



BIOLOGY

BOOKS - UNIVERSAL BOOK DEPOT 1960 BIOLOGY (HINGLISH)

BIOTEHNOLOGY : PRINCIPALES AND PROCESSES

Biotehnology Principales And Processes

1. The linking of antibiotic resistance gene with

the plasmid vector became possible with

A. DNA polymerase

B. Exonucleases

C. DNA ligase

D. Endonucleases

Answer: C

2. Main objective of production of herbicide resistant GM crops is to A. Encourage eco-friendly herbicides B. Reduce herbicide accumulation in food articles for healthy safty C. eliminate weeds from the field without the use of manual labour D. Eliminate weeds from the field without the use of herbicides

Answer: b



3. Which of these is used as vector in gene therapy for SCID

A. Arbovirus

B. Rotavirus

C. Enterovirus

D. Paravovirus

Answer: c





4. Which of the flowing has the ability to transform normal cell into cancerous cell in animal

A. Arbovirus

B. Rotavirus

C. Enterovirus

D. Retrovirus







5. Which one among the following is just a cloning plasmid not an expression plasmid

A. pBAD-18-Cam

B. P BCSK

C. pUC18

D. pET

Answer: c

6. Branch dealing with genetic engineering is

A. Genetic engineering

B. Euthenics

C. Euphenics

D. None of these

Answer: c

7. Genetic engineering means:

A. Manipulation of cell contents

- B. Test tube babies
- C. Manipulation of cytochromes
- D. Manipulation (modification) of genes

Answer: d



8. Who among the following scientists is associated with the discoveries in genetic engineering

A. Khorana

B. Watson

C. Crick

D. Messleson

Answer: a

9. It is now possible to breed plants and animals with desired characters through

A. Genetic engineering

B. Chromosome engineering

C. lkebana technique

D. tissue culture

Answer: a

10. Which of the following organelles is releated with genetic engineering/gene cloning

A. Golgi apparatus

B. Lysosomes

C. Mitochondria

D. Plasmids

Answer: d

11. In genetic engineering, a DNA segment (gene) of interest, is translated to the host cell through a vector. Consider the following four agents (A-D) in this regard and select the correct option about which one or more of these can be used as a vector/vectors Statement (A) A bacterium (B) Plasmid

(C) Plasmodium (D) Bacteriophage

A. (A),(B) and (D) Only

B. (A) only

C. (A) and (C) only

D. (B) and (C) only

Answer: d



12. Recombinant DNA (rDNA) tenchnology is

related with

A. C.darwin

B. Stanley Cohen

C. Herbert Boyer

D. Both (b) and (c)

Answer: D



13. A desirable change in genotype of an organism is obtained by

A. DNA replication

B. Protein synthesis

C. rDNA technology

D. mRNA formation

Answer: C



14. Which of these is widely used in genetic

engineering

A. Anopheles

B. Dragon fly

C. Dragon lizard

D. Fruit fly

Answer: d



15. Identify the plasmid

A. AIU I

B. Hind III

C. Eco RI

D. pBr 322

Answer: d

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16. In recombination vector used is

A. Protein

- B. Agrobacterium tumerfaciens
- C. Nucleic acid
- D. Cellulose

Answer: b



17. First biochemical to be produced commercially by microbial cloning and genetic engineering is

- A. Human insulin
- B. Penicillin
- C. Interferons
- D. Fertility factors





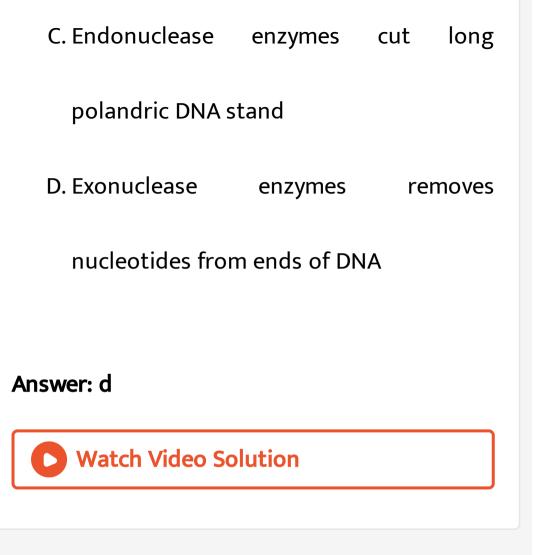
18. Which of the following option is correct for recombinant DNA technology

A. Exonuclease enzyme removes nuclotides

from site within DNA

B. Endonuclease enzyme removes

nucleotides from the ends of DNA



19. Restriction endonucleases are most widely

used in recombinant DNA technology. They are

obtained from

- A. Bacteriphages
- **B. Bacterial cells**
- C. Plasmids
- D. All prokaryotic cells

Answer: b

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20. In recombinant DNA technology, the term

vector refers to

A. Plasmids that can transfer foreign DNA into a living cell B. Cosmids that can cut DNA at specific base sequences C. Plasmids that can join different DNA fragments D. Cosmids that can degrade harmful

proteins

Answer: A

21. An analysis of chromosomal DNA using the southern hybridization technique does not use

A. Autoradiography

B. PCR

C. Electrophoresis

D. Blotting

Answer: a



22. Genetic engineering has been successfully used for producing

A. Animals like bulls for farm work as they have super power

B. Transgenic mice for testing safety of

polio vaccine before use in humans

C. Transgenic models for studying new

treatements for certain cardiac diseases

D. Transgenic Cow-Rosie which produces

high fat milk for making ghee

Answer: b

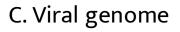
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23. The gene 'rop' present in pBR322 cloning

vector, codes for

A. Original bacterial plasmid

B. Modified bacterial plasmid



D. Transposon

Answer: b



24. Which one of the following techniques made it possible to gnetically engineer living organisms ?

A. Heavier isotope labeling

B. Hybridization

C. Recombinant DNA techniques

D. X-ray diffraction

Answer: c

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25. Which one of the following technique made it possible to genetically engineering living organisms

A. Heavier isotope labeling

B. Hybridization

C. Recombinant DNA techniques

D. X-ray diffraction

Answer: c

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26. Following enzymes /techniques are used in

the process of recombinant DNA technology

A. Eco RI to cut the isolated genome

B. DNA ligase

C. Protease and ribonuclease for removal of proteins and RNA from DNA D. Production of recombination hosts E. lysozyme for isolation of the genetic material (DNA) F. gel electrophoresis for separation and isolation of DNA fragments Mark the correct sequence of their use.

A. Restriction endonucleases and

topoisomerases

B. Endonucleases and polymerases

C. Restriction endonucleases and ligases

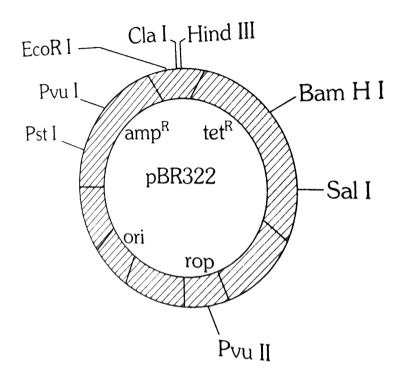
D. Peptidases and ligases

Answer: d

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27. The figure below is the diagrammatic representation of the E.coli vector pBR322. which one of the given options correctly

identifies its certain component (s)



- A. Ori-original restriction enzymes
- B. Ro-reduced osmotic pressure
- C. Hind III, EcoRI -Selectable markers

D. amp^R , tet^R -antibiotic resistance genes

Answer: D



28. PCR and restriction Fragements length Polymorphism are the methods for

A. Study of enzymes

- B. Genetic transformation
- C. DNA sequencing
- D. Genetic -Fingerprinting

Answer: d



29. Fearing that the child to be born may have a genetic disorder, a couple goes to a doctor. Which one of the following techniques is likely to be suggested by the doctor to cure the genetic disorder ?

A. Hybridoma technology

B. Gene therapy

C. rDNA technology

D. Embryo transfer

Answer: b



30. What is the permanent cure of adenosine

deaminase (ADA) deficiency in children ?

A. Hybridoma technology

B. Gene therapy

C. rDNA technology

D. Embryo transfer

Answer: b



31. In genetic engineering, the antibiotics are

used

A. As selectable markers

B. To select healthy vectors

C. As sequences from where replication

starts

D. To keep the cultures free of infection

Answer: a

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32. The colonies of recombinant bacteria appear white in because of

A. Inactivation of glycosidase enzyme in				
recombinant bacteria				
B. Non-recombinant bacteria containing				
beta -galactosidase				
C. Insertional inactivation of alpha-				
galactosidase in non-recombinant				
bacteria				
D. Insertional inactivation of alpha-				
galactosidasea in recombinant bacteria				

Answer: c



33. Precipitates of purified DNA after the addition of chilled ethanol are seen as a collection of the fine threads in suspension. This process is referred as



A. Precipitate DNA

B. Break open the cell to release DNA

C. Facilitate action of restriction enzymes

D. Remove proteins such histones

Answer: a

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34. The below figure refers to recombinant DNA technology indentify A,B,C,D respectively

	Foreign DNA C join foreign DNA C join foreign DNA to plasmid					
	A	В	C	D		
	Restriction Endonuclease	Restriction Endonuclease	DNA ligase	Transformation		
	Exonuclease	Endonuclease	Hydrolase	Transduction		
1	Endonuclease	Exonuclease	DNA ligase	Transformation		

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Endonuclease

(a)

(b)

(c)

(d)

Exonuclease

35. When the chilled ethanol is added in purified DNA, it ultimately precipitates out. The can be show in the figure as collection of

DNA ligase

Transformation

fine threads in the suspension. This process is

known as





A. DNA bands

- **B. DNA recognition**
- C. DNA digestion
- D. DNA spooling

Answer: d





36. Which of the following is not a feature of

the plasmids

A. Independent replication

B. Circular structure

C. Transferable

D. Single-standed

Answer: d

37. Which of the following is a restriction endonulease

A. Hind II

B. protease

C. Dnase l

D. Rnase

Answer: d

38. Which of the following restriction enzymes

produces blunt ends

A. Hind III

B. Sal I

C. Eco RV

D. Xho I

Answer: c

39. A gene whose expression helps to indentify

transformed cell is known as

A. Selectable marker

B. Vector

C. Plasmid

D. Structural gene

Answer: a

40. The correct order of steps in Polymerase Chain Reaction (PCR) is

A. Extension, Denaturation, Annealing

B. Annealing , Extension, Denaturation

C. Denaturation, Extension, Annealing

D. Denaturation, Annealing, Extension

Answer: D

41. The correct order of steps in polymerase chain Reaction (PCR) is

A. Extension, Denaturation, Annealing

B. Annealing , Extension, Denaturation

C. Denaturation, Extension, Annealing

D. Denaturation, Annealing, Extension

Answer: D

42. Rising of dough is due is

A. Multiplication of yeast

B. Production of CO_2

C. Emulsification

D. Hydrolysis of wheat flour starch into

sugars

Answer: b

43. An enzyme catalysing the removal of

nucleotides from the ends of DNA is

A. Endonuclease

B. Exonuclease

C. DNA ligase

D. Hind-II

Answer: b

44. The transfer of genetic material from one bacterium to another through the mediation of a vector like virus is termed as

A. Transduction

B. Conjugation

C. Tranformation

D. Translation

Answer: a

45. Which of the given statements is correct in the context of observing DNA separated by agarose gel electrophoresis ?

A. DNA can be seen in visible light

B. DNA can be seen without staining in

visible light

C. Ethidium bromide stained DNA can be

seen in visible light

D. Ethidium bromide stained DNA can be

seen under exposure to UV light





46. Restriction' in restriction enzyme refers to

A. Cleaving of phsphodiester bond in DNA

by the enzyme

B. Cuttig of DNA at specific position only

C. Prevention of the multiplication of

bacteriophage by the host bacteria

D. All of the above

Answer: C

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47. Which of the following is not required in the preparation of a recombinant DNA molecule ?

A. Restriction endonuclease

B. DNA ligase

C. DNA fragments

D. E coil

Answer: d



48. Which of the following statements does

not hold true for restriction enzyme ?

A. it recognize a palindromic nucleotide

sequence

- B. it is an endonuclease
- C. it is isolated isolated from viruses
- D. It can produce the same kind of sticky

ends in different DNA molecules

Answer: c

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49. The most important feature in a plasmid to

be used as a vector is



B. presence of a selectable marker

C. Presence of sites for restriction

endonuclease

D. it size

Answer: a

50. While isolating DNA from bacteria, which

of the following enzymes is not used ?

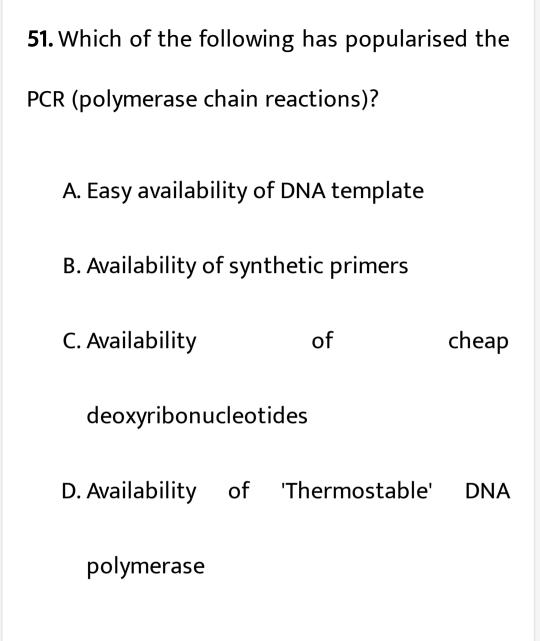
A. Lysozyme

B. Ribonuclease

C. Deoxyribonuclease

D. Protease

Answer: C



Answer: d

52. An antibiotic resistance gene in a vector usually helps in the selection of

A. Competent bacterial cells

B. Transformed bacterial cells

C. Recombinant bacterial cells

D. None of the above

Answer: b

53. Significance of 'heat shock' method in bacterial transformation is to facilitate

- A. Binding of DNA to the cell wall
- B. uptake of DNA through membrane

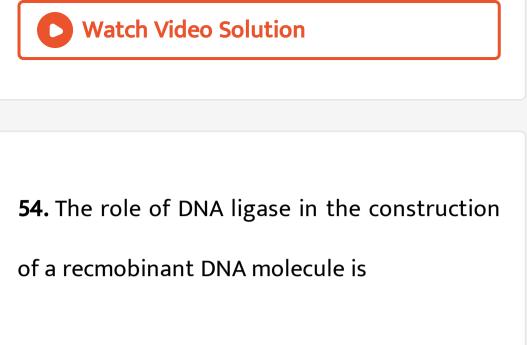
transport proteins

C. Uptake of DNA through transient pores

in the bacterial cell wall

D. Expression of antibiotic resistance gene

Answer: c



A. Formation of phosphodiester bond

between two DNA fragments

B. Formation of hydrogen bonds between

sticky ends of DNA fragments

C. Ligation of all purine and pyrimidine

bases

D. None of the above

Answer: a

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55. Which of the following is not a source of restriction endonuclease ?

A. Haemophilus infuenzae

B. Escherichia coli

C. Entamoeba coli

D. Bacillus amyloliquefaciens

Answer: c

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56. Which of the following steps are catalysed by Taq polymerase in a PCR reaction ?

A. Denaturation of template DNA

B. Annealing of primers to template DNA

C. Extension of primer end of the template

DNA

D. All of the above

Answer: c

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57. A bacterial cell was transformed with a recombinant DNA that was generated using a human gene. However, the transformed cells

did not produce the desired protein. Reason could be

A. Human gene may have intron which

bacteria cannot process

B. Amino acid codons for humans and

bacteria are different

C. Human protein is formed but degraded

by bacteria

D. All of the above

Answer: a



58. Which of the following should be chosen for best yield if one were to produce a recmobinant protein in large amounts ?

A. Laboratory flask of largest capacity

B. A stirred -tank bioreactor without in -lets

and out -lets

- C. A continous culture system
- D. Any of the above

Answer: c



59. Who among the following was awarded the Nobal Prize for the development of PCR technique ?

A. Herbert Boyer

B. Hargovind Khurana

C. Kary mullis

D. Arthur kornberg

Answer: c



60. During DNA purification, which enzymes is used to treat the plant cell

A. Ribonuclease

B. Cellulase

C. Chitinase

D. Ligase





61. The stired-tank reactor is usually

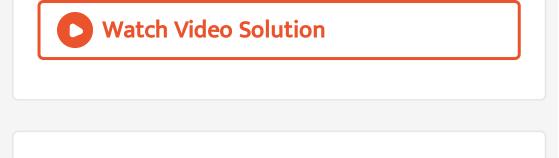
A. Cylinderical

B. Rounded

C. Cup-Shaped

D. Flattened

Answer: A



- 62. A bioreactors refers to
 - A. A device in which substances are treated
 - to stimulate biochemical transformation
 - by living cells
 - B. A nuclear reactor for biological studies
 - C. A tank for biochemical reactions
 - D. Organisms badly reacting to stimuli





- 63. Chimeric DNA is
 - A. A part of recombinant DNA
 - B. In fact passenger DNA
 - C. Recombinant DNA formed by combining

vector DNA and passenger DNA

D. Residual DNA that has no role in genetic

engineering

Answer: c

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64. Which one of the following correctly explains the term 'Chimera'

A. Spontaneously induced deletions

B. Breaking a part of chromosome segment during mutation C. Development of genetically diverse tissues in the same organism D. During mutation at segregation,

chromosome become equally distributed

Answer: c

65. Why is recombinant DNA (rDNA) technology called genetic engineering A. It involves sophisticated technology at microscopic level B. Knowledge of engineering of two DNAs C. It involves manipulation degree in

engineering

D. It includes an authorized degree in engineering

Answer: c



66. In rDNA technology in order to make the bacterial host cells 'component' to accept the rDNA, these are kept in

A. Dilute solutoin of CsCl

B. Divalent anions such as phosphates

C. Chilled ethanol

D. Divalent cations such as calcium

Answer: D



67. Which of the given statements is correct in the context of observing DNA separated by agarose gel electrophoresis ?

A. DNA can be seen in visible light

B. DNA can be seen without staining in

visible light

C. Ethidium bromide stained DNA can be

seen in visible light

D. Ethidium bromide stained DNA can be

seen under exposure to UV light

Answer: d

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68. Recombinant DNA is force to enter the host cells by incubating the cells with rDNA

first

A. On ice followed by heat shock and then

again on ice

B. At low temperature followed by heat

shock and then on ice

C. In ethidium bromide followed by calcium

salts

D. Into chilled ethanol followed by ice and

then into CsCl

Answer: a

69. The desired product of rDNA technology produced through bioreactors on large scale undergoes

A. Elution

- B. Enzymatic action
- C. Biomonitoring
- D. Downstream processing

Answer: d





70. The cloning vectors M13 has genetic material

A. ssRNA

B. dSRNA

C. ssDNA

D. dsDNA

Answer: c



71. Match column I and column II and select

the right option given below

Column I			Column II	
I.	Recombinant DNA technology	A.	Vector	
II.	Cloning Vehicles	В.	Sealing enzyme	
III.	Macromolecular Separation	C.	Electrophores is	
IV.	DNA Ligase	D.	Genetic engineering	

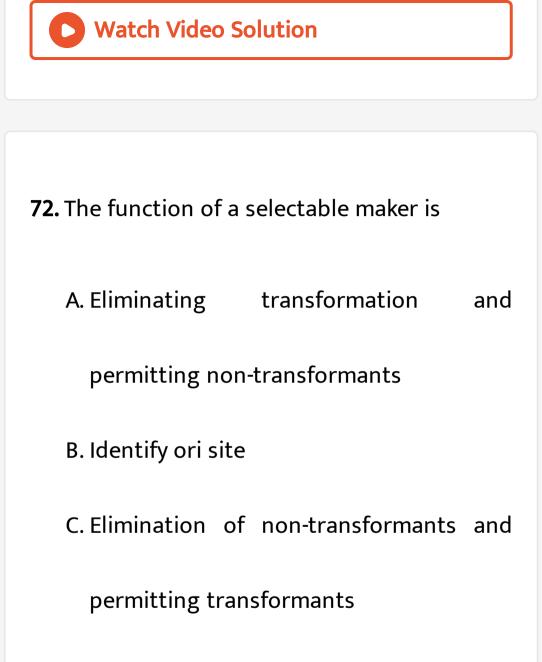
A. I-D,II-A,III-B,IV-C

B. I-A,II-D,III-B,IV-C

C. I-D,II-A,III-C,IV-B

D. I-B,II-A,III-D,IV-C

Answer: c



D. To destroy recognition sites

Answer: C



73. Which of the followijng is/are used in recombinant DNA technology
A. Agarose gel (B). Ethidium bromide
(C) . Plasmid vector (D) Restriction endonuclease

A. A,B

C. C,D

D. A,B,C,D

Answer: d



74. DNA recombinant technology uses

A . Restiction endonucleases B. DNA ligase

C. Cloning vector D. Electrophoresis

B. B,C

C. C,D

D. A,B,C,D

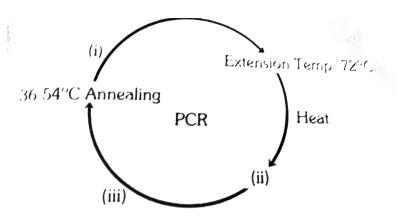
Answer: d

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75. The following cycle refer to the PCR prcess,

name the factors or steps indicated with

numbers



A. (i) Taq polumerase (ii) Extension (iii) Ligation

B. (i) Primer (ii) Denaturation at $94^{\,\circ}C$ (iii)

Taq polymerase

C. (i) Denaturation at $94^{\circ}C$ (ii) Taq

polymerase (iii) Primer

D. (i) Denaturation at $94^{\,\circ}\,C$ (ii) Taq

polymerase (iii) Primer

Answer: d

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76. Observe the figure and select the correct

option out of (a-d)

	A DNA	B DNA	Enzyme recognizing palindrome	Enzyme joining the sticky ends
(a)	Vector	Foreign	DNA ligase	Exonuclease
(b)	Vector	Foreign	Exonuclease	DNA ligase
(c)	Vector	Foreign	EcoRI	DNA ligase
(d)	Vector	Foreign	DNA ligase	EcoRI





77. Genetically engineered bacteria are being

employed for production of

A. Thyroxin

B. Progesterone

C. Insulin

D. Estrogen

Answer: c

78. The first human hormone drug produced by recombinant DNA technology genetic engineering is

A. Thyroxin

B. Progesterone

C. Insulin

D. Estrogen

Answer: C



79. Which vector can clone only a small fragement of DNA

A. Plasmid

B. Cosmid

C. Bacterial artificial chromosomes

D. Yeast artificial chromosomes

Answer: a

80. Which of the following is not a component

of downstream processing

A. Expression

B. Separation

C. Purification

D. Preservation

Answer: A

81. The process of separation and purificaition of expressed protein before marketing is called

A. Upstream processing

B. Downstream processing

C. Bioprocessing

D. Postproduction processing

Answer: b

82. Assertion : in recombinant DNA technology, human genes are often transferred into bacteria (prokaryotes) or yeast (eukaryote). Reason: Both bacteria and yeast multiply very fast to form huge population which express the desired gene.

A. If both the assertion and the reason are

true and the reason is a correct

explanation of the assertion

B. If both the assertion and reason are true

but the reason is not a correct

explanation of the assertion

C. IF the assertion is true but the reason is

false

D. If both the assertion and reason are

false

Answer: a

83. Assertion : Plasmids are extrachromosomal DNA.

Plasmids are found in bacteria and are useful

in genetic engineering

A. If both the assertion and the reason are

true and the reason is a correct

explanation of the assertion

B. If both the assertion and reason are true

but the reason is not a correct

explanation of the assertion

C. IF the assertion is true but the reason is

false

D. If both the assertion and reason are

false

Answer: a

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84. Assertion : Plasmids are single stranded extra chromosomal DNA.

Reason: Plasmids are found in Eukaryotic cells.

A. If both the assertion and the reason are true and the reason is a correct explanation of the assertion B. If both the assertion and reason are true but the reason is not a correct explanation of the assertion C. IF the assertion is true but the reason is false D. If both the assertion and reason are false

Answer: d



85. Assertion : Recognition site should be perfectly single and responsive to commonly used restriction enzymes.
Reason: In pNR 322 Alien DNA is ligated generally in the area of Bam-HI site of tetracycline resistance gene.

A. If both the assertion and the reason are true and the reason is a correct explanation of the assertion B. If both the assertion and reason are true but the reason is not a correct explanation of the assertion C. If the assertion is true but the reason is false D. If both the assertion and reason are false

Answer: B



86. Assertion: A gene from bacillus thuringiensis is incorporated in plant genome to increases yeild. Reason : it is Bt toxin producing genw which

kills larvae of insects.

A. If both the assertion and the reason are

true and the reason is a correct

explanation of the assertion

B. If both the assertion and reason are true

but the reason is not a correct

explanation of the assertion

C. IF the assertion is true but the reason is

false

D. If both the assertion and reason are

false

Answer: a

87. Which of the following tools of recombinant DNA technology is incorrectly paired with its use

A. Restriction enzyme-production of RFLPs B. DNA ligase-enzyme that cuts DNA, creating the stickly end of restriction fragments C. DNA polymerase -used in a polymerase chain reaction to amplify sections of DNA

D. Reverse transciptase -production of c

DNA from mRNA

Answer: b

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88. In recombinant DNA technology, the term

vector refers to

restriction fragments

B. the sticky end of DNA fragments

C. A plasmid used to transger DNA into a

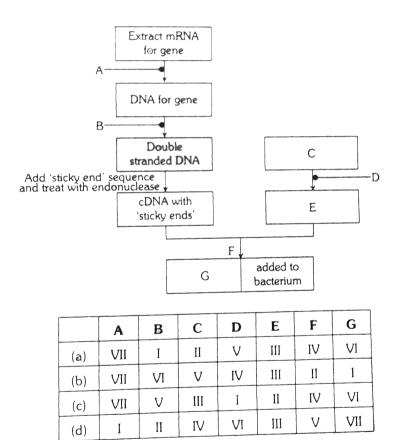
living cell

D. A DNA probe used to identify a

particular gene

Answer: c

89. Identify the labelled iterms A,B,C,D,E,F and G in the diagrame below from the list-I to VII given with



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90. When a recombinant DNA is inserted within the coding sequencing of an enzymes, β -galactosidase

A. This result into inactivation of the enzyme

- B. This is called insertional inactivation
- C. in the presence of insertion, the colonies

do not produce any colour

D. All of these

Answer: D



91. If recombinant DNA is inserted within the coding sequence of enzyme galactosidase, which of the following will occur in case of non -recombinants

A. Insertional inactivation

B. Colonies do not produce any colour

colour

D. Inactivation of enzyme galactosidase

Answer: C

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92. Following enzymes /techniques are used in

the process of recombinant DNA technology

A. Eco RI to cut the isolated genome

B. DNA ligase

C. Protease and ribonuclease for removal of

proteins and RNA from DNA

D. Production of recombination hosts

- E. lysozyme for isolation of the genetic material (DNA)
- F. gel electrophoresis for separation and isolation of DNA fragments

Mark the correct sequence of their use.

A. C,E,B,F,A,D

B. E,C,A,B,F,D

C. E,C,A,F,B,D

D. A,E,C,B,D,F

Answer: c

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93. I. Recombinant DNA technolgy is used to improve crop plants by increasing their productivity, by making them more nutritious and by developing disease resistant
II Bt cotton is resistance to bollworm infestation

III. Bacillus thuringiensis is not harmed by selfcry protein because of its occurrence asprotoxin (inactive)V. Protoxin Cry protein is changed into active

cry protein in the stomach of insects due to alkaline pH in stomach

A. All are correct

- B. I and iV are correct
- C. only III is false
- D. All are false

Answer: c



94. DNA polymerase enzyme is isolated from

which bacteria

A. E.Coli

- B. Thermus aquaticus
- C. Bacillus thuringenesis
- D. Agro bacterium

Answer: a





- 95. Which of the following techniques serve
- the purpose of early diagnosis
- I. r-DNA technology
- II PCR
- III ELISA
- IV Convential method of diagnosis (serum, urine analysis,etc)
 - A. I,II,III
 - B. IV only

C. III only

D. All

Answer: a



96. The DNA molecule to which the gene of

interst is integrated for cloning is called

A. Vector

B. Template

C. carrier

D. Transformer

Answer: a



97. The application of microbial metabolism to

trensform simple raw materials into valuabale

products is

A. Bioactalysis

- B. Genetic engineering
- C. Tissue culture
- D. Fermentation

Answer: d

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98. For rapid production of alcohol , immobilised yeast calles are kept in

A. Silica gel

- B. Wire netting
- C. Porcelain columns
- D. Calcium alginate beads

Answer: d

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99. What is the source of EcoRI

A. Escherichia coli RI

B. Escherichia coli RI 13

C. Escherichia coli 13

D. Escherichia Coli RY 13

Answer: D