

BIOLOGY

BOOKS - MTG BIOLOGY (ENGLISH)

BIOTECHNOLOGY - PRINCIPLES AND PROCESSES



- 1. Who is the father of genetic engineering?
 - A. Steward Linn
 - B. Stanley Cohen
 - C. Paul Berg
 - D. Kary Mullis

Answer: C



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2. Which	of the	following	processes/	techniques'	can	be	included	under
biotechn	ology?							

- (i) In vitro fertilisation
- (ii) Synthesis of a gene
- (iii) Correcting a defective gene
- (iv) Developing a DNA vaccine
 - A. (i) and (ii)
 - B. (ii) and (iii)
 - C. (iii) and (iv)
 - D. (i),(ii),(iii) and (iv)

Answer: D



3. Plasmid used to construct the first recombinant DNA was isolated from which bacterium species?

A. Escherichia coli

B. Salmonella typhimurium

C. Agrobacterium tumefacines

D. Thermus aquaicus

Answer: B



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- **4.** Genetic engineering is possible because
 - A. We can cut DNA at specific sites by restriction endonucleases
 - B. restriction endonucleases purified sites by restriction used in

bacteria

C. the phenomenon of transduction in bacteria is well understood

D. we can see DNA by electron microscope
Answer: A
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5. The term 'molecular scissors' refers to
A. recombinant DNA
B. restriction enzymes
C. Taq polymerase
D. polindromic nucleotide sequences.
Answer: B
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6. The term 'chemical knife' refers to

A. polymerases B. endonucleases C. ribonucleases D. cellulases **Answer: B View Text Solution**



- 7. In recombinant DNA technology, the term vector refers to
 - A. the enzyme that cuts DNA into restriction fragments
 - B. the sticky end of a DNA fragment
 - C. a plasmid used to transfer DNA into a living cell
 - D. a DNA fragment which carries only ori gene

Answer: C



8. One of the key factors, which makes the plasmid the vector in genetic engineering is

A. its resistance to antibiotics

B. its resistance to restriction enzymes

C. its ability to carry a foreign gene

D. its ability to cause infection in the host

Answer: C



- 9. The term 'recombinant DNA' refers to
 - A. DNA of the host cell
 - B. DNA with a piece of foreign DNA
 - C. DNA with selectable marker

D. DNA with more than one recognition sites
Answer: B
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10. The term 'chimeric DNA' refers to
A. DNA with overhanging stretches

B. DNA with palindromic sequences

C. a recombiant DNA

D. molecular scissors

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Answer: C

11. Which of the following contains the key tools for recombinant DNA technology?

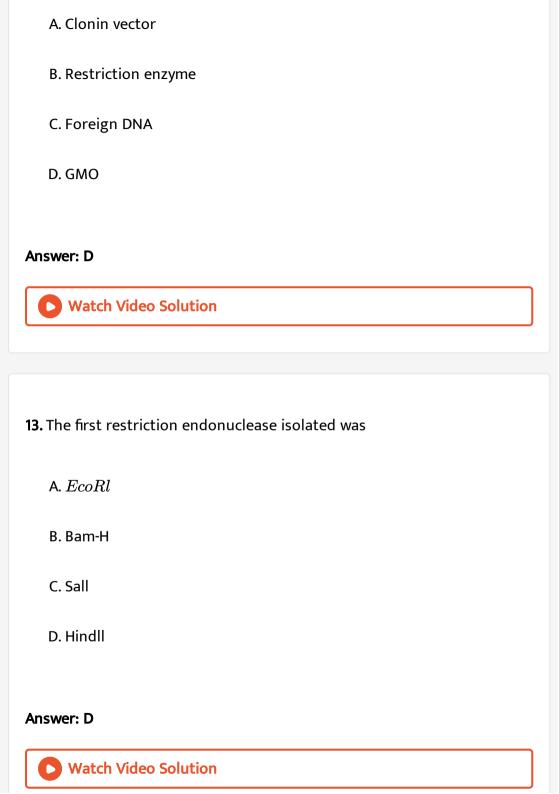
- (i) Restriction endonucleases, ligases, vectors
- (ii) Ligases, host organism, ligases, vectors
- (iii) Vectors, Taq polymerase, primers
- (iv) Restriction exonucleases, ligases, primers, bioreactors
 - A. (i), (ii) and (iii)
 - B. (i) and (ii)
 - C. (i), (iii) and (iv)
 - D. (iii) and (iv)

Answer: B



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12. Which of the following is not a tool of genetic engineering?



14. The letter 'R' in EcoRl is derived from

A. the name of genus

B. the name of strain

C. the name of species

D. the term 'restriction'

Answer: B



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15. Match column I with column II with respect to the nomenclature of restriction enzyme EcoRl and select the correct answer from the given codes.

ColumnII ColumnIII

A. E (i) 1^{st} in order of identification

B. co (ii) Name of genus C. R (iii) Name of species

D. l (iv) Name of strain

B. A-(ii), B-(i), C-(iii), D-(iv)

(a) A - (iii), B - (i), C - (ii), D - (iv)

(b) A - (ii), B - (i), C - (iii), D - (iv)

(c) A - (i), B - (ii), C - (iii), D - (iv)

(d) A - (ii), B - (iii), C - (iv), D - (i)

A. A - (iii), B - (i), C - (ii), D - (iv)

C. A - (i), B - (ii), C - (iii), D - (iv)

D. A-(ii), B-(iii), C-(iv), D-(i)

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Answer: D

A. Escherichia coli RY13

- B. Haemophilus influenzae Rd

16. The source of the restriction enzyme Hindll is

C. Bacillus amyloliuefaciens ${\cal H}$

Answer: B



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- 17. A restriction endonuclease breaks bonds between the
 - A. base pairs of a DNA molecule
 - B. base pairs of a DNA-RNA hybrid molecule
 - C. sugar and phosphate components of a nucleic acid molecule
 - D. exons and introns of a DNA molecule.

Answer: C



18. Read the given statements and select the correct option,

Statement 1 : Restriction endonuclease enzymes recognise a specific palindromic nucleotide sequence in the DNA

Statement 2: Restriction endonuclease enzymes are called as molecular scissors or biological scissors.

- A. (a) Both statements 1 and 2 are correct
- B. (b) statement 1 is correct but statement 2 is incorrect
- C. (c) statement 1 is incorrect but statement 2 is correct
- D. (d) both statements 1 and 2 are incorrect

Answer: A



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19. Study the following figures and identify the enzymes involved in steps I and II.



- A. EcoRl and DNA Ligase
- B. Hind II and DNA Ligase
- C. EcoRi and HindII
- D. Restriction endonuclease and exonuclease

Answer: A



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- 20. Which of the following correctly depicts the recognition site for EcoRI
- ?
- (a) $G-A-A\stackrel{\downarrow}{-}T-T-C$ C-T-T-A-A-G (b) $G-T-C\stackrel{\downarrow}{-}G-A-C$ $C-A-G\stackrel{\uparrow}{-}C-T-G$
- (c) $G \stackrel{\downarrow}{-} T C G A C \ C A G C T G$
- (d) $G \stackrel{\downarrow}{-} A A T T C \\ C T T A A G$

A.
$$G-A-A\stackrel{\downarrow}{-}T-T-C \ C-T-T\stackrel{\uparrow}{-}A-A-G$$

B.
$$G-T-C \stackrel{\downarrow}{-} G-A-C$$
 \uparrow $C-A-G \stackrel{\uparrow}{-} C-T-G$

c.
$$G \stackrel{\downarrow}{-} T - C - G - A - C$$
 $C - A - G - C - T - G$

D.
$$G \stackrel{\downarrow}{-} A - A - T - T - C$$

 $C - T - T - A - A - G$

Answer: D



- 21. The sticky ends of a fragmented DNA molecule are made of
- A. (a) calcium salts
 - B. (b) endonuclease enzyme
 - C. (c) unpaired bases
 - D. (d) methyl groups

Answer: C



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22. The restrcition enzyme responsible for cleavage of following sequence

$$5'-G-T-C\stackrel{\downarrow}{-}G-A-C-3'$$

$$3'-C-A-G - C-T-G-5'$$

A. (a) EcoRl

B. (b) Hindll

C. (c) BamHl

D. (d) EcoRll

Answer: B



23. Identify the palindromic sequence in the following

- A. $\frac{GAATTC}{CTTUUG}$
- $\mathsf{B.} \; \frac{GGATCC}{CCTAGG}$
- c. $\frac{CCTGG}{GGACC}$
- D. $\frac{CDATA}{GCTAA}$

Answer: B



24. Which of the following statements is not correct regarding EcoRl restriction endonuclease enzyme ?

A. It is isolaed from Escherichia coil RY13

B. Its recongition sequence is $5^{\,\prime}-GAATTC-3^{\,\prime}$

3'-CTTAAG-5'

C. It produces complementary blunt ends.

D. None of these

Answer: C



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25. Which of the following sequences is recongnised by restriction enzyme BamHI?

A.
$$5'-G\overset{\downarrow}{A}ATTC-3' \ 3'-CTTAAG-5'$$

B.
$$5' - \stackrel{\downarrow}{AAGCTA} - 3'$$
 $3' - TTCGAT - 5'$

C.
$$5^{\prime}-\overset{\downarrow}{GGATCC}-3^{\prime}$$
 $3^{\prime}-CCTAGG-5^{\prime}$

D.
$$5^{\prime}-CCC\overset{\downarrow}{A}AT-3^{\prime}\ 3^{\prime}-GGGTTA-5^{\prime}$$

Answer: C



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26. If a plasmid vector is digested with EcoRl at a single site, then

A. one sticky end will be produced

B. two sticky ends will be produced

C. four sticky ends will be produced

D. six sticky ends will be produced

Answer: B



27. How many fragments will be generated if you digest a linear DNA molecule with a restriction enzyme having four recognition sites on the

DNA?

(a) 3

(b) 6

(c) 5

(d) 4

28. How many fragments will be generated on the digestion of a closed circular DNA molecule with a restriction enzyme having six recongnition sites on the DNA?

A. 5

B. 7

C. 6

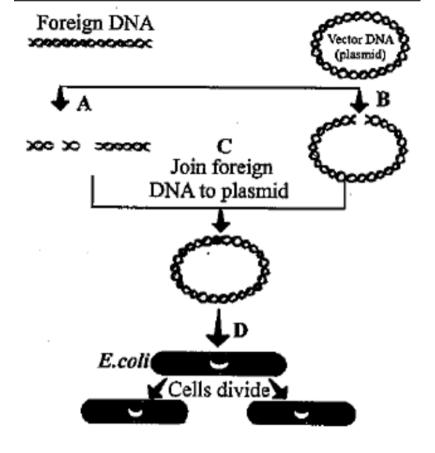
D. 9

Answer: C



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29. The flow chart given below represents the process of recombinant DNA technology. Identify A,B, C and D.



- A. (a) A-Restriction endonuclease, B-Restriction exonuclease, C-DNA ligase, D-Transformation
- B. (b) A-Restriction endonuclease, B-Restriction endonuclease, C-DNA ligase, D-Transformation
- C. (c) A-Restriction endonuclease, B-Restriction endonuclease, C-Hydrolase, D-Transformation

D. (d) A-Restriction endonuclease, B-Restriction endonuclease, C-

Answer: B



Hydrolase, D-Transduction

30. In recombinant DNA technology, a plasmid vector is cleaved by

A. modified DNA ligase

B. a heated alkaline solution

C. the same enzyme that cleaves the donor DNA

D. the different enzyme than that cleaves the donor DNA.

Answer: C



31. Gel electrophoresis is a

A. technique of separation of charged molecules under the influence of magnetic field

B. technique of incorpotation of DNA molecules into the cell through transient pores made due to electrical impulses

C. technique of separation of DNA fragments through the pores of agarose gel under the influence of electric field

D. technique of separation and purification of gene products.

Answer: C



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32. Gel electrophoresis is used for

A. (a) construction of recombinant DNA using cloning vectors

- B. (b) isolation of DNA molecules
- C. (c) cutting of DNA into fragments
- D. (d) separation of DNA fragments according to their size.

Answer: D



- **33.** Having become an expert on gel electrophoresis, you are asked to examine a gel. Where would you find the smallest segments of DNA?
- (a) Near the positive electrode, farthest away from the wells
- (b) Near the negative electrode, close to the wells
- (c) Near the negative electrode, farther away from the wells
- (d) Near the middly, they tend to slow down after the first few minutes
 - A. Near the positive electrode, farthest away from the wells
 - B. Near the negative electrode, close to the wells
 - C. Near the negative electrode, farther away from the wells

D. Near the middly, they tend to slow down after the first few minutes

Answer: A



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34. Which of the following steps performed by person to visualise the DNA bands obtained from gel electrophoresis?

A. (a) Exposure of DNA fragments to UV radiations

B. (b) Staining gel with bromophenol blue followed by exposure to UV radiations

C. (c) Staining gel with ethidium bromide followed by exposure to

D. (d) Person can see the bands without staining.

Answer: C



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UV radiations.

35. Study the given figure carefully and select the incorrect statements regarding this.



- (i) It represents a typical agarose gel electrophoresis in which lane 1 contains undigested DNA
- (ii) Smallest DNA bands are formed at A and largest DNA bands are formed at B
- (iii) The separated DNA fragments can be visualized after staining in the visible light
- (iv) The separated DNA bands are cut out from the agarose gel and extracted from the gel piece. this step is known as elution.
 - A. (i) and (ii)
 - B. (ii) and (iii)
 - C. (ii) and (iv)
 - D. Person can see the bands without staining.

Answer: B

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

A. EcoRI -Production of sticky ends

B. DNA ligase - Multiplication of DNA molecules

C. ori- copy number

D. Selectable market - Identification of transformants

Answer: B



37. If you want of recover many copies of the target DNA, you will choose

a vector

A. Which does not have origin of replication

- B. which has antibiotic resistance gene
- C. whose origin supports high copy number
- D. which has only one restriction site

Answer: C



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- 38. Which of the following statements are correct?
- (i) Restriction enzymes cut the strand of DNA a little away from the centre of the palindrome site, but between the same two bases on the opposite strands.
- (ii) Hind II always cuts DNA molecules at a particular point by reconginisng a specific sequence of six base pairs.
- (iii) Separated DNA fragments cannot be visualised without staining on an agarose gel electrophoresis.
- (iv) 'Ori' is the sequence responsible for controlling the copy number.
- (v) DNA is a positively charged molecule.

- A. (i), (iii) and (v)
- B. (i), (ii),(iii) and (iv)
- C. (iii),(iv) and (v)
- D. (i),(ii),(iii),(iv) and (v)

Answer: B



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- **39.** Which one of the following characteristics is generally not preferred
- for a cloning vector?
- a) An origin of replication
- c)Multiple restriction sites

b) An antibiotic resistance marker

- d) A high copy number
 - A. An origin of replication
 - B. An antibiotic resistance marker

- C. Multiple restriction sites
- D. A high copy number

Answer: C



- **40.** Read the following statements and select the correct ones.
- (i) Same kind of sticky ends are produced when a DNA has been cut by different restriction enzymes.
- (ii) Exonucleases make cuts at specific positions within the DNA.
- (iii) Hind II was the first restriction endonuclease to be isolated.
- (iv) A bacteriophage has the ability to replicate within bacterial cells by integrating its DNA with bacterial DNA.
- (v) Presence of more than one recognition sites for a enzyme within the vector facilitates the gene cloning.
 - A. (i),(iii) and (v)
 - B. (ii) and (iv)

C. (iii) and (iv) D. (ii),(iii) and (iv) **Answer: C Watch Video Solution 41.** Which of the following is not a cloning vector? 1) Cosmid 2) pBR 322 3) Sall 4) Phagemid A. 1) Cosmid B. 2) pBR 322 C. 3) Sall D. 4) Phagemid **Answer: C**

42. Match column I with column II and select the correct answer from the

given codes.

ColumnII ColumnIII

 $A. amp^R$ gene (i)Artificial plasmid

B. Separation of DNA fragments (ii) Selectable marker C. Hindlll (iii) Electrophoresis

D. pBR322 (iv) Haemophilus influenzea

A. 1)
$$A - (iii), B - (ii), C - (i), D - (iv)$$

B. 2)
$$A - (iv), B - (i), C - (iii), D - (ii)$$

C. 3)
$$A - (ii), B - (iii), C - (iv), D - (i)$$

D. 4)
$$A - (iii), B - (iv), C - (i), D - (iii)$$

Answer: C



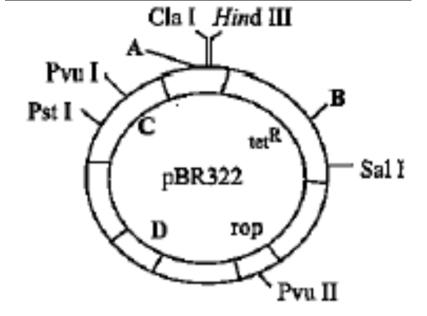
- 43. The gene 'rop' present in pBR322 cloning vector, codes for
- 1) the proteins involved in the translation
- 2) the proteins involved in the replication of the plasmid
- 3) the proteins involved in the synthesis of ampicillin only
- 4) the proteins involved in the synthesis of tetracycline only
 - A. 1) the proteins involved in the translation
 - B. 2) the proteins involved in the replication of the plasmid
 - C. 3) the proteins involved in the synthesis of ampicillin only
 - D. 4) the proteins involved in the synthesis of tetracycline only

Answer: B



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44. Identify A,B,C, and D in the given figure of E. coli cloning vector pBR322 and select the correct option.



- A. $\frac{A}{ ext{Hindl}} \ \frac{B}{EcoRl} \ \frac{C}{amp^R} \ ext{ ori}$
- C. $\frac{A}{BamHl}$ $\frac{B}{Pstl}$ $\frac{C}{ori}$ $\frac{D}{amp^R}$
- D. $\frac{A}{EcoRl}$ $\frac{B}{BamHl}$ $\frac{C}{amp^R}$ ori

Answer: D



45. Read the given statements and select the correct option.

Statement 1: The cloning vector is required to have very few, preferably

single, recongnition sites for the commonly used restriction enzymes.

Statement 2: Presence of more than one recongnition sites within a cloning vector will generate several fragments, which will complicate the process of gene cloining.

- 1) Both statements 1 and 2 are correct
- 2) statement 1 is correct but statement 2 is incorrect
- 3) statement 1 is incorrect but statement 2 is correct
- 4) None of the above
 - A. 1) Both statements 1 and 2 are correct
 - B. 2) statement 1 is correct but statement 2 is incorrect
 - C. 3) statement 1 is incorrect but statement 2 is correct
 - D.

Answer: A



46. pBR322 was the first artificial cloning vector to be constructed. What does "BR" stands for ?

- A. 1) Bacteriophage and Recombinant
- B. 2) Boliver and Rodriguez
- C. 3) Boyer and Replicative
- D. 4) None of these

Answer: B



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- 47. Read the following statements and select the correct ones.
- (i) Electrophoresis is a technique used for the separation of molecules based on their size and charge.
- (ii) Plasmids are extra-chromosomal, self-replicating, usually circular,

double stranded DNA molecules found naturally in many bacteria and

also in some yeast,.

(iii) It is not advisable to use an exonuclease enzyme while producing a recombinant DNA molecule.

(iv) In EcoRI, the roman numberal I indicates that it was the first enzyme isolated from Ecoli

A) (i) and (ii)

B) (iii) and (iv)

C) (i),(ii) and (iv)

D) (i),(ii),(iii) and (iv)

A. A) (i) and (ii)

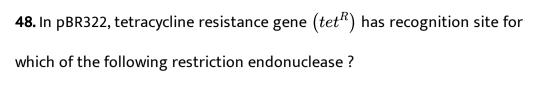
B. B) (iii) and (iv)

C. C) (i),(ii) and (iv)

D. D) (i),(ii),(iii) and (iv)

Answer: D





- 1) Hindlll
- 2) BamHl
- 3) EcoRl
- 4) Pstl
 - A. 1) Hindlll
 - B. 2) BamHl
 - C. 3) EcoRl
 - D. 4) Pstl

Answer: B



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49. Which of the following is not a characteristic of pBR322 vector?

A. 1) It was the first artificial cloning vector constructed in 1977 by Boli-

ver and Rodriguez.

B. 2) It is the most widely used, versatile and easily manipulated vector.

C. 3) It has two antibiotic resistance genes tet^R and amp^R

D. 4) It does not have restriction site for sall.

Answer: D



50. What will be the effect if pBR322, a cloning vector does not carry 'ori' site ?

A. 1) Sticky ends will not produce

B. 2)Transformation will not take place

C. 3) The cell will transform into a tumour cell

D. 4) Replication will not take place

Answer: D



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51. Using recombinant DNA technology, genes from a donor cell can be inserted into a bacterium for DNA replication and protein synthesis. The kind of cells that can be used as gene donors in this technology are

- A. 1) bacteria only
- B. 2) either yeasts or bacteria
- C. 3) eukaryotic cells only
- D. 4) any of these

Answer: D



52. An advantage of using yeasts rather than bacteria as recipient cells for the recombinant DNA of eukaryotes is that yeasts can

- A. 1) produce restriction enzymes
- B. 2) excise introns from the RNA transcript
- C. 3) remove methyl groups
- D. 4) reproduce more rapidly

Answer: B



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53. Read the given statements and select the correct option.

Statement 1: Both bacteria and yeast multiply very fast to form huge populations which express the desired gene.

Statement 2: In recombinant DNA technology, human genes are often transferred into bacteria (prokaryotes) or yeast (eukaryotes).

- A. 1) Both statements 1 and 2 are correct
- B. 2) statement 1 is correct but statement 2 is incorrect
- C. 3) statement 1 is incorrect but statement 2 is correct
- D. 4) Both statements 1 and 2 are incorrect

Answer: A



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54. In the process of insertional inactivation

- 1) a recombinant DNA is inserted within the coding sequence of enzyme $\boldsymbol{\beta}$
- -galactosidase, resulting in inactivation of the enzyme
- 2) a recombinant DNA is inserted within the coding sequence of proteins
- involved in the replication of the plasmid
- 3) a recombinat DNA is inserted within the recongnition site for EcoRl
- 4) none of these

A. 1) a recombinant DNA is inserted within the coding sequence of

enzyme eta-galactosidase, resulting in inactivation of the enzyme

B. 2) a recombinant DNA is inserted within the coding sequence of proteins involved in the replication of the plasmid

C. 3) a recombinat DNA is inserted within the recongnition site for EcoRl

D. 4) none of these

Answer: A



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55. If a person obtains transformants by inserting a recombinant DNA within the coding sequence of enzyme 'beta-galatosidase, he will separate out recombinants from non-recombinats by which of the following observations?

- A. 1) Non-recombinant colonies do not produce any colour whereas
 - reccombinants give blue coloured colonies
- B. 2) Recombinant colonies do not produce any colour whereas nonrecombinants given blue coloured colonies.
- C. 3)Recombinants and non-recombinants both produce blue coloured colonies
- D. 4) No colonies are formed due to insertional inactivation

Answer: B



- **56.** Read the given statements and select the correct option
- Statement 1: In insertional inactivation, blue colour produced by bacterial colonies indicates that the plasmid does not have an insert into the bacterial genome.
- Statement 2: Presence of insert results into insertional inactivation of β -

galactosidase enzyme and the colonies do not produce any colour.

- A) Both statements 1 and 2 are correct
- B) statement 1 is correct but statement 2 is incorrect
- C) statement 1 is incorrect but statement 2 is correct
- D) Both statements 1 and 2 are incorrect
 - A. A) Both statements 1 and 2 are correct
 - B. B) statement 1 is correct but statement 2 is incorrect
 - C. C) statement 1 is incorrect but statement 2 is correct
 - D. D) Both statements 1 and 2 are incorrect

Answer: A



- 57. During insertional inactivation, the presence of a chromogenic substrate gives blue coloured colonies if the plasmid in the bacteria does not have an insert. The blue colour is produced by the enzyme
- 1) α -glucosidase

- 2) restriction endonuclease 3) β -galactosidase
- 4) Taq polymerase
- A. 1) α -glucosidase
 - B. 2) restriction endonuclease
 - C. 3) β -galactosidase
 - D. 4) Taq polymerase

Answer: C



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engineering?

58. Which of the following bacteria is used as a vector for plant genetic

- 1) Agrobacterium tumefaciens
- 2) Bacteriophages
- 3) Thermus aquaticus
- 4) Pyrococcus furiosus

- A. 1) Agrobacterium tumefaciens
- B. 2) Bacteriophages
- C. 3) Thermus aquaticus
- D. 4) Pyrococcus furiosus

Answer: A



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- 59. Which of the following microbes transform normal plant and animal cells to cancerous cells respecively?
 - A. 1) Retroviruses and Rhizobium
 - B. 2) Esherichia coli and Agrobacterium tumefaciens
 - C. 3) Agrobacterium tumefaciens and Retroviruses
 - D. 4) Agrobacterium tumefaciens and A.rhizogenes

Answer: C

60. Read the given statements and select the correct option.

Statement 1: The tumour inducing plasmid (Tiplasmid) acts as a cloning vector in recombinant DNA technology.

Statement 2: The Ti plasmid which is used in the mechanisms of delivering genes to a cell remains pathogenic.

A. 1. Both statements 1 and 2 are correct

B. 2. Statement 1 is correct but statement 2 is incorrect

C. 3. Statement 1 is incorrect but statement 2 is correct

D. 4. Both statements 1 and 2 are incorrect

Answer: B



61. a crown gall bacterium, is called an natural genetic engineer'
of plants.
1) Escherichia coli
2) Streptomyces albus

- 3) Agrobacterium tumefaciens
- A. 1) Escherichia coli

4) Azotobacter

- B. 2) Streptomyces albus
- C. 3) Agrobacterium tumefaciens
- D. 4) Azotobacter

Answer: C



- **62.** DNA cannot pass through a cell membrane as
- 1) it is too big to cross the memebrane

- 2) it is a hydrophilic molecule
- 3) membrane does not have specific proteins to facilitate the transport
- 4) none of these
 - A. 1) it is too big to cross the memebrane
 - B. 2) it is a hydrophilic molecule
 - C. 3) membrane does not have specific proteins to facilitate the

transport

D. 4) none of these

•

Answer: B



- **63.** The term "competent" refers to
 - A. 1) increasing the competition between cells
 - B. 2) making cells impermeable for DNA

C. 3) increasing the efficiency with which DNA enters the bacterium through pores in its cell wall

D. 4) making cells permeable for divalent cations

Answer: C



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64. The correct sequence of making a cell competent is

A. 1. treatment with divalent cation o incubation of cells with recombinant DNA of ice o heat shock $(42^{\circ}C)$ o placing on ice

B. 2. heat shock $(42^\circ C) o$ incubation of cells with recombinant DNA on ice o treatment with divalent cations o placing on ice

C. 3. treatment with divalent cation $\,\rightarrow\,$ placing on ice $\,\rightarrow\,$ incubation

of cells with recombinant DNA on ice $\;
ightarrow\;$ heat shock $(42\,{}^{\circ}\,C)$

D. 4. incubation of cells with recombinant DNA on ice $\,\,
ightarrow\,$ heat shock

 $(42\,{}^{\circ}C)
ightarrow {}$ treatment with divalent cations $ightarrow {}$ placing on ice.

Answer: A



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65. Match the terms given in column I with their definitions in column II and select the correct answer from codes given below.

(iv)Process by which bacteria take up pieces of DN

ColumnII ColumnIII

D. Recombinant DNA

A. Transformation (i) Sequences cut by restriction enzymes

B. Recognition site (ii) Process by which DNA fragments are separate C. Gel electrophoresis (iii) Plasmid DNA that has incorporated human I

A. 1)
$$A-(iii), B-(i), C-(ii), D-(iv)$$

B. 2)
$$A-(iv), B-(i), C-(ii), D-(iii)$$

C. 3)
$$A-(i), B-(ii), C-(iii), D-(iv)$$

D. 4)
$$A-(ii), B-(iii), C-(iv), D-(i)$$

Answer: B



66. Micro-injection is a method used to

A. 1) produce sticky ends of DNA

B. 2) provide protection against pathogen

C. 3) purify the DNA

D. 4) inject recombinant DNA into the nucleus of an animal cell.

Answer: D



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67. Which of the following is required for micro-injection method of gene transfer?

A. 1) micro-particles

B. 2) Micro-pipettes

- C. 3) Divalent cations
- D. 4) UV radiations

Answer: B



- **68.** In biolistic method of gene trasfer, the microparticles coated with foreign DNA are bombarded into target cells at a very high velocity. These microparticles are made up of
- 1) silver or tungsten
- 2) arsenic or silver
- 3) gold or tungsten
- 4) none of these
 - A. 1) silver or tungsten
 - B. 2) arsenic or silver
 - C. 3) gold or tungsten

D. 4) none of these

Answer: C



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- **69.** The different steps of recombinant DNA technology are given below randomly.
- (i) Isolation of the DNA fragments or genes to be cloned
- (ii) Introduction of the recombinant DNA into a suitable cell (usually E. coil) called host (transformation)
- (iii) Multiplication/expression of the introduced gene in the host
- (iv) Selection of the transformed host cells, and identification of the clone containing the desired gene/DNA fragment
- (v) Insertion of the isolated gene in a suitable plasmid vector

Which of the following represents the correct sequences of steps?

A. A)
$$(i)
ightarrow (iii)
ightarrow (iv)
ightarrow (v)$$

B. B)
$$(iii)
ightarrow (ii)
ightarrow (i)
ightarrow (v)
ightarrow (iv)$$

C. C)
$$(i)
ightarrow (v)
ightarrow (ii)
ightarrow (iv)
ightarrow (iii)$$

D. D)
$$(v)
ightarrow (i)
ightarrow (iii)
ightarrow (iv)
ightarrow (ii)$$

Answer: C



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70. The given flow chart depicts the steps to transfer a desirable gene of interest into a plant.



Identify the missing steps (A,B and C) with regard ot following statements and select the correct option.

- (i) Joining of desirable gene to a suitable cloning vector using ligases to create a recombinant DNA molecule.
- (ii) Selection of transformed cells.
- (iii) Transferring the recombinant DNA molecules to teh target cells.

A.
$$A B C$$
 (i) (ii) (iii) .

B. A B C (i) (iii) (ii).

$$egin{array}{lll} {\sf C.} & A & B & C \ (ii) & (iii) & (i). \ & & A & B & C \ (iii) & (i) & (ii). \end{array}$$

Answer: B

Watch Video Solution	or
----------------------	----

respectively , and .

(i)	EcoRl	cuts	the	DNA	between	bases	only	when	the
seq	uence	is p	resent	in the	DNA duple:	х.			

- (ii) Disruption of the cell membranes can be achieved by treating the bacterial cells, plant cells and fungal cells with enzymes
- (iii) Since DNA has a____charge, it moves towards the_____of the electrophoretic chamber.

71. Fill up the blanks and select the correct option.

- A) (i) G and A, GA A T TC (ii) endonuclease, cellulase, chitinase (ii) negative, anode
- B) (i) G and A,G A AT TC(i) lysozyme, cellulase, chitinase (iii) positive, cathode

C) (i) G and A,GA AT C (ii) lysozyme, cellulase, chitinase (ii)negative, anode
D) (i) G and A, GA ATC (ii) lysozyme, cellulase, chitinase (iii) positive,
cathode

A. A) (i) G and A, GA A T TC (ii) endonuclease, cellulase, chitinase (ii) negative, anode

B. B) (i) G and A,G A AT TC(i) lysozyme, cellulase, chitinase (iii) positive, cathode

C. C) (i) G and A,GA AT C (ii) lysozyme, cellulase, chitinase (ii)negative,

anode

D. D) (i) G and A, GA ATC (ii) lysozyme, cellulase, chitinase (iii) positive, cathode

Answer: C



72. In the isolation of DNA, removal of protein and RNA is carried out by							
enzymesandrespectively.							
1) lysozyme ribonuclease							

- 1) Ty302 yTTTC, TIDOTTUCICUSC
- 2) protease, cellulase
- 3) protease, ribonuclease4) ribonuclease, chitinase
 - A. 1) lysozyme, ribonuclease
 - B. 2) protease, cellulase
 - C. 3) protease, ribonuclease
 - D. 4) ribonuclease, chitinase

Answer: C



Watch Video Solution

73. During isolation of genetic material, the chemical used to precipitate out the purified DNA is

A. a) bromophenol blueB. b) chilled ethanolC. c) ethidium bromideD. d) both (a) and (c)

Answer: B



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74. 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. 1) DNA transformation
- B. 2) DNA ligation
- C. 3) DNA spooling
- D. 4) DNA duplication



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75. Match column I with column II and select the correct answer from the given codes.

$$ColumnII$$
 $ColumnIII$

- A. Recombinant DNA (i) Chilled ethanol
- B. Precipitation of DNA (ii)DNAstaining
- C. Transposons (iii) Jumping genes
- D. Ethidium bromide (iv)Genetic engineering

A. 1)
$$A - (iv), B - (i), C - (iii), D - (ii)$$

B. 2)
$$A - (i), B - (iii), C - (ii), D - (iv)$$

C. 3)
$$A - (ii), B - (i), C - (iii), D - (iv)$$

D. 4)
$$A - (iv), B - (ii), C - (i), D - (iii)$$

Answer: A



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- 76. The polymerase chain reaction is a technique used for
- 1) amplification of DNA
- 2) amplification of enzymes
- 3) amplification of proteins
- 4) all of these
 - A. 1) amplification of DNA
 - B. 2) amplification of enzymes
 - C. 3) amplification of proteins
 - D. 4) all of these

Answer: A



77. Process	used	for	amplification	or	multiplication	of	DNA	in	DNA
fingerprintir	ng is								

- 1) polymerase chain reaction
- 2) southern blotting
- 3) northern blotting
- 4) None of these
 - A. 1) polymerase chain reaction
 - B. 2) southern blotting
 - C. 3) northern blotting
 - D. 4) None of these

Answer: A



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78. Primers are

1) chemically synthesised oligonucleotides that are complementary to the

regions of DNA

2) chemically synthesised oligonucleotides that are not complementary

to the regions of DNA

3) chemically synthesised, autonomously replicating circular DNA

molecules

4) specific sequences present on recombinant DNA

A. 1) chemically synthesised oligonucleotides that are complementary

to the regions of DNA

B. 2) chemically synthesised oligonucleotides that are not

complementary to the regions of DNA

C. 3) chemically synthesised, autonomously replicating circular DNA

molecules

D. 4) specific sequences present on recombinant DNA

Answer: A



79. Enzyme 'Taq polymerase' used in PCR, has been isolated from bacterium

- 1) Agrobacterium tumefaciens
- 2)Thermus aquaticus
- 3) Streptomyces albus
- 4) Escherichia coil
 - A. 1) Agrobacterium tumefaciens
 - B. 2)Thermus aquaticus
 - C. 3) Streptomyces albus
 - D. 4) Escherichia coil

Answer: B



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80. Which of the following statements are correct for the enzyme Taq polymerase ?

- (i) It remains active during the high temperature induced denaturation of dsDNA.
 - (ii) It requires primers for carrying out the process of polymerisation.
- (iii) It synthesises the RNA region between the primers, using dNTPs and Mq^{2+} .
 - A. a) (i) and (ii)
 - B.b) (ii) and (iii)
 - C. c) (i),(ii) and (iii)
 - D. d) None of these

Answer: A



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81. Match column I (enzyme) with column II (characteristic/activity) and select the correct answer from the given codes,

ColumnIColumnIIA. Taq DNA polymerase (i) Cleaves the ends of linear DNA B. Exonuclease (ii) Breakdown of fungal cell wall (iii)Stable above $90^{\circ}C$ C. Protease D. Chitinase (iv)Made only by eukaryotic cells (v)Degradation of proteins 1) A - (iii), B - (iv), C - (i), D - (ii) 2) A - (iv) .B - (iii) .C - (i) .D - (ii) 3) A - (ii), B - (i), C - (v), D - (iii)4) A - (iii), B - (i), C - (v), D - (ii) A. 1) A - (iii), B - (iv), C - (i), D - (ii)B. 2) A - (iv), B - (iii), C - (i), D - (ii)

C. 3) A - (ii), B - (i), C - (v), D - (iii)

D. 4) A - (iii), B - (i), C - (v), D - (ii)

Answer: D



- A. a) Tumour inducing Ti plasmid
- B. b) DNA probe Identifies the desired DNA fragment
- C. c) PCR DNA staining
- D. d) Agarose Sea weeds

Answer: C



- **83.** The correct sequence of different steps of polymerease chain reaction is
- 1) annealing \rightarrow denaturation \rightarrow extension
- 2) denaturation \rightarrow extension \rightarrow annealing
- 3) denaturation \rightarrow annealing \rightarrow extension
- 4) extension \rightarrow denaturation \rightarrow annealing
 - A. 1) annealing $\,
 ightarrow \,$ denaturation $\,
 ightarrow \,$ extension
 - B. 2) denaturation \rightarrow extension \rightarrow annealing

C. 3) denaturation \rightarrow annealing \rightarrow extension

D. 4) extension \rightarrow denaturation \rightarrow annealing

Answer: C



Watch Video Solution

2. Apparatus Requirement

84. Given table gives an account of differences between PCR and gene

DNA

Restriction enzyme, ligase, vec

cloning. Which of the following points shows the incorrect difference?

Parameter PCR Gene cloning

1. Efficient More Less

3. Manipulation in vitro in vitro and in vivo

4. cost More Less 5. Automation Yes No

6. Error probability Less More
7. Time for a typical experiment 2-4days 4hours

8. Application More Less

a) 1 and 3

b) 4,5 and 6

c) 4 and 7

d) 4,7 and 8

A. a) 1 and 3 B. b) 4,5 and 6 C. c) 4 and 7 D. d) 4,7 and 8 **Answer: C Watch Video Solution** 85. Which of the following is required to perform polymerase chain reaction? A. a) Primers, dNTPs and DNA polymerase B. b) $DNA, CaCI_2$ and nuclease C. c) Mg^{2+} , DNAD. d) Both (a) and (c) Answer: D

86. In a polymerase chain reaction, temperature required for the steps

- (i) Denaturation,
- (ii) Annealing and
- (iii) Extension are respectively
- A) (i) 94 ° C (ii) 40 ° C (iii) 72 ° C
- B)(i)40 ° C(ii)72 ° C(iii)94 ° C
- C)(i)94 ° C(ii)72 ° C(iii)40 ° C
- D)(i)72 ° C(ii)94 ° C(iii)40 ° C
 - A. A) $(i)94^{\circ}C(ii)40^{\circ}C(iii)72^{\circ}C$
 - B. B) $(i)40^{\circ}\,C(ii)72^{\circ}\,C(iii)94^{\circ}\,C$
 - C. C) $(i)94^{\circ}\,C(ii)72^{\circ}\,C(iii)40^{\circ}\,C$
 - D. D) $(i)72^{\circ}\,C(ii)94^{\circ}\,C(iii)40^{\circ}\,C$

Answer: A



87. In addition to Taq polymerase enzyme which other thermostable DNA polymerases have been isolated to be used in polymerase chain Reaction (PCR)?

- A. a) Pfu polymerase isolated from Pyrococcus furiosus
- B. b) Tli polymerase(vent ploymerase) isolated fromThermococcus litoralis
- C. c) Both (a) and (b)
- D. d) None of these

Answer: C



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88. Given figures represents the steps involved in polymerase chain reaction (PCR). Identify the steps A,B,C and C and select the correct

option.		
Α.		
A	B	
Denaturation at $94-96^{\circ}C$	Extension through Taq polymerase at	
В.		
A	B C	
Denaturation at $94-96^{\circ}C$	Annealing at $40-60^{\circ}C$ Extension	
C.		
A	B C	
Denaturation at $40-60^{\circ}C$	Annealing at $72^{\circ}C$ Extension through	
D.		
A	B	
Extension through Taq polyn	nerase at $72^{\circ}C$ Denaturation at 40°	
Answer: B		
View Text Solution		
89. In a polymerase chain reaction	after the denaturation step why the	
mixture needs to cool down to a lower temperature ?		
	•	

- a) To permit specific annealing of the primers
- b) To give a halt to the reaction mixture
- c) To increase the activity of enzyme Taq polymerase
- d) To obtain the multiple copies of the DNA
 - A. a) To permit specific annealing of the primers
 - B. b) To give a halt to the reaction mixture
 - C. c) To increase the activity of enzyme Taq polymerase
 - D. d) To obtain the multiple copies of the DNA

Answer: A



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90. If a recombinant DNA bearing gene for resistance to antibiotic ampicillin is transferred to E.coli cells, the host cells become transformed into ampicillin resistant cells. If such bacteria are transferred on agar plates containing ampicillin, only transformants will grown and the untransformed recipient cells will die. The ampicillin resistant gene in this

case is called as 1) selectable marker 2) recombinant protein 3) cloning site 4) chemical scalpels A. 1) selectable marker B. 2) recombinant protein C. 3) cloning site D. 4) chemical scalpels Answer: A **Watch Video Solution 91.** Which of the following is not used to transfer the recombinant DNAinto the host? A. 1) Micro-injection method

- B. 2) Gene gun method
- C. 3) Bioreactor
- D. 4) Disarmed pathogen vectors

Answer: C



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- 92. A device in which large volume of living cells are cultured in order to
- get a specific product is called
- 2) agitator

1)PCR

- 3) bioreactor
- 4) assimilator
 - A. 1)PCR
 - B. 2) agitator
 - C. 3) bioreactor

D. 4) assimilator

Answer: C



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- **93.** Which of the following statements are correct with respect to a biorector?
- (i) It can process large volumes of culture.
- (ii) It provides optimum temperature and pH.
- (iii) It is a completely automated tool.
- (iv) It is a compact thermal cycler.
 - A. a) (i) and (ii)
 - B. b) (i),(ii) and (iii)
 - C. c) (iii) and (iv)
 - D. d) (ii) and (iii)

Answer: A

94. Identify the figures (A) and (B) and select the correct option.



- A. $\frac{(A)}{\text{Spargal stirred-tank bioreactor}}$ (B). Spargal stirred-tank bioreactor
- B.
- (A)(B).

Spargal stirred-tank bioreactor Sparged stirred-tank bioreactor (A)(B).

- C. Simple stirred-tank Sparged stirred-tank bioreactor
- (B).(A)Simple stirred-tank bioreactor Simple stirred-tank bioreactor

Answer: C



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95. Stirred-tank bioreactors have advantages over shake flasks because they

- A. a) provide high temperature and pH
- B. b) provide better aeration and mixing properties
- C. c) do not allow the entry of CO_2
- D. d) are easy to operate

Answer: B



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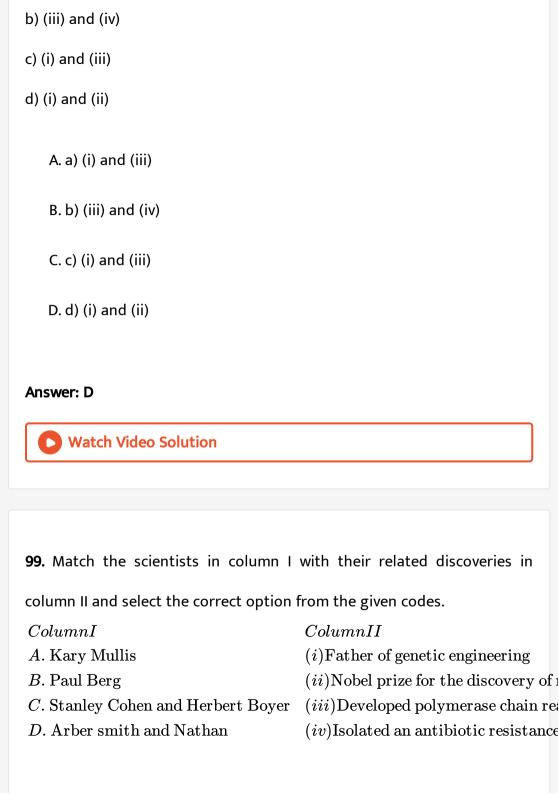
- **96.** After completion of the biosynthetic stage in the bioreactors, the product undergoes separation and purification processes, collectively termed as
 - A. a) transformation
 - B. b) electrophoresis
 - C. c) downstream processing
 - D. d) upstream processing

Answer: C



- **97.** Study the following statements regarding recombinant DNA technology and select the incorrect ones.
- (i) Taq polymerase extends the primers using the nucleotides provided in the reaction.
- (ii) Antibiotic resistance genes are considered as desirable genes in recombinant DNA technology.
- (iii) DNA fragments are separated according to their charge only, in agarose gel electrophoresis.
- (iv) Transformation is a procedure through which a piece of DNA is integrated in to the genome of a host bacterium.
- (v) To produce higher yields of a desired protein, host cells can be multiplied in a continuous culture.
- (vi) Downstream processing is one of the steps of polymerase chain reaction.

A. a) (ii),(iii) and (vi) B. b) (i),(iii) and (v) C. c) (ii),(iii) and (v) D. d) (i),(iv) and (v) Answer: A **Watch Video Solution** 98. Read the following statements and select the incorrect ones. (i) When the transformed cells on agar plates containing ampicillin are spread, both transformed and untransformed cells will grow. (ii) Restriction enzymes are used in isolation and separation of DNA from other macromolecules. (iii) Downstream processing is one of the steps of rDNA technology. (iv) Disarmed pathogen vectors are also used in transfer of rDNA into the host. a) (i) and (iii)



A. 1)
$$A-(iii), B-(i), C-(iv), D-(ii)$$

B. 2) $A-(iii), B-(iv), C-(i), D-(ii)$

C. 3) A - (iv), B - (ii), C - (iii), D - (i)

D. 4) A - (i), B - (iii), c - (iv), D - (ii)

Answer: A



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- (i) _____is a natural polymer extracted from____.
- (ii) The DNA fragments purified by gel electrophoresis are used in constructing_____by joining them with_____.
- (iii) The ligation of alien DNA is carried out at a_____. present in one of

the two_____in a plasmid vector.

(iv)_____enzyme remains active during the high temperature induced denaturation of ds DNA

(v) DNA fragments are resolved according to their_____ through

in agarose gel electrophoresis.

a) (i) Agarose, sea weeds (ii) recombinant DNA, cloning vector (iii) restriction site, antibiotic resistance genes (iv) Taq polymerase (v) size, sieving effect

b) (i) Agarose, sea weeds (ii) Restriction site, antibiotic resistance genes (iii) recombinant DNA, cloning vector (iv) Taq polymerase (v) size, sieving effect

c) (i) Agarose, sea weeds (ii) restriction site, antibiotic resistance genes (iii) recombinant DNA, cloning vector (iv) Taq polymerease (v) size, sieving effect

d) (i) size, sieving effect (ii) agarose, seaweeds (iii) recombinant DNA cloning vector (iv) Taq polymerase (v) restriction site, antibiotic resistance genes

A. a) (i) Agarose, sea weeds (ii) recombinant DNA, cloning vector (iii) restriction site, antibiotic resistance genes (iv) Taq polymerase (v) size, sieving effect

B. b) (i) Agarose, sea weeds (ii) Restriction site, antibiotic resistance genes (iii) recombinant DNA, cloning vector (iv) Taq polymerase (v)

size, sieving effect

C. c) (i) Agarose, sea weeds (ii) restriction site, antibiotic resistance genes (iii) recombinant DNA, cloning vector (iv) Taq polymerease (v) size, sieving effect

D. d) (i) size, sieving effect (ii) agarose, eas weeds (iii) recombinant

DNA cloning vector (iv) Taq polymerase (v) restriction site, antibiotic

resistance genes

Answer: A



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Hots

1. Four mutant strians of bacteria (1-4) all require substance S to grow (each strian is blocked at one step in the S-biosynthesis pathway). Four plates were prepared with minimal medium and a trace of substance, S

to allow a small amount of growth of mutant cells. on plate A, mutant cells of strain 1 were spred over entire surface of tha agar to form a thin law of bacteria. On plate B, the lawn was composed of mutant cells of strain 2, and so on. On each plate, cells of each of the four mutant types were inoculated over the lawn, as indicated in the figure by the circles. dark circles indicae excellent growth. A strain blocked at a later step in the S substance metabolic pathwa accumulates intermediates that can 'feed' a strain blocked at an earlier step.



What is the order of genes (1-4) in the metabolic pathway for synthesis of substance S?

A.
$$2 o 4 o 3 o 1$$

B.
$$2 o 1 o 3 o 4$$

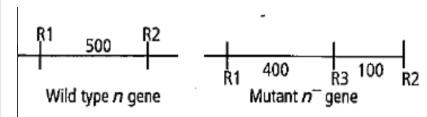
$$\mathsf{C.}\, 1 \to 3 \to 4 \to 2$$

D.
$$1 o 2 o 4 o 3$$

Answer: C

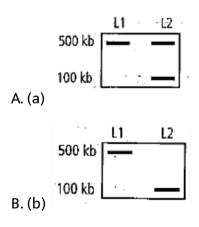


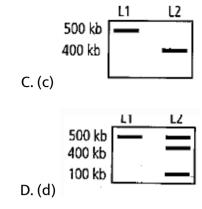
2. The figure shows the restriction enzyme cutting sites (R1-R3) in wild type (n) and mutant (n) gene.



If a radioactively labelled probe (that hybridises at a sequence close to R1) is used for detecting the presence of DNA fragments after gel electrophoresis and Southern blotting, which of the following band patterns will yout expect ?

Note: L1: wild type DNA, L2: mutant DNA





Answer: C



3. Analyse the given diagram which steps involved in the procedure of selecting transformed bacteria.



Identify the bacterial colony which has undergone transformation?

- A. colony 5
 - B. Colony 2
 - C. Colony 4

D. Colony 3

Answer: C



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4. The nucleic acid extracted from animal liver is loaded and run on agarose gel. After staining, it shows following pattern:



If the remaining sample is treated with RNAse and loaded in gel what result would you expect?

A. 📄

B. 📄

C. 📄

D. 📄

Answer: A



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5. The basic procedure involved in the synthesis of recombinanat DNA molecule is depicted below. The mistake in the procedure is



- A. Enzyme polmerase is not included.
- B. The mammalian DNA is shown double stranded
- C. Two different restriction enzymes are used.
- D. Only one fragment is inserted

Answer: C



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Ncert

1. Rising of dough is due is

A. a) multiplication of yeast B. b) production of CO_2 C. c) emulsification D. d) hydrolysis of wheat flour strach into sugars. **Answer: B Watch Video Solution** 2. An enzyme catalysing the removal of nucleotides from the ends of DNA is A. a) endonuclease B. b) exonuclease C. c) DNA ligase D. d) Hind II

Answer: B

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

A. a) transduction

B. b) conjugation

C. c) transformation

D. d) translation

Answer: A



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

A. a) DNA can be seen in visible light

B. b) DNA can be seen without staining in visible light.

C. c) Ethidium bromide stained DNA can be seen in visible light.

D. d) Ethidium bromide stained DNA can be seen under exposure to **UV** light

Answer: D



5. Restriction' in restriction enzyme refers to

A. a) cleaving of phosphodiester bond in DNA by the enzyme

B. b) cutting of DNA at specific position only

C. c) prevention of the multiplication of bacteriophage in bacteria

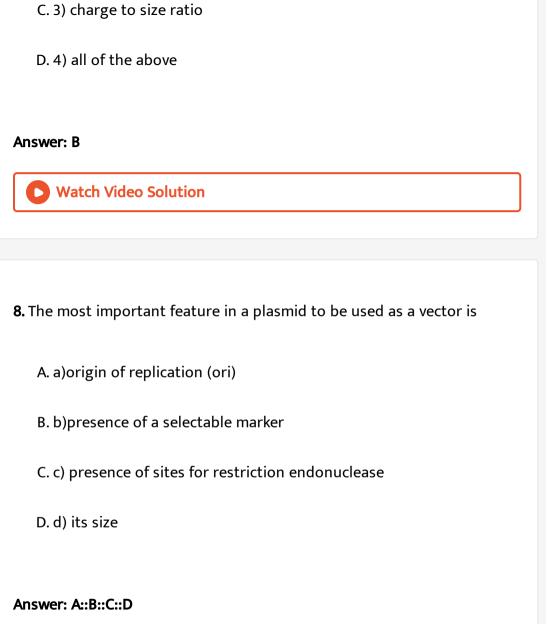
D. d) all of the above

Answer: C



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6. 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
Watch Video Solution
Watch Video Solution
7. In agarose gel electrophoresis, DNA molecules are separated on the basis of their
7. In agarose gel electrophoresis, DNA molecules are separated on the



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9. While isolating DNA from bacteria, which of the following enzymes is not used?

A. a) Lysozyme

B. b) Ribonuclease

C. c) Deoxyribonuclease

D. d) Protease

Answer: C



10. Which of the following has popularised the PCR (polymerase chain reactions)?

A. a) Easy availability of DNA template

B. b) Availability of synthetic primers

C. c) Availability of cheap deoxyribonucleotides

D. d) Availability of 'thermostable' DNA polymerase

Answer: D



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- 11. An antibiotic resistant gene in a vector usually helps in the selection of
 - A. a) competent cells
 - B. b) transformed cells
 - C. c) recombinant cells
 - D. d)none of the above

Answer: B



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

A. a) binding of DNA to the cell wall

B. b) uptake of DNA through membrane transport proteins

C. c) uptake of DNA through transient pores in the bacterial cell wall

D. d) expression of antibiotic resistance gene

Answer: C



13. The role of DNA ligase in the construction of a recmobinant DNA molecule is

A. a) formation of phosphodiester bond between two DNA fragments

B. b)formation of hydrogen bonds between sticky ends of DNA

fragments

- C. c) ligation of all purine and pyrimidine bases
- D. d)none of the above

Answer: A



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- **14.** Which of the following is not a source of restriction endonuclease?
- (a) Haemophilus influenzea
- (b) Escherichia coil
- (c) Entamoeba coil
- (d) Bacillus amyloliquifaciens
- A. Haemophilus influenzea
 - B. Escherichia coil
 - C. Entamoeba coil
 - D. Bacillus amyloliquifaciens

Answer: C

15. 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 and on the template DNA
- (d) All of the above
 - A. Denaturation of template DNA
 - B. Annealing of primers to template DNA
 - C. Extension of primer and on the template DNA
 - D. All of the above

Answer: C



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16. 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

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 degrated by bacteria

D. all of the above

Answer: A



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17. 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 continuous culture system
- (d) Any of the above
 - A. Laboratory flask of largest capcity
 - B. A stirred-tank bioreactro without in-lets and out-lets
 - C. A continuous culture system
 - D. Any of the above

Answer: C



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

(a) Herbert Boyer (b) Hargovind Khurane (c) Kary Mullis (d) Arthur Kornberg A. Herbert Boyer B. Hargovind Khurane C. Kary Mullis D. Athur Kornberg **Answer: C Watch Video Solution** 19. Which of the following statements does not hold true for restriction enzyme? (a) It recongnises a palindromic nucleotide sequence (b) It is an endonuclease.

- (c) It is isolated from viruses
- (d) It produces the same kind of sticky ends in different DNA molecules
 - A. It recongnises a palindromic nucleotide sequence
 - B. It is an endonuclease.
 - C. It is isolated from viruses
 - D. It produces the same kind of sticky ends in different DNA molecules

Answer: C



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Assertion Reason

- **1.** Assertion: In a chemical engineering process, it is necessary to prepare sterile ambience.
- Reason: Sterile ambience inhibits the growth of undesirable microbes during manufacture of product like antibiotics, vaccines and enzymes.

A. a) Both assertion and reason are true and reason is correct explanation of assertion.

B. b) Both assertion and reason are true but reason is not correct explanation for assertion.

C. c) Assertion is true but reason is false.

D. d) Assertion is false but reason is true.

Answer: A



2. Assertion: Asexual reproduction is more important with regard to biotehnology.

Reason: Asexual reproduction preserves the genetic information while sexual reporduction permits variations.

A. a) Both assertion and reason are true and reason is correct explanation for assertion.

B. b) Both assertion and reason are true but reason is not the correct explanation for assertion.

C. c) Assertion is true but reason is false.

D. d) Assertion is false but reason is true.

Answer: A



3. Assertion: Genetic engineering can overcome the drawbacks of traditional hybridisation.

Reason: Genetic engineering can create desired DNA sequences to meet specific requirements.



4. Assertion: A piece of DNA inserted into an alien organism generally does not replicate if not inserted into a chromosome.

Reason: Chromosomes have specific sequences called ori region where DNA replication is initiated.



5. Assertion : Genetic engineering requires both nuclease and ligases.

 $\label{lem:Reason: Ligases produce the nick in the recombinant DNA molecule.}$



6. Assertion: Restriction enzymes Hin and Hpa are produced from two different genera of bacteria.

Reason : Hin is produced from Haemophilus while Hpa is produced from

Hematococcus.



7. Assertion: Restriction enzymes recongise palindromic sequences.Reason: Palindromic sequences read same in both directions of the two strand.A. d) Both assertion and reason are false.B.

D.

C.

Answer: B



8. Assertion: The matrix used in gel electrophoresis should have controllable pore size.

Reason: Agarose concentration can be changed to change pore sizes.

A. a) Both assertion and reason are true and reason is correct explanation for assertion

B. b) Both assertion and reason are true but reason is not correct explanation of assertion

C. c) Assertion is true but reason is false.

D. d) Both assertion and reason are false.

Answer: B



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9. Assertion : All expression vectors are cloning vectors and vise versa.

Reason: Expression vectors have at least the regulatory sequences i.e., promotes, operators, ribosomal binding sites, etc having optimum function in the chosen control but not origin of replication.



10. Assertion: E.coli having pBR322 with DNA insert at BamHI site cannot grow in medium containing tetracyline.

Reason : Recognition site for BamHI is present in ter^R region of pBR22.



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11. Assertion: A bacterial cell with restriction enzymes will be easily infected and lysed by bacteriophages.

Reason: Restriction enzymes catalyse synthesis of protective coat around bacterial cell that prevents bacteriophage attack.



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12. Assertion: Special methods are used for transformation i.e., incorporation of recombinant DNA into host.

Reason: DNA is a hydrophilic molecule.



13. Assertion: Use of chitinase enzyme is necessary for isolation of DNA from yeast cells but not in case of Spirogyra.

Reason: Fungal cell wall is made up of fungal cellulose or chitin.



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14. Assertion: PCR primers must not have self complementary regions.

Reason: Self complementary regions result in hairpin structures adversely affecting the PCR.



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15. Assertion: Downstream processing is generally considered more difficult and costlier in plants than that in microbes.

Reason: Rhizosecretion is used as a method to facilitate easier recovery of recombinant proteins from plants.



Biotechnology Principles And Processes

1. Who is the father of genetic engineering?	

- A. Steward Linn
- B. Stanley Cohen
- C. Paul Berg
- D. Kary Mullis

Answer: C



- 2. Which of the following processes/techniques can be included under biotechnology?
- (i) In vitro fertilisation
- (ii) Synthesis of a gene

(iii) Correcting a defective gene (iv) Developing a DNA vaccine A. (i) and (ii) B. (ii) and (iii) C. (iii) and (iv) D. (i),(ii),(iii) and (iv) Answer: D **Watch Video Solution** 3. Plasmid used to construct the first recombinant DNA was isolated from which bacterium species? A. Escherichia coli B. Salmonella typhimurium C. Agrobacterium tumefacines

D. Thermus aquaicus

Answer: B



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- 4. Genetic engineering is possible because
 - A. We can cut DNA at specific sites by restriction endonucleases
 - B. restriction endonucleases purified sites by restriction used in

bacteria

- C. the phenomenon of transduction in bacteria is well understood
- D. we can see DNA by electron microscope

Answer: A



A. recombinant DNA
B. restriction enzymes
C. Taq polymerase
D. polindromic nucleotide sequences.
Answer: B
View Text Solution
6. The term 'chemical knife' refers to
6. The term 'chemical knife' refers to A. polymerases
A. polymerases
A. polymerases B. endonucleases

5. The term 'molecular scissors' refers to

Answer: B



- 7. In recombinant DNA technology, the term vector refers to
 - A. the enzyme that cuts DNA into restriction fragments
 - B. the sticky end of a DNA fragment
 - C. a plasmid used to transfer DNA into a living cell
 - D. a DNA fragment which carries only ori gene

Answer: C



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8. One of the key factors, which makes the plasmid the vector in genetic engineering is

- A. its resistance to antibiotics
- B. its resistance to restriction enzymes
- C. its ability to carry a foreign gene
- D. its ability to cause infection in the host

Answer: C



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- 9. The term 'recombinant DNA' refers to
 - A. DNA of the host cell
 - B. DNA with a piece of foreign DNA
 - C. DNA with selectable marker
 - D. DNA with more than one recognition sites

Answer: B



10. The term 'chimeric DNA' refers to

A. DNA with overhanging stretches

B. DNA with palindromic sequences

C. a recombiant DNA

D. molecular scissors

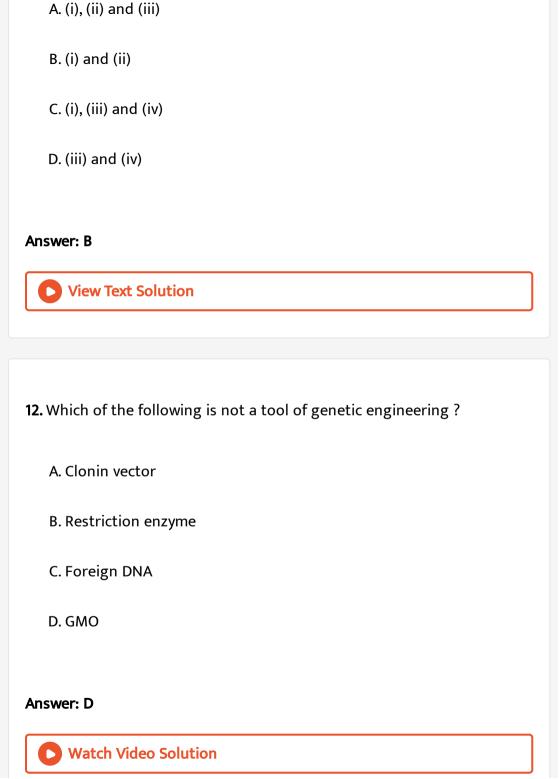
Answer: C



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11. Which of the following contains the key tools for recombinant DNA technology?

- (i) Restriction endonucleases, ligases, vectors
- (ii) Ligases, host organism, ligases, vectors
- (iii) Vectors, Taq polymerase, primers
- (iv) Restriction exonucleases, ligases, primers, bioreactors



13. The first restriction endonuclease isolated was
A. $EcoRl$
B. Bam-H
C. Sall
D. Hindll
Answer: D
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14. The letter 'R' in EcoRl is derived from
A. the name of genus
B. the name of strain
C. the name of species

D. the term 'restriction'

Answer: B



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15. Match column I with column II with respect to the nomenclature of restriction enzyme EcoRl and select the correct answer from the given codes.

ColumnII ColumnII

- A. E (i) 1^{st} in order of identification
- B. co (ii) Name of genus
- C. R (iii) Name of species
- D. l (iv) Name of strain
- (a) A (iii), B (i), C (ii), D (iv)
- (b) A-(ii), B-(i), C-(iii), D-(iv)
- (c) A (i), B (ii), C (iii), D (iv)
- (d) A (ii), B (iii), C (iv), D (i)
 - A. A (iii), B (i), C (ii), D (iv)
 - B. A (ii), B (i), C (iii), D (iv)

C. A - (i), B - (ii), C - (iii), D - (iv)

D. A - (ii), B - (iii), C - (iv), D - (i)

Answer: D



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16. The source of the restriction enzyme Hindll is

A. Escherichia coli RY13

B. Haemophilus influenzae Rd

C. Bacillus amyloliuefaciens ${\cal H}$

D. Streptomyces albus

Answer: B



17. A restriction endonuclease breaks bonds between the

A. base pairs of a DNA molecule

B. base pairs of a DNA-RNA hybrid molecule

C. sugar and phosphate components of a nucleic acid molecule

D. exons and introns of a DNA molecule.

Answer: C



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18. Read the given statements and select the correct option,

Statement 1 : Restriction endonuclease enzymes recognise a specific palindromic nucleotide sequence in the DNA

Statement 2: Restriction endonuclease enzymes are called as molecular scissors or biological scissors.

A. (a) Both statements 1 and 2 are correct

B. (b) statement 1 is correct but statement 2 is incorrect

C. (c) statement 1 is incorrect but statement 2 is correct

D. (d) both statements 1 and 2 are incorrect

Answer: A



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19. Study the following figures and identify the enzymes involved in steps

I and II.



A. EcoRl and DNA Ligase

B. Hind II and DNA Ligase

C. EcoRi and HindII

D. Restriction endonuclease and exonuclease

Answer: A

20. Which of the following correctly depicts the recognition site for EcoRI

(a)
$$G-A-A\stackrel{\downarrow}{-}T-T-C$$
 $C-T-T-A-A-G$

(b)
$$egin{aligned} G-T-C & \stackrel{\downarrow}{-} G-A-C \ C-A-G & \stackrel{\downarrow}{-} C-T-G \end{aligned}$$

(c)
$$G \stackrel{\downarrow}{-} T - C - G - A - C \\ C - A - G - C - T - G$$

(d)
$$G \stackrel{\downarrow}{-} A - A - T - T - C \ C - T - T - A - A - G$$

A.
$$G-A-A\stackrel{\downarrow}{-}T-T-C$$
 \uparrow $C-T-T-A-A-G$

B.
$$G-T-C \stackrel{\downarrow}{-} G-A-C \\ C-A-G \stackrel{\uparrow}{-} C-T-G$$

c.
$$G \stackrel{\downarrow}{-} T - C - G - A - C$$

D.
$$G \stackrel{\downarrow}{-} A - A - T - T - C \ C - T - T - A - A - G$$

Answer: D



- 21. The sticky ends of a fragmented DNA molecule are made of
 - A. (a) calcium salts
 - B. (b) endonuclease enzyme
 - C. (c) unpaired bases
 - D. (d) methyl groups

Answer: C



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22. The restrcition enzyme responsible for cleavage of following sequence

is

$$5'-G-T-C \stackrel{\downarrow}{-} G-A-C-3' \ 3'-C-A-G-C-T-G-5'$$

- A. (a) EcoRl
- B. (b) Hindll

C. (c) BamHl

D. (d) EcoRII

Answer: B



- 23. Identify the palindromic sequence in the following
- A. $\frac{GAATTC}{CTTUUG}$ $\mathsf{B.} \; \frac{GGATCC}{CCTAGG}$
 - c. $\frac{CCTGG}{GGACC}$
 - D. $\frac{CDATA}{GCTAA}$

Answer: B



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24. Which of the following statements is not correct regarding EcoRl restriction endonuclease enzyme ?

A. It is isolaed from Escherichia coil RY13

B. Its recongition sequence is $5^{\,\prime}-GAATTC-3^{\,\prime}$

$$3'-CTTAAG-5'$$

C. It produces complementary blunt ends.

D. None of these

Answer: C



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25. Which of the following sequences is recongnised by restriction enzyme BamHI?

A.
$$5' - \overset{\downarrow}{GAATTC} - 3' \ 3' - CTTAAG - 5'$$

B.
$$\begin{array}{c}
5' - AAGCTA - 3' \\
3' - TTCGAT - 5'
\end{array}$$
C. $\begin{array}{c}
5' - GGATCC - 3' \\
3' - CCTAGG - 5'
\end{array}$
D. $\begin{array}{c}
5' - CCCAAT - 3' \\
3' - GGGTTA - 5'
\end{array}$

c.
$$\frac{5^{7}-GGATCC-3^{7}}{3^{7}-CCTAGG-5^{7}}$$

D.
$$rac{5'-CCCAAT-3'}{3'-GGGTTA-5'}$$

Answer: C



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26. If a plasmid vector is digested with EcoRl at a single site, then

A. one sticky end will be produced

B. two sticky ends will be produced

C. four sticky ends will be produced

D. six sticky ends will be produced

Answer: B



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27. How many fragments will be generated if you digest a linear DNA molecule with a restriction enzyme having four recognition sites on the

DNA?

- (a) 3
- (b) 6
- (c) 5
- (d) 4



28. How many fragments will be generated on the digestion of a closed
circular DNA molecule with a restriction enzyme having six recongnition
sites on the DNA ?

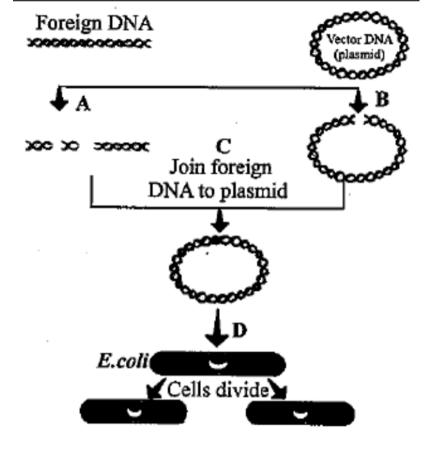
- A. 5
- B. 7
- C. 6
- D. 9

Answer: C



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29. The flow chart given below represents the process of recombinant DNA technology. Identify A,B, C and D.



- A. (a) A-Restriction endonuclease, B-Restriction exonuclease, C-DNA ligase, D-Transformation
- B. (b) A-Restriction endonuclease, B-Restriction endonuclease, C-DNA ligase, D-Transformation
- C. (c) A-Restriction endonuclease, B-Restriction endonuclease, C-Hydrolase, D-Transformation

D. (d) A-Restriction endonuclease, B-Restriction endonuclease, C-

Answer: B



Hydrolase, D-Transduction

30. In recombinant DNA technology, a plasmid vector is cleaved by

A. modified DNA ligase

B. a heated alkaline solution

C. the same enzyme that cleaves the donor DNA

D. the different enzyme than that cleaves the donor DNA.

Answer: C



31. Gel electrophoresis is a

A. technique of separation of charged molecules under the influence of magnetic field

B. technique of incorpotation of DNA molecules into the cell through transient pores made due to electrical impulses

C. technique of separation of DNA fragments through the pores of agarose gel under the influence of electric field

D. technique of separation and purification of gene products.

Answer: C



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32. Gel electrophoresis is used for

A. (a) construction of recombinant DNA using cloning vectors

- B. (b) isolation of DNA molecules
- C. (c) cutting of DNA into fragments
- D. (d) separation of DNA fragments according to their size.

Answer: D



- **33.** Having become an expert on gel electrophoresis, you are asked to examine a gel. Where would you find the smallest segments of DNA?
- (a) Near the positive electrode, farthest away from the wells
- (b) Near the negative electrode, close to the wells
- (c) Near the negative electrode, farther away from the wells
- (d) Near the middly, they tend to slow down after the first few minutes
 - A. Near the positive electrode, farthest away from the wells
 - B. Near the negative electrode, close to the wells
 - C. Near the negative electrode, farther away from the wells

D. Near the middly, they tend to slow down after the first few minutes

Answer: A



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34. Which of the following steps performed by person to visualise the DNA bands obtained from gel electrophoresis?

A. (a) Exposure of DNA fragments to UV radiations

B. (b) Staining gel with bromophenol blue followed by exposure to UV radiations

C. (c) Staining gel with ethidium bromide followed by exposure to

D. (d) Person can see the bands without staining.

Answer: C



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UV radiations.

35. Study the given figure carefully and select the incorrect statements regarding this.



- (i) It represents a typical agarose gel electrophoresis in which lane 1 contains undigested DNA
- (ii) Smallest DNA bands are formed at A and largest DNA bands are formed at B
- (iii) The separated DNA fragments can be visualized after staining in the visible light
- (iv) The separated DNA bands are cut out from the agarose gel and extracted from the gel piece. this step is known as elution.
 - A. (i) and (ii)
 - B. (ii) and (iii)
 - C. (ii) and (iv)
 - D. Person can see the bands without staining.

Answer: B

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

A. EcoRI -Production of sticky ends

B. DNA ligase - Multiplication of DNA molecules

C. ori- copy number

D. Selectable market - Identification of transformants

Answer: B

a vector



37. If you want of recover many copies of the target DNA, you will choose

A. Which does not have origin of replication

- B. which has antibiotic resistance gene
- C. whose origin supports high copy number
- D. which has only one restriction site

Answer: C



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- 38. Which of the following statements are correct?
- (i) Restriction enzymes cut the strand of DNA a little away from the centre of the palindrome site, but between the same two bases on the opposite strands.
- (ii) Hind II always cuts DNA molecules at a particular point by reconginisng a specific sequence of six base pairs.
- (iii) Separated DNA fragments cannot be visualised without staining on an agarose gel electrophoresis.
- (iv) 'Ori' is the sequence responsible for controlling the copy number.
- (v) DNA is a positively charged molecule.

- A. (i), (iii) and (v)
- B. (i), (ii),(iii) and (iv)
- C. (iii),(iv) and (v)
- D. (i),(ii),(iii),(iv) and (v)

Answer: B



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- **39.** Which one of the following characteristics is generally not preferred
- for a cloning vector?
- a) An origin of replication
- b) An antibiotic resistance marker
- c)Multiple restriction sites
- d) A high copy number
 - A. An origin of replication
 - B. An antibiotic resistance marker

- C. Multiple restriction sites
- D. A high copy number

Answer: C



- **40.** Read the following statements and select the correct ones.
- (i) Same kind of sticky ends are produced when a DNA has been cut by different restriction enzymes.
- (ii) Exonucleases make cuts at specific positions within the DNA.
- (iii) Hind II was the first restriction endonuclease to be isolated.
- (iv) A bacteriophage has the ability to replicate within bacterial cells by integrating its DNA with bacterial DNA.
- (v) Presence of more than one recognition sites for a enzyme within the vector facilitates the gene cloning.
 - A. (i),(iii) and (v)
 - B. (ii) and (iv)

C. (iii) and (iv) D. (ii),(iii) and (iv) **Answer: C Watch Video Solution 41.** Which of the following is not a cloning vector? 1) Cosmid 2) pBR 322 3) Sall 4) Phagemid A. 1) Cosmid B. 2) pBR 322 C. 3) Sall D. 4) Phagemid **Answer: C**

42. Match column I with column II and select the correct answer from the

given codes.

ColumnII ColumnIII

 $A. amp^R$ gene (i)Artificial plasmid

B. Separation of DNA fragments (ii) Selectable marker

C. Hindlll (iii)Electrophoresis

D. pBR322 (iv) Haemophilus influenzea

A. 1)
$$A - (iii), B - (ii), C - (i), D - (iv)$$

B. 2)
$$A-(iv), B-(i), C-(iii), D-(ii)$$

C. 3)
$$A - (ii), B - (iii), C - (iv), D - (i)$$

D. 4)
$$A - (iii), B - (iv), C - (i), D - (iii)$$

Answer: C



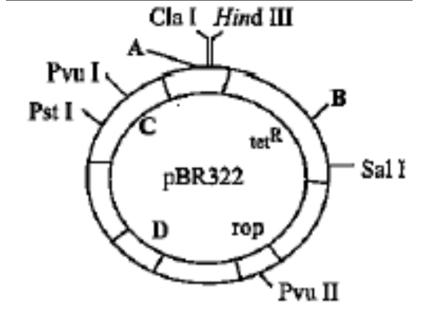
- 43. The gene 'rop' present in pBR322 cloning vector, codes for
- 1) the proteins involved in the translation
- 2) the proteins involved in the replication of the plasmid
- 3) the proteins involved in the synthesis of ampicillin only
- 4) the proteins involved in the synthesis of tetracycline only
 - A. 1) the proteins involved in the translation
 - B. 2) the proteins involved in the replication of the plasmid
 - C. 3) the proteins involved in the synthesis of ampicillin only
 - D. 4) the proteins involved in the synthesis of tetracycline only

Answer: B



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44. Identify A,B,C, and D in the given figure of E. coli cloning vector pBR322 and select the correct option.



- A. $\frac{A}{\text{Hindl}} \ \frac{B}{EcoRl} \ \frac{C}{amp^R} \ \text{ori}$
- C. $\frac{A}{BamHl}$ $\frac{B}{Pstl}$ $\frac{C}{ori}$ $\frac{D}{amp^R}$
- D. $\frac{A}{EcoRl}$ $\frac{B}{BamHl}$ $\frac{C}{amp^R}$ ori

Answer: D



45. Read the given statements and select the correct option.

Statement 1: The cloning vector is required to have very few, preferably

single, recongnition sites for the commonly used restriction enzymes.

Statement 2: Presence of more than one recongnition sites within a cloning vector will generate several fragments, which will complicate the process of gene cloining.

- 1) Both statements 1 and 2 are correct
- 2) statement 1 is correct but statement 2 is incorrect
- 3) statement 1 is incorrect but statement 2 is correct
- 4) None of the above
 - A. 1) Both statements 1 and 2 are correct
 - B. 2) statement 1 is correct but statement 2 is incorrect
 - C. 3) statement 1 is incorrect but statement 2 is correct
 - D.

Answer: A



46. pBR322 was the first artificial cloning vector to be constructed. What does "BR" stands for ?

A. 1) Bacteriophage and Recombinant

B. 2) Boliver and Rodriguez

C. 3) Boyer and Replicative

D. 4) None of these

Answer: B



- 47. Read the following statements and select the correct ones.
- (i) Electrophoresis is a technique used for the separation of molecules based on their size and charge.
- (ii) Plasmids are extra-chromosomal, self-replicating, usually circular, double stranded DNA molecules found naturally in many bacteria and

also in some yeast,.

(iii) It is not advisable to use an evenuslesse enzyme while producing a

(iii) It is not advisable to use an exonuclease enzyme while producing a recombinant DNA molecule.

(iv) In EcoRI, the roman numberal I indicates that it was the first enzyme

isolated from E.coli

A) (i) and (ii)

B) (iii) and (iv)

C) (i),(ii) and (iv)

D) (i),(ii),(iii) and (iv)

A. A) (i) and (ii)

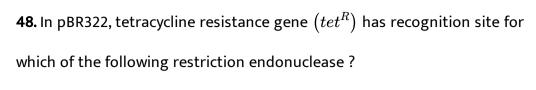
B. B) (iii) and (iv)

C. C) (i),(ii) and (iv)

D. D) (i),(ii),(iii) and (iv)

Answer: D





- 1) Hindlll
- 2) BamHl
- 3) EcoRl
- 4) Pstl
 - A. 1) Hindlll
 - B. 2) BamHl
 - C. 3) EcoRl
 - D. 4) Pstl

Answer: B



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49. Which of the following is not a characteristic of pBR322 vector?

A. 1) It was the first artificial cloning vector constructed in 1977 by Boli-

ver and Rodriguez.

B. 2) It is the most widely used, versatile and easily manipulated vector.

C. 3) It has two antibiotic resistance genes tet^R and amp^R

D. 4) It does not have restriction site for sall.

Answer: D



50. What will be the effect if pBR322, a cloning vector does not carry 'ori'

site?

A. 1) Sticky ends will not produce

B. 2)Transformation will not take place

C. 3) The cell will transform into a tumour cell

D. 4) Replication will not take place

Answer: D



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51. Using recombinant DNA technology, genes from a donor cell can be inserted into a bacterium for DNA replication and protein synthesis. The kind of cells that can be used as gene donors in this technology are

- A. 1) bacteria only
- B. 2) either yeasts or bacteria
- C. 3) eukaryotic cells only
- D. 4) any of these

Answer: D



52. An advantage of using yeasts rather than bacteria as recipient cells for the recombinant DNA of eukaryotes is that yeasts can

- A. 1) produce restriction enzymes
- B. 2) excise introns from the RNA transcript
- C. 3) remove methyl groups
- D. 4) reproduce more rapidly

Answer: B



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53. Read the given statements and select the correct option.

Statement 1: Both bacteria and yeast multiply very fast to form huge populations which express the desired gene.

Statement 2: In recombinant DNA technology, human genes are often transferred into bacteria (prokaryotes) or yeast (eukaryotes).

- A. 1) Both statements 1 and 2 are correct
- B. 2) statement 1 is correct but statement 2 is incorrect
- C. 3) statement 1 is incorrect but statement 2 is correct
- D. 4) Both statements 1 and 2 are incorrect

Answer: A



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54. In the process of insertional inactivation

- 1) a recombinant DNA is inserted within the coding sequence of enzyme $\boldsymbol{\beta}$
- -galactosidase, resulting in inactivation of the enzyme
- 2) a recombinant DNA is inserted within the coding sequence of proteins
- involved in the replication of the plasmid
- 3) a recombinat DNA is inserted within the recongnition site for EcoRl
- 4) none of these

A. 1) a recombinant DNA is inserted within the coding sequence of

enzyme eta-galactosidase, resulting in inactivation of the enzyme

B. 2) a recombinant DNA is inserted within the coding sequence of proteins involved in the replication of the plasmid

C. 3) a recombinat DNA is inserted within the recongnition site for

D. 4) none of these

EcoRl

Answer: A



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55. If a person obtains transformants by inserting a recombinant DNA within the coding sequence of enzyme 'beta-galatosidase, he will separate out recombinants from non-recombinats by which of the following observations?

- A. 1) Non-recombinant colonies do not produce any colour whereas
 - reccombinants give blue coloured colonies
- B. 2) Recombinant colonies do not produce any colour whereas nonrecombinants given blue coloured colonies.
- C. 3)Recombinants and non-recombinants both produce blue coloured colonies
- D. 4) No colonies are formed due to insertional inactivation

Answer: B



- **56.** Read the given statements and select the correct option
- Statement 1: In insertional inactivation, blue colour produced by bacterial colonies indicates that the plasmid does not have an insert into the bacterial genome.
- Statement 2: Presence of insert results into insertional inactivation of β -

galactosidase enzyme and the colonies do not produce any colour.

A) Both statements 1 and 2 are correct

B) statement 1 is correct but statement 2 is incorrect

C) statement 1 is incorrect but statement 2 is correct

D) Both statements 1 and 2 are incorrect

A. A) Both statements 1 and 2 are correct

B. B) statement 1 is correct but statement 2 is incorrect

C. C) statement 1 is incorrect but statement 2 is correct

D. D) Both statements 1 and 2 are incorrect

Answer: A



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57. During insertional inactivation, the presence of a chromogenic substrate gives blue coloured colonies if the plasmid in the bacteria does not have an insert. The blue colour is produced by the enzyme

1) α -glucosidase

- 2) restriction endonuclease 3) β -galactosidase
- 4) Taq polymerase
- A. 1) α -glucosidase
 - B. 2) restriction endonuclease
 - C. 3) β -galactosidase
 - D. 4) Taq polymerase

Answer: C



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engineering?

58. Which of the following bacteria is used as a vector for plant genetic

- 1) Agrobacterium tumefaciens
- 2) Bacteriophages
- 3) Thermus aquaticus
- 4) Pyrococcus furiosus

- A. 1) Agrobacterium tumefaciens
- B. 2) Bacteriophages
- C. 3) Thermus aquaticus
- D. 4) Pyrococcus furiosus

Answer: A



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- 59. Which of the following microbes transform normal plant and animal cells to cancerous cells respecively?
 - A. 1) Retroviruses and Rhizobium
 - B. 2) Esherichia coli and Agrobacterium tumefaciens
 - C. 3) Agrobacterium tumefaciens and Retroviruses
 - D. 4) Agrobacterium tumefaciens and A.rhizogenes

Answer: C

60. Read the given statements and select the correct option.

Statement 1: The tumour inducing plasmid (Tiplasmid) acts as a cloning vector in recombinant DNA technology.

Statement 2: The Ti plasmid which is used in the mechanisms of delivering genes to a cell remains pathogenic.

- A. 1. Both statements 1 and 2 are correct
- B. 2. Statement 1 is correct but statement 2 is incorrect
- C. 3. Statement 1 is incorrect but statement 2 is correct
- D. 4. Both statements 1 and 2 are incorrect

Answer: B



61a crown gall bacterium, is called an natural genetic engineer'						
of plants.						
1) Escherichia coli						
2) Streptomyces albus						

- 3) Agrobacterium tumefaciens

4) Azotobacter

- A. 1) Escherichia coli
- B. 2) Streptomyces albus
- C. 3) Agrobacterium tumefaciens
- D. 4) Azotobacter

Answer: C



- **62.** DNA cannot pass through a cell membrane as
- 1) it is too big to cross the memebrane

- 2) it is a hydrophilic molecule
- 3) membrane does not have specific proteins to facilitate the transport
- 4) none of these
 - A. 1) it is too big to cross the memebrane
 - B. 2) it is a hydrophilic molecule
 - C. 3) membrane does not have specific proteins to facilitate the

D. 4) none of these

transport

Answer: B



- 63. The term "competent" refers to
 - A. 1) increasing the competition between cells
 - B. 2) making cells impermeable for DNA

C. 3) increasing the efficiency with which DNA enters the bacterium through pores in its cell wall

D. 4) making cells permeable for divalent cations

Answer: C



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64. The correct sequence of making a cell competent is

A. 1. treatment with divalent cation o incubation of cells with recombinant DNA of ice o heat shock $(42^{\circ}C)$ o placing on ice

B. 2. heat shock $(42^\circ C) o$ incubation of cells with recombinant DNA on ice o treatment with divalent cations o placing on ice

C. 3. treatment with divalent cation $\,
ightarrow \,$ placing on ice $\,
ightarrow \,$ incubation

of cells with recombinant DNA on ice $\;
ightarrow\;$ heat shock $(42\,{}^{\circ}\,C)$

D. 4. incubation of cells with recombinant DNA on ice $\,\,
ightarrow\,$ heat shock

 $(42\,{}^{\circ}C)
ightarrow {}$ treatment with divalent cations $ightarrow {}$ placing on ice.

Answer: A



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65. Match the terms given in column I with their definitions in column II

(iii) Plasmid DNA that has incorporated human I

ColumnII ColumnIII

C. Gel electrophoresis

A. Transformation (i) Sequences cut by restriction enzymes

B. Recognition site (ii) Process by which DNA fragments are separate

D. Recombinant DNA (iv)Process by which bacteria take up pieces of DN

and select the correct answer from codes given below.

A. 1)
$$A-(iii), B-(i), C-(ii), D-(iv)$$

B. 2)
$$A-(iv), B-(i), C-(ii), D-(iii)$$

C. 3)
$$A-(i),B-(ii),C-(iii),D-(iv)$$

D. 4)
$$A-(ii), B-(iii), C-(iv), D-(i)$$

Answer: B



66. Micro-injection is a method used to

A. 1) produce sticky ends of DNA

B. 2) provide protection against pathogen

C. 3) purify the DNA

D. 4) inject recombinant DNA into the nucleus of an animal cell.

Answer: D



67. Which of the following is required for micro-injection method of gene transfer?

A. 1) micro-particles

B. 2) Micro-pipettes

- C. 3) Divalent cations
- D. 4) UV radiations

Answer: B



- **68.** In biolistic method of gene trasfer, the microparticles coated with foreign DNA are bombarded into target cells at a very high velocity. These microparticles are made up of
- 1) silver or tungsten
- 2) arsenic or silver
- 3) gold or tungsten
- 4) none of these
- A. 1) silver or tungsten
 - B. 2) arsenic or silver
 - C. 3) gold or tungsten

D. 4) none of these

Answer: C



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- **69.** The different steps of recombinant DNA technology are given below randomly.
- (i) Isolation of the DNA fragments or genes to be cloned
- (ii) Introduction of the recombinant DNA into a suitable cell (usually E. coil) called host (transformation)
- (iii) Multiplication/expression of the introduced gene in the host
- (iv) Selection of the transformed host cells, and identification of the clone containing the desired gene/DNA fragment
- (v) Insertion of the isolated gene in a suitable plasmid vector

Which of the following represents the correct sequences of steps?

A. A)
$$(i)
ightarrow (iii)
ightarrow (iv)
ightarrow (v)$$

B. B)
$$(iii)
ightarrow (ii)
ightarrow (i)
ightarrow (v)
ightarrow (iv)$$

C. C)
$$(i)
ightarrow (v)
ightarrow (ii)
ightarrow (iv)
ightarrow (iii)$$

D. D)
$$(v)
ightarrow (i)
ightarrow (iii)
ightarrow (iv)
ightarrow (ii)$$

Answer: C



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70. The given flow chart depicts the steps to transfer a desirable gene of interest into a plant.



Identify the missing steps (A,B and C) with regard ot following statements and select the correct option.

- (i) Joining of desirable gene to a suitable cloning vector using ligases to create a recombinant DNA molecule.
- (ii) Selection of transformed cells.
- (iii) Transferring the recombinant DNA molecules to teh target cells.

A.
$$egin{array}{ccccc} A & B & C \ (i) & (ii) & (iii). \end{array}$$

B. A B C (i) (iii) (ii).

$$egin{array}{lll} {\sf C.} & A & B & C \ (ii) & (iii) & (i). \ & & A & B & C \ (iii) & (i) & (ii). \end{array}$$

Answer: B

0	Watch	Video	Solution
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(i)	EcoRl	cuts	the	DNA	between	bases_	only	when	the
sequence is present in the DNA duplex.						х.			

71. Fill up the blanks and select the correct option.

(ii) Disruption of the cell membranes can be achieved by treating the bacterial cells, plant cells and fungal cells with enzymes respectively , and .

(iii) Since DNA has a charge, it moves towards the of the

- electrophoretic chamber.
- A) (i) G and A, GA A T TC (ii) endonuclease, cellulase, chitinase (ii) negative, anode
- B) (i) G and A,G A AT TC(i) lysozyme, cellulase, chitinase (iii) positive, cathode

C) (i) G and A,GA AT C (ii) lysozyme, cellulase, chitinase (ii)negative, anode
D) (i) G and A, GA ATC (ii) lysozyme, cellulase, chitinase (iii) positive,
cathode

A. A) (i) G and A, GA A T TC (ii) endonuclease, cellulase, chitinase (ii) negative, anode

B. B) (i) G and A,G A AT TC(i) lysozyme, cellulase, chitinase (iii) positive, cathode

C. C) (i) G and A,GA AT C (ii) lysozyme, cellulase, chitinase (ii)negative, anode

D. D) (i) G and A, GA ATC (ii) lysozyme, cellulase, chitinase (iii) positive, cathode

Answer: C



72. In the isolation of DNA, removal of protein and RNA is carried out by						
enzymesandrespectively.						
1) lysozyme, ribonuclease						

- 17 193029IIIC, TIDOTIUCICUSC
- 2) protease, cellulase
- 3) protease, ribonuclease4) ribonuclease, chitinase
 - A. 1) lysozyme, ribonuclease
 - B. 2) protease, cellulase
 - C. 3) protease, ribonuclease
 - D. 4) ribonuclease, chitinase

Answer: C



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73. During isolation of genetic material, the chemical used to precipitate out the purified DNA is

A. a) bromophenol blueB. b) chilled ethanolC. c) ethidium bromide

Answer: B



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D. d) both (a) and (c)

74. 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. 1) DNA transformation
- B. 2) DNA ligation
- C. 3) DNA spooling
- D. 4) DNA duplication



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75. Match column I with column II and select the correct answer from the given codes.

$$ColumnII$$
 $ColumnIII$

- A. Recombinant DNA (i) Chilled ethanol
- B. Precipitation of DNA (ii)DNAstaining
- C. Transposons (iii) Jumping genes
- D. Ethidium bromide (iv)Genetic engineering

A. 1)
$$A - (iv), B - (i), C - (iii), D - (ii)$$

B. 2)
$$A - (i), B - (iii), C - (ii), D - (iv)$$

C. 3)
$$A - (ii), B - (i), C - (iii), D - (iv)$$

D. 4)
$$A - (iv), B - (ii), C - (i), D - (iii)$$

Answer: A



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- 76. The polymerase chain reaction is a technique used for
- 1) amplification of DNA
- 2) amplification of enzymes
- 3) amplification of proteins
- 4) all of these
 - A. 1) amplification of DNA
 - B. 2) amplification of enzymes
 - C. 3) amplification of proteins
 - D. 4) all of these

Answer: A



77. Process	used	for	amplification	or	multiplication	of	DNA	in	DNA
fingerprintir	ng is								

- 1) polymerase chain reaction
- 2) southern blotting
- 3) northern blotting
- 4) None of these
 - A. 1) polymerase chain reaction
 - B. 2) southern blotting
 - C. 3) northern blotting
 - D. 4) None of these

Answer: A



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78. Primers are

1) chemically synthesised oligonucleotides that are complementary to the

regions of DNA

2) chemically synthesised oligonucleotides that are not complementary

to the regions of DNA

3) chemically synthesised, autonomously replicating circular DNA molecules

4) specific sequences present on recombinant DNA

A. 1) chemically synthesised oligonucleotides that are complementary to the regions of DNA

B. 2) chemically synthesised oligonucleotides that are not complementary to the regions of DNA

C. 3) chemically synthesised, autonomously replicating circular DNA molecules

D. 4) specific sequences present on recombinant DNA

Answer: A



79. Enzyme 'Taq polymerase' used in PCR, has been isolated from bacterium

- 1) Agrobacterium tumefaciens
- 2)Thermus aquaticus
- 3) Streptomyces albus
- 4) Escherichia coil
 - A. 1) Agrobacterium tumefaciens
 - B. 2)Thermus aquaticus
 - C. 3) Streptomyces albus
 - D. 4) Escherichia coil

Answer: B



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80. Which of the following statements are correct for the enzyme Taq polymerase ?

- (i) It remains active during the high temperature induced denaturation of dsDNA.
 - (ii) It requires primers for carrying out the process of polymerisation.
- (iii) It synthesises the RNA region between the primers, using dNTPs and Mq^{2+} .
 - A. a) (i) and (ii)
 - B.b) (ii) and (iii)
 - C. c) (i),(ii) and (iii)
 - D. d) None of these

Answer: A



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81. Match column I (enzyme) with column II (characteristic/activity) and select the correct answer from the given codes,

ColumnIColumnIIA. Taq DNA polymerase (i) Cleaves the ends of linear DNA B. Exonuclease (ii) Breakdown of fungal cell wall (iii)Stable above $90^{\circ}C$ C. Protease D. Chitinase (iv)Made only by eukaryotic cells (v)Degradation of proteins 1) A - (iii), B - (iv), C - (i), D - (ii) 2) A - (iv) .B - (iii) .C - (i) .D - (ii) 3) A - (ii), B - (i), C - (v), D - (iii)4) A - (iii), B - (i), C - (v), D - (ii) A. 1) A - (iii), B - (iv), C - (i), D - (ii)B. 2) A - (iv), B - (iii), C - (i), D - (ii)

C. 3) A - (ii), B - (i), C - (v), D - (iii)

D. 4) A - (iii), B - (i), C - (v), D - (ii)

Answer: D



- A. a) Tumour inducing Ti plasmid
- B. b) DNA probe Identifies the desired DNA fragment
- C. c) PCR DNA staining
- D. d) Agarose Sea weeds

Answer: C



is

- **83.** The correct sequence of different steps of polymerease chain reaction
- 1) annealing \rightarrow denaturation \rightarrow extension
- 2) denaturation \rightarrow extension \rightarrow annealing
- 3) denaturation \rightarrow annealing \rightarrow extension
- 4) extension \rightarrow denaturation \rightarrow annealing
 - A. 1) annealing $\,
 ightarrow \,$ denaturation $\,
 ightarrow \,$ extension
 - B. 2) denaturation \rightarrow extension \rightarrow annealing

C. 3) denaturation \rightarrow annealing \rightarrow extension

D. 4) extension \rightarrow denaturation \rightarrow annealing

Answer: C



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2. Apparatus Requirement

6. Error probability

84. Given table gives an account of differences between PCR and gene

DNA

Less

More

Restriction enzyme, ligase, vec

cloning. Which of the following points shows the incorrect difference?

Parameter PCR Gene cloning

1. Efficient More Less

3. Manipulation in vitro in vitro and in vivo

4. cost More Less 5. Automation Yes No

7. Time for a typical experiment 2-4days 4hours

8. Application More Less

a) 1 and 3

b) 4,5 and 6

c) 4 and 7

d) 4,7 and 8

A. a) 1 and 3 B. b) 4,5 and 6 C. c) 4 and 7 D. d) 4,7 and 8 **Answer: C Watch Video Solution** 85. Which of the following is required to perform polymerase chain reaction? A. a) Primers, dNTPs and DNA polymerase B. b) $DNA, CaCI_2$ and nuclease C. c) Mg^{2+} , DNAD. d) Both (a) and (c) Answer: D

86. In a polymerase chain reaction, temperature required for the steps

- (i) Denaturation,
- (ii) Annealing and
- (iii) Extension are respectively
- A) (i) 94 ° C (ii) 40 ° C (iii) 72 ° C
- B)(i)40 ° C(ii)72 ° C(iii)94 ° C
- C)(i)94 ° C(ii)72 ° C(iii)40 ° C
- D) (i) 72 ° C (ii) 94 ° C (iii) 40 ° C
 - A. A) $(i)94^{\circ}C(ii)40^{\circ}C(iii)72^{\circ}C$
 - B. B) $(i)40^{\circ}\,C(ii)72^{\circ}\,C(iii)94^{\circ}\,C$
 - C. C) $(i)94^{\circ}\,C(ii)72^{\circ}\,C(iii)40^{\circ}\,C$
 - D. D) $(i)72^{\circ}\,C(ii)94^{\circ}\,C(iii)40^{\circ}\,C$

Answer: A



87. In addition to Taq polymerase enzyme which other thermostable DNA polymerases have been isolated to be used in polymerase chain Reaction (PCR)?

- A. a) Pfu polymerase isolated from Pyrococcus furiosus
- B. b) Tli polymerase(vent ploymerase) isolated fromThermococcus litoralis
- C. c) Both (a) and (b)
- D. d) None of these

Answer: C



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88. Given figures represents the steps involved in polymerase chain reaction (PCR). Identify the steps A,B,C and C and select the correct

option.	
A.	
A	B
Denaturation at $94-96^{\circ}C$	Extension through Taq polymerase at
В.	
A	B C
Denaturation at $94-96^{\circ}C$	Annealing at $40-60^{\circ}C$ Extension
C.	
A	B C
Denaturation at $40-60^{\circ}C$	Annealing at $72^{\circ}C$ Extension through
D.	
A	B
Extension through Taq polyn	nerase at $72^{\circ}C$ Denaturation at 40°
Answer: B	
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89. In a polymerase chain reaction after the denaturation step why the	
mixture needs to cool down to a lower temperature ?	

- a) To permit specific annealing of the primers
- b) To give a halt to the reaction mixture
- c) To increase the activity of enzyme Taq polymerase
- d) To obtain the multiple copies of the DNA
 - A. a) To permit specific annealing of the primers
 - B. b) To give a halt to the reaction mixture
 - C. c) To increase the activity of enzyme Taq polymerase
 - D. d) To obtain the multiple copies of the DNA

Answer: A



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90. If a recombinant DNA bearing gene for resistance to antibiotic ampicillin is transferred to E.coli cells, the host cells become transformed into ampicillin resistant cells. If such bacteria are transferred on agar plates containing ampicillin, only transformants will grown and the untransformed recipient cells will die. The ampicillin resistant gene in this

case is called as 1) selectable marker 2) recombinant protein 3) cloning site 4) chemical scalpels A. 1) selectable marker B. 2) recombinant protein C. 3) cloning site D. 4) chemical scalpels Answer: A **Watch Video Solution 91.** Which of the following is not used to transfer the recombinant DNAinto the host? A. 1) Micro-injection method

- B. 2) Gene gun method
- C. 3) Bioreactor
- D. 4) Disarmed pathogen vectors

Answer: C



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- **92.** A device in which large volume of living cells are cultured in order to
- get a specific product is called
- 2) agitator

1)PCR

- 3) bioreactor
- 4) assimilator
 - A. 1)PCR
 - B. 2) agitator
 - C. 3) bioreactor

D. 4) assimilator

Answer: C



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- **93.** Which of the following statements are correct with respect to a biorector?
- (i) It can process large volumes of culture.
- (ii) It provides optimum temperature and pH.
- (iii) It is a completely automated tool.
- (iv) It is a compact thermal cycler.
 - A. a) (i) and (ii)
 - B. b) (i),(ii) and (iii)
 - C. c) (iii) and (iv)
 - D. d) (ii) and (iii)

Answer: A

94. Identify the figures (A) and (B) and select the correct option.



- A. $\frac{(A)}{\text{Spargal stirred-tank bioreactor}}$ (B). Spargal stirred-tank bioreactor
- B.
- (A)(B).

Spargal stirred-tank bioreactor Sparged stirred-tank bioreactor (A)(B).

- C. Simple stirred-tank Sparged stirred-tank bioreactor
- (B).(A)Simple stirred-tank bioreactor Simple stirred-tank bioreactor

Answer: C



they

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95. Stirred-tank bioreactors have advantages over shake flasks because

- A. a) provide high temperature and pH
- B. b) provide better aeration and mixing properties
- C. c) do not allow the entry of CO_2
- D. d) are easy to operate

Answer: B



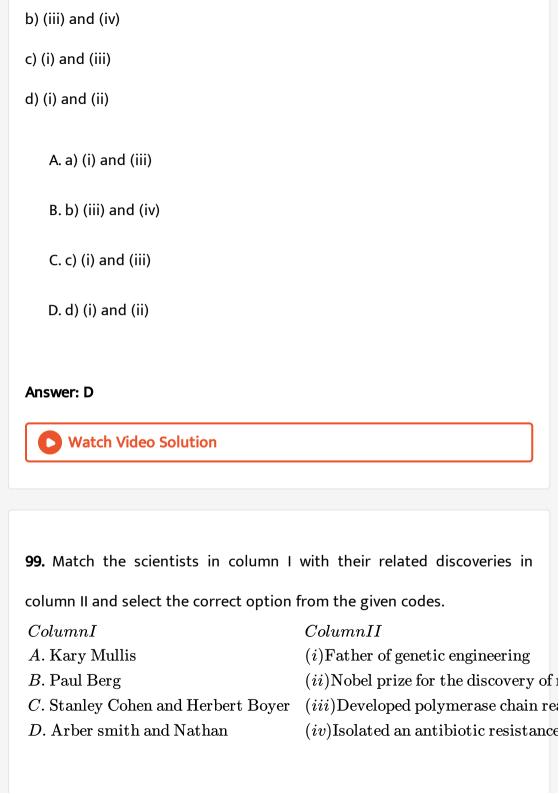
- **96.** After completion of the biosynthetic stage in the bioreactors, the product undergoes separation and purification processes, collectively termed as
 - A. a) transformation
 - B. b) electrophoresis
 - C. c) downstream processing
 - D. d) upstream processing

Answer: C



- **97.** Study the following statements regarding recombinant DNA technology and select the incorrect ones.
- (i) Taq polymerase extends the primers using the nucleotides provided in the reaction.
- (ii) Antibiotic resistance genes are considered as desirable genes in recombinant DNA technology.
- (iii) DNA fragments are separated according to their charge only, in agarose gel electrophoresis.
- (iv) Transformation is a procedure through which a piece of DNA is integrated in to the genome of a host bacterium.
- (v) To produce higher yields of a desired protein, host cells can be multiplied in a continuous culture.
- (vi) Downstream processing is one of the steps of polymerase chain reaction.

A. a) (ii),(iii) and (vi) B. b) (i),(iii) and (v) C. c) (ii),(iii) and (v) D. d) (i),(iv) and (v) Answer: A **Watch Video Solution** 98. Read the following statements and select the incorrect ones. (i) When the transformed cells on agar plates containing ampicillin are spread, both transformed and untransformed cells will grow. (ii) Restriction enzymes are used in isolation and separation of DNA from other macromolecules. (iii) Downstream processing is one of the steps of rDNA technology. (iv) Disarmed pathogen vectors are also used in transfer of rDNA into the host. a) (i) and (iii)



A. 1)
$$A-(iii), B-(i), C-(iv), D-(ii)$$

B. 2) $A-(iii), B-(iv), C-(i), D-(ii)$

C. 3) A - (iv), B - (ii), C - (iii), D - (i)

D. 4) A - (i), B - (iii), c - (iv), D - (ii)

Answer: A



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- (i) _____is a natural polymer extracted from____.
- (ii) The DNA fragments purified by gel electrophoresis are used in constructing____by joining them with_____.
- (iii) The ligation of alien DNA is carried out at a_____. present in one of the two in a plasmid vector.

(iv)_____enzyme remains active during the high temperature induced

denaturation of ds DNA

(v) DNA fragments are resolved according to their_____ through

in agarose gel electrophoresis.

a) (i) Agarose, sea weeds (ii) recombinant DNA, cloning vector (iii) restriction site, antibiotic resistance genes (iv) Taq polymerase (v) size, sieving effect

b) (i) Agarose, sea weeds (ii) Restriction site, antibiotic resistance genes (iii) recombinant DNA, cloning vector (iv) Taq polymerase (v) size, sieving effect

c) (i) Agarose, sea weeds (ii) restriction site, antibiotic resistance genes (iii) recombinant DNA, cloning vector (iv) Taq polymerease (v) size, sieving effect

d) (i) size, sieving effect (ii) agarose, seaweeds (iii) recombinant DNA cloning vector (iv) Taq polymerase (v) restriction site, antibiotic resistance genes

A. a) (i) Agarose, sea weeds (ii) recombinant DNA, cloning vector (iii) restriction site, antibiotic resistance genes (iv) Taq polymerase (v) size, sieving effect

B. b) (i) Agarose, sea weeds (ii) Restriction site, antibiotic resistance genes (iii) recombinant DNA, cloning vector (iv) Taq polymerase (v)

size, sieving effect

C. c) (i) Agarose, sea weeds (ii) restriction site, antibiotic resistance genes (iii) recombinant DNA, cloning vector (iv) Taq polymerease (v) size, sieving effect

D. d) (i) size, sieving effect (ii) agarose, eas weeds (iii) recombinant

DNA cloning vector (iv) Taq polymerase (v) restriction site, antibiotic

resistance genes

Answer: A



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101. Four mutant strians of bacteria (1-4) all require substance S to grow (each strian is blocked at one step in the S-biosynthesis pathway). Four plates were prepared with minimal medium and a trace of substance, S to allow a small amount of growth of mutant cells. on plate A, mutant cells of strain 1 were spred over entire surface of tha agar to form a thin

law of bacteria. On plate B, the lawn was composed of mutant cells of strain 2, and so on. On each plate, cells of each of the four mutant types were inoculated over the lawn, as indicated in the figure by the circles. dark circles indicae excellent growth. A strain blocked at a later step in the S substance metabolic pathwa accumulates intermediates that can 'feed' a strain blocked at an earlier step.



What is the order of genes (1-4) in the metabolic pathway for synthesis of substance S?

A.
$$2 o 4 o 3 o 1$$

B.
$$2 o 1 o 3 o 4$$

$$\mathsf{C.}\, 1 \to 3 \to 4 \to 2$$

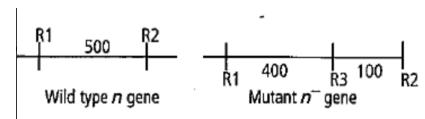
D.
$$1 o 2 o 4 o 3$$

Answer: C



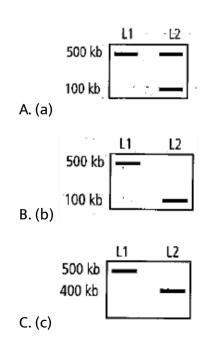
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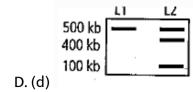
102. The figure shows the restriction enzyme cutting sites (R1-R3) in wild type (n) and mutant (n) gene.



If a radioactively labelled probe (that hybridises at a sequence close to R1) is used for detecting the presence of DNA fragments after gel electrophoresis and Southern blotting, which of the following band patterns will yout expect ?

Note: L1: wild type DNA, L2: mutant DNA





Answer: C



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103. Analyse the given diagram which steps involved in the procedure of selecting transformed bacteria.



Identify the bacterial colony which has undergone transformation?

A. colony 5

B. Colony 2

C. Colony 4

D. Colony 3

Answer: C

104. The nucleic acid extracted from animal liver is loaded and run on agarose gel. After staining, it shows following pattern:



If the remaining sample is treated with RNAse and loaded in gel what result would you expect ?



В. 📄

C. 🔀

D. 📝

Answer: A



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105. The basic procedure involved in the synthesis of recombinanat DNA molecule is depicted below. The mistake in the procedure is



- A. Enzyme polmerase is not included.
- B. The mammalian DNA is shown double stranded
- C. Two different restriction enzymes are used.
- D. Only one fragment is inserted

Answer: C



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- 106. Rising of dough is due is
 - A. a) multiplication of yeast
 - B. b) production of CO_2
 - C. c) emulsification

D. d) hydrolysis of wheat flour strach into sugars.

Answer: B

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107. An enzyme catalysing the removal of nucleotides from the end

107. An enzyme catalysing the removal of nucleotides from the ends of DNA is

- A. a) endonuclease
- B. b) exonuclease
- C. c) DNA ligase
- D. d) Hind II

Answer: B



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

- A. a) transduction
- B. b) conjugation
- C. c) transformation
- D. d) translation

Answer: A



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

- A. a) DNA can be seen in visible light
- B. b) DNA can be seen without staining in visible light.
- C. c) Ethidium bromide stained DNA can be seen in visible light.

D. d) Ethidium bromide stained DNA can be seen under exposure to UV light

Answer: D



110. Restriction' in restriction enzyme refers to

- A. a) cleaving of phosphodiester bond in DNA by the enzyme
- B. b) cutting of DNA at specific position only
- C. c) prevention of the multiplication of bacteriophage in bacteria
- D. d) all of the above

Answer: C



111. 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



112. In agarose gel electrophoresis, DNA molecules are separated on the basis of their

A. 1) charge only

B. 2) size only

C. 3) charge to size ratio

D. 4) all of the above

Answer: B



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- 113. The most important feature in a plasmid to be used as a vector is
 - A. a)origin of replication (ori)
 - B. b)presence of a selectable marker
 - C. c) presence of sites for restriction endonuclease
 - D. d) its size

Answer: A::B::C::D



114. While isolating DNA from bacteria, which of the following enzymes is not used ?

A. a) Lysozyme

B. b) Ribonuclease

C. c) Deoxyribonuclease

D. d) Protease

Answer: C



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115. Which of the following has popularised the PCR (polymerase chain reactions)?

A. a) Easy availability of DNA template

B. b) Availability of synthetic primers

C. c) Availability of cheap deoxyribonucleotides

D. d) Availability of 'thermostable' DNA polymerase

Answer: D



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116. An antibiotic resistant gene in a vector usually helps in the selection of

- A. a) competent cells
- B. b) transformed cells
- C. c) recombinant cells
- D. d)none of the above

Answer: B



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

- A. a) binding of DNA to the cell wall
- B. b) uptake of DNA through membrane transport proteins
- C. c) uptake of DNA through transient pores in the bacterial cell wall
- D. d) expression of antibiotic resistance gene

Answer: C



118. The role of DNA ligase in the construction of a recmobinant DNA molecule is

- A. a) formation of phosphodiester bond between two DNA fragments
- B. b)formation of hydrogen bonds between sticky ends of DNA
 - fragments

C. c) ligation of all purine and pyrimidine bases

D. d)none of the above

Answer: A



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- 119. Which of the following is not a source of restriction endonuclease?
- (a) Haemophilus influenzea
- (b) Escherichia coil
- (c) Entamoeba coil
- (d) Bacillus amyloliquifaciens
- A. Haemophilus influenzea
 - B. Escherichia coil
 - C. Entamoeba coil
 - D. Bacillus amyloliquifaciens

Answer: C

120. 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 and on the template DNA
- (d) All of the above
 - A. Denaturation of template DNA
 - B. Annealing of primers to template DNA
 - C. Extension of primer and on the template DNA
 - D. All of the above

Answer: C



121. 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

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 degrated by bacteria

D. all of the above

Answer: A



122. 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 continuous culture system
- (d) Any of the above
 - A. Laboratory flask of largest capcity
 - B. A stirred-tank bioreactro without in-lets and out-lets
 - C. A continuous culture system
 - D. Any of the above

Answer: C



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

(a) Herbert Boyer (b) Hargovind Khurane (c) Kary Mullis (d) Arthur Kornberg A. Herbert Boyer B. Hargovind Khurane C. Kary Mullis D. Athur Kornberg **Answer: C Watch Video Solution** 124. Which of the following statements does not hold true for restriction enzyme? (a) It recongnises a palindromic nucleotide sequence (b) It is an endonuclease.

- (c) It is isolated from viruses
- (d) It produces the same kind of sticky ends in different DNA molecules
 - A. It recongnises a palindromic nucleotide sequence
 - B. It is an endonuclease.
 - C. It is isolated from viruses
 - D. It produces the same kind of sticky ends in different DNA molecules

Answer: C



- **125.** Assertion: In a chemical engineering process, it is necessary to prepare sterile ambience.
- Reason: Sterile ambience inhibits the growth of undesirable microbes during manufacture of product like antibiotics, vaccines and enzymes.
 - A. a) Both assertion and reason are true and reason is correct explanation of assertion.

B. b) Both assertion and reason are true but reason is not correct explanation for assertion.

C. c) Assertion is true but reason is false.

D. d) Assertion is false but reason is true.

Answer: A



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126. Assertion: Asexual reproduction is more important with regard to biotehnology.

Reason: Asexual reproduction preserves the genetic information while sexual reporduction permits variations.

A. a) Both assertion and reason are true and reason is correct explanation for assertion.

B. b) Both assertion and reason are true but reason is not the correct

explanation for assertion.

C. c) Assertion is true but reason is false.

D. d) Assertion is false but reason is true.

Answer: A



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127. Assertion: Genetic engineering can overcome the drawbacks of traditional hybridisation.

Reason: Genetic engineering can create desired DNA sequences to meet specific requirements.



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128. Assertion: A piece of DNA inserted into an alien organism generally does not replicate if not inserted into a chromosome.

Reason: Chromosomes have specific sequences called ori region where DNA replication is initiated.



129. Assertion: Genetic engineering requires both nuclease and ligases.

Reason: Ligases produce the nick in the recombinant DNA molecule.



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130. Assertion: Restriction enzymes Hin and Hpa are produced from two

Reason : Hin is produced from Haemophilus while Hpa is produced from

Hematococcus.



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different genera of bacteria.

131. Assertion: Restriction enzymes recongise palindromic sequences.

Reason: Palindromic sequences read same in both directions of the two strand.

A. d) Both assertion and reason are false. B. C. D. Answer: B Watch Video Solution 132. Assertion: The matrix used in gel electrophoresis should have controllable pore size. Reason: Agarose concentration can be changed to change pore sizes. A. a) Both assertion and reason are true and reason is correct explanation for assertion B.b) Both assertion and reason are true but reason is not correct explanation of assertion C. c) Assertion is true but reason is false.

D. d) Both assertion and reason are false.

Answer: B



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133. Assertion : All expression vectors are cloning vectors and vise versa.

Reason: Expression vectors have at least the regulatory sequences i.e., promotes, operators, ribosomal binding sites, etc having optimum function in the chosen control but not origin of replication.



134. Assertion: E.coli having pBR322 with DNA insert at BamHI site cannot grow in medium containing tetracyline.

Reason : Recognition site for BamHI is present in ter^R region of pBR22.



135. Assertion: A bacterial cell with restriction enzymes will be easily infected and lysed by bacteriophages.

Reason: Restriction enzymes catalyse synthesis of protective coat around bacterial cell that prevents bacteriophage attack.



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136. Assertion: Special methods are used for transformation i.e., incorporation of recombinant DNA into host.

Reason : DNA is a hydrophilic molecule.



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137. Assertion: Use of chitinase enzyme is necessary for isolation of DNA from yeast cells but not in case of Spirogyra.

Reason: Fungal cell wall is made up of fungal cellulose or chitin.



138. Assertion: PCR primers must not have self complementary regions.

Reason: Self complementary regions result in hairpin structures adversely affecting the PCR.



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139. Assertion: Downstream processing is generally considered more difficult and costlier in plants than that in microbes.

Reason: Rhizosecretion is used as a method to facilitate easier recovery of recombinant proteins from plants.

