

The main functions of the respiratory system include:

1. Exchange of **O<sub>2</sub>** and **CO<sub>2</sub>**.
2. Voice production.
3. Regulation of **plasma pH**.
4. **Olfaction** (sensation of smell)
5. Infection (pathogen invasion) prevention.

**Respiration** is divided into 4 processes:

1. **Pulmonary ventilation** is the movement of air into/out of the **lungs**
2. **External respiration** is the movement of O<sub>2</sub> from the lungs to the **blood** and CO<sub>2</sub> from the blood to the lungs.
3. **Internal respiration** is the movement of O<sub>2</sub> from the blood to the **cell interior** and CO<sub>2</sub> from the cell interior to the blood.
4. **Cellular respiration** is the breakdown of **glucose, fatty acids** and **amino acids** that occurs in **mitochondria** and results in production of **ATP**. It requires O<sub>2</sub> and produces CO<sub>2</sub>. (Note that this type of cellular respiration, which requires O<sub>2</sub>, is known as “**aerobic metabolism**,” whereas breakdown of glucose that produces ATP but does not require O<sub>2</sub> is “**anaerobic metabolism**.”)

The structures of the respiratory system can be divided into the **upper respiratory tract** and the **lower respiratory tract**. The upper respiratory tract refers to the **nose, pharynx**, and their associated structures. The lower respiratory tract includes the **larynx, trachea, bronchi**, and **lungs**.

The respiratory system can also be separated into a **conducting zone** and a **respiratory zone**. The **conducting zone** refers to structures that transport air but play no role in gas exchange. It includes: **nasal cavity, nasopharynx, oropharynx, laryngopharynx, larynx, trachea, bronchi**, and all **bronchioles** except for **respiratory bronchioles**. These structures are involved in transporting, filtering, humidifying, and warming air.

The **respiratory zone** refers to structures where exchange of O<sub>2</sub> and CO<sub>2</sub> occurs. Sites of exchange are known as **alveoli** (sing. **alveolus**). All respiratory zone structures contain alveoli and include: **respiratory bronchioles, alveolar ducts, and alveolar sacs**.

The **nose** is divided into the **external nose** and the **internal nasal cavity**. Air enters the nasal cavity via the **nostrils** or **nares**. The nasal cavity is divided by the **nasal septum**. The nasal septum is formed anteriorly from **septal cartilage**, and posteriorly from the **perpendicular plate** of the **ethmoid bone**, and the **vomer**. The **sphenoid** and **ethmoid** bones form the roof of the nasal cavity. The **hard palate** forms the floor of the nasal cavity. It is composed of the **palatine processes** of the **maxillary bones** and the **horizontal plates** of the **palatine bones**. The lateral walls of the nasal cavity are composed of the maxillae as well as the 3 pairs of **nasal conchae**. The **superior** and **middle nasal conchae** are projections of the **ethmoid bone**. The **inferior nasal conchae** are bones of their own. Nasal conchae increase the surface area of the nasal cavity and make airflow turbulent. The turbulence decreases the velocity of airflow. The resulting

increased amount of time coupled with the large surface area allows the inspired air to be filtered, warmed, and humidified.

The majority of the nasal cavity is lined by **respiratory epithelium**. Respiratory epithelium is **pseudostratified columnar epithelium with goblet cells**. The **mucus** secreted by goblet cells, as well as by mucous glands, helps filter and trap inspired particulate matter. The moist mucus also contributes to the humidification of inspired air. Cilia help sweep mucus to the pharynx where it is swallowed. Respiratory epithelium is underlain by a dense vasculature. The blood helps warm inspired air. Mucus also contains lysozyme as well as immunoglobulins, which help prevent infection. The **olfactory epithelium** is located in the most superior region of the nasal cavity and is involved in olfaction. The nasal cavity is continuous with the nasopharynx via the **posterior nasal apertures**.

The **paranasal sinuses** are cavities within the bones surrounding the nasal cavity. There are a total of 5 sinuses: **ethmoid sinus**, **sphenoid sinus**, **frontal sinus**, and the paired **maxillary sinuses**. Paranasal sinuses contribute to mucus production, lighten the skull, and provide resonance during speech. Openings from the paranasal sinuses empty into the nasal cavity. The 2 **nasolacrimal ducts** also empty (drain tears) into the nasal cavity.

The **pharynx** is the common portion of the respiratory and digestive tracts. It receives air from the nasal cavity and food, drink, and air from the oral cavity. It's continuous with the resp. tract at the larynx and with the digest. tract at the esophagus. The pharynx is divided into 3 sections: **nasopharynx**, **oropharynx**, & **laryngopharynx**. The nasopharynx extends from the posterior nasal apertures to the end of the **soft palate** and is lined by respiratory epithelium. The **soft palate** is the partition btwn the nasopharynx and the oral cavity. It's primarily composed of skeletal muscle. The posterior-most portion that hangs down is the **uvula**. The soft palate and uvula flip up during swallowing and help prevent food/drink from entering the nasopharynx. On the lateral walls of the nasopharynx are the openings to the **auditory tubes** (a.k.a. **Eustachian tubes**). Each auditory tube connects the pharynx to a **middle ear cavity**. They ensure that air pressure within the middle ear cavities is equal to atmospheric pressure. The nasopharynx also contains the **pharyngeal tonsil**.

The oropharynx is inferior to the uvula and superior to the **epiglottis**. It's posterior to the oral cavity and is continuous with it at an arched region known as the **fauces**. 2 sets of tonsils (**palatine** and **lingual**) are located right nearby. It's lined by nonkeratinized stratified squamous epithelium. This provides the necessary protection since this region is a common pathway for food and air.

The laryngopharynx is inferior to the epiglottis and superior to the split between the larynx and the esophagus. It's lined by nonkeratinized stratified squamous epithelium b/c it is also a common pathway for food and air. It's continuous with the larynx inferiorly.

The **larynx** routes food and air down their correct passages. It contains the **vocal cords**, which function in voice production. The larynx is a tube made up of 9 cartilages

connected by membranes, ligaments, and muscles. The cartilages include **thyroid, cricoid, epiglottis** and 3 small, paired cartilages (**arytenoids, cuneiforms, and corniculates**). All the laryngeal cartilages are hyaline cartilage with the exception of the epiglottis, which is elastic cartilage. The thyroid cartilage is the largest and its midline **laryngeal prominence** is the “**Adam’s apple**.” Inferior to the thyroid is the signet ring-shaped cricoid cartilage. It forms the base of the larynx. The arytenoids articulate with the posterior superior border of the cricoid cartilage. The corniculates are attached to the superior edge of the arytenoids. The cuneiforms are found within a membrane just anterior to the corniculates. The epiglottis extends from the superior thyroid cartilage with its free flap abutting the tongue. During swallowing, the epiglottis tips and covers the entrance to the larynx and ensures that food enters the esophagus. Deep to the laryngeal mucosa in the lateral walls of the larynx are the **vocal ligaments**. They extend from the arytenoids cartilages to the posterior surface of the thyroid cartilage. These are the core of the **vocal folds** or **true vocal cords**. Vibration of the vocal ligaments creates sounds that are then modified by the tongue, lips, etc., to produce speech. Skeletal muscles move the arytenoids and other cartilages and change the position and tension of the vocal cords (this modifies the sounds produced). Superior to the vocal folds are the **vestibular folds** or **false vocal cords**. These play no role in voice production. They help prevent food particles from passing thru the larynx as well as provide lubrication for the vocal folds. The space between the vocal folds on the left and those on the right is known as the **glottis**. The larynx is lined by respiratory epithelium below the vocal folds. The remainder is lined by stratified squamous epithelium.

Continuous with the larynx inferiorly is the **trachea**. It is a membranous tube lined by respiratory epithelium and consisting of dense regular CT and smooth muscle reinforced by 15-20 C-shaped rings of cartilage. It extends from the larynx to the mediastinum, where it splits into 2 **primary bronchi**. The trachea is associated with copious mucus secretion. Its rings prevent it from collapsing during inspiration. The open portion of the C is posterior and houses the **trachealis muscle**. Its contraction can alter the radius of the trachea and it plays a large role in coughing. The lack of posterior cartilage is important b/c it provides the esophagus with room to expand when a large bolus of food is swallowed. The last tracheal cartilage is expanded and contains a very sensitive posterior projection known as the **carina**.

The trachea divides into 2 **primary bronchi**. The right primary bronchus is wider, shorter, and more vertical than the left. It’s more likely to be obstructed by a foreign object. Air reaching the bronchi has been significantly filtered, warmed, and humidified. Within the lungs, each primary bronchus divides into **secondary bronchi**. There are 3 secondary bronchi on the right (one for each of the 3 **lobes** of the right lung) and 2 secondary bronchi on the left (one for each of the 2 lobes of the left lung). Secondary bronchi divide to yield **tertiary bronchi** that then divide to yield **quaternary bronchi** and so forth until about 23 branchings have occurred. Once the passageways have a diameter <1mm they are known as **bronchioles**. The **terminal bronchioles** are the last bronchioles without **alveoli**. Bronchioles with alveoli are known as **respiratory bronchioles** and lead into **alveolar ducts**.

As the bronchial tree branches, its histology changes markedly:

1. Cartilage rings are replaced by cartilage plates, and within the bronchioles, cartilage is absent entirely.
2. Epithelium changes from pseudostratified columnar to simple columnar to simple cuboidal.
3. The number of cilia declines.
4. The number of goblet cells declines.
5. The relative amount of smooth muscle increases.

**Respiratory zone** structures are defined by the presence of thin-walled alveoli. Alveoli are the sites of gas exchange. They're first seen w/i respiratory bronchioles. There, the alveoli are not adjacent and are separated by a thicker epithelium. Respiratory bronchioles lead into alveolar ducts. In them, the alveoli are adjacent to one another. Alveolar ducts terminate as **alveolar sacs**, blind clusters of alveoli. There are about 300 million alveoli within the lungs.

The walls of the alveoli are made of simple squamous epithelial cells known as **type I alveolar cells**. Cobwebbing the external surface of the alveoli are **pulmonary capillaries**. These capillaries are lined by endothelium. O<sub>2</sub> and CO<sub>2</sub> are exchanged as they pass through both sets of simple squamous epithelia (alveolar and capillary) as well as the fused basal laminae between the 2. This structure is collectively known as the **respiratory membrane**. Its extreme thinness facilitates the diffusion of O<sub>2</sub> and CO<sub>2</sub>. Interspersed amongst the type I alveolar cells are **type II alveolar cells**. These cells function primarily in the production of **surfactant**, a chemical that helps prevent alveolar collapse. **Alveolar macrophages (dust cells)** monitor the surface of the alveoli. In addition to capillaries, alveoli are also covered by a network of **elastic fibers**.

The **lungs** occupy the entire **thoracic cavity** except for the mediastinum. The thoracic cavity is enclosed and bounded:

- Above by the upper ribs and tissues of the neck.
- At the sides by the ribs and intercostal muscles.
- At the back by the ribs and vertebral column.
- In front by the ribs, costal cartilages, and sternum.
- Below by the **diaphragm** (a strong dome-shaped sheet of skeletal muscle with a central tendon).

Each lung is cone-shaped, associated with its own **pleural cavity**, and connected to the mediastinum by bronchial and vascular attachments known as the **root of the lung**. Anterior, lateral, and posterior surfaces hug the ribs and form the **costal surfaces**. Deep to the clavicle is the **apex**, the narrow superior lung tip. On the medial side of each lung is an indentation known as the **hilum**. Pulmonary vessels, nerves, and lymphatics enter/exit at this point. The left lung is slightly smaller than the right and contains a concavity known as the **cardiac impression** – which much of the heart. The left lung has only 2 lobes – **superior** and **inferior** separated by an **oblique fissure**. The right lung has 3 lobes – **superior**, **middle**, and **inferior** separated by **oblique** and **horizontal fissures** respectively. Lungs largely consist of air space and elastic connective tissue.

The **pleurae** are the thin, double-layered serosa that covers each lung. The **parietal pleura** covers the thoracic wall, the superior surface of the diaphragm, and the mediastinum. It continues around the heart and between the lungs. At the hilum, the parietal pleura is continuous w/ the **visceral pleura**, which covers the external surface of the lungs themselves. The pleurae produce pleural fluid which fills the slit-like **pleural cavity** btwn them. Pleural fluid reduces friction and helps the parietal and visceral pleurae adhere to one another.

The basic mechanism of breathing includes 2 phases – **inspiration** and **expiration**. Air movement occurs when a pressure gradient exists between the air within the lung alveoli and the air in the surrounding atmosphere. There are 3 pressures vital for lung function:

1. **Atmospheric pressure** – pressure exerted by the air surrounding the body.
2. **Intrapulmonary pressure** – pressure exerted by the air within the alveoli.
3. **Intrapleural pressure** – pressure within the pleural cavity.

The lungs are naturally elastic and would have a tendency to collapse if there was not an opposing force keeping them open. The basis of the opposing force is provided by the presence of a pressure gradient between the alveoli and the pleural cavity. Intrapleural pressure is always lower than alveolar pressure. Thus, the air within the alveoli is always “attempting” to leave the alveoli and enter the pleural cavity. This prevents alveolar collapse. If intrapleural pressure equilibrates with alveolar pressure, this gradient is lost and lung collapse can occur. One way the pressure gradient can be lost is if the pleural cavity is opened to the external environment – due to a stab wound perhaps.

The relationship between pressure and volume is given by **Boyle’s Law**, which states that at constant temperature, the pressure of a gas varies inversely with its volume. Thus changes in lung pressure (i.e., the creation of gradients btwn the lungs and the atmosphere) can be achieved by changing lung volume. Changing lung volume is achieved by changing the volume of the thoracic cavity via skeletal muscle contraction.

The following is the sequence of the **quiet inspiratory process**:

1. **Respiratory centers** in the **ventral medulla oblongata** become active.
2. Signals are sent down the **phrenic nerve** to the **diaphragm** and down **intercostal nerves** to the **external intercostal muscles**.
3. Diaphragm and external intercostals contract.
4. Contraction of the diaphragm lengthens the thoracic cavity top to bottom. Contraction of the external intercostals lifts the ribs and sternum increasing the side-to-side and front-to-back dimensions of the thoracic cavity.
5. Volume of the thoracic cavity increases.
6. Lung volume increases.
7. Alveolar pressure decreases. Alveolar pressure is now  $<$  atmospheric pressure.
8. Air flows from the atmosphere into the alveoli until alveolar P = atmospheric P.

In **forced inspiration**, other muscles are involved so as to further increase thoracic volume (and further decrease alveolar pressure). Such muscles include the **scalenes** and **sternocleidomastoids** of the neck, and the **pectoralis minors** of the chest.

**Quiet expiration** is a passive process, (i.e., not powered by skeletal muscle contraction).

1. Phrenic and intercostal nerves cease firing.
2. Diaphragm and external intercostals relax.
3. The thoracic volume decreases.
4. Lung volume decreases.
5. Alveolar pressure increases. Alveolar pressure is now  $>$  atmospheric pressure.
6. Air flows from the alveoli into the atmosphere until alveolar  $P =$  atmospheric  $P$ .

In **forced expiration**, muscles contract in order to further reduce the size of thoracic cavity (and further increase alveolar pressure). Such muscles include the **latissimus dorsi**, **rectus abdominus**, **transverse abdominus**, **obliques**, and **internal intercostals**.

**Airway resistance** can sometimes affect airflow. It's normally insignificant due to the relatively large diameters of the air passages. However, during severe allergic reactions histamine causes contraction of bronchiolar smooth muscle. This decreases airway volume and increase airway resistance. During an asthma attack, vigorous bronchoconstriction can also occur. Mucus or accumulations of infectious material can increase airway resistance. Epinephrine causes relaxation of bronchiolar smooth muscle, increasing bronchiole diameter and decreasing airway resistance.

Another complicating factor is **surface tension**. Water molecules line the inner surfaces of the alveoli. These water molecules have a stronger attraction for one another than for the molecules of gas within the alveolar lumen. This high surface tension can lead to alveolar collapse. Collapsed alveoli require large amounts of energy to inflate during inspiration. Luckily, the type II alveolar cells produce the chemical surfactant. It decreases the cohesiveness of the water molecules and thus reduces alveolar surface tension and decreases the likelihood of alveolar collapse.

**Compliance** refers to the ability of the lungs to expand. The ease with which the lungs can expand facilitates efficient ventilation. Replacement of the elastic lung tissue with inelastic scar tissue as well as reduced surfactant production will decrease lung compliance. Too much compliance is undesirable as well.

Air is made up of 79% nitrogen, 20% oxygen, smaller amounts of carbon dioxide and water vapor, and minute amounts of other gases. The pressure exerted by atmospheric air is a sum of the pressures exerted by each individual gas in the air. Thus each gas in a mixture of gases exerts a certain amount of pressure. This is known as the **partial pressure** for that gas. *Individual gases tend to move from one place to another based on their partial pressure gradient.*

During gas exchange in the lungs the partial pressure of  $O_2$  in the alveoli is 104mmHg. The partial pressure of  $O_2$  in blood entering pulmonary capillaries is 40mmHg. The  $PO_2$

gradient favors flow of O<sub>2</sub> from alveolar air into the pulmonary capillary blood. Meanwhile, the partial pressure of CO<sub>2</sub> in the alveoli is 40mmHg. The partial pressure of CO<sub>2</sub> in blood entering the pulmonary capillaries is 45mmHg. The PCO<sub>2</sub> gradient favors flow of CO<sub>2</sub> from pulmonary capillary blood into the alveolar air.

At the systemic tissues, the situation is reversed. Arterial blood PO<sub>2</sub> is 104mmHg while tissue PO<sub>2</sub> is less than 40mmHg. The PO<sub>2</sub> gradient favors flow of O<sub>2</sub> from the systemic capillary blood into the interstitial fluid and tissue cells. Meanwhile, arterial blood PCO<sub>2</sub> is 40mmHg while tissue PCO<sub>2</sub> is greater than 45mmHg. The PCO<sub>2</sub> gradient favors flow of CO<sub>2</sub> from interstitial fluid and tissue cells into systemic capillary blood.

Notice that the partial pressure gradients for CO<sub>2</sub> are much smaller than the partial pressure gradients for O<sub>2</sub>. CO<sub>2</sub> does not require such a big gradient because it is much more **soluble** in water; thus it enters the plasma much more readily than does O<sub>2</sub>.

O<sub>2</sub> is carried by blood in 2 ways. 1.5% of the O<sub>2</sub> is simply dissolved in plasma. The other 98.5% is bound to **hemoglobin** within red blood cells.

Each Hb molecule can combine with up to 4 oxygen molecules. Hemoglobin with bound O<sub>2</sub> is **oxyhemoglobin**. Hemoglobin w/o bound O<sub>2</sub> is **reduced hemoglobin**. The loading and unloading of O<sub>2</sub> by hemoglobin is given by a single reversible equation: **Hb + O<sub>2</sub> ↔ HbO<sub>2</sub> + H<sup>+</sup>**. In the lungs (highPO<sub>2</sub>), the reaction runs from left to right. In the tissues (low PO<sub>2</sub>), the reaction runs from right to left. When Hb has 4 O<sub>2</sub> molecules bound to it, it's **saturated**. When Hb has less than 4 O<sub>2</sub> molecules bound to it, it's **unsaturated**. In the lungs (PO<sub>2</sub> is 104mmHg), Hb is fully saturated. In the tissues (PO<sub>2</sub> is 40mmHg), Hb is 75% saturated, meaning that, on average, each Hb molecule has 3 molecules of O<sub>2</sub> bound to it. Thus, substantial amounts of O<sub>2</sub> are still available in the venous blood. This provides a reserve for use during increased activity. During aerobic respiration, working muscle cells produce CO<sub>2</sub>, heat, and acid. These factors decrease the affinity that Hb has for O<sub>2</sub>, making Hb more likely to release O<sub>2</sub> to these working cells. Note that carbon monoxide binds to hemoglobin in the same binding site as oxygen, but far more tightly.

CO<sub>2</sub> is transported w/i blood in 3 ways. About 10% is dissolved in plasma. Another 20% is bound to hemoglobin. The equation for this combination is: **Hb + CO<sub>2</sub> ↔ HbCO<sub>2</sub>**. HbCO<sub>2</sub> is known as **carbaminohemoglobin**. CO<sub>2</sub> binds to Hb in a different place than O<sub>2</sub>. 70% is transported as part of the **bicarbonate ion** in plasma.

When CO<sub>2</sub> diffuses out of the tissue fluid, it enters the systemic capillary plasma and then the RBC. Within the RBC, CO<sub>2</sub> combines with water to form **carbonic acid**, which dissociates into a **bicarbonate ion** and a **hydrogen ion**. The equation is: **CO<sub>2</sub> + H<sub>2</sub>O ↔ H<sub>2</sub>CO<sub>3</sub> ↔ HCO<sub>3</sub><sup>-</sup> + H<sup>+</sup>**. This reaction is catalyzed by the enzyme **carbonic anhydrase**. Once bicarbonate is formed it diffuses out of the RBC into the plasma. As this occurs, a **chloride ion** diffuses from the plasma into the RBC. This maintains charge balance and is known as the **chloride shift**. At the pulmonary capillaries, the above events reverse themselves.

The primary control center for respiration is a cluster of neurons in the medulla oblongata known as the **ventral respiratory group (VRG)**. It contains both **inspiratory neurons** and **expiratory neurons**. When the inspiratory neurons fire, signals travel down the phrenic and intercostal nerves and excite the diaphragm and external intercostals – resulting in inspiration. When the expiratory neurons fire, the output to the diaphragm and external intercostals ceases and expiration occurs. The on/off cycle of both these types of neurons creates the basic respiratory rhythm – known as **eupnea**.

There is also a **dorsal respiratory group (DRG)**, which helps to integrate information from peripheral chemoreceptors and stretch receptors. It then inputs this info to the VRG. In the pons, we find the **pontine respiratory group**, which helps modify, breathing rhythm during sleep, talking, and exercise.

The primary factors that influence respiratory rate are the contents of the plasma and the cerebrospinal fluid. The main respiratory stimuli in order of importance are: **CSF pH**; **plasma PCO<sub>2</sub>** and **pH**; and the **plasma PO<sub>2</sub>**. **Central chemoreceptors** on the medulla measure the [H<sup>+</sup>] of the CSF. **Peripheral chemoreceptors** (located primarily in the aortic arch and carotid sinus) measure the PCO<sub>2</sub>, [H<sup>+</sup>], and PO<sub>2</sub> of the plasma.

When plasma CO<sub>2</sub> levels rise, CSF CO<sub>2</sub> levels rise (since CO<sub>2</sub> can easily diffuse thru the blood-brain barrier and enter the CSF from the plasma). Within the CSF, CO<sub>2</sub> combines with H<sub>2</sub>O to form HCO<sub>3</sub><sup>-</sup> and H<sup>+</sup>. Thus, as plasma CO<sub>2</sub> levels rise, CSF pH will decrease. A drop in CSF pH can be quite damaging. Luckily, the medullary chemoreceptors sense the low pH and initiate an ↑ in respiratory rate and depth to rid the body of excess CO<sub>2</sub>. The lowering in pH of the CSF as caused by the rise of CSF PCO<sub>2</sub> as caused by the rise of plasma PCO<sub>2</sub> is the most powerful respiratory stimulus.

When plasma PCO<sub>2</sub> reaches its threshold or when plasma pH drops to its threshold, the peripheral chemoreceptors are activated and they signal the medulla (via the **vagus** and **glossopharyngeal nerves**) to increase respiratory rate and depth. Recall how CO<sub>2</sub> combines with H<sub>2</sub>O to form bicarbonate and a hydrogen ion. This is why ↑ plasma CO<sub>2</sub> results in ↑ plasma H<sup>+</sup> and ↓ plasma pH.

Arterial PO<sub>2</sub> must drop substantially (i.e., to below 60mmHg) before the chemoreceptors sensitive to PO<sub>2</sub> play a role in activating the respiratory centers. In people who chronically retain CO<sub>2</sub>, (perhaps due to **emphysema** or chronic bronchitis) the peripheral PCO<sub>2</sub> receptors become unresponsive. In such individuals, arterial PO<sub>2</sub> levels play a significant role in respiration regulation.

Factors outside the brainstem also influence respiratory rate and depth. The presence of irritants in the respiratory tract can lead to coughing and sneezing as well as cause other changes in rate and depth. The **Hering-Breuer reflex** describes another factor. As forced inhalation proceeds, the lungs stretch. Excess stimulation of lung stretch receptors inhibits the medullary inspiratory neurons and activates medullary expiratory neurons. The **hypothalamus** exerts effects as well. An example is the changes in respiratory rate and depth associated with changes in body T°. In general, as body T° ↑, respiratory rate

and depth  $\uparrow$ ; and as body  $T^\circ \downarrow$ , respiratory rate and depth  $\downarrow$ . Changes in BP affect respiration rate and depth. When BP falls, the respiratory rate increases. When BP rises, the respiratory rate declines. The **cerebral cortex** also influences respiratory rate, e.g., our ability to hold our breath (for a limited time at least). Respiratory rhythm is of course altered by laughing, yawning, crying, speech, hiccupping, etc.

It should be noted that the respiratory system can both cause and correct disturbances in plasma pH. The respiratory system can cause plasma pH disturbances in 2 ways. During **hyperventilation**, plasma  $CO_2$  drops. This causes a decrease in plasma  $H^+$  and thus, an increase in plasma pH. If plasma pH rises above normal levels, it is known as **plasma alkalosis**. In this case, we call it **respiratory alkalosis**. During **hypoventilation**, plasma  $CO_2$  increases. This causes an increase in plasma  $H^+$  and thus, a decrease in plasma pH. If plasma pH drops below normal levels, it is known as **plasma acidosis**. In this case we call it **respiratory acidosis**.

Plasma pH disturbances are not always respiratory in origin. Any alkalosis not caused by respiratory malfunctions is referred to as **metabolic alkalosis**. Causes of metabolic alkalosis include:

1. Vomiting – due to the loss of gastric HCl.
2. Ingestion of excessive antacids (e.g., sodium bicarbonate).
3. Constipation – decreased loss of  $HCO_3^-$  in feces.

In response to metabolic alkalosis, respiratory rate and depth will decrease. This will increase plasma  $CO_2$  levels and decrease plasma pH.

Any acidosis not caused by respiratory malfunctions is referred to as **metabolic acidosis**. Causes of metabolic acidosis include:

1. Severe diarrhea – excess  $HCO_3^-$  loss in feces.
2. Renal disease – failure of kidneys to secrete acids in urine.
3. Excess alcohol ingestion. (Byproducts of alcohol metabolism are acidic.)
4. Starvation – when the body begins to break down fat and muscle protein reserves for energy, acidic metabolites (**ketone bodies**) are produced.
5. Untreated **diabetes mellitus** – fats are broken down and ketone bodies are produced.

In response to metabolic acidosis, respiratory rate and depth will increase. This will decrease plasma  $CO_2$  levels and increase plasma pH.