1-Which of the following is wrong concerning Philadelphia chromosome:

- a -shortening of long arm of chromosome 22
- b -Philadelphia -ve cases have bad prognosis
- c -diagnostic of CML
- d -found in lymphocytes)√(

2 -the commonest cause of jaundice in thalassemia is:

- a -viral hepatitis c
- b -iron deposition in liver
- c -viral hepatitis B
- d -haemolysis)√(

3-Waldeyer' ring does not include:

- a -faucal tonsils
- b -submandibular glands) $\sqrt{(}$
- c -adenoids
- d-lingual tonsils

4-whitch of the following anemias is associated with splenomegaly:

- a -chronic renal failure
- b -aplastic anemia
- c -hereditary spherocytosis)√(
- d -sickle cell anaemia

5 -all may cause abdominal pain in thalassemia major except:

- a -vasculitis)√(
- b -splenic infarction
- c -dragging pain dt huge splenomegaly
- d -pigment stones in gall bladder

6 -Virchow's node receives lymphatics from all except:

- a -testes
- b -stomach
- c-prostate)√(
- d -breast

7 -all produce microcytic anemia except:

- a -sideroblastic an.
- b -thalassemia
- c-pernicious anemia) $\sqrt{}$
- d-lead poisoning

8 -Basophilic stippling is classically seen in:

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- a -CML
- b -myelosclerosis
- c -chronic lead poisoning) $\sqrt{(}$
- d -iron def anemia

9-Increase Fe & normal TIBC are found in:

- a -thalassemia major
- b -haemosidrosis
- c-rheumatoid arthritis)√(
- d-dissiminated malignancy

10 -Non thrombocytopenic purpurais seen in all except:

- a-vasculitis
- b-uraemia
- c-hereditary haemorrhagic telangectasia
- d-SLE)√(

11 -Gum bleeding is characteristic of all except:

- a-chronic phenytoin therapy) $\sqrt{(}$
- b-aplastic an.
- c-scurvy
- d-haemophilia

12-Which of the following is not true in thrombathenia:

- a-prolonged bleeding time
- b-normal plat .Count
- c-plat .Aggregation defect
- d-prolonged clotting time) $\sqrt{(}$

13 -Which is not associated with hypersplenism:

- a-splenomegaly
- b-pancytopenia
- c-hyper cellular bone marrow)√(
- d-reversibility by splenectomy

14-splenectomy is curative in:

- a-G6PD def.
- b-ITP
- c-thalassemia
- d-hereditary sphrocytosis) $\sqrt{(}$

15-Plummer Vinson syndrome is not associated with:

- a-angular stomatitis
- b-splenomegaly
- c -clubbing)√(

d -post cricoid web

16 haemolytic anemia is not produced by:

- a-penicillin
- b-lithium)√(
- c -Quinidine
- d -methyldopa

17-sideroblastic anemia may be treated by all except:

- a-pyridoxine
- b-hydroxy urea)√(
- c -desferroxamine
- d-androgens

18-Henoch schonlien purpura is not associated with:

- a-thrombocytopenia)√(
- b -palpable purpura
- c-intussusception
- d-acute diffuse glomerulonephritis

19 -Outstanding feature of ITP:

- a-fever
- b -gum bleeding)√(
- c -moderate splenomegaly
- d -sterna tenderness

20 -Thrombocytopenia is absent in:

- a-DIC
- b-Wiskottt Aldrich syndrome
- c -Henoch schonlien purpura)√(
- d -myelosclerosis

21 -Cooley's anemia is:

- a-Sickle cell an.
- b-thalassemia major)√(
- c -high ESR
- d-aplastic an.

22 -presence of an .-jaundice -splenomegaly with increase MCH is seen in:

- a-liver cirrhosis
- b-th.major
- c-PNH
- d-herditary spherocytosis)√(

23-all may complicate BM transplantation except:

- a-cataract formation
- b-leucoencephalopathy
- c -cardiomyopathy
- d -emphysema)√(

24-incorrect about pernicious an:

- a-hyperchlorhydria)√(
- b-premature graying of hair
- c -anti intrinsic factor antibody in 60% of pts.
- d -gastric polyp may occur

25-Busulfan therapy lead to all except:

- a-hyperpigmentation
- b-pulm .Fibrosis
- c-optic neuritis)√(
- d -BM suppression

26-increase serum iron —decrease IBC a feature of:

- a-Hookworm infestation
- b-sideroblastic an). $\sqrt{(}$
- c-alcoholic liver dis.
- d-th.major

27 -hepato-splenomegaly with lymphadenopathy occur in all except:

- a-ALL
- b-lymphoma
- c-CML)√(
- d-dissiminated TB

28 - sickle cell an .ls not complicated by:

- A -papillary necrosis
- b-pancreatitis)√(
- c -osteomyelitis
- d -CHF

29 -decrease iron &decrease iron binding capacity are seen in:

- a-recurrent GIT bleeding
- b -intestinal resection
- c -chronic infection) $\sqrt{(}$
- d-menorrhagia

30-Plat .Transfusion is not indicated in:

- a-aplastic an.
- b-uraemia with bleeding

c-DIC

d-immunogenic thrombocytopenia) $\sqrt{(}$

31-best ttt modality in CML is:

a-hydroxyurea

b-allogenic BMT)√(

c-interferon alpha

d-radiotherapy

32-G6PD reflect false normal report in:

a-iron def .an.

b -hypoplastic an.

c-hairy cell leuk.

d-shortly after haemolysis) $\sqrt{(}$

33-half life of plat .ls:

a-1-2days

b-3-4days)√(

c-5-6days

d-7-8days

34-Macrocytosis is a characteristic of all except:

a-an .Of myxedema

b -methotrexate induced

c -chronic alcoholism induced liver dis.

d -SLE)√(

35-which is associated with prolonged bleeding time:

a-polythycemia vera

b-Von Willebrand dis).√(

c -antiphospholipid syndrome

d -haemophilia

36-peripheral bl . Picture is most useful in:

a-NHL

b-multiple myeloma

c-myelodysplastic syndrome

 $d - CML)\sqrt{(}$

37 -splenectomy is contraindicated in:

a-pyruvate kinase def.

b-ITP

c-BM failure)√(

d-angiogenic myeloid metaplasia) myeloproliferative.