**بسم الله الرحمن الرحيم**

**Lecture 10, 11- Neurophysioloy Dr. Noor Jawad**

**2nd stage 2021**

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**Cerebral Blood Flow, Cerebrospinal Fluid, and Brain Metabolism**

***Objective***

***1. What is the effect of Carbon Dioxide and Hydrogen Ions on Cerebral Blood Flow?***

***2. Autoregulation of Cerebral Blood Flow When the Arterial Pressure Changes?***

**Cerebral Blood Flow**

**Normal Rate of Cerebral Blood Flow**

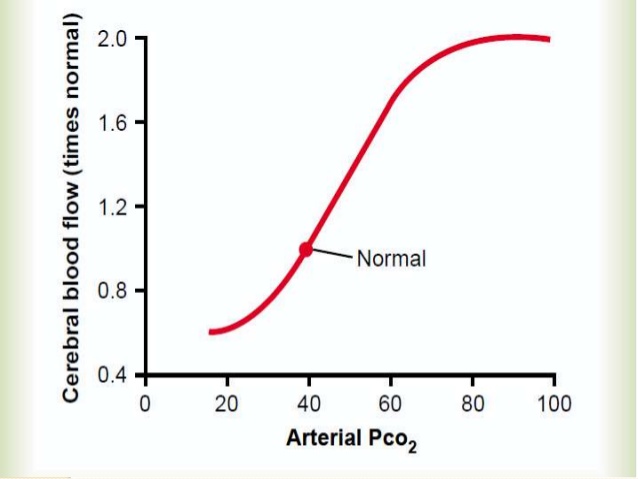
Normal blood flow through the brain of the adult person averages 50 to 65 milliliters per 100 grams of brain tissue per minute. For the entire brain, this amounts to 750 to 900 ml/min, or 15 per cent of the resting cardiac output.

**Regulation of Cerebral Blood Flow**

As in most other vascular areas of the body, cerebral blood flow is highly related to metabolism of the tissue. At least three metabolic factors have potent effects in controlling cerebral blood flow: (1) carbon dioxide concentration, (2) hydrogen ion concentration, and (3) oxygen concentration.

**Increase of Cerebral Blood Flow in Response to Excess Carbon Dioxide or Excess Hydrogen Ion Concentration.**

An increase in carbon dioxide concentration in the arterial blood perfusingthe brain greatly increases cerebral blood flow. This is demonstrated in Figure below, which shows that a 70 per cent increase in arterial PCO2 approximately doubles cerebral blood flow.



**Figure show Relationship between arterial PCO2 and cerebral blood flow**

Carbon dioxide is believed to increase cerebral blood flow by combining first with water in the body fluids to form carbonic acid, with subsequent dissociation of this acid to form hydrogen ions. The hydrogen ions then cause vasodilation of the cerebral vessels—the dilation being almost directly proportional to the increase in hydrogen ion concentration up to a blood flow limit of about twice normal.

Any other substance that increases the acidity of the brain tissue, and therefore increases hydrogen ion concentration, will likewise increase cerebral blood flow. Such substances include lactic acid, pyruvic acid, and any other acidic material formed during the course of tissue metabolism.

**Importance of Cerebral Blood Flow Control by Carbon Dioxide and Hydrogen Ions.**

Increased hydrogen ion concentration greatly depresses neuronal activity. Therefore, it is fortunate that an increase in hydrogen ion concentration also causes an increase in blood flow, which in turn carries hydrogen ions, carbon dioxide, and other acidforming substances away from the brain tissues. Loss of carbon dioxide removes carbonic acid from the tissues; this, along with removal of other acids, reduces the hydrogen ion concentration back toward normal.

Thus, this mechanism helps maintain a constant hydrogen ion concentration in the cerebral fluids and thereby helps to maintain a normal, constant level of neuronal activity.

**Autoregulation of Cerebral Blood Flow When the Arterial Pressure Changes.**

Cerebral blood flow is “autoregulated” extremely well between arterial pressure limits of 60 and 140 mm Hg. That is, mean arterial pressure can be decreased acutely to as low as 60 mm Hg or increased to as high as 140 mm Hg without significant change in cerebral blood flow. And, in people who have hypertension, autoregulation of cerebral blood flow occurs even when the mean arterial pressure rises to as high as 160 to 180 mm Hg.

**Role of the Sympathetic Nervous System in Controlling Cerebral**

**Blood Flow.**

The cerebral circulatory system has strong sympathetic innervation that passes upward from the superior cervical sympathetic ganglia in the neck and then into the brain along with the cerebral arteries. This innervation supplies both the large brain arteries and the arteries that penetrate into the substance of the brain.

However, transection of the sympathetic nerves or mild to moderate stimulation of them usually causes very little change in cerebral blood flow because the blood flow autoregulation mechanism can override the nervous effects.

***Clinical points***

**Cerebral “Stroke” Occurs When Cerebral Blood Vessels Are Blocked**

Almost all elderly people have blockage of some small arteries in the brain, and as many as 10 per cent eventually have enough blockage to cause serious disturbance of brain function, a condition called a “stroke.”

Most strokes are caused by arteriosclerotic plaques that occur in one or more of the feeder arteries to the brain. The plaques can activate the clotting mechanism of the blood, causing a blood clot to occur and block blood flow in the artery, thereby leading to acute loss of brain function in a localized area. One of the most common types of stroke is blockage of the *middle cerebral artery* that supplies the mid portion of one brain hemisphere.

**Cerebrospinal Fluid System**

.The entire cerebral cavity enclosing the brain and spinal cord has a capacity of about 1600 to 1700 milliliters; about 150 milliliters of this capacity is occupied by *cerebrospinal* *fluid* and the remainder by the brain and cord.

**Contents of CSF**

The CSF is derived from [blood plasma](https://en.wikipedia.org/wiki/Blood_plasma) and is largely similar to it, except that CSF is nearly protein-free compared with plasma and has some different [electrolyte](https://en.wikipedia.org/wiki/Electrolyte) levels. Owing to the way it is produced, CSF has a higher [chloride](https://en.wikipedia.org/wiki/Chloride) level than plasma, and an equivalent [sodium](https://en.wikipedia.org/wiki/Sodium) level.

CSF contains approximately 0.3% plasma proteins, or approximately 15 to 40 mg/dL, depending on sampling site. CSF is normally free of [red blood cells](https://en.wikipedia.org/wiki/Red_blood_cell), and at most contains only a few [white blood cells](https://en.wikipedia.org/wiki/White_blood_cell). Any white blood [cell count](https://en.wikipedia.org/wiki/Cell_counting) higher than this constitutes [pleocytosis](https://en.wikipedia.org/wiki/Pleocytosis" \o "Pleocytosis).

**Three Main Functions of CSF**

* CSF protects brain and spinal cord from trauma.
* CSF supplies nutrients to nervous system tissue.
* CSF removes waste products from cerebral metabolism.

**Cushioning Function of the Cerebrospinal Fluid**

A major function of the cerebrospinal fluid is to cushion the brain within its solid vault. The brain and the cerebrospinal fluid have about the same specific gravity (only about 4 per cent different), so that the brain simply floats in the fluid. Therefore, a blow to the head, if it is not too intense, moves the entire brain simultaneously with the skull, causing no one portion of the brain to be momentarily contorted by the blow.

**Formation, Flow, and Absorption of Cerebrospinal Fluid**

Cerebrospinal fluid is formed at a rate of about 500 milliliters each day, which is three to four times as much as the total volume of fluid in the entire cerebrospinal fluid system.

About two thirds or more of this fluid originates as *secretion from the choroid plexuses* in the four ventricles, *mainly in the two lateral ventricles*. Additional small amounts of fluid are secreted by the ependymal surfaces of all the ventricles and by the arachnoidal membranes.

The fluid secreted in the *lateral ventricles* passes first into the *third* *ventricle*; then, after addition of minute amounts of fluid from the third ventricle, it flows downward along the *aqueduct of Sylvius* into the *fourth ventricle*, where still another minute amount of fluid is added.

Finally, the fluid passes out of the fourth ventricle through three small openings, *two lateral foramina of Luschka* and a *midline foramen of Magendie*, entering the *cisterna* *magna*, a fluid space that lies behind the medulla and beneath the cerebellum.

**Secretion by the Choroid Plexus.**

Secretion of fluid into the ventricles by the choroid plexus depends mainly on active transport of sodium ions through the epithelial cells lining the outside of the plexus .

The sodium ions in turn pull along large amounts of chloride ions as well because the positive charge of the sodium ion attracts the chloride ion’s negative charge. The two of these together increase the quantity of osmotically active sodium chloride in the cerebrospinal fluid, which then causes almost immediate osmosis of water through the membrane, thus providing the fluid of the secretion.

**Cerebrospinal Fluid Pressure**

The normal pressure in the cerebrospinal fluid system *when one is lying in a horizontal position* averages 130 millimeters of water (10 mm Hg), although this may be as low as 65 millimeters of water or as high as 195 millimeters of water even in the normal healthy person.

**Regulation of Cerebrospinal Fluid Pressure by the Arachnoidal Villi.**

The normal rate of cerebrospinal fluid formation remains very nearly constant, so that changes in fluid formation are seldom a factor in pressure control. Conversely, the arachnoidal villi function like “valves” that allow cerebrospinal fluid and its contents to flow readily into the blood of the venous sinuses while not allowing blood to flow backward in the opposite direction.

Then, if the cerebrospinal fluid pressure rises still higher, the valves open more widely, so that under normal conditions, the cerebrospinal fluid pressure almost never rises more than a few millimeters of mercury higher than the pressure in the cerebral venous sinuses.

Conversely, in disease states, the villi sometimes become blocked by large particulate matter, by fibrosis, or by excesses of blood cells that have leaked into the cerebrospinal fluid in brain diseases. Such blockage can cause high cerebrospinal fluid pressure, .

**Blood–Cerebrospinal Fluid and Blood-Brain Barriers**

It has already been pointed out that the concentrations of several important constituents of cerebrospinal fluid are not the same as in extracellular fluid elsewhere in the body.

Furthermore, many large molecular substances hardly pass at all from the blood into the cerebrospinal fluid or into the interstitial fluids of the brain, even though these same substances pass readily into the usual interstitial fluids of the body. Therefore, it is said that barriers, called the *blood–cerebrospinal fluid* *barrier* and the *blood-brain barrier*, exist between the blood and the cerebrospinal fluid and brain fluid, respectively.

Barriers exist both at the choroid plexus and at the tissue capillary membranes in essentially all areas of the brain parenchyma *except in some areas of the hypothalamus*, *pineal gland*, where substances diffuse with greater ease into the tissue spaces. The ease of diffusion in these areas is important because they have sensory receptors that respond to specific changes in the body fluids, such as changes in osmolality and in glucose concentration, as well as receptors for peptide hormones that regulate thirst, such as angiotensin II.

The blood-brain barrier also has specific carrier molecules that facilitate transport of hormones, such as leptin, from the blood into the hypothalamus where they bind to specific receptors that control other functions such as appetite and sympathetic nervous system activity.

In general, the blood–cerebrospinal fluid and blood brain barriers are highly permeable to water, carbon dioxide, oxygen, and most lipid-soluble substances such as alcohol and anesthetics; slightly permeable to electrolytes such as sodium, chloride, and potassium; and almost totally impermeable to plasma proteins and most non–lipid-soluble large organic molecules.

**Brain Metabolism**

Like other tissues, the brain requires oxygen and food nutrients to supply its metabolic needs.

**Total Brain Metabolic Rate and Metabolic Rate of Neurons.**

Under resting but awake conditions, the metabolism of the brain accounts for about 15 per cent of the total metabolism in the body, even though the mass of the brain is only 2 per cent of the total body mass. Therefore, under resting conditions, brain metabolism per unit mass of tissue is about 7.5 times the average metabolism in non–nervous system tissues.

Most of this excess metabolism of the brain occurs in the neurons, not in the glial supportive tissues. The major need for metabolism in the neurons is to pump ions through their membranes, mainly to transport sodium and calcium ions to the outside of the neuronal membrane and potassium ions to the interior.

Each time a neuron conducts an action potential, these ions move through the membranes, increasing the need for additional membrane transport to restore proper ionic concentration differences across the neuron membranes. Therefore, during excessive brain activity, neuronal metabolism can increase as much as 100 to 150 per cent.

**Special Requirement of the Brain for Oxygen—Lack of Significant Anaerobic Metabolism.**

Most tissues of the body can live without oxygen for several minutes and some for as long as 30 minutes. During this time, the tissue cells obtain their energy through processes of anaerobic metabolism, which means release of energy by partially breaking down glucose and glycogen but without combining these with oxygen.

This delivers energy only at the expense of consuming tremendous amounts of glucose and glycogen. However, it does keep the tissues alive.

The brain is not capable of much anaerobic metabolism. One of the reasons for this is the high metabolic rate of the neurons, so that most neuronal activity depends on second-by-second delivery of oxygen from the blood.

Putting these factors together, one can understand why sudden cessation of blood flow to the brain or sudden total lack of oxygen in the blood can cause unconsciousness within 5 to 10 seconds.

**Under Normal Conditions Most Brain Energy Is Supplied by Glucose.**

Under normal conditions, almost all the energyused by the brain cells is supplied by glucose derivedfrom the blood. As is true for oxygen, most of this isderived minute by minute and second by second fromthe capillary blood, with a total of only about a 2-minutesupply of glucose normally stored as glycogen in theneurons at any given time.

A special feature of glucose delivery to the neurons is that its transport into the neurons through the cell membrane is not dependent on insulin, even though insulin is required for glucose transport into most other body cells.

Therefore, in patients who have serious diabetes with essentially zero secretion of insulin, glucose still diffuses readily into the neurons—which is most fortunate in preventing loss of mental function in diabetic patients.

Yet, when a diabetic patient is overtreated with insulin, the blood glucose concentration can fall extremely low because the excess insulin causes almost all the glucose in the blood to be transported rapidly into the vast numbers of insulin-sensitive non-neural cells throughout the body, especially into muscle and liver cells.

When this happens, not enough glucose is left in the blood to supply the neurons properly, and mental function does then become seriously deranged, leading sometimes to coma and even more often to mental imbalances and psychotic disturbances— all caused by overtreatment with insulin.

**Thank you**

***References : Guyton and Hall textbook of medical physiology, thirteen edition***