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## **BRAINWARE UNIVERSITY**

Term End Examination 2024-2025
Programme – M.Sc.(BT)-2024
Course Name – Genetics and Biostatistics
Course Code - MBT20204
( Semester II )

Full Marks : 60 Time : 2:30 Hours

[The figure in the margin indicates full marks. Candidates are required to give their answers in their own words as far as practicable.]

## Group-A

(Multiple Choice Type Question)

1 x 15=15

- Choose the correct alternative from the following :
- (i) Select the correct criterion to decide whether to use a histogram or a bar chart.
  - a) Use a histogram for discrete data and a bar chart for continuous data
  - c) Both can be used interchangeably
- b) Use a bar chart for discrete data and a histogram for continuous data
- d) Neither is suitable for data visualization
- (ii) Select the most suitable graph you would recommend for visualizing the correlation between two variables.
  - a) Bar chart

b) Histogram

c) Scatter plot

- d) Box plot
- (iii) Write the most appropriate statistical measure you would recommend to determine the middle value of a dataset.
  - a) Mean

b) Median

c) Mode

- d) Variance
- (iv) Write the correct classification for a dataset with a large difference between Q3 and Q1.
  - a) High variance

b) High standard deviation

c) High interquartile range

- d) Low variability
- (v) Select the appropriate method to decide whether a dataset is positively skewed.
  - a) Check if the median is greater than the mean
- b) Check if the mean is greater than the median
- c) Look at the left tail of the distribution
- d) Use a pie chart
- (vi) Classify the different modes of inheritance using pedigree analysis.
  - a) Autosomal dominant, autosomal recessive, X-linked dominant, X-linked recessive
- b) Genetic drift, gene flow, mutation, selection

	c) Monohybrid, dihybrid, polygenic, epistatic	d) Cladogram, phylogram, dendrogram, unrooted tree	
(vii)	Recommend a method to measure gene flow bet	ween populations.	RARY
(viii)	<ul><li>a) Hardy-Weinberg equation</li><li>c) Pedigree analysis</li><li>Decide which concept explains the movement of</li></ul>	d) Sex ratio determination garasat, K	re University olkata -700125
	a) Genetic drift     c) Natural selection     Classify the different inheritance patterns observe	b) Gene flow d) Inbreeding	
	<ul> <li>a) Autosomal dominant, autosomal recessive,</li> <li>X-linked dominant, X-linked recessive</li> <li>c) Cladogram, phylogram, dendrogram,</li> <li>unrooted tree</li> <li>Decide whether correlation implies causation in s</li> </ul>	<ul> <li>b) Mutations, recombination, gene flow, solinked</li> <li>d) Phylogenetic trees, Hardy-Weinberg equilibrium, recombination</li> </ul>	sex-
(xi)	<ul><li>a) Yes</li><li>c) Only if p &lt; 0.05</li><li>Decide which test is most appropriate for analyzing groups.</li></ul>	b) No d) Only for parametric tests ng variance among multiple independent	
	a) t-test	b) ANOVA	
	c) Poisson regression	d) Correlation	
(xii)	Classify correlation coefficients based on the stre		
(xiii	a) Weak, moderate, strong     c) High, low, average     Classify different types of regression models.	b) Small, medium, large d) Positive, negative, neutral	
(xiv	<ul><li>a) Simple, multiple, logistic</li><li>c) Univariate, bivariate, multivariate</li><li>) Label which test is most suitable for analyzing ca</li></ul>	b) Linear, exponential, logarithmic d) All Itegorical survey data.	L. Chea
	a) Chi-square test	b) ANOVA	
	c) Regression	d) t-test	
(xv	) Select the name of the statistical test used for co		
	a) t-test	b) Chi-square test	
	c) ANOVA	d) Correlation analysis	
	Grou	un-B	
	(Short Answer T		3 x 5=15
2. (	Calculate the skewness of a dataset where the mea	an is greater than the median.	(3)
3. Classify different types of gene interactions that influence phenotypic expression.			(3)
4. Classify different types of linear and nonlinear regression models.			(3)
6. (	Ilustrate how error propagation affects experimen Create a box plot for the given dataset: 15, 21, 30, Q1, Q3, and IQR.	tal data analysis. 42, 50, 55, 62, 70, 80. Identify the median,	(3)
	C	DR .	(=)
	Construct a frequency distribution table for the fol 12, 18, 22, 25, 22, 18, 12.	llowing dataset: 8, 12, 15, 12, 18, 20, 15, 15	5, (3)
	Gro	up-C	
		Type Questions)	5 x 6=30

7.	7-2 - State of type is and type in citors in trypotitesis testing and tricil	(5)
	consequences in biomedical research.	
8.	Report the effects of genetic bottlenecks on conservation efforts.	(5)
9.	Construct a frequency distribution table for the following dataset: 5, 10, 10, 15, 15, 20, 20, 25, 30, 35.	(5)
10.	. Illustrate the concept of genetic correlation between two traits with a relevant example.	(5)
	. Classify different types of factorial experiment designs.	(5)
12.	. Illustrate the role of linkage disequilibrium in GWAS studies.	(5)
	OR	-
	Summarize quantitative genetics models in predicting phenotypic traits.	(5)

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