R19

Code: BP505T

B.Pharm III Year I Semester (R19) Regular & Supplementary Examinations January 2023 PHARMACEUTICAL JURISPRUDENCE

Time: 3 hours Max. Marks: 75

PART - A

(Compulsory Question)

1		Answer the following: (10 X 02 = 20 Marks)		
	(a)	Give offences and penalties about import of drugs.	2M	
	(b)	Give two examples of permitted colours as per Drugs and Cosmetics Act 1940.	2M	
	(c) (d)	What is Drug Consultative Committee (DCC)? Give the labelling requirements and write specimen label for schedule G	2M 2M	
	(e)	Define London proof spirit under M&TP Act	2M	
	(f)	Mention the ex-officio members of PCI.	2M	
4	(~)	Mention the objectives of D&MR Act	2M	
	(h)	Write a short on retail price of formulations.	2M	
	(i)	Mention the functions of Right to Information Act.	2M	
	(j)	Give the criteria for patenting of inventions.	2M	
		PART – B		
		(Answer any two questions: 02 X 10 = 20 Marks)		
2		Describe the classes of drugs and cosmetics which are prohibited from import and import under License.	10M	
3	(a)	Write the constitution and functions of Drug Technical Advisory Board (DTAB)	7M	
	(b)	Describe about restricted license.	ЗМ	
4	(0)	Write in detail about Education Description of Otata and Joint Otata Discussion On the	71.4	
-	(a) (b)	Write in detail about Education Regulation of State and Joint State Pharmacy Councils. Write a note on patent preparation.	7M 3M	
	(0)	while a note on patent preparation.	Sivi	
PART – C				
		(Answer any seven questions: 07 X 05 = 35 Marks)		
5	a)		3M	
^	b)	What are qualifications of Drug Inspector?	2M	
6	a)	= April 11 strot about alcottono proparationo.	3M	
	b)	Mention the offences and penalties in contravention of Pharmacy Act.	2M	
7		Give the constitution and functions of Institutional Animal Ethical Committee (IAEC).	5M	
8	a)	Mention the role of pharmacist in relation to his medical profession. What is Hathi committee.	3M	
9	b) a)	Write a note on list of permitted colors as per Drugs and Cosmetics Act 1940.	2M 3M	
Ü	b)			
10		Describe schedule P, U & V under Drugs and Cosmetics Act 1940	2M 5M	
11			3M	
		Pharmacy Act	A044000000	
12	b)		2M	
12	a) b)	3	3M	
13	2		2M 2M	
	b)		3M	
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Code: BP505T

12

13

B.Pharm III Year I Semester (R19) Regular Examinations February 2022 PHARMACEUTICAL JURISPRUDENCE

Time: 3 hours Max. Marks: 75

PART - A

		(Compulsory Question)
1	(a) (b) (c) (d) (e) (f) (g) (h)	Answer the following: (10 X 02 = 20 Marks) What is drugs and cosmetics act? How many times has it been modified? Which classes of drugs are prohibited from import? How are the colours used in drug products coded? What is the role of drug analysts? What are patented preparations? Why is the production of opium and poppy regulated? Why was the prevention of cruelty to animals act 1960 formulated? "The price of drugs need to be regulated" - Why?
	(i) (j)	What are the various types of IPR? What is the role of ethics in pharmaceutical industry?
	U/	PART – B (Answer any two questions: 02 X 10 = 20 Marks)
2	(a) (b)	Write a note on the labelling requirements of drugs and cosmetics. Write a note on the role of licensing authorities.
3	(a) (b)	What are the regulations for the export of ayurvedic and homeopathic preparations? What are the objectives and scope of pharmacy act?
4	(a) (b)	What are magic remedies? Why are they not advised as per the drug and magic remedies act? Write a note on the code of pharmaceutical ethics.
5		PART – C (Answer any seven questions: 07 X 05 = 35 Marks) Write a note on the import of drugs.
6		What are the various schedules mentioned in the drugs and cosmetics act 1940?
7		Write a few salient features about the top regulatory authority for the sale of drugs in India.
8		What is the function of narcotic and psychotropic consultative committee?
9	ı	What are the guidelines for breeding and stocking of animals as per CPCSEA?
1)	What are bulk drugs? How are they priced and sold?
1	1	Write a note on the mudaliar committee.

What are the pros and cons of patented technology over generics?

Why transparency is important in the sale and manufacturing of drugs?

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B.Pharm III Year I Semester (R19) Regular Examinations February 2022

PHARMACEUTICAL JURISPRUDENCE

Time: 3 hours

Max. Marks: 75

PART - A

(Compulsory Question)

- Answer the following: (10 X 02 = 20 Marks) 1
 - What is drugs and cosmetics act? How many times has it been modified? (a)
 - Which classes of drugs are prohibited from import? (b)
 - How are the colours used in drug products coded? (c)
 - What is the role of drug analysts? (d)
 - What are patented preparations? (e)
 - Why is the production of opium and poppy regulated? (f)
 - Why was the prevention of cruelty to animals act 1960 formulated? (g)
 - "The price of drugs need to be regulated" Why? (h)
 - What are the various types of IPR? (i)
 - What is the role of ethics in pharmaceutical industry? (i)

PART - B

(Answer any two questions: 02 X 10 = 20 Marks)

- Write a note on the labelling requirements of drugs and cosmetics. 2 (a)
 - Write a note on the role of licensing authorities. (b)
- What are the regulations for the export of ayurvedic and homeopathic preparations? 3 (a)
 - What are the objectives and scope of pharmacy act? (b)
- What are magic remedies? Why are they not advised as per the drug and magic remedies act? (a)
 - Write a note on the code of pharmaceutical ethics. (b)

PART - C

(Answer any seven questions: 07 X 05 = 35 Marks)

- Write a note on the import of drugs. 5
- What are the various schedules mentioned in the drugs and cosmetics act 1940? 6
- Write a few salient features about the top regulatory authority for the sale of drugs in India. 7
- What is the function of narcotic and psychotropic consultative committee? 8
- What are the guidelines for breeding and stocking of animals as per CPCSEA? 9
- What are bulk drugs? How are they priced and sold? 10
- Write a note on the mudaliar committee. 11
- What are the pros and cons of patented technology over generics? 12
- Why transparency is important in the sale and manufacturing of drugs? 13