

1) The following contractile proteins participate on the process of myocardial muscle contraction: p 94

- ☒ a. actin, myosin, sarcoplasmic reticulum
- ☒ b. myosin, mitochondria and sarcoplasmic reticulum
- ☒ c. actin, myosin, tropomyosin, troponin
- d. actin, myosin, troponin, tropomyosin and sarcoplasmic reticulum
- e. troponin and calcium
- ☒ f. actin, myosin, sarcoplasmic reticulum and sarcolemma

2) Under the aerobic conditions, the heart muscle gains energy mainly by:

- ☒ a. the beta-oxidation of the fatty acids
- b. anaerobic glycolysis
- ☒ c. aerobic glycolysis
- d. pentose cycle
- e. glycogenolysis
- f. aerobic glycolysis and glycogenolysis

3) Myocardium uses ATP:

- a. exclusively for contraction
- b. exclusively for relaxation
- ☒ c. mainly for transmembrane ion transport
- d. mainly for basal metabolism
- ☒ e. mainly for the action of contraction and relaxation
- f. mainly for preservation of subcellular structures

4) Afterload is determined by following factors: p 99

- a. preload and volume of blood in venous bed
- b. preload and peripheral arterial resistance
- c. preload and contractility
- d. the length of the muscular fibers and peripheral venous resistance
- ☒ e. arterial compliance, arterial system blood volume and peripheral arterial resistance
- f. volume of blood in the left ventricle at the end of diastole

5) The most frequent causes of heart failure are: p 104

- a. hypertension and valvular diseases
- b. hypertension and heart hypertrophy
- ☒ c. hypertension and ischemic heart disease
- d. ischemic heart disease and cardiomyopathy
- e. myocardial infarction and diastolic dysfunction
- f. myocardial infarction and hypovolemic shock

6) Forward failure means: p 104

- a. failure of the arterial bed
- b. failure of the venous bed
- ☒ c. signs and symptoms developing as a result of insufficient tissue perfusion
- d. breathlessness
- e. the disturbance of the skeletal muscle function (mainly fatigue)
- ☒ f. failure of the left ventricular afterload

7) Heart failure is: p.103

- a. the state, shortening the life expectancy in 50%
- b. the state, shortening the life to five years
- ☒ c. the syndrome, being the result of variable cardiac diseases
- d. the most frequent pathologic unit of the cardiovascular system
- e. the disease, which always causes death
- f. the arrest of the heart

8) In central cyanosis: p.110

- a. the acral parts of the body are cold
- b. the upper extremities are cold, lower are warm
- c. the lower extremities are cold, upper are warm
- d. the extremities are cold but mucous layers are warm
- ☒ e. the temperature of the acral parts of the body is normal
- ~~f.~~ the amount of the the reduced hemoglobin in the capillary blood is more than 50%

9) The gallop rhythm means that: p.111-112

- ☒ a. above the heart, we hear not two but three sounds
- b. above the heart, we hear not four but five sounds
- c. that the heart rate is above 100 beats per minute
- d. that the heart sounds mimics the gallop of quagaa (zebra)
- e. that the patient has „pulses alternans“
- ☒ f. we hear all four sounds of the heart

10) The most frequent cause of mitral stenosis is: (wiki) + p.112

- a. heart failure with „relative“ mitral stenosis
- b. breathlessness
- c. embolization into the arterial system
- d. atrial fibrillation and cyanosis
- ☒ e. rheumatic fever
- ~~f.~~ bacterial endocarditis

11) One of the consequences of mitral stenosis is: p.127

- ☒ a. arterial system embolism
- ~~b.~~ pulmonary embolism
- c. cor pulmonale acutum
- d. left ventricular hypertrophy
- e. relative mitral insufficiency
- f. foramen ovale apertum

12) The typical manifestations of the aortic stenosis are as follows: p.132

- ☒ a. collapse, stenocardia, sudden death
- b. shock and excentric hypertrophy of the left ventricle
- c. palpitation and peripheral cyanosis
- d. diastolic murmur
- e. cyanosis and repeated tromboembolism
- f. surprisingly good toleration of the physical effort

13) In aortic stenosis following pathology occur: p.132

- ☒ a. concentric hypertrophy of the left ventricle
- ☒ b. excentric hypertrophy of the left ventricle
- c. excentric hypertrophy of the left and right ventricles
- d. concentric hypertrophy of the left atria and ventricle
- e. septal defect
- f. dilatation of both ventricles

14) Great systolic-diastolic amplitude is typical for:

- ☒ a. aortic insufficiency
- b. mitral insufficiency
- c. ventricular septal defect
- d. aortic and mitral insufficiency
- e. aortic stenosis
- f. aortic and mitral stenosis

15) The filling pressure of the left ventricle in cardiogenic shock is: p. 180

- a. decreased
- b. extremely decreased- resulting in hypoperfusion of the periphery
- c. almost normal
- ☒ d. increased
- e. to speak about filling pressure in shock is a non-sense
- f. lower then filling pressure of the right ventricle

16) In the beginning of the septic shock the minute volume is:

- a. decreased
- ☒ b. normal or increased
- c. extremely decreased
- d. decreased as a result of endotoxin
- e. indefinable
- f. accelerated

17) Following states are considered to be the complication of myocardial infarction: p. 199-204

- ☒ a. general adaptive syndrome
- ☒ b. ventricular dysrhythmias
- c. myocarditis
- ☒ d. Frank-Starling mechanism
- ☒ e. ischemic degeneration of the brain
- ☒ f. atherosclerosis of the coronary arteries

18) In the earliest period of atherosclerosis following pathology is present: p. 191

- ☒ a. endothelial cells
- b. fibose plates
- ☒ c. fibromuscular plates
- d. foam cells
- e. plaque fissures
- f. thrombus

19) The basic features of neurons involve:

- a. membrane excitability
- ☒ b. transmission of signals

- ☒ c. release of neurotransmitters
- ☒ d. release of hormones
- ☒ e. release of digestive enzymes
- ☐ f. phagocytosis
- ☒ g. processing of signals
- ☒ h. regulation of motor functions

20) The cells having supporting or helping function in the nervous system are:

- ☒ a. neurons
- ☐ b. Schwann cells
- ☐ c. lymphocytes
- ☒ d. satellite cells
- ☒ e. astrocytes
- ☒ f. oligodendrocytes
- ☒ g. microglial cells
- ☒ h. ependymal cells

21) Glial cells:

- ☒ a. exert supporting function
- ☒ b. exert nutritional function
- ☒ c. produce and increase the fibrin formation
- ☒ d. exert regenerate or restorative function
- ☒ e. participate on the transmission of signals
- ☒ f. are less vulnerable than the neurons
- ☒ g. participate on the transfer of information from the peripheral tissues
- ☒ h. separate cerebrospinal fluid and blood from the brain tissue

22) Main source of energy for neurons is:

- ☒ a. glucose
- ☐ b. amino acids
- ☐ c. lipids
- ☐ d. fructose
- ☐ e. triacylglycerols
- ☐ f. cholesterol
- ☐ g. eikosanoids
- ☐ h. galactose

23) On the neuroendocrine stress reaction participates:

- ☒ a. the sympathoadrenal system
- ☒ b. the hypothalamic-pituitary-adrenal axis
- ☒ c. the parasympathetic nervous system
- ☒ d. catecholamines
- ☒ e. glucocorticoids
- ☒ f. vasopressin
- ☐ g. erythropoietin
- ☒ h. angiotensin II

24) Diseases of the nervous system with mainly intrinsic causes are characterized by the following:

- ☒ a. changes of the chromosomal number

- b. CNS infection
- c. are the genetically conditioned diseases
- d. are neuromuscular diseases
- e. the decisive factors remain unknown
- f. are metabolic disturbances
- g. changes of the chromosomal structures
- h. changes resulting from the destruction of the nervous system

25) Diseases of the nervous system with mixed or possibly unknown etiology can be subdivided into these subgroups:

- a. infection of the CNS
- b. tumors of the CNS
- c. the neuroimmunological diseases
- d. the neuromuscular diseases
- e. the metabolic disturbances
- f. the degenerative diseases
- g. some neuropathies
- h. the genetically conditioned diseases

26) Diseases of the congenital disturbances of the amino acid metabolism involve:

- a. phenylketonurea
- b. maple syrup disease
- c. generalized gangliosidosis
- d. Tay-Sachs amaurotic familiar idiocy
- e. cystationinurea
- f. citrulinurea
- g. Gaucher disease
- h. Gierke disease

27) Diseases with lipid metabolism disturbances include:

- a. phenylketonurea
- b. maple syrup disease
- c. generalized gangliosidosis
- d. Tay-Sachs amaurotic familiar idiocy
- e. galactosemia
- f. Fabry syndrome
- g. Gaucher disease
- h. Gierke disease

28) Diseases with disturbance of sacharide metabolism include:

- a. Cori disease
- b. maple syrup disease
- c. McArdle disease
- d. Tay-Sachs amaurotic familiar idiocy
- e. galactosemia
- f. citrulinurea
- g. Pompe disease
- h. Gierke disease

29) Neuronal injury might be of many degrees:

- a. interruption of the nerve cords
- b. interruption of the commissures of the brain
- c. interruption of the nerve fibers
- d. functional injury caused for example by pressure
- e. interruption of the prefrontal cortex
- f. death of an axon without interruption to the endoneural tubes
- g. axonal death with interruption of the endoneural tubes

30) Apart from the direct outcomes of head injury there might commonly be some complications that occur after few hours and days: **P446**

- a. insomnia
- b. hemorrhage
- c. infection
- d. peripheral cyanosis
- e. tumors
- f. brain edema
- g. leak of the cerebrospinal fluid
- h. dyspnea

31) The vasogenic edema is caused by: **P447**

- a. direct disturbance of the active transport of brain cells
- b. transependymal shift of the cerebrospinal fluid
- c. osmotic disturbances
- d. changes in the membrane permeability
- e. disturbance of released acetylcholine
- f. disturbance of the Na^+ pump
- g. disturbances of the amino acid metabolism
- h. the toxic effect of free O^- radicals

32) We distinguish 4 stages of intracranial hypertension: **P449**

- a. the stage of the total compensation
- b. the stage of the partial compensation
- c. the stage of the stabilization
- d. the stage of the destabilization
- e. the stage of the decompensation
- f. the stage of the partial decompensation
- g. the vasomotor paralysis

33) Hydrocephalus might be caused by: **P450**

- a. an excessive production of cerebrospinal fluid
- b. a decrease production of cerebrospinal fluid
- c. cerebrospinal fluid flow obstruction
- d. disturbance of the cerebrospinal fluid absorption
- e. a increase resorption of the cerebrospinal fluid
- f. congenital stenosis of aqueductus Sylvii
- g. atresia of aqueductus Sylvii
- h. atresia of foramen Magendie and Luschle

34) The neurodegenerative diseases of the CNS include:

- a. Parkinson's disease

- ☒ b. Alzheimer's disease
- c. demyelization disease
- d. epilepsy
- e. multiple sclerosis
- f. myasthenia gravis
- g. Fabry syndrome
- h. the acute haemorrhagic leukoencephalopathy

35) For prionosis is valid:

- ☒ a. they are contagious diseases
- ☒ b. they are neurodegenerative diseases
- c. they are exclusively genetically based
- d. they are not transmitted between different species
- ☒ e. they are transmitted between different species
- f. accumulation of a gama-amyloid
- g. accumulation of a tau proteins
- h. accumulation of alpha-synuclein and amyloid

36) Diseases of motor neuron include:

- ☒ a. Huntington's disease, chorea
- b. Alzheimer's disease
- ☒ c. amyotrophic lateral sclerosis
- ☒ d. progressive supranuclear palsy
- e. tabes dorsalis
- ☒ f. myasthenia gravis
- ☒ g. epilepsy
- h. hydrocephalus

37) Neuromuscular diseases include:

- ☒ a. Huntington's disease, chorea
- b. Lambert-Eaton myasthenic syndrome
- ☒ c. amyotrophic lateral sclerosis
- ☒ d. progressive supranuclear palsy
- ☒ e. tabes dorsalis
- ☒ f. myasthenia gravis
- ☒ g. epilepsy
- h. hydrocephalus

38) Neurodegenerative disorders include: (wiki) category

- ☒ a. Huntington's disease, chorea
- ☒ b. Alzheimer's disease
- ☒ c. amyotrophic lateral sclerosis
- d. progressive supranuclear palsy
- ☒ e. Parkinson's disease
- f. myasthenia gravis
- g. epilepsy
- h. hydrocephalus

39) Microscopic features of Alzheimer disease include: patho

- ☒ a. accumulation of a neurofibrillary tangles

- ~~b.~~ accumulation of a prion proteins
- c. accumulation of a beta-amyloid
- d. accumulation of an alpha-amyloid
- e. accumulation of a gama-amyloid
- f. accumulation of parkin proteins
- g. accumulation of a tau proteins
- h. occurrence a senile plaques

40) **Parkinson's disease is characterized by:** (wiki)

- ~~a.~~ accumulation of alpha-synuclein
- b. accumulation of a prion proteins
- c. accumulation of a beta-amyloid
- d. accumulation of an alpha-amyloid
- e. accumulation of a gama-amyloid
- ~~f.~~ accumulation of parkin proteins
- ~~g.~~ accumulation of tau proteins
- h. loss of dopaminergic cells

41) **The Parkinson's disease is clinically manifested by:** (p.455)

- a. the typical finger movements
- b. the typical head tremor
- c. the typical hand tremor
- d. a mask face
- e. hypertrophy of heart sympathetic innervation
- f. a muscular tonus disturbance
- g. generalized seizures
- h. the abnormal postural tonus

42) **Demyelination diseases include:** (wiki) (p.451, 453)

- a. Huntington's disease/chorea
- b. multiple sclerosis
- c. amyotrophic lateral sclerosis
- d. progressive supranuclear palsy
- e. Parkinson's disease
- f. myasthenia gravis
- g. acute disseminated encephalopathy
- h. acute hemorrhagic leukoencephalopathy

? 43) **Common features of neurodegenerative diseases include:** (wiki)

- a. accumulation of aberrant proteins
- b. neuroinflammation
- ~~c.~~ mitochondrial abnormality
- d. deficit of glutamate
- e. deficit of serine
- f. deficit of glycine
- g. relatively selective loss of neurons of certain phenotype
- h. deficit of aspartate

44) **Neurotransmitters are release predominantly from:**

- a. presynaptic nerve ending

- b. muscle cells
- c. postsynaptic nerve endings
- d. endoplasmatic reticulum
- e. mitochondria
- f. sarcoplasmatic reticulum
- g. cell nucleus
- h. dendrites

45) For ionotropic receptor is valid: (wiki)

- a. inside ion channel is localised Mg^{2+}
- b. is also ion channel
- c. is localized exclusively in mitochondrial membranes
- d. is always bound with G-protein
- e. is localized in cell nucleus
- f. is responsible for slow changes of electrical charge on postsynaptic membrane
- g. is responsible for fast changes of electrical charge on postsynaptic membrane
- h. is represented by nicotinic receptors

46) For metabotropic receptor is valid: (wiki)

- a. is also ion channel
- b. is localized exclusively in cytoplasm of neurons
- c. is usually bound to system of G-proteins
- d. is localized in cell nucleus
- e. is represented by muscarinic receptor
- f. is responsible for slow changes of electrical charge on postsynaptic membrane
- g. is responsible for fast changes of electrical charge on postsynaptic membrane
- h. is represented by nicotine receptors

47) Nicotine receptor: (wiki)

- a. belongs into the group of ionotropic receptors
- b. belongs into the group of metabotropic receptors
- c. its agonist is norepinephrine
- d. its agonist is acetylcholine
- e. its agonist is dopamine
- f. its agonist is glutamate
- g. its agonist is aspartate
- h. its agonist is epinephrine

48) Muscarine receptor:

- a. belongs into the group of ionotropic receptors
- b. belongs into the group of metabotropic receptors
- c. its agonist is serotonin
- d. its agonist is dopamine
- e. its agonist is ATP
- f. its agonist is acetylcholine
- g. its agonist is angiotensin

h. its agonist is GABA

49) The result of transnuclear signalization is:

anything

- a. synthesis of new receptors
- b. synthesis of new ion channels
- c. complete termination of translation
- d. synthesis of new enzymes
- e. changes in reactivity of neuron
- f. changes of neuronal plasticity
- g. immediate changes in electrical charges on neuronal membrane
- h. immediate opening of ion channels in neuronal membrane

50) Classical small molecule neurotransmitters include:

- a. acetylcholine
- b. dopamine
- c. enkephalin
- d. ACTH
- e. endorphin
- f. angiotensin II
- g. glutamate
- h. GABA

51) Neurotransmitters/neuromodulators with large molecule represent:

- a. enkephalin
- b. serotonin
- c. substance P
- d. glutamate
- e. GABA
- f. CRH
- g. angiotensin II
- h. dopamine

52) The main excitatory neurotransmitter in the nervous system is:

- a. glutamate
- b. GABA
- c. glycine
- d. dopamine
- e. serotonin
- f. epinephrine
- g. histamine
- h. ACTH

53) The main inhibitory neurotransmitter in the nervous system is:

- a. GABA
- b. glutamate
- c. epinephrine
- d. serotonin
- e. histamine
- f. ATP
- g. ACTH