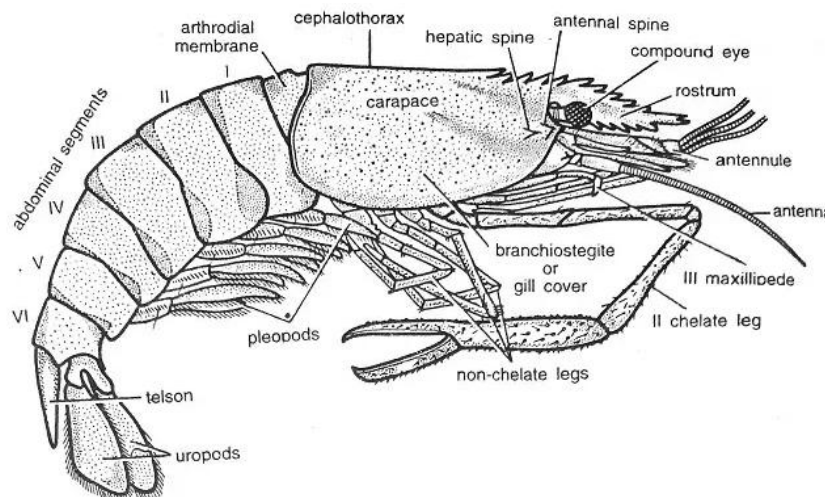
- **Body Shape:** Deep-bodied and laterally compressed, often robust. Body depth can be quite variable depending on environmental conditions.
- **Coloration:** Highly variable. "Pure form" females and non-breeding males are silvery-grey to pale olive with 2-5 mid-lateral blotches. Breeding males become much darker, often dark olive to black.
- **Head**
 - **Snout:** Rounded in females and non-breeding males. Sexually mature males often develop an enlarged, concave upper profile of the head and robust, protruding jaws, sometimes described as "duckbill-like."
 - **Mouth:** Terminal and wide, with thick lips.
 - **Barbels:** Absent.
- **Scales:** Large, cycloid scales.
 - **Lateral line:** Interrupted lateral line (a key characteristic of Cichlids). It typically consists of two parts: an upper section running along the dorsal profile and a lower section running along the mid-body. Scales count on the lateral line series is 30-32.
- **Fins**
 - **Dorsal fin:** Long and continuous, extending over most of the back, with both spiny and soft rays. The posterior part of the dorsal and anal fins often has elongated, pointed extensions, especially in males.
 - **Caudal fin:** Rounded or truncate (not typically forked like carps), often with a red margin in breeding males.
 - Anal fin has three spines.
- **Distinguishing feature:** The combination of an interrupted lateral line, a long dorsal fin with spiny and soft rays, and often a rounded caudal fin, along with the distinct color patterns and head shape changes in breeding males, are characteristic of Tilapia.

STUDY OF ECONOMIC IMPORTANCE OF FOLLOWINGS

1. PRAWN (e.g., *Penaeus monodon* - Giant Tiger Prawn,

Macrobrachium rosenbergii - Giant Freshwater Prawn)

Prawns are decapods (10 legs) belonging to the order Decapoda, suborder Dendrobranchiata. They are distinct from shrimp, often being larger and having branching gills.



Morphological Characters to Observe

- **Body:** Elongated, cylindrical, segmented. Divided into a cephalothorax (fused head and thorax covered by a carapace) and an abdomen (tail).
- **Rostrum:** A prominent, serrated spine projecting forward from the carapace. The number and arrangement of dorsal and ventral teeth on the rostrum are crucial for species identification.
- **Antennules & Antennae:** Two pairs of sensory appendages on the head.
- **Eyes:** Stalked compound eyes.
- **Pereiopods (Walking Legs):** Five pairs on the cephalothorax. The first three pairs often end in small pincers (chelae), used for feeding.
- **Pleopods (Swimmerets):** Five pairs of paddle-like appendages on the abdomen, used for swimming, brood care (in some species), and distinguishing sexes (e.g., petasma in males, thelycum in females for *Penaeus*).
- **Uropods & Telson:** The last abdominal segment bears a central **telson** and two pairs of lateral **uropods**, forming a tail fan used for rapid backward escape (tail flip).
- **Gills:** Branching gills (dendrobranchiate) are a key internal feature distinguishing them from true shrimp.

Economic Importance

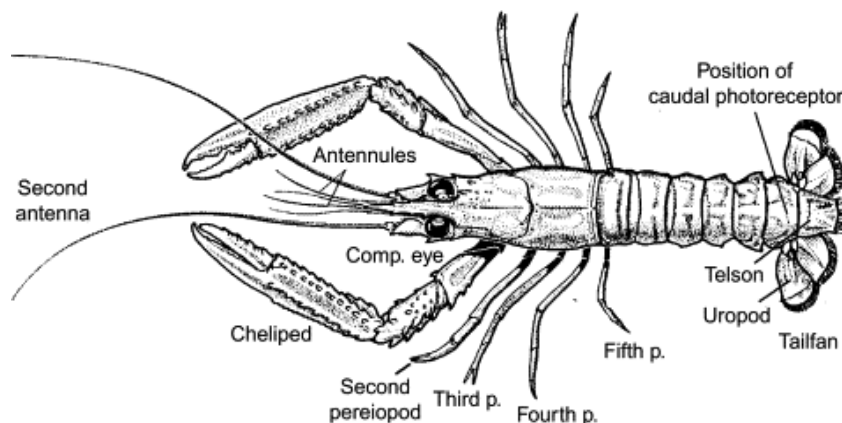
- **Primary Food Source:** Prawns are a highly prized seafood globally, consumed fresh, frozen, cooked, or processed. Their sweet, firm flesh is in high demand.

- **Aquaculture:** Globally, prawn farming is a massive industry. Species like *Penaeus monodon* and *Litopenaeus vannamei* (Whiteleg Shrimp) are extensively farmed in brackish water, while *Macrobrachium rosenbergii* is farmed in freshwater. This provides livelihoods and food security.
- **Wild Capture Fisheries:** Significant wild catches occur, though many stocks are under pressure due to overfishing.
- **Export Commodity:** Major export earner for many coastal developing nations.
- **Bait:** Smaller prawns can be used as live or dead bait for other fish.

2. LOBSTER (e.g., *Nephrops norvegicus* - Norway Lobster

Homarus americanus - American Lobster)

True lobsters are large marine decapods known for their large, asymmetric claws. Spiny lobsters (Palinuridae) lack large claws but have prominent antennae.



Morphological Characters to Observe

- **Body:** Robust, elongated, cylindrical, heavily armored with a thick exoskeleton. Also divided into a cephalothorax (covered by a large, rigid carapace) and a muscular abdomen (tail).
- **Claws (Chelae):** A defining feature. Usually two large, unequal claws on the first pair of pereiopods
 - **Crusher claw:** Larger, more robust, with blunt teeth for crushing.
 - **Pinch/Cutter claw:** More slender with sharper teeth for cutting and tearing. (Note: Spiny lobsters lack these large claws).
- **Antennae:** Long and thick, used for sensing.
- **Rostrum:** Short, sharp spine on the carapace, less prominent than in prawns.
- **Pereiopods (Walking Legs):** Five pairs. The first pair forms the large claws; the next two pairs have small pincers; the last two pairs are simple walking legs.
- **Pleopods:** Smaller than those of prawns, located on the underside of the abdomen.
- **Tail Fan:** Consisting of a central telson and broad uropods, forming a powerful paddle for rapid backward escape.

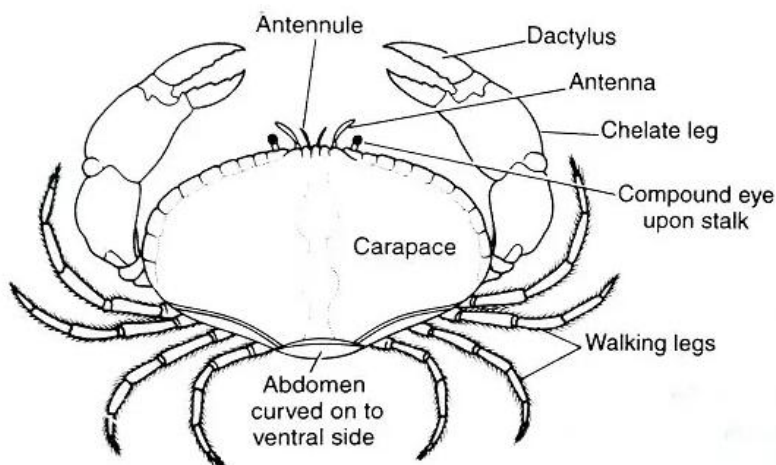
- **Coloration:** Varies by species and habitat (e.g., blue, green, brown, often turning bright red when cooked).

Economic Importance

- **Luxury Seafood:** Lobsters are considered a premium and highly desirable seafood item globally, commanding high prices.
- **Wild Fisheries:** Primarily sourced from wild capture fisheries, often using traps (pots). These fisheries are economically vital to many coastal communities.
- **Tourism:** "Lobster tours" and culinary tourism around lobster are significant in some regions (e.g., New England, Canada).
- **Limited Aquaculture:** While research is ongoing, large-scale commercial aquaculture of true lobsters is challenging due to slow growth rates, cannibalism, and complex life cycles. Spiny lobster aquaculture is developing in some areas.
- **Export Value:** Major export commodity for countries with significant lobster populations.

3. CRAB (e.g., *Portunus pelagicus* - Blue Swimmer Crab, *Scylla serrata* - Mud Crab)

Crabs are decapods characterized by a short, broad body and a reduced abdomen tucked underneath the cephalothorax.



Morphological Characters to Observe

- **Body:** Primarily consists of a broad, flattened, and often spiny carapace (cephalothorax). The abdomen is small and folded tightly underneath the carapace (visible as a flap on the underside).
- **Claws (Chelae):** The first pair of pereopods forms prominent, powerful claws, used for defense, feeding, and courtship. Size and shape vary greatly by species and sex.
- **Pereopods (Walking Legs):** Typically four pairs of walking legs (besides the claws), adapted for walking, swimming (in swimming crabs, the last pair are flattened paddles), or climbing.

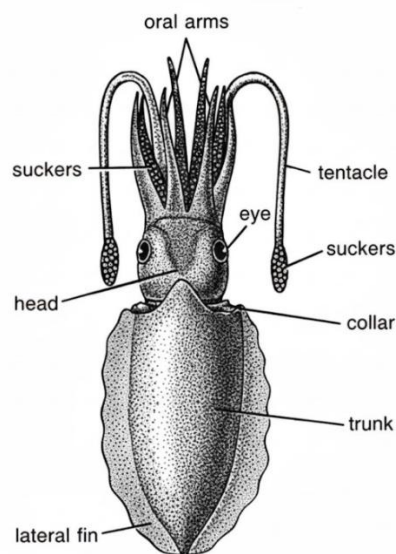
- **Eyes:** Stalked eyes, often retractable.
- **Mouthparts:** Complex mouthparts hidden under the front of the carapace.
- **Gill Slits:** Located under the lateral margins of the carapace.
- **Sexual Dimorphism:** The shape of the abdominal flap is key for sex identification:
 - **Male:** Narrow, inverted 'T' or 'V' shape.
 - **Female:** Broad, rounded, 'U' shape (to carry eggs).
- **Coloration:** Extremely varied (e.g., blue, green, brown, red), often cryptic to blend with the environment.

Economic Importance

- **Major Food Source:** Crabs are a highly popular and valuable seafood, consumed fresh, frozen, canned, or as crab meat.
- **Fisheries:** Extensive wild capture fisheries exist worldwide, using pots, nets, or lines.
- **Aquaculture (Crab Fattening/Grow-out):** While full life-cycle aquaculture is complex for many species, "crab fattening" (holding wild-caught, lean crabs in ponds until they fill with meat) and grow-out of juvenile crabs are significant. Mud crabs (*Scylla serrata*) are a prime example.
- **By-products:** Crab shells can be processed for chitin and chitosan, valuable in industries like pharmaceuticals, cosmetics, and water treatment.
- **Tourism:** Crab festivals and crab-fishing tours are popular.

4. SEPIA (Cuttlefish - e.g., *Sepia pharaonis*)

Cuttlefish are cephalopods, related to octopuses and squids, characterized by an internal shell called a cuttlebone.



Morphological Characters to Observe

- **Body:** Oval, flattened, and robust, with a distinctive fringed fin (lateral fin) running along most of the body margin.

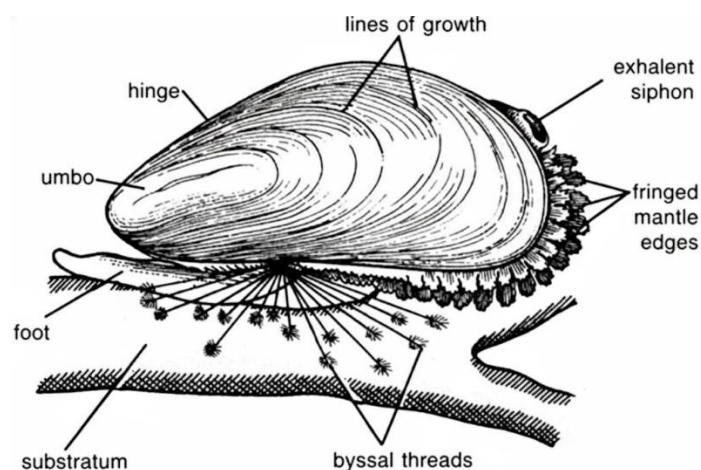
- **Head:** Large, with prominent, W-shaped pupils.
- **Arms & Tentacles:** **Eight arms** with suckers, arranged around the mouth, and **two long, retractable tentacles** with suckered clubs at their ends (used for catching prey). The tentacles are usually hidden in pouches.
- **Mantle:** The main body sac, containing the internal organs.
- **Siphon/Funnel:** A muscular tube under the head used for jet propulsion (expelling water).
- **Cuttlebone:** An internal, porous, calcified shell (phragmocone) running along the dorsal side of the mantle. This is crucial for buoyancy control and is often preserved separately. Its size and shape are species-specific.
- **Chromatophores:** Specialized pigment cells in the skin allowing for rapid and complex color changes for camouflage, communication, and display.
- **Ink Sac:** Produces dark ink for defense, expelled through the siphon.

Economic Importance

- **Food Source:** Cuttlefish are a popular seafood, especially in Mediterranean, Asian, and Southeast Asian cuisines. They are consumed fresh, frozen, dried, or as calamari (though squid is more common for this). Their firm, white meat is highly valued.
- **Fisheries:** Caught by trawling, jigging, and traditional methods.
- **Cuttlebone:** The internal shell (cuttlebone) is highly valued:
 - **Pet Industry:** Used as a calcium supplement and beak conditioner for pet birds.
 - **Polishing Agent:** Historically used as a polishing agent for metals.
 - **Art:** Used for molds in jewelry casting.
- **Ornamental:** Some smaller species are kept in specialized aquariums.
- **Ink:** Historically used as a pigment (sepia ink in art).

5. MUSSEL (e.g., *Mytilus edulis* - Blue Mussel, *Perna viridis* - Green Mussel)

Mussels are bivalve molluscs, meaning their soft body is enclosed within two hinged shells (valves).



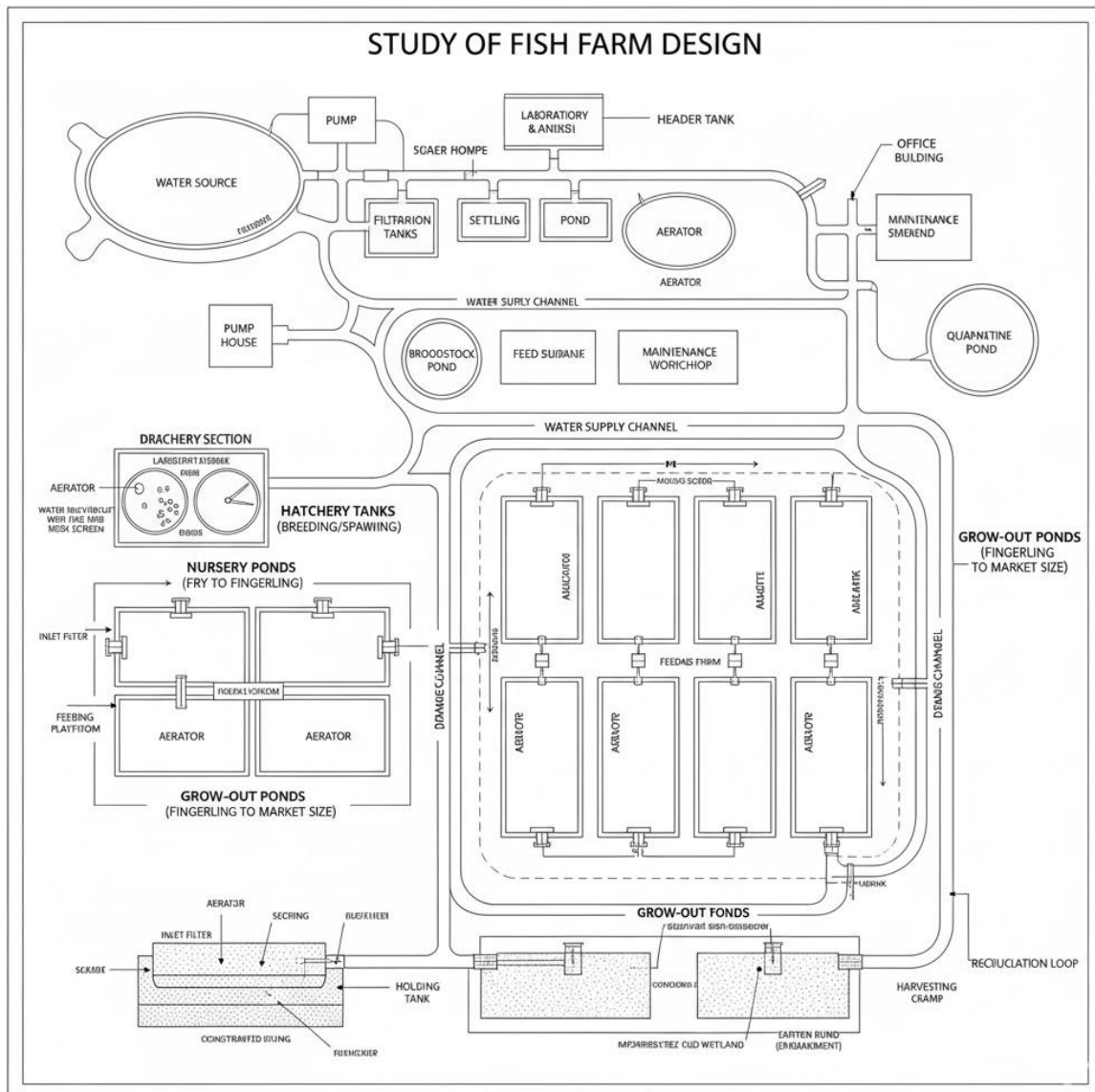
Morphological Characters

- **Shell (Valves):** Two equal or nearly equal shells, typically elongated, wedge-shaped, or oval. The **umbo** (oldest part of the shell) is usually at the pointed anterior end.
- **Hinge:** The dorsal margin where the two valves are joined by a ligament, allowing them to open and close.
- **Growth Lines:** Concentric lines on the shell surface indicating growth periods.
- **Byssal Threads:** A bundle of strong, fibrous threads secreted by a gland in the foot, used to anchor the mussel to hard substrates (rocks, ropes, other shells). This is a unique and key feature.
- **Adductor Muscles:** (Internal, but seen in open specimens/charts). Powerful muscles that close the shells.
- **Mantle:** The fleshy tissue lining the inside of the shells, responsible for secreting the shell.
- **Gills:** Large, sheet-like structures inside the mantle cavity, used for filter feeding (extracting food particles from water) and respiration.
- **Foot:** A small, tongue-like organ used for limited movement and byssal thread secretion.
- **Siphons:** Incurrent and excurrent siphons (sometimes visible as openings) used to draw in and expel water for feeding and respiration.

Economic Importance

- **Primary Food Source:** Mussels are a highly popular and sustainable seafood. They are rich in protein, minerals, and omega-3 fatty acids. Consumed fresh, steamed, boiled, in stews, or processed.
- **Aquaculture: Mussel farming (mussel culture or mariculture)** is a massive and growing industry worldwide. They are typically cultivated on ropes, rafts, or poles, as they are sessile filter feeders. This method is environmentally friendly as it requires no feed input.
- **Ecosystem Services:** As filter feeders, mussels play a vital role in water purification, removing suspended particles and improving water quality in estuaries and coastal areas.
- **Bio-indicators:** Their ability to accumulate pollutants makes them useful as bio-indicators for environmental monitoring programs.
- **By-products:** Shells can be crushed and used as a source of calcium carbonate, or for decorative purposes.
- **Pearl Culture (Limited):** While not their primary economic importance, some mussel species can produce pearls, though less valuable than those from oysters.

STUDY OF FISH FARM DESIGN (CHART/MODEL)



Ideal Fish Farm

1. Structure and Layout of Fish Pond

This refers to the overall arrangement and characteristics of the ponds within the farm.

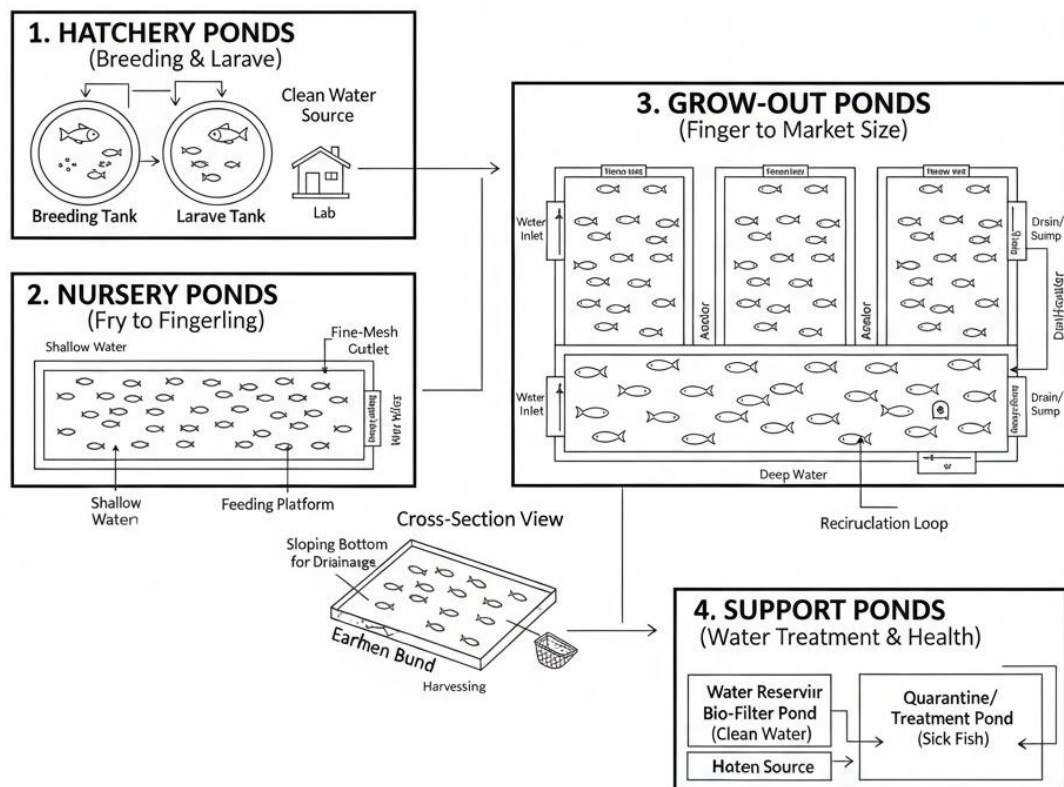
A. Pond Types Based on Function

A well-designed fish farm often segregates different stages of fish growth into specialized ponds to optimize conditions and management.

- **Hatchery (or Breeding/Spawning Pond)**

- **Purpose:** Where broodstock (parent fish) are kept for breeding, and eggs are fertilized and hatched into larvae/fry.
- **Structure/Layout on Chart/Model**
 - **Size:** Typically, *smaller* than nursery or grow-out ponds. They might be relatively shallow or even indoor tanks, depending on the species and breeding method (e.g., haplo breeding, circular tanks for induced breeding).

- **Shape:** Often circular, rectangular, or square tanks for controlled breeding and easy collection of eggs/larvae.
- **Features:** Might show specific installations for egg collection, aeration, water circulation, and temperature control (e.g., heaters).
- **Location:** Often close to a water source and a laboratory/observation area for close monitoring.
- **Importance:** Crucial for initial fish production, controlling reproduction, and ensuring genetic quality.



Types of Ponds used for fishery practices

- **Nursery Pond**
 - **Purpose:** Where newly hatched fry is reared to fingerling size. This stage is critical for high survival rates.
 - **Structure/Layout on Chart/Model**
 - **Size:** *Intermediate* in size, larger than hatchery tanks but smaller than grow-out ponds.
 - **Shape:** Usually rectangular or square, allowing for easy netting and management.
 - **Depth:** Often *shallower* than grow-out ponds (e.g., 0.5 to 1.5 meters) to promote natural food production and easier monitoring of young fish.
 - **Features:** Might show fine mesh screens at inlets/outlets to prevent escape of small fish, and provision for frequent water exchange.

- **Location:** Situated to allow easy transfer of fingerlings to grow-out ponds.
- **Importance:** Provides a protected environment for vulnerable fry, allowing them to grow larger and stronger before being exposed to the rigors of a grow-out system, significantly increasing survival.
- **Grow-out Pond (Production Pond)**
 - **Purpose:** The main production units where fingerlings are grown to market size. This is where the bulk of the economic value is generated.
 - **Structure/Layout on Chart/Model**
 - **Size:** *Largest* ponds on the farm, often several acres in size for extensive or semi-intensive systems, or smaller for intensive systems.
 - **Shape:** Typically, rectangular, square, or irregular based on topography. Rectangular ponds are common for ease of management and harvesting.
 - **Depth:** Deeper than nursery ponds (e.g., 1.5 to 3 meters), providing stable temperatures and more water volume.
 - **Features:** Will prominently feature water inlets/outlets, possibly aerators, feeding areas, and harvesting sumps (channels to collect fish during drainage).
 - **Location:** Usually the most numerous ponds, strategically placed for efficient water flow and access.
 - **Importance:** The core of the farm's production, where fish attain their marketable size and weight.

B. Pond Shape and Orientation

- **Rectangular/Square:** Most common for ease of construction, feeding, sampling, and harvesting (especially with seining). Often oriented with the long axis perpendicular to the prevailing wind for better aeration.
- **Circular:** Used for high-intensity culture, allowing for self-cleaning due to circular flow, but more complex to construct.
- **Irregular:** Can be used in extensive systems where ponds conform to natural topography, reducing construction costs.

C. Pond Bottom and Sides

- **Bottom Slope:** A slight slope towards the drain facilitates complete drainage during harvesting and cleaning. Charts would show cross-sectional views indicating this slope.
- **Lining:** Can be natural earth, clay (for impermeability), or artificial liners (e.g., HDPE, PVC) for intensive systems to prevent seepage and facilitate cleaning. Charts would show cross-sections illustrating the lining.
- **Bunds (Embankments):** Explained below.

2. Water Inlet/Outlet

These are critical for water management, controlling water quality, and enabling harvesting.

- **Water Inlet**
 - **Function:** Brings fresh water into the pond.
 - **Structure/Layout on Chart/Model**
 - **Location:** Typically placed at one end of the pond, often at a higher elevation than the pond bottom to allow gravity flow.
 - **Design:** Can be a simple pipe, a sluice gate, or a channel. Often includes a **screen or filter** to prevent the entry of wild fish, predators, or unwanted debris/vegetation.
 - **Pump/Gravity:** Charts would show if water is supplied by pumps (for lifting water from a source) or by gravity (if the source is higher than the farm).
 - **Importance:** Essential for maintaining good water quality, replenishing dissolved oxygen, diluting waste products, and controlling temperature.
- **Water Outlet/Drainage System**
 - **Function:** Removes water from the pond, allowing for controlled water exchange, drainage for harvesting, and cleaning.
 - **Structure/Layout on Chart/Model**
 - **Location:** Typically, at the *deepest part* of the pond, usually at the opposite end from the inlet, allowing for complete drainage.
 - **Design**
 - **Monk (or Sluice Gate/Standpipe):** A common, vertical structure with removable boards or a gate to control water level. It usually has a screen to prevent fish escape during water exchange. Charts/models would show its cross-section and its position relative to the pond bottom.
 - **Drain Pipe:** A pipe leading from the monk to a common drainage channel or directly to a discharge point.
 - **Harvesting Sump:** Often a small, deeper collection pit at the base of the monk/drain inside the pond. When the pond is drained, fish accumulate here, making harvesting easier.
 - **Importance:** Crucial for water quality management (flushing out metabolites), disease control (draining and drying ponds), and efficient harvesting.

3. Bunds (Embankments)

These are the raised earth walls that define and contain the ponds.

- **Importance:**
 - **Water Retention:** The primary function, holding water within the pond.
 - **Accessibility:** Provide access for farm operations (feeding, sampling, maintenance, harvesting).
 - **Protection:** Act as barriers against predators and prevent fish escape.
 - **Partitioning:** Separate different ponds or sections of the farm.

4. Aerators

Devices used to increase the dissolved oxygen (DO) concentration in the pond water.

- **Purpose:** Fish require sufficient dissolved oxygen to survive and grow. In intensive systems, biological oxygen demand (BOD) from fish waste and feed can deplete DO rapidly.
- **Types (on Chart/Model):** Charts/models would show different types of aerators and their placement.
 - **Paddle Wheel Aerators:** Most common. A motor drives paddles that agitate the water surface, incorporating atmospheric oxygen. Charts would show their position (often in corners or along sides) and the splash they create.
 - **Jet Aerators:** Force water through nozzles, creating fine bubbles and mixing.
 - **Diffusers (Air Blowers):** Air is pumped through diffusers at the pond bottom, creating small bubbles that rise and transfer oxygen. Charts would show tubing laid on the pond bottom and an air compressor unit.
 - **Aspirators:** Draw air into a water stream, creating bubbles.
- **Placement:** Usually strategically placed to ensure uniform oxygen distribution throughout the pond, especially in areas of high fish density or poor circulation.
- **Importance**
 - **Preventing Fish Mortality:** Critical for preventing fish kills due to low oxygen.
 - **Optimizing Growth:** Adequate DO improves feed conversion ratio and growth rates.
 - **Supporting Biota:** Essential for beneficial aerobic bacteria that break down waste.
 - **Increasing Stocking Density:** Allows for higher stocking densities, increasing farm productivity.

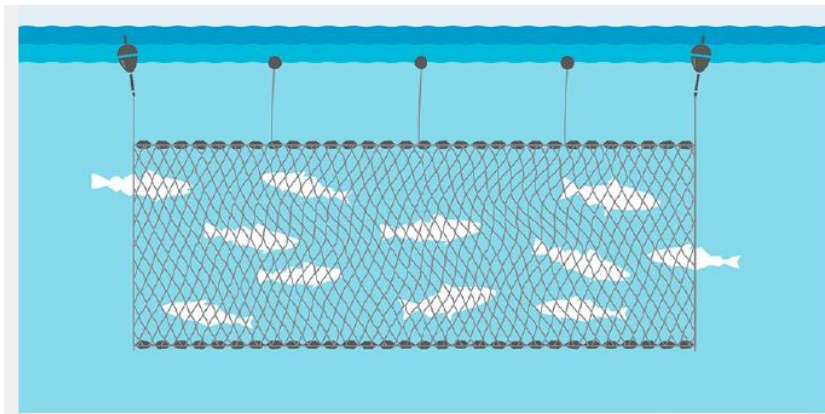
FISHING CRAFTS: GILL NET, CAST NET, TRAWL NET, LONG LINES, SEINE NET, DRIFT NET, PURSE SEINES

FISHING CRAFTS (FISHING GEAR)

Fishing gear refers to the tools and equipment used to capture aquatic organisms. The selection of gear depends on the target species, their behavior, the fishing environment (freshwater, marine, depth), and regulatory frameworks.

1. GILL NET

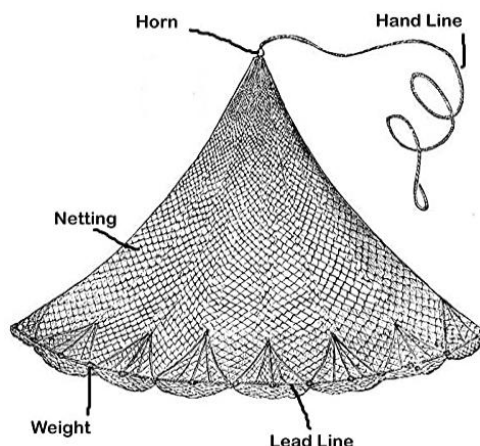
A gill net is a wall of netting, usually set vertically in the water column, designed to entangle fish by their gills as they try to swim through it. The mesh size is crucial; fish of a certain size range will be caught by their gill covers, getting stuck in the mesh.



- **Structure/Components**
 - **Netting:** Made of monofilament or multifilament nylon.
 - **Floatline (Corkline):** The top rope, with floats attached, that keeps the net upright in the water.
 - **Leadline (Footline):** The bottom rope, weighted with leads, that keeps the net stretched vertically.
- **Method of Operation**
 - **Set Gill Net:** Anchored to the bottom or attached to buoys at the surface, remaining stationary.
 - **Drift Gill Net:** Allowed to drift freely with the current, often fished at the surface or mid-water, attached to a vessel. (Note: "Drift net" in your list refers to a type of gill net, often large-scale, which has faced controversy).
- **Target Species:** Highly selective for fish within a specific size range. Used for a wide variety of species, including salmon, tuna, cod, mackerel, and various freshwater fish.
- **Advantages:** Relatively simple to operate, low fuel consumption, size-selective.
- **Disadvantages:** Can be indiscriminate if not properly sized, potential for ghost fishing (lost nets continue to catch), and bycatch of non-target species (including marine mammals and birds). Large-scale drift nets have been banned in many international waters due to high bycatch.

2. CAST NET

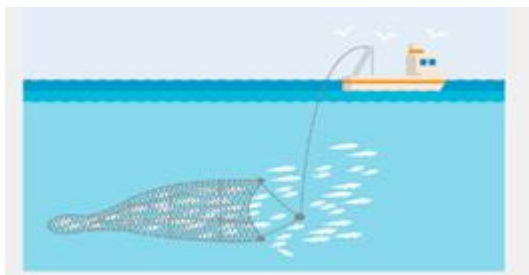
A small, circular net with weights around its perimeter, designed to be cast by hand.



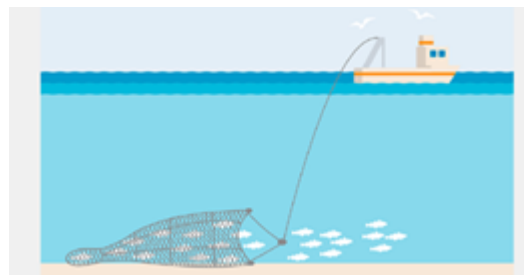
- **Structure/Components**
 - **Circular Netting:** Usually with a small mesh.
 - **Lead Line:** A weighted rope around the circumference.
 - **Hand Line (Rope):** Attached to the center of the net, held by the user.
 - **Brail Lines (Tuck Lines):** Lines running from the lead line to a central point, which pull the net closed when the hand line is retrieved.
- **Method of Operation:** Thrown over a school of small fish from a boat or shore. As it sinks, the weights cause it to spread into a circle. Once it reaches the bottom or over the fish, the user pulls the hand line, closing the net and trapping the fish inside.
- **Target Species:** Primarily small baitfish (sardines, anchovies, mullet) or small fish for subsistence fishing in shallow waters.
- **Advantages:** Simple, inexpensive, effective for small schooling fish, environmentally friendly when used judiciously.
- **Disadvantages:** Limited to shallow water, requires skill to cast effectively, small catches.

3. TRAWL NET

A large, cone-shaped net is towed through the water by one or two boats. It's designed to catch fish by filtering large volumes of water.



PELAGIC TRAWL

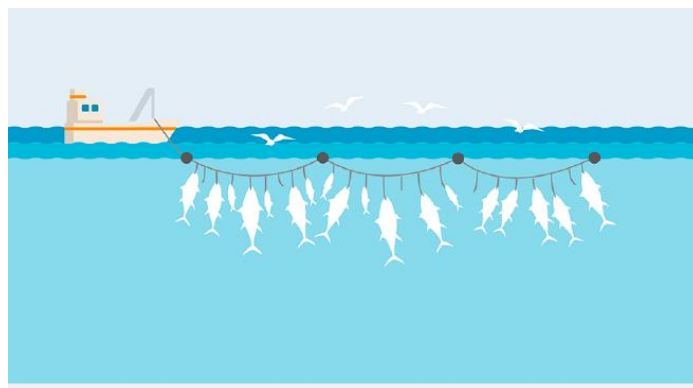


BOTTOM TRAWL

- **Structure/Components**
 - **Wings:** The wide, open front sections.
 - **Body:** The main conical part of the net.
 - **Codend:** The narrow, closed end where the fish accumulate.
 - **Doors (Otter Boards):** Large, rectangular or curved plates that spread the mouth of the net horizontally.
 - **Floats & Weights:** To keep the net open vertically.
- **Method of Operation**
 - **Bottom Trawling:** Towed along the seabed, often catching demersal (bottom-dwelling) fish and shellfish. This can cause significant habitat disturbance.
 - **Midwater Trawling (Pelagic Trawling):** Towed in the mid-water column, targeting schooling pelagic (open-ocean) fish.
- **Target Species:** Wide variety of demersal fish (cod, haddock, flatfish, shrimp, crabs) and pelagic fish (herring, mackerel, hake).
- **Advantages:** Highly efficient for catching large quantities of fish, relatively low labor per unit of catch.
- **Disadvantages:** High potential for bycatch, can be damaging to seabed habitats (bottom trawling), high fuel consumption, less selective.

4. LONGLINES

A long main line, often many kilometers long, to which numerous shorter lines (gangions or snoods) with baited hooks are attached at regular intervals.

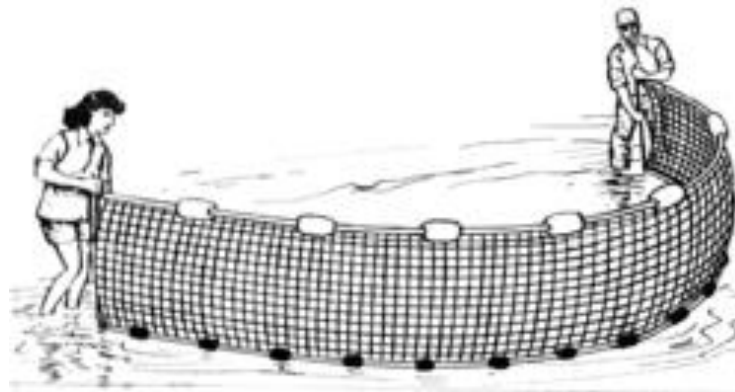


- **Structure/Components:**
 - **Main Line:** A strong, long line.
 - **Gangions/Snoods:** Shorter lines attached to the main line.
 - **Hooks:** Baited hooks at the end of each gangion.
 - **Floats & Anchors:** To set the line at a specific depth (surface, mid-water, or bottom).

- **Method of Operation:** Set in the water and left to soak for several hours or overnight, allowing fish to find and take the bait. The line is then retrieved, and fish are unhooked.
- **Target Species:** Large predatory fish like tuna, swordfish, halibut, cod, sharks.
- **Advantages:** Highly selective for target species and size (depending on hook size and bait), relatively low fuel consumption, less habitat impact than trawling (unless set on sensitive bottoms).
- **Disadvantages:** Can have significant bycatch of seabirds (if not properly mitigated), marine mammals, and non-target fish. Managing and setting/retrieving very long lines can be labor-intensive.

5. SEINE NET

A long wall of netting, often with a bag or "bunt" in the center, is designed to encircle a school of fish. It's operated by pulling the ends of the net together, often from shore or a boat.

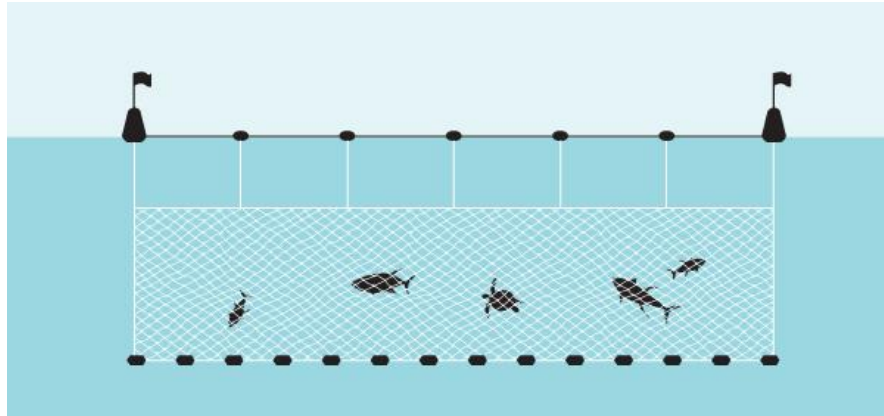


- **Types**
 - **Beach Seine (Shore Seine):** Pulled from shore.
 - **Danish Seine/Scottish Seine (Fly Shooting):** A boat deploys the net in a large circle and then pulls the ropes and net from the boat, herding fish into the bag.
- **Structure/Components:** Similar to a long gill net but designed to enclose, not entangle.
 - **Netting:** Can vary in mesh size.
 - **Floatline & Leadline:** To keep it vertical.
 - **Wings:** The outer parts of the net.
 - **Bag/Bunt:** The central, deeper part where fish accumulate.
- **Method of Operation:** Deployed to encircle fish. The net is then gradually hauled in, trapping the fish in the bag.
- **Target Species:** Schooling fish in coastal areas (mullet, sardines, small pelagic fish) and some demersal species (Danish seine).
- **Advantages:** Can be effective for specific schooling fish, less damaging to habitat than trawling if operated correctly.

- **Disadvantages:** Requires specific bottom conditions (for beach seines), can still have bycatch, dependent on fish schooling behavior.

6. DRIFT NET

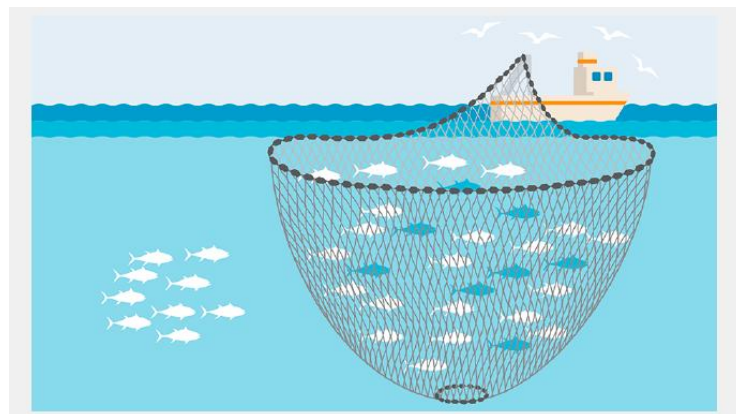
As mentioned under Gill Nets, a drift net is a type of gill net that is not anchored but allowed to drift freely with the ocean currents, typically attached to a fishing vessel. They can be very long.



- **Structure/Components:** Same as a gill net (netting, floatline, leadline).
- **Method of Operation:** Deployed at night and retrieved in the morning.
- **Target Species:** Highly migratory pelagic species like tuna, marlin, swordfish, and salmon.
- **Advantages:** Effective for wide-ranging species.
- **Disadvantages:** **Highly controversial and largely banned internationally** due to "ghost fishing" (lost nets continue to catch indiscriminately) and massive bycatch of marine mammals (dolphins, whales), seabirds, and non-target fish. The term "large-scale high seas driftnet fishing" refers to nets over 2.5 km in length.

7. PURSE SEINE

A very large net designed to encircle a school of fish, and then its bottom is drawn together (pursed) to enclose the fish completely, like a drawstring purse.



- **Structure/Components**
 - **Netting:** A long, deep wall of netting.
 - **Floatline:** Top line with numerous floats.
 - **Leadline:** Bottom line with weights.
 - **Purse Rings:** Rings attached to the leadline through which a "purse line" runs.
 - **Purse Line:** A strong cable that runs through the purse rings, allowing the bottom of the net to be closed.
- **Method of Operation:** A vessel (seiner) encircles a school of fish (often located using sonar or spotter planes). The net is rapidly deployed around the school. Once the circle is complete, the purse line is hauled in, drawing the bottom of the net closed and forming a "bag" that traps the fish. The net is then brought alongside the vessel, and the fish are pumped or scooped onboard.
- **Target Species:** Highly efficient for large schools of pelagic fish such as tuna, sardines, mackerel, anchovies, herring, and salmon.
- **Advantages:** Extremely efficient for large schools, can catch huge volumes of fish, relatively low bycatch if targeting pure schools and using methods to avoid non-target species (e.g., FAD-free tuna purse seining).
- **Disadvantages:** Can result in large bycatch if set on mixed schools or around Fish Aggregating Devices (FADs) that attract various species, requires large vessels and specialized equipment, can lead to overfishing if not managed properly.

GOAT FARMING AND DAIRY SCIENCE

STUDY OF GOAT BREEDS:

SIROHI, JAMUNAPARI, OSMANABADI, BOER, BEETAL, SAANEN

Goat Farming and Dairy Science: A. Study of Goat Breeds

This study focuses on identifying and understanding the unique characteristics and economic utility of different goat breeds. For each breed, you'd typically examine:

- **Origin and Distribution:** Where the breed comes from and where it's commonly found.
- **Morphological Characteristics:** Distinctive physical features like size, color, horn presence/shape, ear type, hair coat.
- **Production Traits:** Primary use (meat, milk, fiber, or dual-purpose), average milk yield, growth rate, prolificacy (number of kids per kidding).
- **Adaptability:** Resistance to diseases, tolerance to different climates/environments.

1. SIROHI



- **Origin and Distribution:** Native to the Sirohi district of Rajasthan, India, and also found in parts of Gujarat. It's well-adapted to the arid and semi-arid regions of these states.
- **Morphological Characteristics**
 - **Size:** Medium-sized breed.
 - **Color:** Predominantly **brown** in various shades, often with **light brown patches or spots** (piebald markings). Sometimes entirely brown or white.
 - **Ears:** Flat, leaf-like, and **medium-sized**, generally drooping downwards.
 - **Horns:** Both sexes possess **short, curved horns**, generally directed upwards and backwards.
 - **Coat:** Short, coarse hair.
 - **Face:** Slightly convex (Roman nose) is sometimes observed.

- **Production Traits**
 - **Primary Use: Dual-purpose (meat and milk).** Highly valued for meat due to good growth rate and feed conversion. Also provides a moderate amount of milk.
 - **Milk Yield:** Moderate, typically 0.5-1.0 kg/day.
 - **Kidding:** Good prolificacy, often producing twins.
- **Adaptability:** Known for its hardiness, resistance to common diseases, and ability to thrive in harsh, hot, and dry conditions with limited fodder availability.

2. JAMUNAPARI



- **Origin and Distribution:** Indigenous to the Yamuna riverine tracts (hence "Jamunapari") of Uttar Pradesh, India (especially Etawah district). Also popular in other parts of India and widely exported.
- **Morphological Characteristics**
 - **Size:** **Very large** and majestic, one of the largest Indian goat breeds.
 - **Color:** Variable, but typically **white or cream**, with patches of tan, black, or brown on the head and neck.
 - **Ears:** Very long, flat, and **pendulous (drooping)**, often folded, resembling a "leaf" or "banana" shape. They can touch the ground if the goat lowers its head.
 - **Horns:** Short, flat, and curved, generally pointing backward. Some animals may be polled (hornless).
 - **Face:** Distinctive **highly convex or Roman nose**, giving it a regal appearance.
 - **Legs:** Long, often with **long hair on the hind legs** (known as "feathering" or "hairy breeches").
 - **Udder:** Well-developed udder with long teats.

- **Production Traits**
 - **Primary Use: Dual-purpose (milk and meat).** Renowned as a dairy goat, often called "poor man's cow." Also good for meat due to its large size.
 - **Milk Yield:** High, typically 2.0-3.0 kg/day, with some individuals yielding more.
 - **Kidding:** Often single kids, sometimes twins.
- **Adaptability:** Adapted to extensive grazing systems. While hardy, they prefer warmer climates and good fodder.

3. OSMANABADI



- **Origin and Distribution:** Native to the Osmanabad district of Maharashtra, India. Widely distributed across Maharashtra and neighboring states.
- **Morphological Characteristics**
 - **Size:** Medium to large size.
 - **Color:** Predominantly **black**, but variations like brown, white, or spotted are also seen.
 - **Ears:** Medium-sized, flat, and drooping downwards, but not as long as Jamunapari.
 - **Horns:** Both sexes possess **backward-curving horns** of medium length.
 - **Coat:** Short and coarse hair.
- **Production Traits**
 - **Primary Use:** Primarily a **meat breed**, known for its good body weight and high dressing percentage. Also provides a moderate amount of milk, making it dual-purpose.
 - **Milk Yield:** Moderate, around 0.5-1.0 kg/day.
 - **Kidding:** Highly prolific, known for **high twinning and even triplets**, which contribute to its meat production potential.

- **Adaptability:** Very hardy and well-adapted to various climatic conditions, including semi-arid and tropical regions. Known for good disease resistance and scavenging ability.

4. BOER



- **Origin and Distribution:** Developed in South Africa. It's a globally recognized and highly popular **meat goat breed**, widely introduced to many countries, including India, for commercial meat production.
- **Morphological Characteristics**
 - **Size:** **Very large** and heavily muscled, renowned for its substantial build.
 - **Color:** Distinctive coloration: **white body with a reddish-brown head and a white blaze** on the face. Color variations (e.g., solid red, pied) exist but are less common for purebreds.
 - **Ears:** Long, broad, and **pendulous** (drooping).
 - **Horns:** Strong, thick, and backward-curving horns.
 - **Face:** Strong, often Roman nose.
 - **Body:** Very well-fleshed, compact, and muscular, reflecting its primary purpose.
- **Production Traits**
 - **Primary Use:** **Exclusively a meat breed.** Exceptional growth rate, superior feed conversion efficiency, and excellent carcass quality. Often used for cross-breeding with indigenous goats to improve meat production traits.
 - **Milk Yield:** Low, only sufficient for raising its kids.
 - **Kidding:** Good prolificacy, typically twins.
- **Adaptability:** Relatively hardy, but generally requires better management and nutrition compared to indigenous breeds, especially in intensive farming systems.

5. BEETAL



- **Origin and Distribution:** Originates from the Punjab region of India and Pakistan. It's one of the largest and most prominent Indian dairy goat breeds.
- **Morphological Characteristics:**
 - **Size:** **Large-sized** breed, comparable to Jamunapari but generally slightly less robust.
 - **Color:** Highly variable, common colors include **black with white spots, brown, red, or mottled (mix of colors)**.
 - **Ears:** Long, broad, and **pendulous (drooping)**, similar to Jamunapari but often less curled.
 - **Horns:** Medium-sized, spirally twisted, and directed backward and upward. Some individuals can be polled.
 - **Face:** Prominent, often **Roman nose**.
 - **Udder:** Well-developed and capacious udder, indicating good milk production.
- **Production Traits**
 - **Primary Use:** **Dual-purpose (milk and meat)**, but primarily valued as a **dairy breed**. It's often referred to as the "dairy goat of India." Also has good meat qualities due to its size.
 - **Milk Yield:** High, typically 1.5-2.5 kg/day, making it one of the best milk producers among Indian breeds.
 - **Kidding:** Good prolificacy, often twins.
- **Adaptability:** Well-adapted to plains and semi-arid regions. Relatively hardy but thrives best under semi-intensive or intensive management with good feed.

6. SAANEN



- **Origin and Distribution:** Originates from the Saanen Valley in Switzerland. It is the world's most widely recognized and highest milk-producing dairy goat breed, found in almost every country with a dairy goat industry.
- **Morphological Characteristics:**
 - **Size:** Large-sized and elegant, with a refined appearance.
 - **Color:** Distinctive **pure white or creamy white** (no spots or patterns). Occasionally, a "biscuit" patch may be seen.
 - **Ears:** Medium-sized, erect, and alert, pointing slightly forward. This is a key distinguishing feature from pendulous-eared Indian breeds.
 - **Horns:** Usually **polled (hornless)**, but horned individuals can occur.
 - **Face:** Straight or slightly dished face.
 - **Hair:** Short, fine hair, sometimes with a fringe of longer hair on the back and thighs.
 - **Udder:** Exceptionally large, well-attached, and capacious udder with well-placed teats.
- **Production Traits**
 - **Primary Use:** **Exclusively a dairy breed.** Unmatched milk production globally.
 - **Milk Yield:** Very high, typically 3.0-5.0 kg/day, with top producers exceeding this significantly. Known for persistent lactation.
 - **Milk Quality:** Milk has a lower fat content compared to some other breeds.
 - **Kidding:** Good prolificacy, often twins or triplets.
- **Adaptability:** Prefers cooler climates. Requires excellent management, high-quality feed, and clean housing for optimal production. Can be more susceptible to heat stress in hot, humid climates.

STUDY DAIRY PRODUCTS: MILK, CURD, GHEE, PANEER, CHEESE, KHOA

Study of Dairy Products

This study aims to provide a detailed understanding of common dairy products, covering their raw material (milk), processing methods, physical and chemical properties, and economic significance.



MILK



CURD



GHEE



PANEER



CHEESE



KHOA

1. MILK

- **Description:** The primary raw material for all other dairy products. It's a complex biological fluid secreted by the mammary glands of mammals. In dairy science, it typically refers to milk from cows, buffalo, goats, or sheep.
- **Composition**
 - **Water:** Approximately 87-88%
 - **Lactose (Milk Sugar):** Around 4.5-5.0% (primary carbohydrate)
 - **Fat:** Highly variable (e.g., 3.5-6.0% for cow milk, 6.0-9.0% for buffalo milk). Present as tiny globules in an emulsion.
 - **Proteins:** Around 3.2-3.5% (Caseins ~80%, Whey proteins ~20%). Caseins are responsible for curd formation.
 - **Minerals:** (Calcium, Phosphorus, Potassium, etc.) ~0.7%
 - **Vitamins:** (A, D, B vitamins, etc.)
- **Processing (for liquid consumption)**
 - **Chilling:** Rapid cooling to 4°C to inhibit microbial growth.
 - **Clarification/Filtration:** Removal of suspended impurities.

- **Standardization:** Adjusting fat content to meet legal requirements (e.g., skim, toned, full cream milk).
- **Homogenization:** Breaking down fat globules into smaller, uniform sizes to prevent cream separation (creaming up) and improve mouthfeel.
- **Pasteurization:** Heating milk to specific temperatures for a set time to kill pathogenic microorganisms and extend shelf life (e.g., LTLT: 63°C for 30 min; HTST: 72°C for 15 sec; UHT: 135-150°C for a few seconds).
- **Packaging:** Aseptic packaging for UHT, regular packaging for pasteurized milk.
- **Economic Importance:** Staple food globally, source of essential nutrients, massive industry providing livelihoods from farm to processing.

2. CURD (DAHI IN INDIA)

- **Description:** A fermented milk product, semi-solid in consistency, produced by the action of lactic acid bacteria on milk.
- **Process**
 1. **Heating Milk:** Milk is typically boiled and then cooled to a lukewarm temperature (around 40-45°C).
 2. **Inoculation (Setting):** A small amount of previous curd (starter culture) or a commercial bacterial culture (containing *Lactococcus lactis*, *Streptococcus thermophilus*, *Lactobacillus bulgaricus*, etc.) is added.
 3. **Incubation:** The milk is left undisturbed at a warm temperature (30-45°C) for several hours (e.g., 6-10 hours).
 4. **Coagulation:** Lactic acid bacteria ferment lactose into lactic acid, which lowers the pH. This causes the casein proteins to coagulate and form a gel, resulting in the semi-solid curd.
 5. **Cooling:** Once set, the curd is refrigerated to slow down further fermentation.
- **Composition:** Similar to milk but with reduced lactose, increased acidity, and altered protein structure. Contains live beneficial bacteria.
- **Economic Importance:** Widely consumed as a food item, base for many traditional dishes (Raita, Lassi), probiotic benefits, significant part of dairy consumption in many cultures.

3. GHEE

- **Description:** Clarified butterfat, anhydrous (almost water-free). A traditional fat popular in South Asia and the Middle East.
- **Process (Traditional)**
 1. **Butter/Cream Source:** Starts with butter or cream (often from curd /dahi).
 2. **Slow Heating:** Butter/cream is gently heated.

3. **Separation:** As heating continues, water evaporates, and the milk solids (proteins, lactose) separate and settle at the bottom or collect on the surface as foam.
4. **Clarification:** Heating continues until all moisture is removed and the milk solids turn golden brown. The clear fat is then carefully decanted or filtered.
5. **Cooling & Granulation:** The liquid ghee is cooled, forming characteristic granules upon solidification (a sign of good quality).
- **Composition:** Almost 99.5% fat. Contains fat-soluble vitamins (A, D, E, K).
- **Economic Importance:** Valued for its distinct flavor, high smoke point (suitable for frying), long shelf life, and traditional/cultural significance in cooking and rituals. High commercial value.

4. PANEER

- **Description:** A fresh, unripe cheese, common in Indian cuisine. It's a non-melting, non-fermented, acid-coagulated cheese.
- **Process**
 1. **Heating Milk:** Full-fat milk is heated to near boiling (90-95°C).
 2. **Coagulation:** An acidic coagulant (lemon juice, vinegar, citric acid solution, or sour whey) is added to the hot milk. The acid lowers the pH, causing the casein proteins to coagulate into curds.
 3. **Separation:** The curds separate from the greenish-yellow liquid whey.
 4. **Draining & Pressing:** The curds are collected in a cheesecloth, excess whey is drained, and the curds are pressed underweight to form a firm block.
 5. **Chilling:** The block of paneer is chilled to firm up its texture.
- **Composition:** High in protein and fat, low in lactose.
- **Economic Importance:** Versatile ingredient in Indian vegetarian dishes (curries, snacks, desserts). Good source of protein, especially important in vegetarian diets.

5. CHEESE

- **Description:** A diverse group of fermented and ripened milk products, produced by coagulating milk protein (casein) and separating it from the whey. There are thousands of varieties worldwide.
- **General Process (variations exist for each type):**
 1. **Milk Preparation:** Milk (cow, goat, sheep, and buffalo) is often pasteurized, standardized, and sometimes homogenized.
 2. **Starter Culture Addition:** Specific lactic acid bacteria cultures are added to ferment lactose and produce lactic acid.
 3. **Rennet Coagulation:** Rennet (an enzyme, traditionally from calf stomachs, now often microbial) is added to coagulate the casein proteins, forming a solid curd.

4. **Cutting the Curd:** The soft curd is cut into small pieces to release more whey.
 5. **Cooking (Heating):** The curds are gently heated to further expel whey and firm up the curd.
 6. **Draining/Pressing:** Whey is drained, and the curds are often pressed to remove more whey and form a solid mass.
 7. **Salting:** Salt is added for flavor, preservation, and to control microbial growth.
 8. **Shaping:** The cheese is formed into its characteristic shape (blocks, wheels, etc.).
 9. **Ripening/Aging (Maturation):** The most critical stage. Cheese is stored under controlled temperature and humidity for weeks, months, or even years. During this time, enzymes (from rennet, milk, and microbes) break down proteins and fats, developing the unique flavor, aroma, and texture of the specific cheese variety. (Fresh cheeses like cottage cheese or cream cheese skip this step).
- **Composition:** Highly variable depending on type, but generally concentrated in protein and fat, with very low or no lactose in aged cheeses.
 - **Economic Importance:** Global staple food, vast culinary applications, major export commodity for many countries, provides significant economic activity from dairy farming to processing and retail.

6. KHOA (MAWA)

- **Description:** A concentrated milk product, traditionally made by slowly evaporating milk in an open pan until it reaches a semi-solid consistency. It forms the base for numerous Indian sweets.
- **Process**
 1. **Milk Heating:** Full-fat milk (often buffalo milk due to higher solids) is taken in a large, shallow, heavy-bottomed pan (karahi).
 2. **Slow Evaporation:** The milk is continuously stirred over a moderate flame to prevent scorching. Water evaporates, and the milk thickens.
 3. **Concentration:** Stirring becomes more vigorous as the milk thickens, preventing lumps and ensuring uniform evaporation.
 4. **Final Stage:** Heating continues until the milk solids reach a pasty, semi-solid, crumbly consistency that leaves the sides of the pan.
 5. **Cooling:** The khoa is removed from heat and cooled.
- **Composition:** Highly concentrated milk solids. Typically contains 25-30% moisture, 20-25% fat, and 20-25% protein, and about 30-35% lactose.
- **Economic Importance:** Essential ingredient in a vast array of Indian sweets (e.g., gulab jamun, peda, barfi), major component of the traditional Indian dairy market.

PARASITOLOGY

ASCARIS

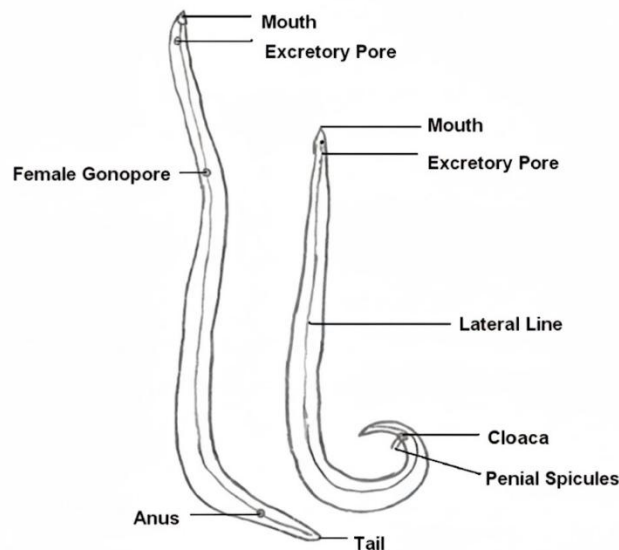
- Morphology (Male, Female, Sexual dimorphism)
- Life cycle
- Parasitic adaptations are explained in detail

Parasitology: A. Ascaris (*Ascaris lumbricoides*)

Ascaris lumbricoides is the largest nematode (roundworm) parasitizing the human intestine. It causes a condition known as ascariasis, which is a significant public health problem, especially in tropical and subtropical regions with poor sanitation.

a. Morphology (Male, Female, Sexual Dimorphism)

Ascaris worms are cylindrical, elongated, and have a smooth, tough cuticle (outer covering). They taper slightly at both ends.



FEMALE AND MALE ASCARIS

1. MALE ASCARIS

- **Size:** Generally **smaller** than the female. Typically measures about 15-30 cm in length and 2-4 mm in diameter.
- **Posterior End:** The most distinctive feature is its **ventrally curved (hooked) posterior end**.
- **Spicules:** Possesses **two chitinous copulatory spicules** (penial spicules) that protrude from the cloaca (a common opening for digestive and reproductive tracts) at the posterior end. These are used to hold open the female's genital pore during copulation.
- **Cloaca:** A common opening for the digestive and reproductive systems.
- **Genital Papillae:** Small, sensory structures (pre-anal and post-anal papillae) are present near the cloaca.

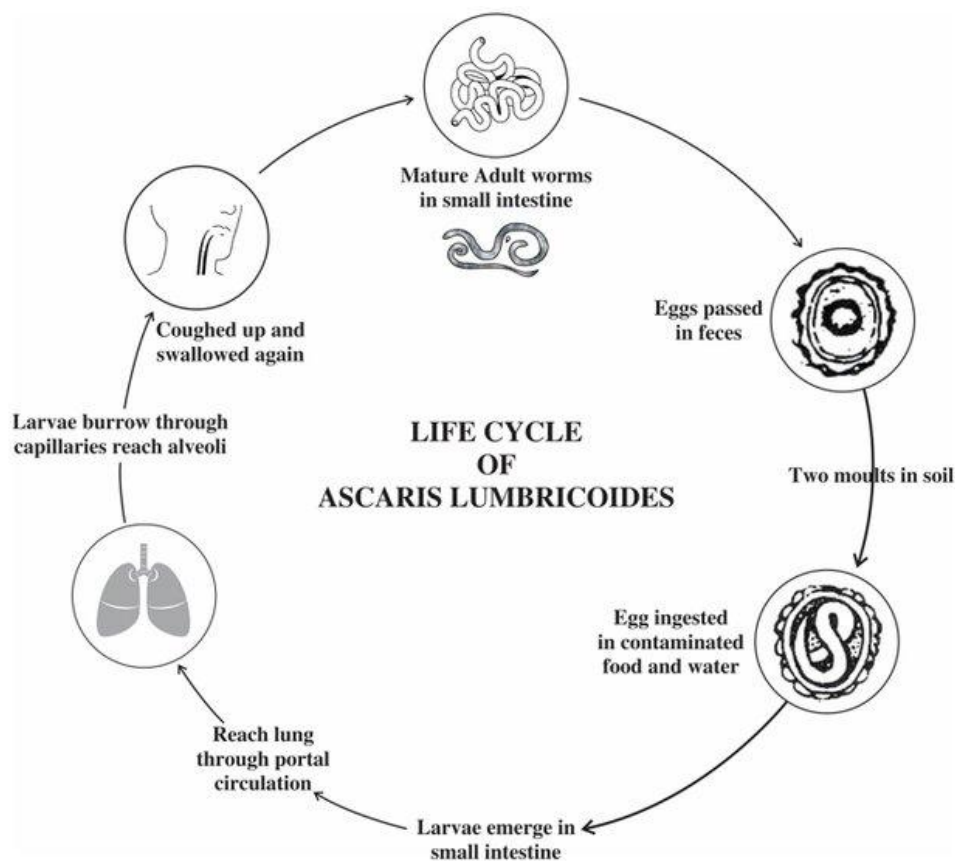
2. Female *Ascaris*

- **Size:** Larger and more robust than the male. Measures about 20-49 cm in length and 3-6 mm in diameter.
- **Posterior End:** The posterior end is **straight and conical**, not curved.
- **Vulva (Genital Aperture):** A distinct transverse slit located approximately one-third of the body length from the anterior end. This is where the female receives sperm and lays eggs.
- **Anus:** A separate opening at the posterior end, distinct from the vulva.
- **Reproductive System:** Highly prolific, with two long, convoluted uteri that fill most of the body cavity, containing millions of eggs.

3. Sexual Dimorphism (Key Differences between Male and Female)

- **Size:** Female is significantly larger and thicker than the male.
- **Posterior End:** Male's posterior end is curved ventrally with spicules; female's is straight and conical.
- **Genital Opening:** Male has a cloaca at the posterior end; female has a vulva one-third down the body and a separate anus at the posterior end.
- **Reproductive Structures:** Male has testes and spicules; female has ovaries, oviducts, and a large, egg-filled uterus.

b. Life Cycle (*Ascaris lumbricoides*)



Ascaris lumbricoides has a direct life cycle, meaning it does not require an intermediate host. It's a complex cycle involving migration through the human body.

1. Egg Stage (in Environment)

- **Unembryonated Eggs:** Adult female worms in the human intestine lay a prodigious number of unembryonated eggs (200,000 per day!). These eggs are passed in the feces of an infected person.
- **Embryonation/Larvation:** When passed in feces, the eggs are not immediately infective. They require specific environmental conditions (warmth, moisture, shady soil) to develop. Within 2-4 weeks (depending on conditions), a first-stage larva (L1) develops inside the egg, which then molts to a second-stage larva (L2), which is the infective stage. The egg containing the L2 larva is called an embryonated egg.

2. Infection of Host (Human)

- Humans become infected by ingesting infective embryonated eggs from contaminated soil, water, or food (e.g., unwashed vegetables).

3. Hatching and Larval Migration (in Human)

- **Hatching:** Once ingested, the embryonated eggs hatch in the small intestine, releasing **L2 larvae**.
- **Penetration & Hepatic Portal System:** The L2 larvae penetrate the intestinal wall and enter the **hepatic portal system** (blood vessels leading to the liver).
- **Liver & Heart:** They are carried to the **liver**, where they remain for 3-4 days and undergo a molt to L3 larvae. From the liver, they travel through the bloodstream to the heart.
- **Lungs:** From the heart, they are carried to the lungs via the pulmonary arteries. Here, they break out of the capillaries into the alveoli (air sacs).
- **Alveolar Molt & Migration:** In the alveoli, the larvae grow, feed on blood, and undergo another molt (to **L4 larvae**). They then migrate up the bronchial tree, trachea, and into the pharynx.
- **Swallowing:** The larvae are coughed up and then swallowed back down the esophagus.

4. Adult Stage (in Intestine)

- **Intestinal Development:** The swallowed L4 larvae pass through the stomach and reach the **small intestine**, where they mature into adult male and female worms.
- **Sexual Maturation & Reproduction:** Within 6-10 weeks after infection, the adult worms become sexually mature, mate, and the female begins laying eggs, completing the cycle.

c. Parasitic Adaptations

Ascaris lumbricoides exhibits several remarkable adaptations that allow it to survive and thrive as an endoparasite

(a) Morphological adaptations

1. The body is long and cylindrical, pointed at both ends.
2. The mouth is bounded by three lips which help the parasite to attach with mucous membrane of the host's intestine.
3. The parasite is devoid of Locomotory organs as the parasite lives in the intestine where protection from enemies and food supply are ensured.
4. The body wall of *Ascaris* is covered with cuticle, resistant to the digestive enzymes of the host.

(b) Anatomical adaptations

1. The pharynx is muscular that facilitates ingestion of food by sucking action. There are no digestive glands.
2. Sense organs are ill-developed, being found only on lips in the form of papillae.
3. The digestive tract is simple without provision for storage, as there is constant supply of food.

(c) Physiological adaptations

1. The body wall is covered with tough, thick and resistant cuticle, shields against the digestive enzymes of the host and antitoxins.
2. Ingested food of this parasite is pre- digested, so that there are no elaborate digestive glands.
3. The respiration is almost entirely anaerobic. Extremely low metabolic rate and anaerobic respiration enable the worm to live inside the host's intestine, where the free oxygen is negligible.
4. Reproductive system of *Ascaris* is well- developed and numerous eggs are produced to make up for the poor chances of the right host being reached.
5. The eggs are covered with resistant covering or chitinous shell which provides safety to the zygote and embryonated eggs from unfavorable environmental factors.
6. The minute size and resistant nature of eggs make them to withstand prolonged dryness and cold. The minute size eggs afford far and wide dispersal of the parasite.

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