National Board of Examinations

Question Paper Name :	DrNB Medical Genetics Paper1
Subject Name :	DrNB Medical Genetics Paper1
Creation Date :	2023-10-15 14:11:36
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Share Answer Key With Delivery Engine :	No
Actual Answer Key :	No

DrNB Medical Genetics Paper1

Group Number :	1
Group Id :	327187789
Group Maximum Duration :	0
Group Minimum Duration :	180
Show Attended Group? :	No
Edit Attended Group? :	No
Group Marks :	100
Is this Group for Examiner? :	No
Examiner permission :	Cant View
Show Progress Bar? :	No

DrNB Medical Genetics Paper1

Section Id :	327187792
Section Number :	1
Section type :	Offline

Mandatory or Optional :	Mandatory
Number of Questions to be attempted :	10
Section Marks :	100
Enable Mark as Answered Mark for Review and Clear Response :	Yes
Maximum Instruction Time :	0
Sub-Section Number :	1
Sub-Section Id :	327187796
Question Shuffling Allowed :	No
Is Section Default? :	null

Question Number : 1 Question Id : 32718718322 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

Please write your answers in the answer booklet within the allotted pages as follows:-

Question Number	Answer to be attempted within	Question Number	Answer to be attempted within
Q. 1	Page 1-5	Q. 6	Page 26-30
Q. 2	Page 6-10	Q. 7	Page 31-35
Q. 3	Page 11-15	Q. 8	Page 36-40
Q. 4	Page 16-20	Q. 9	Page 41-45
Q. 5	Page 21-25	Q. 10	Page 46-50

1. What are Poly ADP-Ribose Polymerase (PARP) and PARP inhibitors (PARPi)? Briefly describe their mechanism of action and clinical uses. [4+6]

Question Number : 2 Question Id : 32718718323 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

a) List various approaches for gene sequencing. [4]

b) Define variants and outline variant discovery workflow. [2+4]

Question Number : 3 Question Id : 32718718324 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

- a) Genomic Databases. [3]
- b) Genomic Browsers. [2]
- c) Write down their clinical applications. [5]

Question Number : 4 Question Id : 32718718325 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

Explain Epigenetics. Describe the mechanism behind Epigenetics and give three examples as applicable to genetic disorders. [3+(4+3)]

Question Number : 5 Question Id : 32718718326 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

Name a database using Artificial Intelligence (AI) for dysmorphological diagnosis. Write down the challenges of using Articial Intelligence (AI) and Machine Learning (ML) in human dysmorphology. [1+9]

Question Number : 6 Question Id : 32718718327 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

Discuss the molecular basis of Down's syndrome. How will you do a rapid diagnosis of Down's syndrome in a prenatal setting? [5+5]

Question Number : 7 Question Id : 32718718328 Question Type : SUBJECTIVE Consider As

Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

What are pseudo-deficiency alleles? Explain with at least two examples. Mention its role in genetic counseling. [4+4+2]

Question Number : 8 Question Id : 32718718329 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

- a) Role of bio-banks for genetic conditions. [5]
- b) Recent guidelines for establishment of bio-banks in India. [5]

Question Number : 9 Question Id : 32718718330 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

a) Homeobox genes and their relevance in human genetic disorders. [5]b) How they are studied? Give examples. [5]

Question Number : 10 Question Id : 32718718331 Question Type : SUBJECTIVE Consider As Subjective : Yes Calculator : None Response Time : N.A Think Time : N.A Minimum Instruction Time : 0

Correct Marks : 10

Discuss the causes of aneuploidy and explain by a diagram the segregation of chromsomes during gamete formation in a balanced reciprocal translocation carrier. [5+5]