

**UKA TARSADIA UNIVERSITY**  
**M. Pharm. (Pharmaceutical Analysis) (3<sup>rd</sup> Semester)**  
**040060302: Pharmaceutical and Cosmetic Analysis**

**Duration: 3 hours**

**Maximum marks: 70**

**Instructions:**

1. Attempt all questions
2. Write each section in a separate answer book.
3. Make suitable assumptions wherever necessary.
4. Figures to the right indicate full marks allocated to that question.
5. Draw diagrams/figures wherever necessary.

**SECTION-1**

**Q.1] (a) Answer the following: [7 x 1 = 7]**

- 1) What is photolysis?
- 2) What do you mean by stabilizers?
- 3) What is biological sample?
- 4) Enlist biological matrices.
- 5) What is hypersensitivity?
- 6) What do you mean by repeated insult irritant?
- 7) What is polymorphic transformation?

**(b) Attempt any four: [4 x 2 = 8]**

- 1) What is mean kinetic temperature?
- 2) How is selectivity determined in bioanalytical method development?
- 3) What do you mean by oxidative autocatalysis?
- 4) What is an open patch test? How is it performed?
- 5) What do you mean by long term stability?
- 6) Enlist the preservatives commonly used in pharmaceuticals. Explain assay principle of any one preservative.

**Q.2] (a) Discuss physical stability testing of tablets. [5]**

**OR**

**(a) Describe principle, procedure, advantages and disadvantages of liquid-liquid extraction. [5]**

**(b) The interaction of moisture with the drug substance and excipients can significantly affect the physical stability of the final drug product. Explain giving suitable examples. [5]**

**OR**

**(b) Name the pathways of chemical degradation. Enumerate the factors affecting chemical stability of drugs. Discuss the significance of hydrolytic degradation study of drugs. [5]**

**Q.3] Attempt any two [2 x 5 = 10]**

- (a) What is the significance of skin irritation test? How is it performed?
- (b) Describe US-FDA guidelines for bioanalytical method development.
- (c) Discuss physical stability testing of suspensions.

## SECTION-2

**Q.4] (a) Answer the following: [7 x 1 = 7]**

- 1) What is a Gross sample?
- 2) What do you mean by Acceptable Quality Level (AQL)?
- 3) Define: Iodine value.
- 4) Name the methods for determination of moisture and volatile matter.
- 5) What is GRAS list?
- 6) Enlist the types of cosmetic pencils.
- 7) What do you mean by softening point of lipstick?

**(b) Attempt any four: [4 x 2 = 8]**

- 1) Write classification of cosmetics.
- 2) Explain the principle of any one method for determination of rosin.
- 3) How will you prepare composite samples for liquid soaps?
- 4) Describe packing and marking requirements for skin powder for infants.
- 5) How will you perform the test for stability of smell for after shave lotions?
- 6) Enlist the requirements for shaving creams as per BIS specifications.

**Q.5] (a) Describe the principle and procedure involved in determination of foaming power. Name the cosmetics for which determination of foaming power is required. [5]**

**OR**

**(a) Name the types of shampoos. Enlist the ideal properties of shampoos. Enumerate the requirements for both types of shampoos. [5]**

**(b) Describe the principle and procedure involved in determination of combined alkali and total anhydrous soap. [5]**

**OR**

**(b) Describe the requirements for tooth paste as per BIS specification. [5]**

**Q.6] Attempt any two [2 x 5 = 10]**

- (a) Name the types of hair oils. Describe the requirements for all types of hair oils as per BIS specification.**
- (b) Describe the scale of sampling for henna powder. Enumerate the requirements for henna powder as per Indian standard specification.**
- (c) Discuss the quality control of raw materials used in cosmetic industry.**