

# Digestive System Processes: Chemical and Physical

## **Objectives**

- List the digestive system enzymes involved in the digestion of proteins, fats, and carbohydrates; state their site of origin; and summarize the conditions promoting their optimal functioning.
- ☐ Name the end products of protein, fat, and carbohydrate digestion.
- ☐ Define enzyme, catalyst, control, substrate, and hydrolase.
- Describe the different types of enzyme assays, and the appropriate chemical tests to determine if digestion of a particular foodstuff has occurred.
- ☐ Discuss the role of temperature and pH in the regulation of enzyme activity.
- ☐ State the function of bile in the digestive process.
- Explain why swallowing is both a voluntary and a reflex activity, and discuss the role of the tongue, larynx, and gastroesophageal sphincter in swallowing.
- ☐ Compare and contrast segmentation and peristals is as mechanisms of mixing and propulsion in digestive tract organs.

## **Materials**

#### Part I: Enzyme Action

#### General Supply Area

- Hot plates
- 250-ml beakers
- Boiling chips
- Test tubes and test tube rack
- Wax markers
- Water bath set at 37°C (if not available, incubate at room temperature and double the time)
- Ice water bath
- · Chart on board for recording class results

#### Activity 1: Starch Digestion

Dropper bottle of distilled water

Text continues on next page. →

# **Mastering** A

For related exercise study tools, go to the Study Area of MasteringA&P. There you will find:

- Lab PAL
- Practice Anatomy
   A&PFlix
- PhysioEx PEx
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## Pre-Lab Quiz

- 1. Circle the correct underlined term. Enzymes are catalysts / substrates that increase the rate of chemical reactions without becoming a part of
- 2. Circle True or False. Breakdown products of fats are absorbed by the lymphatic system and are then transported into the systemic circulation by lymph.
- 3. A(n) is a specimen or standard against which all experimental samples are compared.
  - **b.** control
- c. substrate
- d. trial
- 4. One enzyme that you will be studying today, produced by the salivary glands and secreted into the mouth, hydrolyzes starch to maltose. It is
- 5. Circle True or False. When you use iodine to test for starch, a color change to blue-black indicates a positive starch test.
- **6.** If Benedict's test in the starch assay produces a precipitate, then your test will be recorded as positive for maltose.
  - a. blue to black
- b. green to orange
- **7.** The enzyme , produced by the pancreas, is responsible for breaking down proteins.
  - **b**. kinase c. lipase d. trypsin **a.** amylase
- 8. Circle the correct underlined term. The enzyme pancreatic lipase / pepsin hydrolyzes neutral fats to their component monoglycerides and fatty acids.
- 9. Circle True or False. Both smooth and skeletal muscles are involved in the propulsion of foodstuffs along the alimentray canal.
- movements are local contractions that mix foodstuffs with digestive juices and increase the rate of absorption.
  - a. Deglutition
- **b**. Elimination
- c. Peristaltic
- d. Segmental

- Dropper bottles of the following: 1% alpha-amylase solution\* 1% boiled starch solution, freshly prepared 1% maltose solution Lugol's iodine solution (IKI) Benedict's solution
- Spot plate

#### Activity 2: Protein Digestion

 Dropper bottles of 1% trypsin and 0.01% BAPNA solution

#### Activity 3: Bile Action and Fat Digestion

- Dropper bottles of 1% pancreatin solution, litmus cream (fresh cream to which powdered litmus is added to achieve a deep blue color), 0.1 NHCl, and vegetable oil
- Bile salts (sodium taurocholate)
- Parafilm® (small squares to cover the test tubes)

#### Part II: Physical Processes

#### Activity 5: Observing Digestive Movements

- Water pitcher
- Paper cups
- Stethoscope
- Alcohol swab
- Disposable autoclave bag
- · Watch, clock, or timer

#### Activity 6: Video Viewing

- Television and VCR or DVD player for independent viewing of video by student
- Interactive Physiology<sup>®</sup>, Digestive

PEx PhysioEx™ 9.1 Computer Simulation Ex. 8 on p. PEx-119

he food we eat must be processed so that its nutrients can reach the cells of our body. First the food is mechanically broken down into small particles, and then the particles are chemically (enzymatically) digested into the molecules that can be absorbed. Food digestion is a prerequisite to food absorption. (You have already studied mechanisms of passive and active absorption in Exercise 5 and/or PhysioEx Exercise 1. Before proceeding, review that material.)

### **Digestion of Foodstuffs: Enzymatic Action**

**Enzymes** are large protein molecules produced by body cells. They are biological catalysts, meaning that they increase the rate of a chemical reaction without themselves becoming part of the product. The digestive enzymes are hydrolytic enzymes, or hydrolases. Their substrates, or the molecules on which they act, are organic food molecules which they break down by adding water to the molecular bonds, thus cleaving the bonds between the chemical building blocks, or monomers.

Each enzyme hydrolyzes only one or a small group of substrate molecules, and specific environmental conditions are necessary for it to function optimally. Since digestive enzymes actually function outside the body cells in the digestive tract, their hydrolytic activity can also be studied in a test tube.

Figure 39.1 is a flowchart depicting the progressive digestion of carbohydrates, proteins, fats, and nucleic acids. It summarizes the specific enzymes involved, their site of formation, and their site of action. Acquaint yourself with the flowchart before beginning this experiment, and refer to it as necessary during the laboratory session.

#### General Instructions for Activities 1–3

Work in groups of four, with each group taking responsibility for setting up and conducting one of the following experiments. In each activity, you are directed to boil the contents of one or more test tubes. To do this, obtain a 250-ml beaker, boiling chips, and a hot plate from the general supply area. Place a few boiling chips into the beaker, add about 125 ml of water, and bring to a boil. Place the test tube for each specimen in the water for the number of minutes specified in the directions. You will also be using a 37°C bath and an ice water bath for parts of these experiments.

Upon completion of the experiments, each group should communicate its results to the rest of the class by recording them in a chart on the board. Each assay contains one or more controls, the specimens against which experimental samples are compared. All members of the class should observe the controls as well as the experimental results and be able to explain the tests used and the results observed and anticipated for each experiment.



## Activity 1

#### **Assessing Starch Digestion by Salivary Amylase**

- 1. From the general supply area, obtain a test tube rack, 10 test tubes, and a wax marking pencil. From the Activity 1 supply area, obtain a dropper bottle of distilled water and dropper bottles of maltose, amylase, and starch solutions.
- 2. In this experiment you will investigate the hydrolysis of starch to maltose by salivary amylase. You will need to be able to identify the presence of starch and maltose, the breakdown product of starch, to determine to what extent the enzymatic activity has occurred. Thus controls must be prepared to provide a known standard against which comparisons can be made. Starch decreases and sugar increases as hydrolysis occurs, according to the following formula:

Starch + watermaltose

<sup>\*</sup>The alpha-amylase must be a low-maltose preparation for good results.

<sup>&</sup>lt;sup>†</sup>Prepare by adding 1 g starch to 100 ml distilled water; boil and cool; add a pinch of salt (NaCl). Prepare fresh daily.

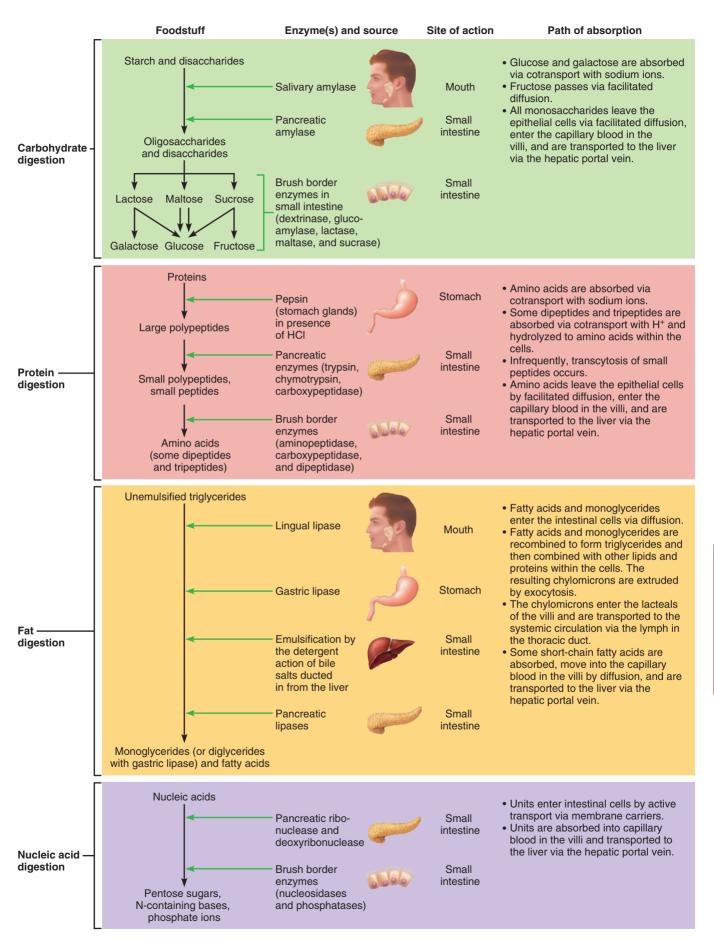


Figure 39.1 Flowchart of digestion and absorption of foodstuffs.

Two students should prepare the controls (tubes 1A to 3A) while the other two prepare the experimental samples (tubes 4A to 6A).

- Mark each tube with a wax pencil and load the tubes as indicated in the Activity 1 chart below, using 3 drops (gtt) of each indicated substance.
- Place tubes in the incubation conditions listed in the Activity 1 chart below for approximately 1 hour. Shake the rack gently from time to time to keep the contents evenly mixed.
- At the end of the hour, perform the amylase assay described below.
- While these tubes are incubating, proceed to Physical Processes: Mechanisms of Food Propulsion and Mixing (p. 606). Be sure to monitor the time so as to complete this activity as needed.

#### Amylase Assay

- 1. After one hour, obtain a spot plate and dropper bottles of Lugol's iodine solution (for the IKI, or iodine, test) and Benedict's solution from the Activity 1 supply area. Set up your boiling water bath using a hot plate, boiling chips, and a 250-ml beaker.
- 2. While the water is heating, mark six depressions of the spot plate 1A–6A (*A* for amylase) for sample identification.
- 3. Using a pipet, transfer a drop of the sample from each of the tubes 1A–6A into the appropriately numbered spot. Into each sample drop, place a drop of Lugol's iodine (IKI) solution. A blue-black color indicates the presence of

starch and is referred to as a **positive starch test**. If starch is not present, the mixture will not turn blue, which is referred to as a **negative starch test**. Record your results (+ for positive, – for negative) in the Activity 1 chart and on the board.

4. Into the remaining mixture in each tube, place 3 drops of Benedict's solution. Put each tube into the beaker of boiling water for about 5 minutes. If a green-to-orange precipitate forms, maltose is present; this is a **positive sugar test**. A **negative sugar test** is indicated by no color change. Record your results in the Activity 1 chart and on the board.

## WHY THIS Lactose Intolerance

Lactose is a disaccharide found in milk and, to a lesser extent, in other dairy products. Lactose is hydrolyzed into glucose and galactose by the enzyme lactase. Infants usually produce lactase, but many individuals become lactose intolerant (lactase nonpersistent) as they mature. When lactose-intolerant individuals ingest dairy products, the bacteria in their colon react with lactose, producing gas, which causes bloating, flatulence, abdominal cramps, and diarrhea. Individuals who continue to produce lactase into adulthood are lactase-persistent and are able to metabolize lactose normally. Ingestion of probiotic bacteria has been found to reduce the symptoms of lactose intolerance, possibly because probiotic bacteria that produce their own lactase replace the gas-producing bacterial flora in the colon. ■

Activity 1: Salivary Amylase Digestion of Starch						
Tube no.	1A	2A	3A	4 <b>A</b>	5 <b>A</b>	6A
Additives (3 gtt ea)		starch.	maltose,	Boil amylase 4 min, then add starch		
	amylase, water	water	water	amylase boiled amylase, starch	amylase, starch	amylase, starch
Incubation condition	37°C	37°C	37°C	37°C	37°C	0°C
IKI test (color change)						
Result: (+) or (-)						
Benedict's test (color change)						
Result: (+) or (-)						

#### Additive key:

- = Amylase
  - se = Starch
- = Maltose
- = Water

## **Protein Digestion by Trypsin**

**Trypsin,** an enzyme produced by the pancreas, hydrolyzes proteins to small peptides. BAPNA (*N*-alpha-benzoyl-arginine-*p*-nitroanilide) is a synthetic trypsin substrate consisting of a dye covalently bound to an amino acid. Trypsin

hydrolysis of BAPNA cleaves the dye molecule from the amino acid, causing the solution to change from colorless to bright yellow. The color change from clear to yellow is direct evidence of hydrolysis by trypsin.

## Activity 2

#### **Assessing Protein Digestion by Trypsin**

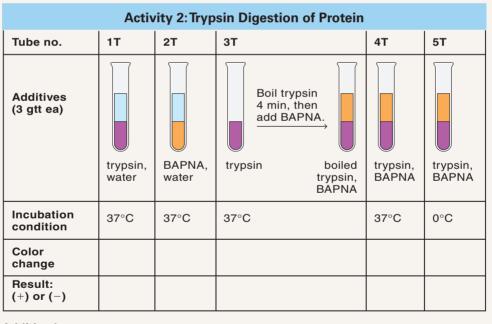
- 1. From the general supply area, obtain five test tubes and a test tube rack, and from the Activity 2 supply area get a dropper bottle of trypsin and one of BAPNA. Bring these items to your bench.
- 2. Two students should prepare the controls (tubes 1T and 2T) while the other two prepare the experimental samples (tubes 3T to 5T).
- Mark each tube with a wax pencil, and load the tubes as indicated in the **Activity 2 chart**, using 3 drops (gtt) of each indicated substance.
- Place all tubes in a rack in the appropriate water bath for approximately 1 hour. Shake the rack occasionally to keep the contents well mixed.

At the end of the hour, examine the tubes for the results of the trypsin assay (detailed below).

#### Trypsin Assay

Since BAPNA is a synthetic color-producing substrate, the presence of yellow color indicates a **positive hydrolysis test**; the dye molecule has been cleaved from the amino acid. If the sample mixture remains clear, a **negative hydrolysis test** has occurred.

Record the results in the Activity 2 chart and on the board.



#### Additive key:

= Trypsin =

= BAPNA

= Water

# Pancreatic Lipase Digestion of Fats and the Action of Bile

The treatment that fats and oils go through during digestion in the small intestine is a bit more complicated than that of carbohydrates or proteins—pretreatment with bile to physically emulsify the fats is required. Hence, two sets of reactions occur.

First:

Then:

Fat/oil droplets  $\frac{lipase}{(digestion)}\!\!>\!$  monoglycerides and fatty acids

The term **pancreatin** describes the enzymatic product of the pancreas, which includes enzymes that digest proteins, carbohydrates, nucleic acids, and fats. It is used here to investigate the properties of **pancreatic lipase**, which hydrolyzes fats and oils to their component monoglycerides and two fatty acids.

Fats/oils bile (emulsification) minute fat/oil droplets

Since fatty acids are organic acids, they acidify solutions, decreasing the pH. An easy way to recognize that digestion is ongoing or completed is to test pH. You will be using a pH indicator called *litmus blue* to follow these changes; it changes from blue to pink as the test tube contents become acidic.

## Activity 3

#### **Demonstrating the Emulsification Action of Bile and Assessing Fat Digestion by Lipase**

- 1. From the general supply area, obtain nine test tubes and a test tube rack, plus one dropper bottle of each of the solutions in the Activity 3 supply area.
- 2. Although *bile*, a secretory product of the liver, is not an enzyme, it is important to fat digestion because of its emulsifying action. It physically breaks down large fat particles into smaller ones. Emulsified fats provide a larger surface area for enzymatic activity. To demonstrate the action of bile on fats, prepare two test tubes and mark them 1E and 2E (*E* for emulsified fats).
- To tube 1E, add 20 drops of water and 4 drops of vegetable oil.
- To tube 2E, add 20 drops of water, 4 drops of vegetable oil, and a pinch of bile salts.
- Cover each tube with a small square of Parafilm, shake vigorously, and allow the tubes to stand at room temperature.

After 10 to 15 minutes, observe both tubes. If emulsification has not occurred, the oil will be floating on the surface of the water. If emulsification has occurred, the fat droplets will be suspended throughout the water, forming an emulsion.

In which tube has emulsification occurred? \_\_\_\_\_

- **3.** Two students should prepare the controls (1L and 2L, *L* for lipase) while the other two students in the group set up the experimental samples (3L to 5L, 4B, and 5B, where *B* is for bile), as illustrated in the **Activity 3 chart**.
- Mark each tube with a wax pencil and load the tubes using 5 drops (gtt) of each indicated solution.
- Place a pinch of bile salts in tubes 4B and 5B.
- Cover each tube with a small square of Parafilm, and shake to mix the contents of the tube.
- Remove the Parafilm, and place all tubes in a rack in the appropriate water bath for approximately 1 hour.
   Shake the test tube rack from time to time to keep the contents well mixed.
- At the end of the hour, perform the lipase assay below.

#### Lipase Assay

Fresh cream provides the fat substrate for this assay; add litmus powder to it to make litmus cream. The basis of this assay is a pH change that is detected by the litmus powder indicator. Alkaline or neutral solutions containing litmus are blue but will turn reddish in the presence of acid. If digestion occurs, the fatty acids produced will turn the litmus cream from blue to pink. Because the effect of

Activity 3: Pancreatic Lipase Digestion of Fats							
Tube no.	no. 1L 2L 3L		4L	5L	4B	5B	
Additives (5 gtt ea)			Boil pancreatin 4 min, then add litmus cream.				
	pancreatin, water	litmus cream, water	pancreatin boiled pancreatin, litmus cream	pancreatin, litmus cream	pancreatin, litmus cream	pancreatin, litmus cream, bile salts	pancreatin, litmus cream, bile salts
Incubation condition	37°C	37°C	37°C	37°C	0°C	37°C	0°C
Color change							
Result: (+) or (-)							

#### Additive key:

= Pancreatin

= Litmus cream





hydrolysis by lipase is directly seen, additional assay reagents are not necessary.

1. To prepare a color control, add 0.1 N HCl drop by drop to tubes 1L and 2L (covering the tubes with a square of

Parafilm after each addition and shaking to mix) until the cream turns pink.

2. Record the color of the tubes in the Activity 3 chart and on the board.

## Activity 4

### **Reporting Results and Conclusions**

- 1. Share your results with the class as directed in the General Instructions (p. 600).
- 2. Suggest additional experiments, and carry out experiments if time permits.
- 3. Prepare a lab report for the experiments on digestion. (See Getting Started, on MasteringA&P.)



## Group Challenge

#### **Odd Enzyme Out**

The following boxes each contain four digestive enzymes. One of the listed enzymes does not share a characteristic that the other three do. Working in groups of three, discuss the characteristics of the enzymes in each group. On a separate piece of paper, one student will record the characteristics for each enzyme for the group. Discuss the possible candidates for the "odd enzyme". Once the group

has come to a consensus, circle the enzyme that doesn't belong with the others, and explain why it is singled out. Sometimes there may be multiple reasons why the enzyme doesn't belong with the others. Include as many as you can think of, but make sure it does not have the key characteristic shared by the other three.

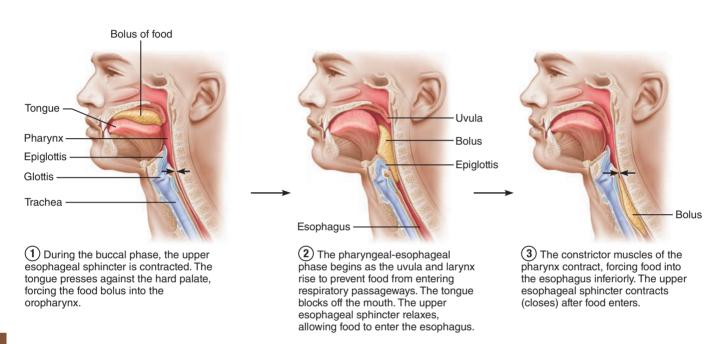
1. Which is the "odd enzyme"?	Why is it the odd one out?
Trypsin	
Carboxypeptidase	
Pepsin	
Chymotrypsin	
2. Which is the "odd enzyme"?	Why is it the odd one out?
Lactase	
Pepsin	
Aminopeptidase	
Trypsin	
3. Which is the "odd enzyme"?	Why is it the odd one out?
Maltase	
Pancreatic lipase	
Nucleosidase	
Dipeptidase	
4. Which is the "odd enzyme"?	Why is it the odd one out?
Sucrase	
Dextrinase	
Glucoamylase	
Chymotrypsin	

## Physical Processes: Mechanisms of Food Propulsion and Mixing

Although enzyme activity is a very important part of the overall digestion process, foods must also be processed physically (by chewing and churning) and moved by mechanical means along the tract if digestion and absorption are to be completed. Muscles are involved in producing the movements of foodstuffs along the gastrointestinal tract. Although we tend to think only of smooth muscles when visceral activities are involved, both skeletal and smooth muscles are involved in the physical processes. This fact is demonstrated by the simple activities that follow.

### **Deglutition (Swallowing)**

**Swallowing,** or **deglutition,** is largely the result of skeletal muscle activity and occurs in two phases: *buccal* (mouth) and *pharyngeal-esophageal*. The buccal phase (**Figure 39.2** step ①) is voluntarily controlled and initiated by the tongue. Once begun, the process continues involuntarily in the pharynx and esophagus through peristalsis, resulting in the delivery of the swallowed contents to the stomach (Figure 39.2 steps (2)–(3)).



**Figure 39.2 Swallowing.** The process of swallowing consists of voluntary (buccal) (step 1) and involuntary (pharyngeal-esophageal) phases (steps 2-3).

## Activity 5

#### **Observing Movements and Sounds of the Digestive System**

- 1. Obtain a pitcher of water, a stethoscope, a paper cup, an alcohol swab, and an autoclave bag in preparation for making the following observations.
- 2. While swallowing a mouthful of water, consciously note the movement of your tongue during the process. Record your observations.
- 3. Repeat the swallowing process while your laboratory partner watches the externally visible movements of your larynx. (This movement is more obvious in a male, since males have a larger Adam's apple.) Record your observations.

- What do these movements accomplish?
- 4. Before donning the stethoscope, your lab partner should clean the earpieces with an alcohol swab. Then, he or she should place the diaphragm of the stethoscope over your abdominal wall, approximately 2.5 cm (1 inch) below the xiphoid process and slightly to the left, to listen for sounds as you again take two or three swallows of water. There should be two audible sounds—one when the water splashes against the gastroesophageal sphincter,

and the second when the peristaltic wave of the esophagus arrives at the sphincter and the sphincter opens, allowing water to gurgle into the stomach. Determine, as accurately as possible, the time interval between these two sounds and record it below.

Interval between arrival of water at the sphincter and the

opening of the sphincter: \_\_\_\_\_ sec

This interval gives a fair indication of the time it takes for the peristaltic wave to travel down the 25 cm (10 inches) of the esophagus. (Actually the time interval is slightly less than it seems, because pressure causes the sphincter to relax before the peristaltic wave reaches it.)



Dispose of the used paper cup in the autoclave bag.

## **Segmentation and Peristalsis**

Although several types of movements occur in the digestive tract organs, peristalsis and segmentation are most important as mixing and propulsive mechanisms (**Figure 39.3**).

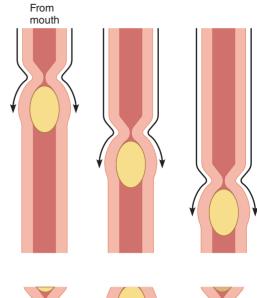
**Peristaltic movements** are the major means of propelling food through most of the alimentary canal. Essentially, they are waves of contraction followed by waves of relaxation that squeeze foodstuffs through the alimentary canal, and they are superimposed on segmental movements.

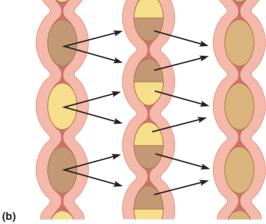
**Segmental movements** are local constrictions of the organ wall that occur rhythmically. They serve mainly to mix the foodstuffs with digestive juices and to increase the rate of absorption by continually moving different portions of the chyme over adjacent regions of the intestinal wall. However, segmentation is also an important means of food propulsion in the small intestine, and slow segmenting movements called haustral contractions are frequently seen in the large intestine.

## Activity 6

#### **Viewing Segmental and Peristaltic Movements**

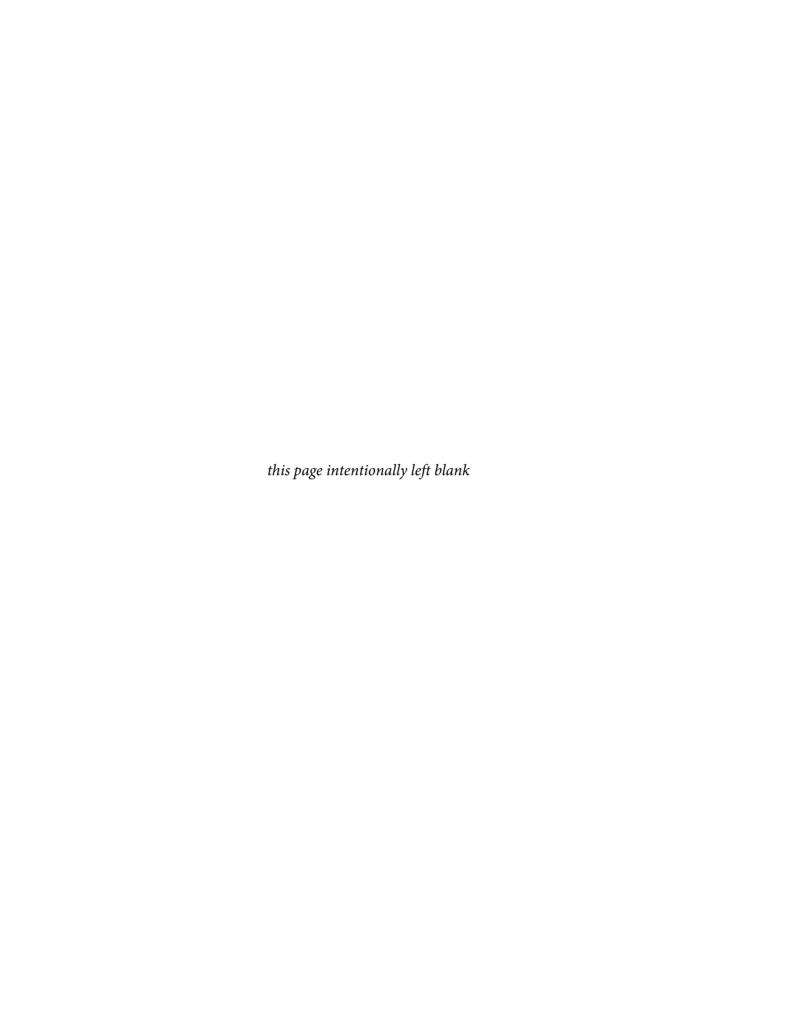
If a video showing some of the propulsive movements is available, go to a viewing station to view it before leaving the laboratory. Alternatively, use the *Interactive Physiology®* module on the Digestive System to observe gut motility.





(a)

Figure 39.3 Peristaltic and segmental movements of the digestive tract. (a) Peristalsis: neighboring segments of the intestine alternately contract and relax, moving food along the tract. (b) Segmentation: single segments of intestine alternately contract and relax. Because inactive segments exist between active segments, food mixing occurs to a greater degree than food movement. Peristalsis is superimposed on segmentation movements.





## **REVIEW SHEET**

# Digestive System Processes: Chemical and Physical

Name		L	_abTime/Date				
Digestion of Fo	oodstuffs: Enzy	matic Action	1				
	definitions with the pro						
Key: a. catalyst	b. control	c. enzyme d.	substrate				
	1. substance on w	vhich a catalyst works	6				
	2. biologic catalyst; protein in nature						
	3. increases the r	ate of a chemical read	ction without becoming	ing part of the product			
	4. provides a star	ndard of comparison	for test results				
2. List the three charac	cteristics of enzymes						
	digestive system are cla						
Enzyme	Organ producing it	Site of action	Substrate(s)	Optimal pH			
Salivary amylase							
Trypsin							
Lipase (pancreatic)							
	ucts of digestion for the						
fats:		and					
WHY THIS   6 How door							
MATTERS 6. How does	s the substrate for amyla	ase differ from the su	bstrate for lactase?				
How are	the substrates similar? _						

	7. Where does lactose hydrolysis occur for lactase-persistent individuals?						
	Where does lactose hydrolysis occur for lactose-intolerant individuals who have consumed probiotic bacte-						
	rial microflora?						
8.	You used several indicators or tests in the laboratory to determine the presence or absence of certain substances. Choose the correct test or indicator from the key to correspond to the condition described below.						
	Key: a. Lugol's iodine (IKI) b. Benedict's solution c. litmus d. BAPNA						
	1. used to test for protein hydrolysis, which was indicated by a yellow color						
	2. used to test for the presence of starch, which was indicated by a blue-black color						
	3. used to test for the presence of fatty acids, which was evidenced by a color change from blue to pink						
	4. used to test for the presence of reducing sugars (maltose, glucose) as indicated by a blue to green or orange color change						
9.	What conclusions can you draw when an experimental sample gives both a positive starch test and a positive maltose						
	test after incubation?						
	Why was 37°C the optimal incubation temperature?						
	Why did very little, if any, starch digestion occur in test tube 4A?						
	When starch was incubated with amylase at 0°C, did you see any starch digestion?						
	Why or why not?						
	Accume you have said to a group of your poors that amulass is capable of starch by dralysis to malters. If you had						
	Assume you have said to a group of your peers that amylase is capable of starch hydrolysis to maltose. If you had						
	not done control tube 1A, what objection to your statement could be raised?						
	What if you had not done tube 2A?						
10.	In the exercise concerning trypsin function, why was an enzyme assay such as Benedict's or Lugol's iodine (IKI), which						
	test for the presence of a reaction product, not necessary?						
	Why was tube 1T necessary?						
	Why was tube 2T necessary?						
	Trypsin is a protease similar to pepsin, the protein-digesting enzyme in the stomach. Would trypsin work well in the						
	stomach? Why?						

	/onospodaridos	•							
S	Substance	Mechanism of absorption	Blood	Lymph					
15.		absorption (passive or active $(\checkmark)$ whether the absorption wo teals).							
	Why?								
	this interval?								
14.	Assume you have been c	hewing a piece of bread for 5 or	r 6 minutes. How would you ex	spect its taste to change during					
13.		nzymes operate optimally at a pl acid. How is the proper pH for th							
	What specific experimen	tal conditions resulted in denat	ured enzymes?						
12.	The three-dimensional structure of a functional protein is altered by intense heat or nonphysiological pH even though peptide bonds may not break. Such inactivation is called denaturation, and denatured enzymes are nonfunctional. Explain why.								
	What role does bile play	What role does bile play in fat digestion?							
	How did the tubes contain	How did the tubes containing bile compare with those not containing bile?							
	Why is bile not considere	Why is bile not considered an enzyme?							
	Why did the litmus indicator change from blue to pink during fat hydrolysis?								
	Explain the reason for the difference.								
	In the procedure concerning pancreatic lipase digestion of fats and the action of bile salts, how did the appearance of tubes 1E and 2E differ?								
TI.	In the procedure concern	ing pancreatic lipase digestion	of fats and the action of bile	salts, how did the appearance					

Substance	Mechanism of absorption	Blood	Lymph
Monosaccharides			
Fatty acids and monoglycerides			
Amino acids			
Water			
Na+, Cl-, Ca <sup>2+</sup>			

1 <b>6</b> . P	eople on a strict diet to lose weigh	t begin to metabolize stored f	ats at an acc	elerated rate. How does this condition	
а	ffect blood pH?				
				t; bread = starch) from the mouth to the nd what specific enzymes are involved	
Γ					
L					
e e	ffect on the digestive process by ac	ting on other cells or structure f the digestive tract. For each l	es and causi hormone be	rmones into the blood. These exert aring them to release digestive enzymes low, note the organ producing the horsed.	
Hormone		Produced by		Target organ(s) and effects	
Sec	cretin				
Ga	strin				
Ch	olecystokinin				
-	vsical Processes: Mec	hanisms of Food F	Propulsi	on and Mixing	
1 <b>9</b> . C	complete the following statements.  Swallowing, or _1_, occurs in tw	o phases—the <u>2</u>	1.		
	nd <u>3</u> . One of these phases, the <u>4</u> During the voluntary phase, the <u>5</u>		2.		
	ood into the back of the throat. Dur 6_rises to ensure that its passagev				
	ne epiglottis so that the ingested sune respiratory passageways. It is po				
	vater while standing on your head l arried along the esophagus involur				
		andstuffs on the 8	5		
	f <u>7</u> . The pressure exerted by the f phincter causes it to open, allowing		0.		
s		g the foodstuffs to			
s e o	phincter causes it to open, allowing nter the stomach.	y the foodstuffs to we movements that and <u>10</u> . One of these	6.		
s e o n ir	phincter causes it to open, allowing nter the stomach. The two major types of propulsi ccur in the small intestine are <u>9</u> a	the foodstuffs to  ve movements that and 10. One of these y mix the foods and to ing different parts of	6. 7.		
s e o n ir th	phincter causes it to open, allowing nter the stomach.  The two major types of propulsiccur in the small intestine are _9_ anovements, _11_, acts to continually ncrease the absorption rate by movements.	the foodstuffs to  ive movements that and _10 One of these ive mix the foods and to ing different parts of mucosa, but it has less	6. 7. 8.		