Rajiv Gandhi University of Health Sciences

First Semester M. Pharm Degree Examination - MAY-2018

[Time: 3 Hours]

Pharmaceutical Validation Q.P. CODE: 5113

Your answers should be specific to the questions asked. Draw neat, labeled diagrams wherever necessary.

LONG ESSAY (Answer any Three)

- 1. Discuss the history and various phases of Drug development and Drug approvals such as IND, NDA and ANDA.
- 2. Elaborate on URS, FAT, SAT and Requalification of a manufacturing equipment.
- 3. Explain the significance of Qualification for analytical instruments? Describe the qualification of FTIR Spectrometer.
- 4. Write in detail the validation of utility systems in Pharmaceutical Industry with emphasis on HVAC system.

SHORT ESSAY (Answer any Nine)

9 X 5 = 45 Marks

- 5. Define Validation and write about Validation Master Plan.
- 6. How is qualification of Disintegration apparatus done?
- 7. Explain the qualification of volumetric flask.
- 8. Write types of sampling methods in Cleaning Validation.
- 9. Write the qualification of Pharmaceutical water system utility.
- 10. How is accuracy and precision determined in Analytical Method Validation?
- 11. Discuss the regulatory aspects of Pharmaceutical drug manufacture.
- 12. Write the salient features of Consumer Protection Act.
- 13. Explain types of Validation in Pharmaceutical Industry.
- 14. Write a note on maintaining status for Calibration, preventive maintenance and change management of manufacturing equipments.

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3 X 10 = 30 Marks

[Max. Marks: 75]