



قائمة الاسئلة

خضاب الدم-مختبرات -الثالث- درجة الاختبار (60)

أ.د.محمد النزيلي

- 1) If iron in hemoglobin is oxidized to 3+, what disease may occur ?
 - 1) - Sickle cell anemia
 - 2) - Cystic fibrosis
 - 3) + Methemoglobinemia
 - 4) - Carbon monoxide poisoning
- 2) Hemoglobin molecule is composed of:
 - 1) - Two globin chains and two heme molecules
 - 2) + Four globin chains and four heme molecules
 - 3) - Four globin chains and ferrous iron
 - 4) - Two globin chains and ferric iron
- 3) A point mutation in the beta-globin gene changing the codon number 6 from glutamate to valine will likely cause what disease?
 - 1) - Methemoglobinemia
 - 2) - Carbon monoxide poisoning
 - 3) + Sickle cell anemia
 - 4) - Thalassemia
- 4) All of the followings are true about iron metabolism EXCEPT :
 - 1) - Iron is constituent of Hb, Mb, heme enzymes and transferrin
 - 2) - Iron is transported from blood to tissues by transferrin
 - 3) - Iron is stored in macrophages as ferritin and hemosiderin
 - 4) + Iron is regulated by erythropoietin
- 5) What has the highest affinity for oxygen ?
 - 1) - Hemoglobin F
 - 2) - Hemoglobin A
 - 3) + Myoglobin
 - 4) - Hemoglobin A2
- 6) The most common anemia in the world is:
 - 1) + Iron deficiency anemia
 - 2) - Megaloblastic anemia
 - 3) - Sideroblastic anemia
 - 4) - Sickle cell anemia
- 7) ATP is generated in RBC by:
 - 1) - Methemoglobin reductase shunt
 - 2) - Hexose-monophosphate pathway
 - 3) + Embden-Meyerhof pathway
 - 4) - Lubering-Rapaport shunt
- 8) The function of the methemoglobin reductase shunt is to:
 - 1) - provide cellular energy
 - 2) - produce methemoglobinemia
 - 3) + prevent oxidation of heme iron
 - 4) - control the rate of glycolysis
- 9) Glucose 6-phosphate dehydrogenase (G6PD) deficiency is characterized by:
 - 1) - it is the most common genetic disease in human (X-linked disease)
 - 2) - It causes a hemolytic anemia associated with Heinz bodies
 - 3) - It induced by some drugs, ingestion of fava beans or by heavy infection





- 4) All Answers are Correct
- 10) Extravascular hemolysis is characterized by:
- 1) - Occurs in macrophages of RES of liver, spleen and BM
 - 2) - The most common mechanism in normal RBC destruction
 - 3) - Initiate by signals to macrophages to ingest damaged or senescent RBC
 - 4) All Answers are Correct
- 11) What is the Hb that results when glutamate in the 6th position of beta chain is replaced by lysine ?
- 1) - Hb S
 - 2) Hb C
 - 3) - Hb E
 - 4) - Hb Yakima
- 12) All of the followings are true about vitamin B12 metabolism EXCEPT :-
- 1) - The daily dietary intake is 5-30 μ g, daily needs is 5 μ g.
 - 2) - It presents in foods of animal origin e.g. liver, meat, eggs and milk
 - 3) The total body stores 2-3mg sufficient for 2-5 months.
 - 4) - Presents in two coenzyme forms as methyl B12 and adoB12
- 13) All of the followings are true about folate metabolism EXCEPT :
- 1) - The absorption takes place in duodenum and jejunum
 - 2) - It is absorbed as methyl THF
 - 3) - It is transported in blood weakly bound to albumin
 - 4) None of the above
- 14) Hb Bart's consists of:
- 1) - Four β globin chains
 - 2) Four γ globin chains
 - 3) - Four α globin chains
 - 4) - Four delta globin chains
- 15) Hb H consists of:
- 1) Four β globin chains
 - 2) - Four γ globin chains
 - 3) - Four α globin chains
 - 4) - Four delta globin chains
- 16) The activity of hexose-monophosphate pathway increases the RBC source of :
- 1) - Glucose and lactic acid
 - 2) - 2,3-DPG and methemoglobin
 - 3) NADPH and reduced glutathione
 - 4) - ATP and other purine metabolites
- 17) All of the followings are true about vitamin B12 metabolism EXCEPT :
- 1) - The absorption takes place in Ileum
 - 2) - Intrinsic factor (IF) is important for B12 absorption
 - 3) - It is transported by 3 types of Transcobalamins (TCI, II, III)
 - 4) None of the above
- 18) All of the followings are true about folic acid metabolism EXCEPT :
- 1) - The daily dietary intake is 200-250 μ g, daily needs 100-150 μ g.
 - 2) - It presents in most foods especially liver, meat, green vegetables and fruits
 - 3) Total body stores 10-12 mg sufficient for 2-4 years.
 - 4) - It is Important for DNA synthesis
- 19) In α -thalassemia, deletion of three α -genes produces:
- 1) - Hb S
 - 2) Hb H





- 3) - Hb D
4) - Hb M
- 20) A qualitative abnormality in Hb may involve all of the followings EXCEPT:
- 1) - Replacement of one or more amino acids in a globin chain
 - 2) - Addition of one or more amino acids in a globin chain
 - 3) - Deletion of one or more amino acids in a globin chain
 - 4) Decreased production of a globin chain
- 21) If a defect in the oxidative pathway occurs, what will result?
- 1) - Insufficient amounts of reduced glutathione
 - 2) - Denaturation of globin
 - 3) - Precipitation of Heinz bodies
 - 4) All Answers are Correct
- 22) Patients with SCD usually do not exhibit symptoms until 6 months of age because:
- 1) - The mother's blood has a protective effect
 - 2) - Hemoglobin levels are higher in infants at birth
 - 3) Higher levels of Hb F are present
 - 4) - The immune system is not fully developed
- 23) Through routine screening, prospective parents discover that they are both heterozygous for Hb S. What percentage of their children potentially could have SCD?
- 1) 25%
 - 2) - 50%
 - 3) - 75%
 - 4) - 100%
- 24) When binding with oxygen, what type of binding curve does hemoglobin have ?
- 1) - Linear
 - 2) Sigmoidal
 - 3) - Hyperbolic
 - 4) - All Answers are Correct
- 25) Which of the following statements describes the oxygen binding curve of haemoglobin ?
- 1) - Each of the four oxygens bind with equal facility
 - 2) The binding of the first oxygen molecule enhances the binding of the other three oxygen molecules
 - 3) - The binding of the first oxygen molecule makes the binding of the other three oxygen molecules more difficult
 - 4) - The binding of the first oxygen molecule has no effect on the binding of the remaining three oxygen molecules
- 26) The Embden-Meyerhof pathway net gain of ATP provides high energy phosphates to:
- 1) - maintain membrane lipids
 - 2) - power the cation pump needed for Na⁺/K⁺ pump and calcium flux
 - 3) - preserve the shape and flexibility of the cellular membrane
 - 4) All Answers are Correct
- 27) Thalassemia is caused by:
- 1) - Increase in α chain production
 - 2) - Increase in β chain production
 - 3) Decrease in α or β chain production
 - 4) - All Answers are Correct
- 28) Disorders of heme synthesis include the followings EXCEPT:
- 1) - Iron deficiency anemia
 - 2) - Sideroblastic anemia
 - 3) Megaloblastic anemia





- 4) - Porphyria
- 29) All of the following factors favoring iron absorption EXCEPT:
- 1) - Ferrous form (Fe+2)
 - 2) + Tea
 - 3) - Reduced serum hepcidin
 - 4) - Increased expression of DMT-1 in duodenal enterocytes
- 30) Iron overload is caused by:
- 1) - Excessive absorption of iron
 - 2) - Excessive repeated blood transfusions
 - 3) - Genetic mutation of the HFE gene causing C282Y protein change
 - 4) + All Answers are Correct

