

Registration No :

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Total Number of Pages : 01

**M.Pharm.  
M.PH2G.6**

**2<sup>nd</sup> Semester Regular Examination 2017-18  
ADVANCED PHARMACEUTICAL TECHNOLOGY  
BRANCH : PHARMACEUTICAL TECHNOLOGY**

**Time : 3 Hours**

**Max Marks : 70**

**Q.CODE : C725**

**Answer Question No.1 which is compulsory and any five from the rest.**

**The figures in the right hand margin indicate marks.**

**Answer all parts of a question at a place.**

- Q1 Answer the following questions: *Short answer type* : (2 x 10)**
- a) What is fluidization technique?
  - b) What is HEPA filter? How to determine its efficiency?
  - c) What is film coating? Give examples of aqueous coating materials.
  - d) What is metered dose inhaler?
  - e) Double cone blender is suitable equipment for mixing of lubricants with dried granules. True or false: Justify.
  - f) What is retrospective validation?
  - g) Name the statistical designs used in optimization of formulations.
  - h) Name the propellants used in development of aerosol dosage form.
  - i) Name the equipments used in manufacturing of suspensions and emulsions.
  - j) Name different type of dryers used in drying of granules.
- Q2 a) Write in detail about recent developments in diluents used in the formulation of tablets. (5)**
- b) Discuss in detail about physics of tablet compression. (5)**
- Q3 Write in detail about improvements in equipments used for granulation technology. (10)**
- Q4 a) What is scale up of in formulation? Discuss briefly about effect of scale up on process parameters like mixing and granulation. (5)**
- b) Discuss critically about validation of solid dosage forms. (5)**
- Q5 Write in detail about advances in formulation and manufacturing of aerosols. (10)**
- Q6 Write about advances in materials and production techniques for parenteral dosage forms. (10)**
- Q7 a) Write in about application of statistical techniques in product development. (5)**
- b) Discuss about validation of analytical methods. (5)**
- Q8 Write short notes on the following : (2.5 x 4)**
- a) Computerization for in process quality control of tablets
  - b) Parts of tablet compression press
  - c) Microbiological testing of water
  - d) Types of validation