CHAPTER 11

Biotechnology: Principles and Processes

11.1 Principles of Biotechnology

- 1. The DNA molecule to which the gene of interest is integrated for cloning is called
 - (a) template
- (b) carrier
- (c) transformer
- (d) vector.
- (2015)
- 2. The cutting of DNA at specific locations became possible with the discovery of
 - (a) selectable markers (b) ligases
 - (c) restriction enzymes (d) probes.
- (2015)
- **3.** Which one of the following is a case of wrong matching?
 - (a) Somatic hybridization
- Fusion of two diverse cells
- (b) Vector DNA
- Site for *t*RNA synthesis
- (c) Micropropagation *In vitro* production of plants in large numbers
- (d) Callus
- Unorganised mass of cells produced in tissue culture (201
- **4.** Which one of the following techniques made it possible to genetically engineer living organisms?
 - (a) Recombinant DNA techniques
 - (b) X-ray diffraction
 - (c) Heavier isotope labelling
 - (d) Hybridization

(Mains 2011)

- 5. Which of the following are used in gene cloning?
 - (a) Nucleoids
- (b) Lomasomes
- (c) Mesosomes
- (d) Plasmids
- (2010)
- **6.** Manipulation of DNA in genetic engineering became possible due to the discovery of
 - (a) restriction endonuclease
 - (b) DNA ligase
 - (c) transcriptase
 - (d) primase.

(2002)

- 7. The bacteria generally used for genetic engineering is
 - (a) Agrobacterium
- (b) Bacillus
- (c) Pseudomonas
- (d) Clostridium. (2000)

- **8.** Which of the following is related to genetic engineering?
 - (a) Heterosis
- (b) Mutation
- (c) Plastid
- (d) Plasmid (1999)
- 9. Genetic engineering is possible, because
 - (a) we can cut DNA at specific sites by endonucleases like DNase I
 - (b) restriction endonucleases purified from bacteria can be used *in vitro*
 - (c) the phenomenon of transduction in bacteria is well understood
 - (d) we can see DNA by electron microscope.

(1998)

- When scientists make an animal superior by view of genotype, introducing some foreign genes in it, is called
 - (a) immunization
- (b) genetic engineering
- (c) tissue culture
- (d) biotechnology. (1996)
- **11.** Which of the following organelles is related with genetic engineering?
 - (a) Mitochondria
- (b) Plasmids
- (c) Golgi bodies
- (d) Lysosomes

(1994)

11.2 Tools of Recombinant DNA Technology

- **12.** Identify the wrong statement with regard to restriction enzymes.
 - (a) Each restriction enzyme functions by inspecting the length of a DNA sequence.
 - (b) They cut the strand of DNA at palindromic sites.
 - (c) They are useful in genetic engineering.
 - (d) Sticky ends can be joined by using DNA ligases. (NEET 2020)
- 13. Choose the correct pair from the following.
 - (a) Ligases
- Join the two DNA molecules
- (b) Polymerases
- Break the DNA into fragments
- (c) Nucleases
- Separate the two strands of
- DNA
- (d) Exonucleases Make cuts at specific positions within DNA

(NEET 2020)

	0,									
14.	The specific palindromic	sequence which is recognised	20.	Which of the following is commonly used a						
	by <i>Eco</i> RI is			introducing a DNA fr	agment in human ly					
	(a) 5' - GAATTC - 3'	(b) 5' - GGAACC - 3'		(a) Retrovirus	(b) Ti plasmid					
	3' - CTTAAG - 5'	3' - CCTTGG - 5'		(c) λ phage	(d) pBR322 (
	(c) 5' - CTTAAG - 3' 3' - GAATTC - 5'	(d) 5' - GGATCC - 3' 3' - CCTAGG - 5'.	21.	The DNA fragments separated on an agabe visualised after staining with						
		(NEET 2020)		(a) acetocarmine	(b) aniline blu					

- 15. The sequence that controls the copy number of the linked DNA in the vector, is termed
 - (a) selectable marker (b) Ori site
 - (c) palindromic sequence (d) recognition site.

(NEET 2020)

- **16.** In gel electrophoresis, separated DNA fragments can be visualized with the help of
 - (a) acetocarmine in bright blue light
 - (b) ethidium bromide in UV radiation
 - (c) acetocarmine in UV radiation
 - (d) ethidium bromide in infrared radiation.

(NEET 2020)

- 17. Following statements describe the characteristics of the enzyme restriction endonuclease. Identify the incorrect statement.
 - (a) The enzyme recognises a specific palindromic nucleotide sequence in the DNA.
 - (b) The enzyme cuts DNA molecule at identified position within the DNA.
 - (c) The enzyme binds DNA at specific sites and cuts only one of the two strands.
 - (d) The enzyme cuts the sugar-phosphate backbone at specific sites on each strand. (NEET 2019)
- 18. A selectable marker is used to
 - (a) help in eliminating the non-transformants, so that the transformants can be regenerated
 - (b) identify the gene for a desired trait in an alien organism
 - (c) select a suitable vector for transformation in a specific crop
 - (d) mark a gene on a chromosome for isolation using restriction enzyme. (Odisha NEET 2019)
- 19. Given below are four statements pertaining to separation of DNA fragments using gel electrophoresis. Identify the incorrect statements.
 - (i) DNA is negatively charged molecule and so it is loaded on gel towards the anode terminal.
 - (ii) DNA fragments travel along the surface of the gel whose concentration does not affect movement of DNA.
 - (iii) Smaller the size of DNA fragment larger is the distance it travels through it.
 - (iv) Pure DNA can be visualized directly by exposing UV radiation.

Choose correct answer from the options given below.

- (a) (i), (iii) and (iv)
- (b) (i), (ii) and (iii)
- (c) (ii), (iii) and (iv)
- (d) (i), (ii) and (iv) (Odisha NEET 2019)

- as a vector for mphocytes?
 - (NEET 2018)
- arose gel can
- (c) ethidium bromide
- (d) bromophenol blue.

(NEET 2017)

- 22. DNA fragments are
 - (a) negatively charged (b) neutral
 - (c) either positively or negatively charged depending on their size
 - (d) positively charged.

(NEET 2017)

- 23. A gene whose expression helps to identify transformed cell is known as
 - (a) vector
- (b) plasmid
- (c) structural gene
- (d) selectable marker.

(NEET 2017)

- **24.** What is the criterion for DNA fragments movement on agarose gel during gel electrophoresis?
 - (a) The smaller the fragment size, the farther it moves.
 - (b) Positively charged fragments move to farther end.
 - (c) Negatively charged fragments do not move.
 - (d) The larger the fragment size, the farther it moves. (NEET 2017)
- 25. A foreign DNA and plasmid cut by the same restriction endonuclease can be joined to form a recombinant plasmid using
 - (a) EcoRI
- (b) Taq polymerase
- (c) polymerase III
- (d) ligase. (NEET-II 2016)
- **26.** Which of the following restriction enzymes produces blunt ends?
 - (a) SalI
- (b) EcoRV (c) XhoI
- (d) HindIII (NEET-II 2016)
- 27. Which of the following is not a feature of the plasmids?
 - (a) Transferable
- (b) Single-stranded
- (c) Independent replication
- (d) Circular structure

(NEET-I 2016)

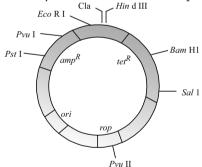
- **28.** Which of the following is a restriction endonuclease?
 - (a) DNase I
- (b) RNase
- (c) Hind II
- (d) Protease

(NEET-I 2016)

- **29.** The introduction of T-DNA into plants involves
 - (a) exposing the plants to cold for a brief period
 - (b) allowing the plant roots to stand in water
 - (c) infection of the plant by Agrobacterium tumefaciens
 - (d) altering the pH of the soil, then heat-shocking the plants. (2015)

- **30.** Which vector can clone only a small fragment of DNA?
 - (a) Bacterial artificial chromosome
 - (b) Yeast artificial chromosome

 - (c) Plasmid (d) Cosmid (2014)
- 31. Commonly used vectors for human genome sequencing
 - (a) T DNA
- (b) BAC and YAC
- (c) expression vectors
- (d) T/A cloning vectors. (2014)
- 32. The colonies of recombinant bacteria appear white in contrast to blue colonies of non-recombinant bacteria because of
 - (a) insertional inactivation of alpha galactosidase in recombinant bacteria
 - (b) inactivation of glycosidase enzyme in recombinant bacteria
 - (c) non-recombinant bacteria containing beta galactosidase
 - (d) insertional inactivation of alpha galactosidase in non-recombinant bacteria. (NEET 2013)
- 33. DNA fragments generated by the restriction endonucleases in a chemical reaction can be separated by
 - (a) electrophoresis
- (b) restriction mapping
- (c) centrifugation
- (d) polymerase chain reaction. (NEET 2013)
- **34.** The given figure is the diagrammatic representation of the *E. coli* vector pBR322. Which one of the given options correctly identifies its certain component(s)?



- (a) *ori*-original restriction enzyme
- (b) rop-reduced osmotic pressure
- (c) HindIII, EcoRI selectable markers
- (d) amp^R , tet^R -antibiotic resistance genes (2012)
- 35. A single strand of nucleic acid tagged with a radioactive molecule is called
 - (a) vector
- (b) selectable marker
- (c) plasmid
- (d) probe.
- (2012)

(2012)

- **36.** For transformation, micro-particles coated with DNA to be bombarded with gene gun are made up of
 - (a) silver or platinum (b) platinum or zinc

 - (c) silicon or platinum (d) gold or tungsten.

- - **37.** Biolistics (gene-gun) is suitable for
 - (a) disarming pathogen vectors
 - (b) transformation of plant cells
 - (c) constructing recombinant DNA by joining with
 - (d) DNA fingerprinting.

(Mains 2012)

- **38.** In genetic engineering, the antibiotics are used
 - (a) as selectable markers
 - (b) to select healthy vectors
 - (c) as sequences from where replication starts
 - (d) to keep the cultures free of infection.

(Mains 2012)

- 39. Which one of the following represents a palindromic sequence in DNA?
 - (a) 5' GAATTC 3' (b) 5' - CCAATG - 3'
 - 3' CTTAAG 5' 3' - GAATCC - 5'
 - (c) 5' CATTAG 3' (d) 5' - GATACC - 3'
 - 3' GATAAC 5' 3' - CCTAAG - 5'

(*Mains 2012*)

40. Given below is a sample of a portion of DNA strand giving the base sequence on the opposite strands. What is so special shown in it?

5' GAATTC 3'

- 3' _____ CTTAAG _____ 5'
- (a) Replication completed
- (b) Deletion mutation
- (c) Start codon at the 5' end
- (d) Palindromic sequence of base pairs (2011)
- **41.** There is a restriction endonuclease called *Eco*RI. What does "co" part in it stand for?
 - (a) colon
- (b) coelom
- (c) coenzyme
- (d) coli (2011)
- **42.** Agarose extracted from sea weeds is used in
 - (a) spectrophotometry (b) tissue culture
 - (c) PCR
- (d) gel electrophoresis.

(2011)

43. Which one of the following palindromic base sequences in DNA can be easily cut at about the middle by some particular restriction enzyme?

- 5' CGTTCG 3' ATGGTA –
- 5' GATATG — CTACTA -
- GAATTC -
- CACGTA -
 - (2010)- CTCAGT -
- 44. Which one of the following is used as vector for cloning genes into higher organisms?
 - (a) Baculovirus
 - (b) Salmonella typhimurium
 - (c) Rhizopus nigricans (d) Retrovirus (2010)

45. DNA or RNA segment tagged with a radioactive 53. Two microbes found to be very useful in genetic molecule is called engineering are (b) probe (a) vector (a) crown gall bacterium and Caenorhabditis elegans (c) clone (d) plasmid. (2010)(b) Escherichia coli and Agrobacterium tumefaciens (c) Vibrio cholerae and a tailed bacteriophage **46.** Restriction endonucleases are enzymes which (d) *Diplococcus sp.* and *Pseudomonas sp.* (a) make cuts at specific positions within the DNA (2006)molecule (b) recognize a specific nucleotide sequence for **54.** Restriction endonucleases binding of DNA ligase (a) are present in mammalian cells for degradation (c) restrict the action of the enzyme DNA of DNA when the cell dies polymerase (b) are used in genetic engineering for ligating two (d) remove nucleotides from the ends of the DNA DNA molecules molecule. (2010)(c) are used for in vitro DNA synthesis 47. In genetic engineering, a DNA segment (gene) of (d) are synthesized by bacteria as part of their interest, is transferred to the host cell through a defense mechanism. vector. Consider the following four agents (i-iv) in **55.** The *Ti* plasmid, is often used for making transgenic this regard and select the correct option about which plants. The plasmid is found in one or more of these can be used as a vector/vectors. (a) Azotobacter (i) Bacterium (ii) Plasmid (b) *Rhizobium* of the roots of leguminous plants (iv) Bacteriophage (iii) Plasmodium (c) Agrobacterium (a) (i), (ii) and (iv) (b) (i) only (d) Yeast as a 2 mm plasmid. (2004)(c) (i) and (iii) (d) (ii) and (iv) **56.** The most thoroughly studied of the known bacteria-(Mains 2010) plant interactions is the **48.** Polyethylene glycol method is used for (a) cyanobacterial symbiosis with some aquatic ferns (a) biodiesel production (b) gall formation on certain angiosperms by (b) seedless fruit production Agrobacterium (c) energy production from sewage (c) nodulation of Sesbania stems by nitrogen fixing (d) gene transfer without a vector. (2009)bacteria **49.** Which one of the following is commonly used in (d) plant growth stimulation by phosphatetransfer of foreign DNA into crop plants? solubilising bacteria. (2004)(a) Meloidogyne incognita 57. Which one of the following bacteria has found (b) Agrobacterium tumefaciens extensive use in genetic engineering work in plants? (c) Penicillium expansum (a) Clostridium septicum (d) Trichoderma harzianum (2009)(b) Xanthomonas citri **50.** Gel electrophoresis is used for (c) Bacillus coagulens (a) construction of recombinant DNA by joining (d) *Agrobacterium tumefaciens* (2003)

with cloning vectors

(b) isolation of DNA molecules

(c) cutting of DNA into fragments

(d) separation of DNA fragments according to their (2008)

51. The linking of antibiotic resistance gene with the plasmid vector became possible with

(a) DNA polymerase

(b) exonucleases

(c) DNA ligase

(d) endonucleases. (2008)

(2006)

52. Restriction endonuclease

- (a) synthesizes DNA
- (b) cuts the DNA molecule randomly
- (c) cuts the DNA molecule at specific sites
- (d) restricts the synthesis of DNA inside the nucleus.

- **58.** Which of the following enzymes are used to join bits of DNA?
 - (a) Ligase

(b) Primase

(c) DNA polymerase

(d) Endonuclease

(2002)

- **59.** A mutant strain of T₄ Bacteriophage, R-II, fails to lyse the *E. coli* but when two strains R-IIX and R-IIY are mixed then they lyse the *E. coli*. What may be the possible reason?
 - (a) Bacteriophage transforms in wild.
 - (b) It is not mutated.
 - (c) Both strains have similar cistrons.
 - (d) Both strains have different cistrons.

(2001)

60. Which of the following cut the DNA from specific 11.3 Processes of Recombinant DNA places? **Technology** (a) E.coli restriction endonuclease I **68.** Match the organism with its use in biotechnology. (b) Ligase (A) Bacillus (i) Cloning vector (c) Exonuclease thuringiensis (d) Alkaline phosphate (2001)(B) Thermus (ii) Construction of first 61. Maximum number of bases in plasmids discovered rDNA molecule aquaticus so far (C) Agrobacterium (iii) DNA polymerase (a) 50 kilo base (b) 500 kilo base tumefaciens (c) 5000 kilo base (d) 5 kilo base. (2001)(D) Salmonella (iv) Cry proteins **62.** Plasmid has been used as vector because typhimurium (a) it is circular DNA which have capacity to join to Select the correct option from the following. eukaryotic DNA (A) (B) (C) (D) (b) it can move between prokaryotic and eukaryotic (a) (ii) (iv) (iii) (i) cells (b) (iv) (iii) (i) (ii) (c) both ends show replication (c) (iii) (ii) (iv) (i) (d) it has antibiotic resistance gene. (2000)(d) (iii) (iv) (i) (ii) (NEET 2020) **69.** DNA precipitation out of a mixture of biomolecules 63. The process of replication in plasmid DNA, other than initiation, is controlled by can be achieved by treatment with (a) chilled chloroform (a) mitochondrial gene (b) plasmid gene (b) isopropanol (c) chilled ethanol (c) bacterial gene (d) none of these. (d) methanol at room temperature. (1999)(NEET 2019) **70.** Which one of the following equipments is essentially 64. Recombinant DNA is achieved by cleaving the required for growing microbes on a large scale, for pro-DNAs by industrial production of enzymes? (a) ligase (a) Bioreactor (b) BOD incubator (b) restriction endonuclease (c) Sludge digester (d) Industrial oven (c) primase (NEET 2019) (d) exonucleases. (1998)71. The correct order of steps in Polymerase Chain 65. Two bacteria found to be very useful in genetic Reaction (PCR) is engineering experiments are (a) extension, denaturation, annealing (a) Nitrobacter and Azotobacter (b) annealing, extension, denaturation (b) Rhizobium and Diplococcus (c) denaturation, extension, annealing (c) Nitrosomonas and Klebsiella (d) denaturation, annealing, extension. (d) Escherichia and Agrobacterium. (1998)(NEET 2018) 66. Restriction endonucleases are 72. The process of separation and purification of (a) used for in vitro DNA synthesis expressed protein before marketing is called (b) used in genetic engineering (a) downstream processing (c) synthesized by bacteria (b) bioprocessing (d) present in mammalian cells for degradation of (c) postproduction processing DNA. (1998)(d) upstream processing. (NEET 2017) **67.** The restriction enzymes are used in genetic engineering, 73. Stirred-tank bioreactors have been designed for (a) purification of product (a) they can cut DNA at specific base sequence

(b) they are nucleases that cut DNA at variable sites

(c) they can degrade harmful proteins

(d) they can join different DNA fragments.

(b) addition of preservatives to the product

vessel.

(1995)

(c) availability of oxygen throughout the process

(d) ensuring anaerobic conditions in the culture

(NEET-II 2016)

,	(a) S	epara	_	cessin	(b)	Purifi Expre	ession		2016)	81.	(c) (d)	genet DNA genet sich o	seque ic fing	ncing erprii	nting.		ıt reg	arding	(2012) g DNA
75.	The Taq polymerase enzyme is obtained from (a) Bacillus subtilis (b) Pseudomonas putida (c) Thermus aquaticus (d) Thiobacillus ferroxidans. (NEET-I 2016)										(a) (b) (c)	A in re	ecipient						
76.	An analysis of chromosomal DNA using the Southern hybridization technique does not use (a) electrophoresis (b) blotting (c) autoradiography (d) It remains acceptable to the figure below Polymerase Characteristics (d) It remains acceptable to the figure below Polymerase Characteristics (d) It remains acceptable to the figure below Polymerase Characteristics (d) It remains acceptable to the figure below Polymerase Characteristics (d) It remains acceptable to the figure below Polymerase Characteristics (a) electrophoresis (b) blotting (c) autoradiography (d) PCR. (2014)											ow sh nain correc	v shows three steps (A, B, C) of ain Reaction (PCR). Select the orrect identification together with						
77.	by (a) P (b) N (c) el	what it represents? PCR and RAPD Northern blotting electrophoresis and HPLC microscopy. what it represents? A. 5' Region to be amplified A. 5' Region to be amplified A. 5' S'																	
78.	Which of the following is not correctly matched for the organism and its cell wall degrading enzyme? (a) Algae - Methylase (b) Fungi - Chitinase (c) Bacteria - Lysozyme (d) Plant cells - Cellulase (NFET 2013)												ut 50°C						
79.	During the process of isolation of DNA, chilled ethanol is added to (a) precipitate DNA (b) break open the cell to release DNA (c) facilitate action of restriction enzymes (d) remove proteins such as histones. (Karnataka NEET 2013)									83.	 (c) C - extension in the presence of heat stable DNA polymerase (d) A - annealing with two sets of primers								
80.	PCR are th		estricti hods		igmen	t leng	th pol	ymorp	ohism			availa	0						
								—(/ER KE	_								
1.	(d)	2.	(c)	3.	(b)	4.	(a)	5.	(d)	6.	(a)	7.	(a)	8.	(d)	9.	(b)	10.	(b)
11.	(b)	12.	(d)	13.	(a)	14.	(a)	15.	(b)	16.	(b)	17.	(c)	18.	(a)	19.	(d)	20.	(a)
21.	(c)	22.	(a)	23.	(d)	24.	(a)	25. 35	(d)	26. 36	(b)	27.	(b)	28.	(c)	29.	(c)	30.	(c)
31. 41.	(b) (d)	32. 42.	(c) (d)	33.43.	(a) (c)	34.44.	(d) (d)	35. 45.	(d) (b)	36. 46.	(d) (a)	37.47.	(b) (d)	38. 48.	(a) (d)	39. 49.	(a) (b)	40. 50.	(d) (d)
51.	(c)	52.	(a) (c)	53.	(b)	54.	(d)	45. 55.	(b)	56.	(a) (b)	57.	(d)	58.	(a)	49. 59.	(d)	60.	(a)
61.	(b)	62.	(a)	63.	(c)	64.	(b)	65.	(d)	66.	(b)	67.	(a)	68.	(b)	69.	(c)	70.	(a)
71.	(d)	72.	(a)	73.	(c)	74.	(d)	75.	(c)	76.	(d)	77.	(a)	78.	(a)	79.	(a)	80.	(d)
81.	(d)	82.	(c)	83.	(d)		. ,		. ,		. ,		. /		. /		. /		

(a) study of enzymes

74. Which of the following is not a component of |