

School of Biomedical Science

Bachelor of Science in Clinical Nutrition and Dietetics Semester End Examination - Jun 2024

Duration: 180 Minutes Max Marks: 100

Sem II - Q1UF203T - Introduction to Clinical Research

General Instructions

Answer to the specific question asked
Draw neat, labelled diagrams wherever necessary
Approved data hand books are allowed subject to verification by the Invigilator

1)	Interpret the objective of Phase I clinical Trial	K1(2)
2)	Interpret the importance of protocol in clinical trial	K2(4)
3)	Distinguish between Bioavailability and Bioequivalence	K2(6)
4)	Discuss the informed consent process.	K3(9)
5)	Discuss different areas of clinical research	K3(9)
6)	Distinguish between randomization and blinding. Discuss the scope	K5(10)
	of randomization and blinding in clinical trail	
7)	Discuss the function of Ethic committee	K4(12)
8)	Analyze and elaborate the importance of source documents in clinical trial	K5(15)
9)	Discuss the bioavailability and bioequivalence studies in detail	K5(15)
10)	Discuss the different areas and scope of Clinical Research.	K6(18)