

Fourth Prof. A/2015
Examination: Doctor of Pharmacy (Pharm.D.)

Roll No. .....

Subject: Pharmaceutics-VI (Industrial Pharmacy)

PAPER: 3

TIME ALLOWED: 3 hrs. MAX. MARKS: 100

## NOTE: Attempt any FIVE questions. All Questions carry equal marks.

- Q.1. a) What is filtration. Write down the types of filtration used in pharmaceutical industry.(10)
  - b) What are defects of tablets? (10)
- Q.2. a) Define the milling and elaborate the types of milling? (8)
  - b) Write down a not on Hammer mill? (6)
  - c) What is Ball milling? (6)
- Q.3. a) What are semisolid preparations and write note their packaging techniques? (10)
  - b) What are suspending agents? (5)
  - c) Write a short note on ophthalmic preparations? (5)
- Q.4. a) What are emulsions and there types? (8)
  - b) Write down methods for preparations of emulsions? (6)
  - c) What are suspending agent and give few examples? (6)
- Q.5 a). What are Excipients of tablets? (8)
  - b). Define mixing. (6)
  - c). which instruments are used for solid mixing? (6)
- Q.6. a) What are different methods of tablet preparations, describe dry granulation method? (12)
  - b) Discuss spray drying and freeze drying. (8)
- Q.7. Write note on
  - a) Rotary dryers (5)
  - b) Quality assurance (5)
  - c) Micro-Emulsions (5)
  - d) Compartment drying (5)



Fourth Prof. A/2016

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Subject: Pharmaceutics-VI (Industrial Pharmacy)

PAPER: 3

TIME ALLOWED: 3 hrs.

MAX. MARKS: 100

## NOTE: Attempt any FIVE questions. All Questions carry equal marks.

### Question 1:

- (a) Define Hazard, Safety and Risk. Discuss different detailed aspects of chemical hazards? (10)
- (b) Write a detailed note on the Fire as an industrial hazard and explain the detection and prevention of fire in an industry. (10)

Question 2: Define Good Manufacturing Practices (GMP) and Current Good Manufacturing Practices (cGMP) and discuss the significance and different aspects of GMP for Finished pharmaceutical products (FPP). (20)

### Question.3:

- (a) Discuss in detail the principal and working of Ball mill and colloidal mill. (10)
- (b) Briefly discuss spray drying technique and its application in detail? (10)

### Question.4:

- (a) Describe the aqueous solvent systems used for the parenteral preparations (12)
- (b) Describe the method of production for a terminally sterilized formulation. (8)

### Question: 5

- (a) Discuss the methods used for ampules sealing? (6)
- (b) What is clean-in-place concept? How equipment are cleaned for sterile manufacturing? (74)

Question: 6: write a short note on four (04) of the followings where each part contains equal (05) marks.

- 1- Filter press
- 2- Clarification
- 3- Mechanism of freeze drying
- 4- Mechanism of milling
- 5- Mass transfer

### Question: 7:

- (a) Discuss the problems associated with the manufacturing of tablets and suggest the solutions to overcome these problems
- (b) Define angle of Repose and write in detail the method of its measurement (10)



Fourth Prof. 2nd A/2016 Examination: Doctor of Pharmacy (Pharm.D.)

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Subject: Pharmaceutics-VI (Industrial Pharmacy)

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MAX. MARKS: 100

## NOTE: Attempt any FIVE questions. All Questions carry equal marks.

### Ouestion 1:

- (a)- What are different theories of drying? Write down the classification of dryers? (10)
- (b)- Discuss emulsions and their stability? Write down mechanical equipments used for emulsions preparation? (10)

### **Ouestion 2:**

Define quality assurance. Write detailed note on Good Manufacturing Practices and Current Good Manufacturing Practices used in pharmaceutical industry? (20)

### Question 3:

- (a)- Discuss in detail the principal and working of cutter mill and fluid energy mill? (10)
- (b)- Define mixing. What are the reasons for size reduction and factors affecting size reduction?(10)

## Question 4:

- (a)- Define sterile area. Write detailed note on Inprocess Quality Control of parenterals? (10)
- (b)- What is filter media? Discuss leaf filter and rotary filter in detail. (10)

### Ouestion 5:

- (a)- What is solid-air interface and angle of repose? (6)
- (b)- Discuss in detail the tablet coating, and problems involved in tableting? (12)

Question 6: Write a short note on 04 of the followings. All parts carry equal (05) marks. (20)

- 1- Heat transfer
- Evaporation under reduced pressure
- 3- Granulation
- 4- Inflammable gases and dusts
- 5- Size analysis and sieving.

### Question 7:

- (a)- Define packaging. Discuss packaging area and influence of packaging materials on pharmaceuticals' products? (10)
- (b)- How suspensions are formulated? And which type of equipments are employed for their preparation? (10)

Fourth Prof: A/2017 Examination: Doctor of Pharmacy (Pharm.D.) Roll No. .....

Subject: Pharmaceutics-VI (Industrial Pharmacy)

PAPER: 3

TIME ALLOWED: 3 hrs. MAX. MARKS: 100

## NOTE: Attempt any FIVE questions. All Questions carry equal marks.

### Question 1:

- (a) Briefly discuss the different types of hazards and their impact on the life of employees working in a Pharmaceutical manufacturing unit.
- (b) Write a detailed note on the fire hazards with reference to identification and prevention fire hazards in the pharmaceutical industry.

### Question 2:

Discuss in detail the aspects of Good Manufacturing Practices (GMP) and its applications for producing quality Finished pharmaceutical products (FPP) in an industry.

### Question.3:

- (a)- Discuss the following four only (05 marks each)
- 1- Heat transfer
- 2 Mass transfer
- 3- Types of dryers
- 4- Pin Mill
- Q.4: (a) Discuss the details of manual size reduction method and write the major differences between manual and mechanical size reduction
- (b) Define the different terms used to explain the filtration process (08).
- Q. 5. (a) Discuss the principle, working and advantages of the colloid mill, fland mill and fluid energy mill.
- (b). Discuss the different methods of size reduction and with the help of examples.
- Q. 6 (a) Define compression and consolidation and describe phenomenon of Angle of Repose (10)
- (b) -Give the classification of various industrial dryers and discuss the details of fluidized bed dryer. (10).
  - Q. 7. (a) Discuss the equipment used in the preparation of semisolid dosage forms (10)
    - (b) Discuss the formulation aspects of emulsions in a Pharmaceutical industry (10)



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TIME ALLOWED: 3 hrs. MAX. MARKS: 100

## NOTE: Attempt any FIVE questions. All Questions carry equal marks.

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Question	1	٠
Ancarion		

- (a) Define Hazard, Give the different classes of hazards and discuss in detail the different aspects of fire hazards?
  (10)
- (b)- Write a detailed note on the chemicals as an industrial hazard and explain the prevention of chemical hazards in industry. (10)

### Question 2:

Define Good Manufacturing Practices (GMP) and Current Good Manufacturing Practices (cGMP) and discuss the significance and different aspects of GMP for Finished pharmaceutical products (FPP). (20)

### Question.3:

(a)- Discuss in detail the types of heat transfer mechanism and its applications

(10)

(b)- Briefly discuss the principles and mechanism of mass transfer and its applications in pharmaceutical industrial? (10)

### Question.4:

- (a) Describe the ideal characteristics of sterile products and explain the requirements for the preparation of parenteral preparations in pharmaceutical industry (12)
- (b) Describe the method of production for sterilized products in pharmaceutical industry (8
- Q.5 (a) What is comminution? Discuss the advantages, disadvantages and the factors affecting the particle size reduction (10)
- (b) Define filtration, clarification, sedimentation and decantation and also discuss the mechanism of filtration with reference to Darcy's Law.
- Q.6 a) Define & discuss the properties of an ideal filter media and also discuss the different factors affecting the selection of filter media. (06)
  - b) Write a detail note on the different types of filter media used in an industry.

(14)

### Q.7. Write note on four of the following (5 marks each)

- a) Compression and Compaction
- b) Angle of Repose
- c) Ball mill
- d) Hussner's ratios
- e) Problems of tablets manufacturing



# Fourth Prof: Annual – 2018 Examination: Doctor of Pharmacy (Pharm.D.)

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Subject: Pharmaceutics-IV (Industrial Pharmacy) (New Course) PAPER: 3 (Part – II)

TIME ALLOWED: 2 Hrs. & 30 min. MAX. MARKS: 80

# Attempt this Paper on Separate Answer Sheet provided. Attempt any 4 questions. Each question carry equal marks.

		1944 (1946)	
Q 2	a)	Describe theory of heat transfer, what are the application	10
	15.1	of heat transfer in pharmaceutical industry?	
	b)	Give classification of dryers used in pharma industry,	
		discuss Fluidized bed dryer in detail.	10
Q3	a)	Why powder volume is difficult to measure? What is the	10
		solution then? How the porosity affects the tablets	
		properties?	
	b)	Describe some approaches to avoid chemical hazards in	10
		pharmaceutical industry.	
Q4	a)	How sterility is maintained in the clean room of a	10
		pharmaceutical industry?	20
	b)	Why granulation is important? Name its various types and	10
	1112-181	describe the steps in wet granulation process.	
		the steps in wee grantatation process.	
Q 5		Give brief answers of following.	C. Etherales
			5 Marks
		The state of the s	each
		a tubici manufacturing	
		with reasons and remedy to avoid.	
		iii. Importance of HVAC system in pharmaceutical	
		industry.	
		<ol> <li>Different tablet coating techniques</li> </ol>	
1000100			
0,6	a)	What are the different mechanism for mechanical size	10
		reduction? Discuss in construction and operation of	
		Planetary Ball mill.	
	b)	List down different characteristics of a packaging materials	10
		and describe the reasons why glass is preferred over the	
		plastics for the packaging of injectable.	
Q 7	a)	Discuss the mechanism of liquids/liquids mixing and	10
		describe the construction and application of silver son	10
		homogenizer	
	b)	Can we dry solution/suspension? How? Describe the	10
		principal of such methods	10
		principal of such methods	



Fourth Prof: Annual – 2018

Examination: Doctor of Pharmacy (Pharm.D.)

Subject: Pharmaceutics-IV (Industrial Pharmacy) (New Course)

PAPER: 3 Part - I (Compulsory)

TIME ALLOWED: 30 min. MAX. MARKS: 20

### Attempt this Paper on this Question Sheet only.

Please encircle the correct statement. Each MCQ carries 1 Mark. This Paper will be collected back after expiry of time limit mentioned above.

Q.1 MCQ's (Select the most appropriate option)

- Slicing and cutting principles are used for the size reduction of -----type of drugs
  - A) Animal origin
  - B) Mineral origin
  - C) Vegetable origin
  - D) Synthetic drugs
- 2. The porcelain mortar and pestle, used for the small scale bruising process, is not suitable for the-----
  - A) Tannins
  - B) Acidic drug
  - C) Basic drugs
  - D) None of the above
- 3. ----- is a process in which coarse emulsion is converted into a colloidal state of uniform composition by reducing particle size.
  - A) Homogenization
  - B) Sedimentation
  - C) Grinding
  - D) Both A and B are correct options
- 4. Rate of filtration is ----- to the viscosity of liquid to be filtered.
  - A) Inversely proportional
  - B) Directly proportional
  - C) None of them
  - D) Equal to
- 5. Which is one of the most common granulating agent?
  - A) Acacia
  - B) Gelatin
  - C) Tragacanth
  - D) Starch

B) Effervescent tablets
C) Dispensing tablets
D) Hypodermic tablets
<ul> <li>7. Randomization of dissimilar particles within a system is called</li></ul>
D) Drying
8. The method most widely used for measuring particle size distribution in pharmaceutical industry is?
A) Sieving B) Microscopy C) Sedimentation D) Laser diffraction
9. Tablet coating adds an approximate increase in weight of A) 3-5 % B) 1-2% C) 2-5% D) 10-20%
10. In coating of tablets pan speeds for non-aqueous film coating is  A) 10-15 rmp  B) 3-10 rmp  C) 40 rmp  D) 100 rmp
11. During coating when atomization is very fine it may lead to effect called  A) Orange peel  B) Bridging  C) Spray drying  D) Capping
<ul> <li>12. If average weight of one tablet is 250 mg what would be the weight of API batch for 1 lac tablets <ul> <li>A) 2500 kg</li> <li>B) 25 kg</li> <li>C) 250 kg</li> <li>D) 250000 kg</li> </ul> </li> <li>13. Spray dried lactose is prone to in presence of excess moisture.</li> </ul>
A) Darkening

6. Which tablets are designed to produce solution rapidly?

A) Implantation tablets

B) OxidationC) HydrationD) Hydro-oxidation

14. Percentage of binders used in tablet formulation	
A) 5-10 %	
B) 10 - 20%	
C) 20 - 30%	
D) 0.01 - 1%	
15. In sugar coating, material is used for polishing	
A) Carnauba wax	
B) Cocoa powder	
C) Oleic acid	
D) Gelatin	
16. Glass apparatus used in the production of sterile products can be sterilized	
16. Glass apparatus used in the production of sterile products can be sterilized by	
A) Dry heat 140 degrees for 30 minutes	
B) Dry heat 120 degrees for 1 hour	
C) Saturated steam 121 degrees for 15 minutes	
D) Using Suitable Disinfectant	
17is used as biological indicator in radiation sterilization.	
A) Bacillus subtilis	
B) Clostridium sporogenes	177
C) Bacillus pumilis	
D) Bacillus stereo-thermophilous	
18. Mesh size is the number of opening per	
A) Inch	
B) Square inch	
C) Linear inch	
D) Centimeter	
19. Empty capsule has moisture content in the range of	
A) 5%-10%	
B) 12%-15 %	
C) 30%-45%	
D) 50%-70%	
20. If average weight of one tablet is 250 mg what would be the weight of API	
batch for 1 lac tablets	
A) 2500 kg	
B) 25 kg	
C) 250 kg	
111 Z50000 kg	



Fourth Prof: 2<sup>nd</sup> Annual - 2018 Examination: Doctor of Pharmacy (Pharm.D.)

Subject: Pharmaceutics-IV (Industrial Pharmacy) (New Course)

MAX. TIME: 2 Hrs. 30 Min.

PAPER: 3 Part - II MAX. MARKS: 80

## ATTEMPT THIS (SUBJECTIVE) ON THE SEPARATE ANSWER SHEET PROVIDED

Note: Attempt any four questions, all questions carry equal marks. a) Describe theory of mass transfer, what are the application 10 of mass transfer in pharmaceutical industry? b) Discuss principle, operation and application of Tray dryer in 10 detail. 10 03 a) Define compression, consolidation and compaction, discuss their mechanisms in detail. 10 b) Name the methods of tablet manufacturing, make flow chart to describe these, how the choice is made between a) How you will differentiate the sterile area of a 10 04 pharmaceutical industry into different classes? What type of operation are carried out in each area? b) Milling of APIs affect their flow, mixing and compressional 10 properties, discuss this argument with reasons. 5 Marks Q5 Give brief answers of following terms. Steps in the manufacturing of hard gel capsules each ii. Sugar coating of tablets Picking and mottling with reasons and remedy to avoid. Bulk and terminal sterilization of injectables a) Discuss the principle, working and applications of colloid 5+5 Q6 mill and fluid energy mill. b) Discuss different types of material used to prepare the 10 containers for pharmaceutical packaging. a) Describe different mechanism of milling with suitable 10 07 examples, which type of milling instrument is suitable for the size reduction of waxy materials 10 b) Write a note on different types of impeller Mixers used for liquid mixing?

PAPER: 3

## UNIVERSITY OF THE PUNJAB

Fourth Prof: 2<sup>nd</sup> Annual – 2018 Examination: Doctor of Pharmacy (Pharm.D.)

`\	Roll	No.	in	Words.	

Roll No. in Fig. .....

Subject: Pharmaceutics-IV (Industrial Pharmacy)

Part - I (Compulsory)

MAX. TIME: 30 Min.

MAX. MARKS: 20

Signature of Supdt.:

Attempt this Paper on this Question Sheet only.

Please encircle the correct option. Division of marks is given in front of each question.

This Paper will be collected back after expiry of time limit mentioned above.

(New Course)

## Q.1. Encircle the right answer cutting and overwriting is not allowed.

(1x20=20)

- Mechanical process of size reduction of solid substances is called as
  - A) Levigation
  - B) Pulverization
  - C) Milling
  - D) Trituration
- 2. The ball mill consists of a horizontally rotating hollow vessel of cylindrical shape with-----
  - A) The length equal to its diameter
  - B) The length slightly greater than its diameter
  - C) The length slightly shorter than its diameter
  - D) The diameter is much shorter
- 3. The size reduction process a pin mill follows the mechanism of
  - A) Cutting
  - B) Compression
  - C) Grinding
  - D) Impact and attrition
- 4. Solid collected on filter medium is the desired product then the process is known as
  - A) Cake filtration ·
  - B) Sedimentation
  - C) Filter aids
  - D) Filter press
  - The solids retained on a filter during filtration process is known as------.
    - A) Filtrate
    - B) Mixture
    - C) Residue
    - D) Filter Aid
  - 6. Clarification is applied when the solids do not exceed-----percent.
    - A) 2.0%
    - B) 1.0%
    - C) 3.0%
    - D) 10%

7.	Which t	est is useful for tes	sting membra	ane efficiency?		
	15	A) Diffusion testin	g			
		B) Forward Flow to	T10			
		C) Bubble point te				
		D) Integrity Testin				
8.	Mesh s	creen of basket rac	ck assembly o	of disintegration a	apparatus is	
	A)	15 mesh screen				
	B)	10 mesh screen				
		20 mesh screen				
	D)	25 mesh screen				
9.	Cappin	g of tablets during	manufacturi	ng occurs due to-		
		Air entrapment				
	100	Plastic deformation				
		Improper adjustme		off blades		
		All of these option				
10.				ects of shock & ab	rasion is evaluated	
	by usin	g				
	0.0	Hardness tester				
	110000	Disintegration test	apparatus			
	,	Friabilator				
		Screw gauge		72		
11.	Which	of the following is	not added in	lozenges?		
	A)	Binder				
	B)	Disintegrant				
	C)	Sweetener				
	D)	All				
12.	Sub co	ating is given to th	e tablets dur	ing sugar coating		
		To Increase the bu				
		To avoid deteriors				
		To prevent the sol		lic medium		
	D)	To avoid stickiness	S			
13.		egrity of HEPA filte		ecked by	test.	
	1777	TOT aerosol challe				
		DOP aerosol challe	enge			
	500	Killer-Killani				
	D)	Fehling's				
14		the following are fi	lter-aids exce	ept		
	7.00	Diatomite				
		Carbon				
		Cellulose		_	D-w- 1	062
	D)	Polyvinyle chloride	e		Page 2	01 3

15. The	non	-aqueous vehicle used in injections is	
A)	Alco	phol	
B)	Gly	cerol	
C)	Alc	pholic sodium stearate	
D)	Veg	etable oils	
10.50		nules high angle of repose indicates	,
16 10		High porosity	
		High bulk density	
		Smooth surface	
	D)	Rough surface	
17 su	perdi	sintegrant are essential component of	
		200	
	A)	Buccal Tablets	
	B)	Gastroretentive drug delivery system	
		Lozenges	
		Fast dissolving tablets	
18. G	ass a	pparatus used in the production of sterile produc	cts can be sterilized
by			
2.50	A)	Dry heat 140 degrees for 30 minutes	
	BI	Dry heat 120 degrees for 1 hour	
	C)	Saturated steam 121 degrees for 15 minutes	
	D)	Using Suitable Disinfectant	
10 1/	thich	statement is NOT correct?	
19. V	Λì	Buccal route avoids first pass metabolism	
	(2)	Parenteral route avoids first pass metabolism	
	C)	Sublingual route avoids first pass metabolism	
	(0)	Oral route avoids first pass metabolism	
20.1	n par	coating technique , the core material is	
	A)	Solid	
		Liquid	
		Gas	
		Both A and B are correct	



UNIVERSITY OF THE PUNJAB Fourth Prof: 2<sup>nd</sup> Annual - 2018 Examination: Doctor of Pharmacy (Pharm.D.)

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Subject: Pharmaceutics-VI (Industrial Pharmacy) (Old Course) PAPER: 3

MAX. TIME: 3 Hrs. MAX. MARKS: 100

## NOTE: Attempt any FIVE questions. All questions carry equal marks.

		All all and the language and the languag	
Q		Describe theory of mass transfer, what are the application of mass transfer in pharmaceutical industry?	10
	b	dryer in detail.	10
Q 2	a	What is the difference b/w manual and mechanical size reduction? Write a detail note on manually used method.	10
	b)	What is unit pack? Discuss the blister packaging and material used to prepare it.	10
Q 3		their mechanisms in detail.	10
	b)	chart to describe these, how the choice is made between these methods?	10
Q 4	a)	How you will differentiate the sterile area of a pharmaceutical industry into different classes? What type of operation are carried out in each area?	10
	b)	Milling of APis affect their flow, mixing and compressional properties, discuss this argument with reasons.	10
Q.5		Give brief answers of following terms.  i. Steps in the manufacturing of hard gel capsules ii. Sugar coating of tablets iii. Picking and mottling with reasons and remedy to avoid.	5 Marks each
		iv. Bulk and terminal sterilization of injectables	
Q 6	a)	Discuss the principle, working and applications of colloid mill and fluid energy mill.	5+5
	b)	Discuss different types of material used to prepare the containers for pharmaceutical packaging.	10
Q7	a)	mixing, describe Silverson mixer in detail	
	D	Describe some approaches to avoid mechanical hazards in pharmaceutical industry?	10

Doctor of Pharmacy (Pharm.D.) Fourth Prof: Annual-2019 Roll No. in Words. .....

Subject: Pharmaceutics-IV (Industrial Pharmacy) (New Course)

Paper: 3 Part - I (Compulsory) Time: 30 Min. Marks: 20

HEET ONLY. Signature of Supdt.:

` Roll No. in Fig. .....

## ATTEMPT THIS PAPER ON THIS QUESTION SHEET ONLY.

Division of marks is given in front of each question.

This Paper will be collected back after expiry of time limit mentioned above.

Q.1.	Enci	rcle the correct option.		(20x1=20)
i.	Mech a) c)	nanical process of size re Levigation Milling	duction of solid b) d)	d substances is called as: Pulverization Trituration
ii.	Most a) c)	commonly used disinteg Calcium phosphate Mannitol	rant in tablet de b) d)	osage form is: Lactose Starch
iii.	The s a) c)	size reduction process in Cutting Grinding	a pin mill follov b) d)	vs the mechanism of Impact Impact and attrition
iv.	Solid a) c)	collected on filter mediun Cake Filter aids	n is the desired b) d	d product then the process is known as Sedimentation Filter press
V.	Which a) c)	n is one of the most comr Acacia Tragacanth	non granulatin b) c)	g agent? Gelatin Starch
vi.	Which a) c)	tablets are designed to Implantation tablets Dispensing tablets	produce solution b) d)	on rapidly? Effervescent tablets Hypodermic tablets
/ii.	Rando a) c)	omization of dissimilar pa Milling Mixing	rticles within a b) d)	
/iii.	The m pharm a) c)	nethod most widely used to acceutical industry is? Sieving Sedimentation	for measuring b) d)	particle size distribution in  Microscopy Laser diffraction
Χ.	Which a) c)	term is used for separati Capping Picking	ion of a tablet i b) d)	nto two or more distinct layers?  Lamination  Sticking
i.	a)	is the problem when ma Hopper Dyes		9
d.	a)	coating when atomizatio Orange peel Spray drying		t may lead to effect called Bridging Capping

xii.	If aver	rage weight of one tablet is 25 olets:	0 mg what	would be the weight of API batch for 1
	a) c)	2500 kg 250 kg	b) d)	25 kg 250000 kg
xiii.	a) c)	fficiency of HEPA filter in remo 99.0% 99.99%	d)	99.97%
xiv.	a)	entage of binders used in table 5 - 10 % 20 - 30%	d)	0.01 - 1%
xv.	In sug a) c)	gar coating, Carnauba wax Oleic acid	d)	Gelatin
xvi.	a) c)	ranules high angle of repose i High porosity Smooth surface	d)	Rough surface
xvii.	a)	non-aqueous vehicle used in i Alcohol Alcoholic sodium stearate	d)	Vegetable oils
xviii.	Glas a) b) c) d)	Dry heat 140 degrees for 30 Dry heat 120 degrees for 1 Saturated steam 121 degree Using Suitable Disinfectant	hour es for 15	erile products can be sterilized by minutes
xix.	a) b) c)	ch statement is NOT correct?  Buccal route avoids first pa Parenteral route avoids first Sublingual route avoids first Oral route avoids first pass	st pass me st pass me s metabolis	etabolism sm
XX.	Dur a) c)	ring sugar coating, following co Sub-coating Syrup coating	pating prev b) d)	vent the moisture penetration Seal-coating Polishing

Doctor of Pharmacy (Pharm.D.) Fourth Prof: Annual-2019

Roll	No.					
		-				

Subject: Pharmaceutics-IV (Industrial Pharmacy) (New Course)

Paper: 3 Part - II

Time: 2 Hrs. 30 Min. Marks: 80

## ATTEMPT THIS (SUBJECTIVE) ON THE SEPARATE ANSWER SHEET PROVIDED

Note: Attempt any FOUR questions. Each question carries equal marks.

Q2	a)	What are different mechanisms of heat transfer, Describe the factors affecting the drying process.	10
	b)	Discuss principle, operation and application of fluid bed	
	٠,	dryer in detail.	10
Q 3	a)	How solid state properties affect the Tablet manufacturing process? How these are settled then?	10
	b)	Name different components of tablet formulation, describe different stages of granulation process.	10
Q4	a)	Describe the classification of clean room of a	10
		pharmaceutical industry. How laminar air flow helps in	
		maintaining the sterility of clean room?	
	b)	What are the components of sterile products? How	10
		sterilization of such products is achieved?	
Q5		Give brief answers of following.	5 Marks
		Cive bile: dilaweis or romoving.	J IVIGI KS
		i. Filter Aid	each
		i. Filter Aid	
		Filter Aid     Steps in manufacturing of Hard gelatin capsule	
Q6	a)	<ul> <li>i. Filter Aid</li> <li>ii. Steps in manufacturing of Hard gelatin capsule</li> <li>iii. Sticking and Picking with reasons and remedy.</li> <li>iv. Sugar coating of tablets</li> <li>Milling of APIs affect their flow, mixing and compressional</li> </ul>	
Q6	72.22	<ul> <li>i. Filter Aid</li> <li>ii. Steps in manufacturing of Hard gelatin capsule</li> <li>iii. Sticking and Picking with reasons and remedy.</li> <li>iv. Sugar coating of tablets</li> <li>Milling of APIs affect their flow, mixing and compressional properties, discuss this argument with reasons.</li> </ul>	each
Q6	a) b)	<ul> <li>i. Filter Aid</li> <li>ii. Steps in manufacturing of Hard gelatin capsule</li> <li>iii. Sticking and Picking with reasons and remedy.</li> <li>iv. Sugar coating of tablets</li> <li>Milling of APIs affect their flow, mixing and compressional properties, discuss this argument with reasons.</li> <li>Compare glass as pharmaceutical packaging material, list</li> </ul>	each
	b)	<ul> <li>i. Filter Aid</li> <li>ii. Steps in manufacturing of Hard gelatin capsule</li> <li>iii. Sticking and Picking with reasons and remedy.</li> <li>iv. Sugar coating of tablets</li> <li>Milling of APIs affect their flow, mixing and compressional properties, discuss this argument with reasons.</li> <li>Compare glass as pharmaceutical packaging material, list down the advantages and disadvantages of this material.</li> </ul>	each
Q 6 Q 7	72.22	<ul> <li>i. Filter Aid</li> <li>ii. Steps in manufacturing of Hard gelatin capsule</li> <li>iii. Sticking and Picking with reasons and remedy.</li> <li>iv. Sugar coating of tablets</li> <li>Milling of APIs affect their flow, mixing and compressional properties, discuss this argument with reasons.</li> <li>Compare glass as pharmaceutical packaging material, list down the advantages and disadvantages of this material.</li> <li>Describe various instruments used in the manufacturing of</li> </ul>	each 10 10
	b) a)	<ul> <li>i. Filter Aid</li> <li>ii. Steps in manufacturing of Hard gelatin capsule</li> <li>iii. Sticking and Picking with reasons and remedy.</li> <li>iv. Sugar coating of tablets</li> <li>Milling of APIs affect their flow, mixing and compressional properties, discuss this argument with reasons.</li> <li>Compare glass as pharmaceutical packaging material, list down the advantages and disadvantages of this material.</li> <li>Describe various instruments used in the manufacturing of Emulsions? Discuss any two of them</li> </ul>	each 10 10
	b)	<ul> <li>i. Filter Aid</li> <li>ii. Steps in manufacturing of Hard gelatin capsule</li> <li>iii. Sticking and Picking with reasons and remedy.</li> <li>iv. Sugar coating of tablets</li> <li>Milling of APIs affect their flow, mixing and compressional properties, discuss this argument with reasons.</li> <li>Compare glass as pharmaceutical packaging material, list down the advantages and disadvantages of this material.</li> <li>Describe various instruments used in the manufacturing of</li> </ul>	each 10 10

Roll No. in Fig. .....

Roll No. in Words. .....

Signature of Supdt.:

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Doctor of Pharmacy (Pharm.D.) Fourth Prof: Annual-2019

Subject: Pharmaceutics-VI (Industrial Pharmacy) (Old Course)

Paper: 3 Part - I (Compulsory)

Time: 30 Min. Marks: 20

## ATTEMPT THIS PAPER ON THIS QUESTION SHEET ONLY.

Division of marks is given in front of each question.

This Paper will be collected back after expiry of time limit mentioned above.

### Q.1. Encircle the correct option.

(20x1=20)

i.	a)	Antioxidants	b)	stainer into the product during storage: Stabilizer				
		olorants	d)	Diluents				
ii.	The	mixing mechanism analogous to		nsport is referred to as:				
		Convective mixing		Shear mixing				
	200	Diffusive mixing		Turbulent mixing				
iii.	If average weight of one tablet is 250 mg what would be the weight of API batch for							
		ablets						
	a)	2500 kg	b)	25 kg				
	c)	250 kg		250000 kg				
iv.		melting point substances causes	3	problem in tablet:				
	a)	Picking	b)					
	c)	Sticking	d)	Mottling				
٧.	The	reduction of bulk volume of the n	naterial	as a result of displacement of the				
	gaseous phase is called:							
	a)	Compaction	b)	Compression				
	c)	Slugging		D. Consolidation				
vi.	Clar	Clarification is applied when the solids do not exceed percent.						
	a)	2.0%		1.0%				
	c)	3.0%	7.5	10%				
vii.	Which test is used for testing membrane efficiency?							
				Forward Flow test				
	c)	Bubble point test	d)	Integrity Testing				
viii.	Warehouse is the area of industry where							
	a)							
	b) Production is carried out							
	c) Maintenance work is carried out							
	d)	Raw material and finished goo	ds are s	tored				
ix.	Capping of tablets during manufacturing occurs due to							
	a)	Air entrapment						
	b)	Plastic deformation						
	c) Improper adjustment of sweep-off blades							
	d) All of these options are correct							
X.	Air locks made in different parts of pharma industry help in							
	a)	Filtration of air	b)	Preventing cross contamination				
	c)	Maintaining sterility of area	d)	Helps in ventilation				

xi.	Which	of the following excipient is not	added i	n lozenges?					
	a)	Binder	b)	Disintegrant					
	c)	Sweetener	d)	Diluent					
xii.	During	g sugar coating of tablets, the S	ub coatir	ng is applied in order to					
ZIII.	a)	To increase the bulkiness							
	b) To avoid deterioration due to microbial attack								
	c)	To prevent the solubility in acid							
	d)								
xiii.	Sodiu	Sodium starch glycolate is classed under							
	a)	Glidant	b)	Super disintegrants					
	c)	Lubricants	d)	Diluents					
xiv.	Most commonly used diluent in tablet dosage form is:								
	a)	Tricalcium phosphate	b)	Mannitol					
	c)	Starch	d)	Lactose					
XV.		on-aqueous vehicle used in inje	ections is	··					
	a)	Alcohol	b)	Glycerol					
	c)	Alcoholic sodium stearate	d)	Vegetable oils					
xvi.	Glass apparatus used in the production of sterile products can be sterilized by:								
	a) Dry heat 140°C for 30 minutes								
	b) Dry heat 120°C for 1 hour								
	c)								
	d)	d) Using Suitable Disinfectant							
xvii.		is the area where mate	erials are	stored until tested for quality.					
	A)	Clean area	b)	Compounding section					
	c)	Dispensing area	d)	Quarantine area					
xviii.	Mesh	size is the number of opening	per	<del></del>					
	a)	Inch	b)	Square inch					
	c)	Linear inch	d)	Centimeter					
xix.	Hard gelatin capsule shells have moisture content in the range of:								
	a.	5%-10%	b)	12%-15 %					
	c)	30%-45%	d)	50%-70%					
XX.	The	particle size directly affects the		property of tablets.					
	a)	Weight	b)	Weight variation					
	c)	Disintegration time	d)	Dissolution rate					

Doctor of Pharmacy (Pharm.D.) Fourth Prof: Annual-2019

Roll No. .....

Subject: Pharmaceutics-VI (Industrial Pharmacy) (Old Course)

Paper: 3 Part - II

Time: 2 Hrs. 30 Min. Marks: 80

# ATTEMPT THIS (SUBJECTIVE) ON THE SEPARATE ANSWER SHEET PROVIDED

Note: Attempt any FOUR questions. Each question carries equal marks.

Q2	a)	Describe theory of mass transfer, what are the application of mass transfer in pharmaceutical industry?	10
	b)	How we can dry temperature sensitive products? Give a	
	-,	detailed account of dryer used for such operation.	10
Q3	a)		10
		powders? how these properties affect the tableting process.	
	b)	Name the methods of tablet manufacturing, make flow	10
	•	chart to describe tablet manufacturing by wet granulation method?	
Q4	a)	200700 Table	10
33 <b>3</b> 200		pharmaceutical industry? What is the role of Laminar flow hood in this process?	
		nood in this process:	
	b)	Define sterilization? Describe different methods used for the terminal sterilization of pharmaceutical formulation	10
Q5		Give brief answers of following terms.	5 Marks
ų s		i. Bubble point test of filter integrity	each
		ii. Film coating of tablets	cucii
		iii. Capping and lamination with reasons and remedy.	
		iv. Parts of single punch tablet machine	
		14. Tarts of single parter tablet machine	
Q 6	a)	Discuss the mechanism of size reduction during milling process. Discuss the principle, working and applications of	5+5
	212	ball mill.	
	b)	Briefly describe different packaging materials used in	10
		pharmaceuticals, what are different factors that affect the	
	62	selection of packaging material?.	
Q7	a)	- Made (1919) - Te 20 및 발생 14.1 및 14.1 및 14.1 및 14.1 를 14.1 를 14.1 및	
		mixing, describe the operation of Silverson homogenizer	10
	b)		10
		focusing on its sources, causes, types of fire, detection and prevention	

### UNIVERSITY OF THE PUNJAB Doctor of Pharmacy (Pharm.D.) Fourth Prof: Annual-2021 Subject: Pharmaceutics-IV (Industrial Pharmacy) (New Course) Paper: 3 Part - I (Compulsory) Time: 30 Min. Marks: 20', ..... ATTEMPT THIS PAPER ON THIS QUESTION SHEET ONLY. Division of marks is given in front of each question. This Paper will be collected back after expiry of time limit mentioned above. Encircle the correct option. Q.1. 1. Subcoating is given to the tablets: (A) To avoid deterioration due to microbial attack (B) To avoid stickiness (C) To prevent the solubility of in acidic media (D) To increase the bulk 2. Sigma blade mixers are commonly used in: (A) Dry granulation (B) Powder mixing (C) Wet granulation (D) Crude fibre mixing 3. Poorly manufactured tablets may have small pinholes on the surface. This phenomenon is known as: (A) Leaching (B) Picking (C) Mottling (D) Chipping 4. Which of the following industrial dryer is used to dry tablet granules? (A) Drum dryer (B) Fluidized bed dryer (C) Spray dryer (D) Freeze dryer 5. One of the following is used as a pH dependent controlled release excipient: (A) Carnauba wax (B) Hydroxy propyl methyl cellulose phthalate (C) Methyl cellulose (D) Glyceryl monostearate 6. In the preparation of multilayer tablets one of the substances listed is used for hydrophilic matrix coating: (A) CMC (B) Shellac (C) Stearyl alcohol (D) Beeswax 7. Durability of a tablet to combined effects of shock and abrasion is evaluated by using: (A) Hardness tester (B) Disintegration apparatus (C) Friabilator (D) Screw Gauge 8. For good flow properties, angle of repose should be: (A) 10 - 20 (B) 20 (C) 20 - 30(D) 30-40

(A) Starch (B) Gelatin

(A) Starch

(D) HPMC

(C) Sodium alginate (D) Tragacanth

10. Identify superdisintegrant:

(B) Sodium starch glycolate (C) Sodium alginate

9. Which of the following polymer used to prepare mucoadhesive tablets: Page 1 of 2

`. Roll No. in Fig. .....

(20x1=20)

Roll No. in Words. .....

Signature of Supdt.:

the homogentical industry
11. Glass is used as one of the major packing materials in pharmaceutical industry,
its internal structure is based on the network of
A. Silicon atoms
B. Polymer
C. Silicon and oxygen atoms
D. Diluents
12. Air locks installed in different parts of pharma industry help in
A. Filtration of air
B. Preventing cross contamination
C. Maintaining sterility of area D. Helps in ventilation
13. Drug dissolution might be faster be faster from a tablet prepared by?
A. direct compression
B. dry granulation
C. wet granulation
D. both dry and wet granulation
14. Methyl cellulose is used as?
A. wet binder
B. dry binder
C. filler
D. Dilyant
15. Which type of mixtures are used for batch mixing of solids?
A. planetary mixers
B. zig-zag mixers
C. sigma blade mixers
D. turbines
16. Swirl is a problem with such arrangement of impellers?
A. side-entering
B. off-centre
C. inclined entering
D. declined entering  17. Charring of cellulose materials/sugars can occur with which of the following
methods of sterilization?
A. Dry heat sterilization
B. Moist heat sterilization
C. Filtration
D. Dediction
18. Air locks installed in different parts of pharma industry help in
A. Filtration of air
B. Preventing cross contamination
C. Maintaining sterility of area
- wy 1 !atilation
1 I MAGAGO THOT INCOME THAILY HILLIGADOLIC VILLAND VII
19. Milling is a rigorous mechanical process that leaves many approximately for the product, however one the following effect can be used as an opportunity for
useful actions
A Agglomerations of particles
B. Charge development of particle surfaces
C. Amorphization
D. Chemical instability  20. Glass apparatus used in the production of sterile products can be sterilized by
A Dry heat 140 °C for 30 minutes
B. Dry heat 120 °C for 1 hour
C. Saturated steam 121 °C for 15 minutes  D. Using Suitable Disinfectant
D Using Suitable Districtant



Doctor of Pharmacy (Pharm.D.) Fourth Prof: Annual-2021

Roll No. ....

Subject: Pharmaceutics-IV (Industrial Pharmacy) (New Course)

aper: 3 Part – II

Time: 2 Hrs. 30 Min. Marks: 80

## ATTEMPT THIS (SUBJECTIVE) ON THE SEPARATE ANSWER SHEET PROVIDED

Note: Attempt any FOUR questions. Each question carries equal marks.

Q.2. Write note on the following

 $(4 \times 5 = 20)$ 

- a) Mass volume relationship
- b) Porosity, voids and Bulk Volume
- c) Heckle plot
- d) Compressibility
- Q.3. A) Define Compaction and discuss the different stages involved in powder compression. (10)
- B) Define Consolidation and discuss the concept of cold welding and asperitic melting. (10)
- Q.4. Define Physics of tablet and discuss the seven different events occur during compression. (20)
- Q.5. Draw the diagram and discuss the advantages and disadvantages of (20)
  - 1) Rotary filter
  - 2) Meta filter
- Q.6. A) Different mechanisms of milling, describe construction and working of ball mill. (10)
- B) Briefly describe the followings:

 $(5 \times 2 = 10)$ 

- i. Composition sterile products.
- ii. Indicators used for dry heat sterilization.
- iii. Specification of class A Clean room.
- iv. Importance of leakage tests of injectables
- v. Advantages of using rubber as packing material
- Q.7. A) What are sterile products? Classify them. Describe the "environment control" in respect of production of such products. (10)
- B) Briefly describe the followings:

 $(5 \times 2 = 10)$ 

- i. Effect of moisture on milling process
- ii. Materials used in pharmaceutical packaging
- iii. Leaching and sorption in respect of plastic containers
- iv. Applications of size reduction in pharmacy
- v. Disadvantages of milling