Total Pages-6 UG/III/MICB/H/VI/18 (New)

2018

MICROBIOLOGY

[Honours]

PAPER - VI

Full Marks: 90

Time: 4 hours

The figures in the right hand margin indicate marks

Candidates are required to give their answers in their own words as far as practicable

Illustrate the answers wherever necessary

[NEW SYLLABUS]

GROUP - A

Answer any two questions:

 15×2

1. (a) What is genotype and phenotype? $1\frac{1}{2}+1\frac{1}{2}$

(b) Give the structure of prokaryotic genome. 4

			2
	(c)	What is backcross?	2,
	(d)	Write short notes on:	2 + 2 + 2
		(i) Incomplete dominance	
		(ii) Co-dominance	
		(iii) Epistasis.	
2.	(a)	What is the operon concept?	2
	(b)	What is the difference between lac	c and 3
40		tryptophan operon?	73.00
	(c)	Give the life sketch of T ₄ phage.	5
	(d)	What is theta (θ) replication?	2.
	(e)	Write note on transcription.	3
3.	(a)) What is spontaneous and induced muta	ation? 2+2
	(b)	What is the difference between perecand paracentric inversion?	entric 2
	(c) What is microbial mutant?	2
	(a	f) What way you will isolate the mic mutant?	crobial 3

	(e)	What is site directed mutagenesis?	2
×	(f)	What is "Delbruck's fluctuation test?	2
4.	(a)	Describe the steps of Agrobact mediated gene transfer.	erium 4
	(b)	Mention the application of generating in medicine, agriculture environmental pollution control.	
	(c)	What do you mean by "bioethics of g engineering"?	enetic 2
,	(d)	Write the steps of Recombinant technology.	DNA 3
		GROUP - B	
		Answer any five questions:	8 x 5
5.	(a)	A DNA molecule was digested by restrenzyme and analysed by gel electrophe. Only one band appear — Explain.	
	(b)	How R-plasmid helps in the developm drug resistance?	nent of
UG/	шим	CB/H/VI/18 (New)	(Turn Over)

UG/III/MICB/H/VI/18 (New)

6.	(a)	What is Real Time PCR?	4
	(b)	Mention the drawbacks of genetic engineering.	4
7.	(a)	Write a note on SOS repair.	4
	(b)	Compare between Western blotting and Southern blotting.	4
8.	(a)	Briefly describe the steps of protein sequencing.	4
	(b)	Briefly describe nif gene.	4
9,	(a)	Why P^{BR322} is considered as suitable cloning vector?	3
	(b)	Briefly describe the types of restriction endonuclease with example.	5
10.	(a)	Distinguish between co-transduction and abortive transduction.	4
	(b)	Distinguish between genomic library and cDNA library.	đ,

(Continued)

UG/III/MICB/H/VI/18 (New)

11.	(a)	State the protocol used for developing transgenic plant.	4
	(b)	Briefly discuss the process of replica plating technique to detect mutation in bacteria.	4
12.	(a)	Briefly mention the steps of isolation of DNA in the laboratory.	4
1	(b)	Write notes on cosmid and phagemid.	4
12		GROUP - C	
	3	Answer any five questions: 4>	< 5
13.		ention the process of mismatch repair and base cision repair.	4
14	. Wi	hat are okazaki fragments? Mention the role topoisomerase and DNA gyrase.	4
15	dit	hy DNA fingerprinting is done? State the fference between inducible and repressable eron.	
16		ention the molecular mechanism of lysogenic cle.	:

UG/III/MICB/H/VI/18 (New)

(Turn Over)

17.	Give an example of Base analogue and state its mode of action.	4
18.	Write note on central dogma.	4
19.	Write note on Interrupted mating experiment.	4
20.	Briefly mention about semiconservative	
	replication.	4