Roll No.

Sig. of Candidate.

Answer Sheet No	_
Sig. of Invigilator	

HSSC-II

HAEMATOLOGY AND BLOOD BANKING HSSC-II SECTION - A (Marks 10)

A. (ii) Which granu A. C. (iii) What A. C. (iv) Which anem A. C. (v) Accor A. C. (vi) The tr chrom A. (vii) What A. B. C. (viii) A pati	lies, with a few primary dark granules? Band Promyelocyte is the composition of normal adult haen 92-95% HbA, 5-8% Hb A_2 , 1-2% Hb 80-85% HbA, 2-3% Hb A_2 , 1-2% Hb h of the following laboratory results would	C. nucleus v B. D. noglobin? o F B. o F D. dd NOT b B. D. of anemic B. D.	Bone Marrow D. All of these with clumped chromatin and small pink secon Myelocyte Meta myelocyte? 90-92% HbA, 2-3% Hb A_2 , 2-5% Hb F 95-97% HbA, 2-3% Hb A_2 , 1-2% Hb F be a usual diagnostic criterion for a patient with Decreased hematocrit level Decreased RBC count its, megaloblastic anemia is a Macrocytic, hyperchromic anemia
(ii) Which granu A. C. (iii) What A. C. (iv) Which anem A. C. (v) Accor A. C. (vi) The tr chrom A. (vii) What A. B. C. D. (viii) A pati most A.	h granulocytic cell has a kidney shaped alles, with a few primary dark granules? Band Promyelocyte is the composition of normal adult haen 92-95% HbA, 5-8% Hb A_2 , 1-2% Hb 80-85% HbA, 2-3% Hb A_2 , 1-2% Hb h of the following laboratory results would hia? Decreased haemoglobin level Decreased platelet count riding to the morphological classification Macrocytic, hypochromic anemia Macrocytic, normo-chromic anemia ranslocation that results in the formation	B. D. noglobin? D.	with clumped chromatin and small pink second Myelocyte Meta myelocyte $ \begin{array}{ccccccccccccccccccccccccccccccccccc$
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C. (iii) What A. C. (iv) Which anem A. C. (v) Accor A. C. (vi) The tr chrom A. (vii) What A. B. C. D. (viii) A pati most A.	Promyelocyte : is the composition of normal adult haem $92\text{-}95\%$ HbA, $5\text{-}8\%$ Hb A_2 , $1\text{-}2\%$ Hb $80\text{-}85\%$ HbA, $2\text{-}3\%$ Hb A_2 , $1\text{-}2\%$ Hb of the following laboratory results would also in the following laboratory results would be creased haemoglobin level Decreased platelet count right of the morphological classification Macrocytic, hypochromic anemia Macrocytic, normo-chromic anemia ranslocation that results in the formation	D. noglobin? o F B. o F D. ld NOT b B. D. of anemic B. D.	Meta myelocyte ? $90\text{-}92\% \text{ HbA, } 2\text{-}3\% \text{ Hb } A_2 \text{ , } 2\text{-}5\% \text{ Hb F} $ $95\text{-}97\% \text{ HbA, } 2\text{-}3\% \text{ Hb } A_2 \text{ , } 1\text{-}2\% \text{ Hb F} $ be a usual diagnostic criterion for a patient with
(iii) What A. C. (iv) Which anem A. C. (v) Accor A. C. (vi) The tr chrom A. (vii) What A. B. C. D. (viii) A pati most A.	is the composition of normal adult haem 92-95% HbA, 5-8% Hb A_2 , 1-2% Hb 80-85% HbA, 2-3% Hb A_2 , 1-2% Hb h of the following laboratory results would hia? Decreased haemoglobin level Decreased platelet count rding to the morphological classification Macrocytic, hypochromic anemia Macrocytic, normo-chromic anemia ranslocation that results in the formation	noglobin? o F B. o F D. ld NOT b B. D. of anemic B. D.	90-92% HbA, 2-3% Hb A_2 , 2-5% Hb F 95-97% HbA, 2-3% Hb A_2 , 1-2% Hb F be a usual diagnostic criterion for a patient with Decreased hematocrit level Decreased RBC count its, megaloblastic anemia is a Macrocytic, hyperchromic anemia
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chrom A. (vii) What A. B. C. D. (viii) A pati most A.			Normocytic, normochromic anemia
A. (vii) What A. B. C. D. (viii) A pati most A.	acamas	of the ph	hiladelphia chromosome(Ph) involves
(vii) What A. B. C. D. (viii) A pati most A.	HUSUITIES.		
A. B. C. D. (viii) A pati	21 and 22 B. 22 and 9	C.	8 and 14 D. 21 and 9
B. C. D. (viii) A pati most A.	is the purpose of the APTT test in moni	toring he	emostasis?
C. D. (viii) A pati most A.	Measures factors of the intrinsic path	way	
D. (viii) A pati most A.	Detects the deficiency of factors for b	oth intrin	nsic and extrinsic pathways
(viii) A pati most A.	Measures circulating FDP's.		
most A.	Detects platelet dysfunctions		
A.	tient presents with a platelet count of 212	2x 10 ⁹ /L	and a bleeding time of 12 mins. These resu
	probably suggest	_	
C.	Decreased platelet production	B.	Increased platelet destruction
	Increased platelet function	D.	None of these
(ix) Which	h of the following conditions would most	likely she	ow a decrease in factor VIII?
A.	Haeamophilia A	B.	Haemophilia B
C.	Myeloproliferative disorder	D.	Carriers of Haemophilia A
(x) Bason	philic stippling is composed of		
A.	DNA	B.	Precipitated stain
C.	Denatured haemoglobin	D.	RNA

HAEMATOLOGY AND BLOOD BANKING HSSC-II

Time allowed: 2:20 Hours

Total Marks Sections B and C:

Student Bounty.com Answer any twelve parts from Section 'B' and any two questions from Section 'C' on the separately provided answer book. Use supplementary answer sheet i.e. Sheet-B if required. Write your answers neatly and legibly.

SECTION - B (Marks 24)

- Attempt any TWELVE parts. The answer to each part should not exceed 2 to 4 lines. (12 x 2 = 24) Q. 2
 - Name the different stages of Red Blood Cell production. (i)
 - Write down the names of different acute Leukemias. (ii)
 - Name the tests to check the extrinsic system of coagulation. What coagulation factors are involved (iii) in this system?
 - Write about the subgroups of Blood group A. (iv)
 - Name different haemoparasites. (V)
 - Name different blood products commonly used. (vi)
 - Define the following terms: (vii)
 - Hypochromia
 - b. Poikilocytosis
 - Anisocytosis
 - How can you classify anemias? (viii)
 - Write down the main clotting factors involved in the intrinsic system of coagulatian. (ix)
 - What is meant by thromobocytopenia? What is the common cause of thrombocytopenia in (x) children?
 - At what temperature do you store the following Blood products: (xi)
 - Red Cell concentrate
 - Platelets concentrate b
 - Fresh frozen plasma
 - Name the different stages of white blood cell development. (xii)
 - What is Reticulocyte count? Name the stain used for its staining. (xiii)
 - (xiv) Write down the causes of Eiosinophilia.
 - Name different malarial parasites. (xv)
 - Write down the names of different types of haemophilia. What is the commonest type? (xvi)

SECTION - C (Marks 16)

Attempt any TWO questions. All questions carry equal marks. Note:-

 $(2 \times 8 = 16)$

- Write the complications of Blood Transfusion. Q. 3
- What is the principle of Bleeding time? List the methods of performing it. Also describe any one of the methods Q. 4
- Write down the different stages of Granulopoiesis. Briefly describe each stage. Q. 5