

Q23 a) The function of Taq Polymerase) is to produce multiple copies of DNA in PCR. we used Taq polymerase because of heat tolerant so it can work at high temperature.

b) loading dyes is mixed with DNA samples for used in gel electrophoresis dyes generally show us how fast our gel is running. → while agarose gel separate DNA fragment at a specific size.

c) primary antibody.  
it is antibody which is directly attached or bind to the antigen the region of primary antibody recognize the epitope on the antigen.

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Secondary antibody:

It binds with the primary antibody to help in detection. Sorting and purification of target antigens (primary and secondary both are added in indirect Elisa.)

D) Blotting paper is used in Southern blotting for finding the specific DNA fragment.

OR:

It is a hybridization technique for the identification of specific nucleic acid and gene and it is mainly used during different stages of gene expression (DNA, RNA)



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Q2 (a) (1) The diagnosis of Corona virus occurs through a Special PCR known as RT-PCR.

It is a nuclear derived technique through which the Corona virus can be found in even a small sample from the patient and it allows for the rapid detection which is vital to limit the spread of Corona virus.

Originally in this method radioactive isotope markers are used to detect genetic materials. more frequently fluorescence dyes are used now a days.

② Specimens Collection from throat, nasal nasopharyngeal, Sputum and bronchial fluids.

R.T-PCR are used in expression profiling to determine the expression of gene it is also to identify the sequence of RNA transcript including transcription start and termination sites.

→ ⑤ R.T-PCR can be carried out by one step R.T-PCR protocols or two step R.T-PCR protocols.

1) One step R.T-PCR take mRNA targets and subjects them to reverse transcription and then PCR amplification in a single test tube.

2) Two step R.T-PCR as the name implies occur in two step first the reverse transcription and then the PCR.



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(c) USE:-

- ① N95 mask.    ② lab coat.
- ③ Hand washing facilities  
Such as Sanitizers.
- ④ Gloves.
- ⑤ Avoid to touch face  
nose eyes during experiment.
- ⑥ protection of eyes from  
lab goggles.
- ⑦ head cover.    ⑧ Foot wear.
- ⑨ wastes should be thrown  
dustbin or bin.
- ⑩ Avoid eating or drinking.  
Smoking were the experiment occur.

Procedure:-

All procedure should be performed in the way that minimizes the formation of aerosols and droplets.

-mouth pipetting must be strictly prohibited.

• All these infectious items which can cause splashes, droplets.

• use of BCS mainly (class II BCS).

Q1 (a) 1) most people are diagnosed with CML through blood test known as CBC Complete blood count.

→ CBC is done for a regular checkup and in CML it is used for WBCs finding because CML patients have high number of WBCs.

Platelets counts

2) Bone marrow aspiration and biopsy. its is a process or procedure in which a hollow needle is inserted in the hip bone and reached that needle into bone marrow and collect a sample for the diagnosed of CML.

3) A CT scan show if any lymph node in your ~~body~~ body become enlarged.

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(34) MRI is mainly used for brain and spinal cord.

(5) PCR in CML is used to look for the BCR-ABL genes in leukemia cell.

(6) FISH test is used for chromosomal study (BCR-ABL)

(7) Genetic test.

(8) Blood chemistry test.

(9) Bone marrow test.

(10) Imaging test which is not usually needed.

But some time necessary for your liver picture.

(11) Biochemical finding.

→ Q1 (b) The main cause of CML is unknown but it is believed that it is caused from the genetic mutation.

During the swapping of DNA between the chromosome (22 and 9) lead the formation of genes which is known as BCR-ABL.

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Now this gene produced a specific proteins called Tyrosine kinase, this proteins cause cml cells to grow and divide it out of control as a results of cml occur.

→ Q1 (c) Philadelphia occur due to the translocation of chromosome 22 and 9 respectively. So it mean that it is mutation which only occurs due to normals chromosomes which is not inheritable but a small amount of BCR-ABL oncogen are present in cml patients which may or may not inherited to their child or offspring. Supply we can say that it is a somatic mutation which is not inheritable to ~~off~~ offspring.