

FirstRanker.com 9204_2010_2 Firstranker!s choice Rajiv Gandhiw Infivensity of Health Sciencesom First Year M.Pharm Degree Examination – October 2010

[Max. Marks: 100] [Time: 3 Hours]

GOOD MANUFACTURING AND LABORATORY PRACTICE

(Revised Scheme 2)

Q.P. CODE: 9204

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

LONG ESSAY (Answer any TWO)

2 X 20 = 40 Marks

- Discuss the organizational set up of Quality Control unit, their responsibilities and various activities of the organization.
- a) What is the design of the air control system adopted in the manufacturing units of 2. antibiotics?
 - b) Discuss the procedure to handle the complaints, its evaluation and rectification.
- a) List out the ICH guidelines to qualify and quantify impurities in new drug substances. 3.
 - b) What are precautions to be taken in handling the cytotoxic substances in the manufacturing unit?

SHORT ESSAY (Answer any FIVE)

5 X 10 = 50 Marks

- What are the measures to be considered before constructing a manufacturing unit? 4.
- List out the Good Laboratory Practice regulations to be followed in caring animals. 5.
- What do you mean by compliance of policy guidelines? 6.
- 7. What is statutory certificate of suitability? Give its significance.
- 8. Discuss the various warehousing procedure for drug products
- 9. Write about validation equipments and documentation procedure to be followed in Industry.

SHORT NOTES 2 X 5 = 10 Marks

- What are the merits and short coming of using computer in maintaining Good Laboratory Practice and good Manufacturing practice?
- International Standards Organization standards 14642-14644

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