

THE TAMIL NADU DR. M.G.R. MEDICAL UNIVERSITY

[LO 987]

MAY 2019

Sub. Code: 2987

**M.PHARM. DEGREE EXAMINATION**  
**(PCI New regulations 2016)**  
**SEMESTER-II**  
**BRANCH-VI – PHARMACOLOGY – MPL**  
**PAPER III – PRINCIPLES OF DRUG DISCOVERY**

*Q.P. Code : 262987*

**Time : Three hours**

**Maximum : 75 Marks**

**I. Elaborate on:**

**(2 x 20 = 40)**

1. a) What are the different statistical methods applied to validate the developed QSAR model? Explain PLS model.  
b) Differentiate SAR & QSAR.
2. Explain the following:
  - a) One Target validation techniques.
  - b) Anti sense technologies.
  - c) Briefly explain the cost involvement and feasibility of new drug discovery in India.

**II. Write notes on:**

**(7 x 5 = 35)**

1. What are micro arrays? Write its importance in target discovery.
2. What are the parameters to be considered in lead optimization procedure?
3. What is meant by:
  - a)  $\alpha$ Helix
  - b) Zinc finger proteins
  - c) Pharmacophore
  - d) Scaffold
  - e) Nucleotide
4. Write a note on rationale drug design.
5. Name the programs used in docking process. What is meant by G score?
6. Why solubility has been considered as one of the important parameters in drug development? Explain.
7. Explain the term COMFA & COMSIA.

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