



P-MPH203-T

Seat No. _____

M. Pharm. (Sem. II) (CBCS) Examination

July/August – 2018

MPH-203T : Computer Aided Drug Development

Time : 3 Hours]

[Total Marks : 75

Instructions :

- (1) Figure to the right indicates marks.
- (2) Draw neat and clean diagrams as and when required.

1 Answer all the questions, each carry 2 marks : 10×2=20

- (1) What is the role of P-gp ?
- (2) What do you mean by population modelling ?
- (3) What are statistical parameters ?
- (4) Define drug absorption and active transport.
- (5) How parameter sensitivity analysis performed ?
- (6) What do you mean by simulation ?
- (7) Describe innovative uses of computers in R & D.
- (8) How simulation of protein and genes help in drug development ?
- (9) Differentiate descriptive versus mechanistic modelling.
- (10) How drug permeate through intestine ?

2 Answer any 2 out of 3, each carry 10 marks : 2×10=20

- (1) Describe the ICH Q8 guideline in detail.
- (2) Write a note on computer simulation of whole animal and organs.
- (3) Describe in detail clinical data collection and management.

3 Answer any 7 out of 9, each carry 5 marks : 7×5=35

- (1) Describe the computer involvement in pharmaceutical drug development.
- (2) Write a detailed note on optimization parameters.
- (3) Describe in detail artificial intelligence.

- (4) Write a note on IVIVC.
 - (5) Describe advantages and disadvantages of automation in pharmaceuticals.
 - (6) What do you mean by ethics of computing in pharmaceutical research ?
 - (7) Write a detailed note on biowaiver.
 - (8) Describe in detail ASBT.
 - (9) Write a note on BBB-choline transporter models.
-