REV-01 BPH/02/05

B. PHARM. SIXTH SEMESTER QUALITY ASSURANCE BP606T [REPEAT]
[USE OMR SHEET FOR OBJECTIVE PART] 2023/06



Marks: 20

 $1 \times 20 = 20$ 

Full Marks: 75

Duration: 3 hrs.

(PART-A: Objective)

Time: 30 min.	

Choose the correct answer from the following:

- GMP consists of how many components?
  - a. 5 c. 4

- b. 6 d. 3
- The 'S' in QSEM stands for:
  - a. Stability

b. Safety

- c. Sanitization

- d. None of these
- QA is a \_\_\_
- a. managerial

b. Corrective d. None of these

- c. safety
- QC is oriented.
- a. process

b. product

- c. provision
- The Q<sub>7</sub> guidelines describes about: a. Quality risk management

b. Pharmacopoeia

d. performance

- d. Impurities c. GMP
- The long term testing conditions for temperate zones is:

a. 25°C, 60% RHc. 30°C, 45% RH

- b. 21°C, 45% RH d. None of these
- In powdered glass test which indicator is used?
  - a. Phenolphthalein
- b. Methyl orange

c. Methyl red

- d. None of these
- The water attack test is used for which type of glass?
  - a. Type I

b. Type II

c. Both a and b

- d. None of these
- The arsenic test is conducted for glass containers intended for what purpose?
  - a. All aqueous products
- b. Oral dosage forms
- c. Aqueous parenteral products
- d. None of these

10.	The self-sealability test is an important test products?  a. All parenteral  c. Multi-dose parenteral	<ul><li>performed for closures of which</li><li>b. Single-dose parenteral</li><li>d. All of these</li></ul>	h type of
11.	<ul> <li>What is the full form of OECD?</li> <li>a. Organization for Economic Control &amp; Drugs</li> <li>c. Organization for Economic Cooperation &amp; Development</li> </ul>	<ul><li>b. Organization for Excise Co Duty</li><li>d. None of the above</li></ul>	ontrol &
12.	GLP is designated as a new part of Chapter a. 21 CFR Part 58 c. 22 CFR Part 58	<ul><li>21 of the CFR as:</li><li>b. 21 CFR Part 56</li><li>d. 22 CFR Part 56</li></ul>	
13.	Communal wastes include:  a. Cardboard  c. Needles	b. Tissue d. None of these	
14.	Incineration method refers to a:  a. Sterilization method  c. Both a and b	<ul><li>b. Waste disposal method</li><li>d. None of these</li></ul>	
15.	<ul><li>E . For chemical disinfection which compou</li><li>a. Chlorine compounds</li><li>c. Both a and b</li></ul>		
16.	SOP should be written in:  a. active voice, present tense c. passive voice, present tense	<ul><li>b. active voice, past tense</li><li>d. None of these</li></ul>	
17.	a. Internal Audit c. Regulatory Audit	udit. b. External Audit d. All of the above	
18.	are the specifications a manufa equipment.  a. Installation qualification  c. Performance qualification	<ul><li>cturer uses to describe a device</li><li>b. Design qualification</li><li>d. Operational qualification</li></ul>	or
19.	The format of Validation Master Plan should a. Documentation format c. Both a and b		
	GLP is a formal regulation created by USFD on:		roposed
	<ul><li>a. Nov 20, 1978</li><li>c. Nov 22, 1976</li></ul>	<ul><li>b. Nov 19, 1976</li><li>d. Nov 21, 1978</li></ul>	
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## PART-B: Descriptive

Tin	ne: 2 hrs. 30 min.	ks: 35
	[ Answer any seven (7) questions ]	
1.	Describe about the design and construction of Premises.	
2.	Explain about the objectives and elements of TQM.	5
3.	Explain the various tests for plastic containers.	5
4.	Mention the various steps for handling and evaluation of complaints.	5
5.	Write short notes on the following: SOPs Quality Audits	5
6.	Explain the various tests for paper and boards as secondary packaging materials.	5
7.	Describe about the ICH guidelines.	5
8.	Write a detailed note on Plant Layout.	5
9.	What are the various measures taken in industries for the control of contamination?	5
	[Answer any two (2) questions]	
1.	Write a detailed note on the various categories of GMP.	10
2.	Describe the various GLP Practices.	10
3.	Write in details about the handling of waste materials and their	10