



AJEENKYA

D Y PATIL UNIVERSITY

End Term Examination (December 2019)

School: SOM

Program: MSc CLINICAL RESEARCH

Course: CLINICAL TRIAL
AND DATA MANAGEMENT

Course Code: MGT684

Semester: 3rd

Max Marks: 50 marks

Duration (mins): 150 mins

Instructions to the students

- 1) **The Question paper consists of two sections- Section I and Section II.**
- 2) **All questions are compulsory.**
- 3) **Neat flow charts or diagrams must be drawn wherever necessary**
- 4) **Figures on the right indicate full marks**

SECTION 1

Q 1) Answer the following:

[2x5=10]

- a) Explain the meaning of project in clinical trial process.
- b) What are the characteristics of project management?
- c) Describe in brief adaptive studies.
- d) Describe – Review of ambulatory or hospital medical records.
- e) What are the various methods for collection of data from secondary sources?

Q 2) Attempt any two of the following:

[5x2=10]

- a) **Explain the phases of clinical trials in detail.**
- b) What are the advantages and disadvantages of secondary data?
- c) What are the roles and responsibilities of a project manager?
- d) Explain in detail various methods used for collection of primary data.

Q 3) Attempt any one of the following:

[5x1=5]

- a) Describe in detail the recruitment and retention of study subjects in clinical trial process.
- b) Explain the various stages involved in project management.

SECTION II

Q 1) Answer the following:

[2x5=10]

- a) What is EMA?
- b) Describe the meaning of manual query writing.
- c) Describe participants as key stakeholders in clinical trial process.
- d) Elaborate CRC.
- e) Explain confidentiality in clinical trial process.

Q 2) Attempt any two of the following:

[5x2=10]

- a) Explain in detail Clinical Trial Agreement (CTA) Process.
- b) What are the roles and responsibilities of clinical research associate (CRA).
- c) Describe in detail about the key stakeholders involved in clinical trial process
- d) Differentiate between CRO and SMO.

Q 3) Attempt any one of the following:

[5x1=5]

- a) Describe Investigator's** outlined in Good Clinical Practices and regulations.
- b) Explain in detail logistics management organization in clinical research.

***** ALL THE BEST *****