

UNIVERSITY EXAMINATIONS - 2014/2015

LEVEL III EXAMINATIONS FOR THE DEGREES OF BACHELOR OF MEDIC AND BACHELOR OF SURGERY (6 YEAR PROGRAM)

HHP 300: GENERAL PATHOLOGY

SAQ/MCQ PAPER

DATE: TUESDAY 19TH MAY, 2015

TIME 9.00 A.M. - 12.00 N

INSTRUCTIONS:

- There will be 5 minutes reading time
- 2. Enter your Registration Number in all your answer books and scripts
- 3 The examination consists of 2 parts

Part A: - MCQ

- i) Each question has only one correct answer
- ii) Answer the question in the answer sheet provided
- iii) If you do correction do so very clearly

Part B: - SAQ

- Answer each question in a separate book
- in Number all your questions clearly

PART A: MULTIPLE CHOICE QUESTIONS

- An AIDS defining malignancy
 - Primary Central Nervous system lymphoma B)
 - Squamous cell carcinoma of the conjunctiva
 - C) Gastric adenocarcinoma
 - Di Hepatocellular carcinoma
 - Squamous cell carcinoma of the skin
- 2 An AIDS associated malignancy
 - Primary central nervous system lymphoma
 - Kaposi sarcoma
 - C) Burkitt lymphoma
 - D) Diffuse large B cell
 - Squamous cell carcinoma of conjunctiva
- HIV virus drives carcinogenesis via all EXCEPT:
 - A) Mutations
 - B) Cytokines surge
 - C) Micro RNAs
 - D) Co-infective agents
 - Large pool of proliferating B cells
- 4. Cellular Features of malignant cells EXCEPT
 - A) Hyperchromasia
 - Increased nuclei cytoplasmic ratio
 - C) Irregular nuclear membrane
 - D) Prominent nucleoli
 - E) invasion
- 5. True of sarcomas
 - Benign mesenchymal tumours A)
 - Growth pattern shows nests, cords and trabeculae B)
 - Usually spreads through haematogenous route C)
 - TNM staging is not applicable D)
 - Does not metastasise to lymphnodes E)
- An example of soft tissue tumour EXCEPT 6.
 - Firbroepithelial polyp A)
 - Lipoma B)
 - Uterine fibroids C)
 - Osteosarcoma D)
 - Rhabdomyoma E)
- Familial tumour with a suppressor gene EXCEPT 7.
 - WILMS tumour A)
 - Neuro fibromatoses-1 B)
 - Retino blastoma C)
 - Burkitts lymphoma D)
 - Familial polyposis coli



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- Special stain used to demonstrate amyloid deposition
 - A) Congo Red
 - B) Haematoxylin and eosin
 - () Giemsa stain
 - D) Grocott stain
 - E) Pap stain
- 90% of the amyloid deposits are composed of
 - A) Glycosaminoglycans
 - B) Apolipoprotein I
 - C) Serum amyloid P
 - D) Serum amyloid E
 - E) Aggregation of misfolded proteins
- Cellular adaptive response EXCEPT: 10.
 - Hypertrophy A)
 - Hyperplasia B)
 - C) Atrophy
 - Neoplasia D)
 - Metaplasia E)
- Tissue composed of stable cells EXCEPT
 - Intestinal mucosa
 - B) Liver
 - Kidney C)
 - Pancreas D)
 - Endothelial cells E)
- Not a role of extra cellular matrix in tissue repair
 - Provide mechanical support for cell anchorage
 - Control all growth by signalling through links with intracellular integrins A) B)
 - Limit capillary proliferation within site of repair C)
 - Establishment of tissue micro-environment D)
 - Storage and presentation of regulatory molecules E)
- Following a myocardial infarction, healing occurs by 13.
 - Regeneration A)
 - Inflammation B)
 - **Fibrosis** C)
 - Metaplasia D)
 - hyperplasia E)
- TRUE regarding malignant hypertension 14.
 - 75% recover with no loss of renal function, A)
 - Is associated with abnormal renin levels
 - B) Is more common compared to benign hypertension C)
 - Affects 1-5% of hypertensive patients D)
 - Has an insidious onset« E)

- 15 Not a major criterion in diagnosis of Rheumatic fever
 - A) Sydenham's chorea
 - B) Carditis
 - €) Polyarthralgia
 - D) Erythemas nodesum
 - E) Subcutaneous nodules
- Causes for Lymphocytosis include all EXCEPT
 - A) Chronic lymphocytic leukaemia
 - B) Infectious mononucleosis
 - C) Brucellosis
 - D) Haemolysis
 - E) Tuberculosis
- Peripheral blood features of neutrophil leucocytosis with a left shift and toxic granulation would be consistent with
 - A) Parasitic infection
 - B) Bacterial infection
 - C) Virial infection.
 - Bone marrow hypoplasia
 - E) Acute myeloid leukaemia,
- Infestation with the following parasite is associated with a microcytic hypochromic blood picture
 - A) Necator americanus
 - B) Plasmodium falciparum
 - C) Leishmania donovani
 - D) Diphilopbothrium latum
 - E) Ascaris lumbricoides
- 19. The following one is classified amongst the early acting growth factors
 - A) Erythropoietin
 - B) Thrombo poietin
 - C) Gm CSF
 - D) G-CSF
 - E) m-CSF
- Microcytic red cell changes occurs in the following conditions EXCEPT
 - A) Iron deficiency anemia
 - B) Thalassemias
 - C) Lead poisoning
 - D) Hypothyroidism
 - E) Sideroblastic anaemia
- The following changes are associated with iron def anaemia
 - A) Elevated serum iron levels >
 - B) Elevated serum ferritin
 - C) Reduced MCH
 - D) Raised MCV «
 - E) Reduced total iron binding capacity +

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- Vitamin B₁₂ 22.
 - Rich sources constitute plants -A)
 - B) Has no role in haemopoiesis .
 - C) Normal serum levels 160-923 ng/L -
 - D) Is absorbed in the stomach using intrinsic factor
 - E) Destroyed during cooking > 70%
- . 23 The serum folate levels in adults is
 - A) 160 - 640 µg/L×
 - B) $3 - 15 \mu g/L$
 - C) 160 - 500 μg/dL
 - D) 4 20 µg/dL
 - E) 60 - 80 µg/L
- Non megaloblastic macrocytic anemia are associated with the following condition EXCEPT
 - A) Liver disease
 - B) Drugs
 - Vitamin B₁₂ deficiency-C)
 - Aplastic anemia D)
 - Hypotheyrodism E)
- A mother is blood group A+ve and the father blood group AB+ve, the offspring will t 25. blood group EXCEPT
 - 0 + veA)
 - B +ve B)
 - A+ve C)
 - AB +ve D)
 - Any of the above E)
- The shelf life of platelets concentrates is
 - 35 days in room temperature A)
 - 3 5 days in room temperature B)
 - 21 days C)
 - 3 5 days at refrigeration D)
 - 28 days E)
- An ideal blood donor concentrate's 27.
 - 16 year old or younger . A)
 - Regularly donates blood. B)
 - A multiparous female C)
 - Bisexual D)
 - Commercial donor ' E)

- Adhesion, aggregation and release are functions of the
 - A) Fibrinolysis
 - B) Anticoagulation
 - C) Coagulation system
 - D) Blood vessel
 - E) platelet
- The inheritance pattern of haomophilia B is
 - Double Autosomal
 - B) X-linked recessive
 - C) Unknown
 - D) Autosomal dominant
 - Autosomal recessive
- In haemophilia A laboratory tests manifest with
 - A prolonged prothrombin time ye &
 - B) Abnormal platelet function in presence of antibodies of factor VII
 - C) Normal thrombin time
 - D) A prolonged bleeding time
 - Prolongation of the activated partial thromboplastin time
- A known haemophilia patient presents to the casualty with a minor bleeding. appropriate investigation is
 - A) Serum blood sugar
 - Serum creatinine
 - C) Blood slide for malaria parasite
 - D) Coagulation screen
 - E) Serum lactate dehydrogenase
- Which one of the following is the **ODD** one out
 - Hereditary elliptocytosis
 - Defect in the hexose monophosphate shunt -B)
 - C) Hereditary stomatocytosis
 - Disorders of permeability of membrane D)
 - Abnormal membrane lipid composition -
- Which of the following is NOT included in immune haemolytic anaemia
 - IgM mediated A)
 - Warm antibody B)
 - C) Paroxysmal nocturnal haemoglobinuria
 - IgG mediated D)
 - E) Cold antibody,
- One of the following regions does NOT have a high prevalence of HbS
 - Mediterranean. A)
 - B) West Africa
 - C) East Africa
 - D) Saudi Arabian peninsula
 - South East Asia -

- 35. The definitive diagnostic test in sickle cell disease is
 - A) Sickling test
 - B) Hb electrophoresis
 - () Hb solubility test
 - D) Peripheral blood picture
 - E) Positive family history for sickle cell disease
- Excess ADH production may lead to:
 - A) Polyuria
 - B) Hypernatraemia
 - Increased plasma osmolarity
 - D) Increased urine osmolarity
 - E) protemuria
- Iron overload is usually associated with 37.
 - Increased plasma ferritin A)
 - Decreased plasma iron * B)
 - Increased total iron binding capacity C)
 - Increased plasma transferrin-D)
 - Pernicious anaemia -E)
- Causes of hypercalcemia include the following: exter
 - Thiazide diuretics
 - Sarcoidosis . B)
 - Tuberculosis C)
 - Thyrotoxicosis D)
 - Chronic kidney disease E)
- The total amount of iron in the body is about 39.
 - 3-4 grams A)
 - 2 10 grams B)
 - 15 30 milligrams C)
 - 100 2000 grams D)
 - 1-2 grams E)
- Causes of Hyperphosphatemia include the following EXCEPT: 40.
 - Renal FAILURE A)
 - Vitamin D excess B)
 - Tumour lysis syndrome C)
 - Hypoparathyroidism. D)
 - Alcohol withdrawal E)
- High Cortisol levels and low ACTH levels indicate: 41.
 - Addison's Disease A)
 - Acromegaly B)
 - Pheochromocytoma C)
 - Cushing's Disease D)
 - Prolactinona E)

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	-1	g. om oon yurare make						
	E)	High carbohydrate intake						
	D)	Anorexia						
		Malabsorption						
	B)	Bacteriuria						
	A)	Proteinuria						
48.	A po	ositive test for ketone bodies in urinalysis may be indicative of						
	E)	Catheter urine						
		Random urine						
	0.022.53	Early morning urine						
	10.00	3 hour urine						
	A)	A) 24 hour urine						
47.	Whi	Which of the following specimen is preferred for emergency tests?						
	E)	48 hours						
	D)	72 hours						
	C)	24 hours						
	B)	12 hours						
	,	Thou						
46.	For l	now long does sodium fluoride stabilize glucose in blood at room temperature						
	E)	Glucagon deficiency						
	D)	Hypothyroidism						
	,C)	Pheochromocytoma -						
	B)	Addison's Disease						
		hese endocrinopathies lead to hyperglyeaemia EXCEPT:						
45.	Allt	hese endocrinonathies lead to hypogycemes						
	E) .	> 11.0 mmol/L						
	D)	3.2 - 7.8 mmol/L 4						
	(C)	3.2 - 6.1 mmol/L-6/51 VB3						
	B)	2.2 – 3.2 mmol/L						
		· L.Z minor L						
44.	The r	eference range for fasting plasma blood sugar is approximately:						
	-/	120 Brains						
	E)	75 grams 120 grams						
	C) D)	50 grams						
	B)	45 grams						
		B						
43.	How	much glucose is administered in an oral glucose tolerance test?						
	1000	•						
	E)	Hypergonadotrophic hypegonadism						
	D)	Secondary hypothyroidism						
	C)	Secondary adrenal cortical insufficiency Lack of lactation						
. •		D Will Hall						
4.2	A)	Dwarfism						

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•49	Which of the following is the major paraprotein associated with multiple myeloma?						
	•A)	IgA					
•	B)	IgD					
	(C)	Ig D €					
	D)	IgM -					
	E)	IgG					
50.	Wh	Which of the following urinary metabolite is associated with catecholamine metabolism?					
	A)	Globulins					
	B)	Bence jones					
	C)	Dopa					
	D)	Vanillyl mandelic acid					
	E)	creatinine					
51.	In serum protein electrophoresis, an abnormal ceruloplasmin (CER) protein can be						
	dete	ected in					
	A)	The albumin band					
385	B)	The alpha one globulin band					
	C)	The alpha two globulin band					
	D)	Beta globulin band					
	E)	Gamma globulin region					
52.	Dec	reased total and LDL-cholesterol with normal triglycerides suggest					
	A)	Tangier's disease					
•	B)	Hypo-lipoproteinaemia					
	C)	Abetalipoproteinaemia					
	D)	Lp(a) disease					
	E)	Hypo-β-lipoproteinaemia					
	T., els	e differential diagnosis of hypercalcaemia the following should not be included					
53.		Vitamin D dependent rickets					
	A)	the state of the s					
	B)	Excess absorption secondary to "milk alkali syndrome"					
	C)	Multiple myeloma					
	D)	Primary hyperparathyroidism					
	E)	Primary hyperparamy					
54.	Of th	e enzymes listed which one is least useful for reflecting hepatobiliary disease					
54.		0.07					

- - A)
 - Glutamate dehydrogenase B)
 - S' nucleotidase C)
 - ALP D)
 - Leucine amino peptidase (arylamidase) E)
- True of a specimen for measurement of lipids:

 A) Use of fluoride is mandatory. 55.

 - Should be frozen immediately B)
 - Should be a fasting specimen C)
 - Only serum should be used D)
 - None of the above E)

Describe the mechanisms involved in the termination of an acute inflammatory (2.5 marks)

Discuss paraneoplastic syndromes under the following

(12.5 marks)

- Definition
- Discuss the clinical importance of paraneoplastic syndrome 11)
- 111) Describe the pathogenesis of one endocrine paraneoplastic syndrome

A 29 year old has the following blood count results. Hb - 5.9 gldlJune WBC - 6.2 x 109/L - 150

MCV - 55 fl MCH - 18 pc. 24 5 Platelets - 400 x 10⁹/L -

1) Interpret the above results.

(3 marks)

Give the most likely differential diagnosis. 11)

(2 marks)

Outline relevant investigations to confirm diagnosis for this patient (7 marks) 111)

Define haemolytic red cell disorders.

(2 marks)

- Outline the laboratory investigations of a patient with haemolytic red cell. ii)
 - (10 marks)
- With the aid of a diagram outline the pathway of the Coagulation System. iii)

(12 marks)

- A 58 year old male patient presented at the medical clinic and the attending physician 4. suspected diabetes mellitus. Answer the following questions:
 - List five signs and symptoms the patient might be having. i)

(2.5 marks)

- List the biochemical tests in sequential order the physician will order to make a ii) diagnosis of diabetes mellitus. (6 marks)
- List three (3) complications of diabetes mellitus. iii)

(1.5 marks)

What is the most likely type of diabetes mellitus is in this patient. (2 marks)

For each hormone listed above, give its overall function. (5 marks)

ofing

List five endocunopathies arising from the hyper functioning anterior pituitary. (5 marks)

b) i) Discuss causes of acute renal failure. (6 marks) -

ii) Describe biochemical investigations in this condition. (6½ marks)