

Name:	
Enrolment No:	

<b>UPES</b>	
<b>End Semester Examination, December 2023</b>	
Course: Pharmaceutical Jurisprudence	Semester: V
Program: B. Pharma	Duration: 03 Hours
Course Code: BP 505T	Max. Marks: 75
<b>Instructions: Attempt all the sections</b>	

<b>SECTION A</b>			
<b>(20Qx1M=20 Marks)</b>			

S. No.		Marks	COs
Q. 1.	Schedule C/C1 is regarding.....	1	CO1
Q. 2.	Drugs and cosmetics act is related to.....	1	CO1
Q. 3.	Enlist committees formed for the Pharmaceutical Legislation in India?	1	CO1
Q. 4.	Medical Termination of Pregnancy Act and Rules were introduced in the year _____	1	CO3
Q. 5.	Give full form of CPCSEA	1	CO3
Q. 6.	DPCO came into existence in which year?	1	CO3
Q. 7.	Define “hospital” as per MTP Act?	1	CO3
Q. 8.	MRP of scheduled formulation is calculated as .....	1	CO3
Q.9.	What % of margin is fixed for retail price?	1	CO2
Q. 10.	In which year Mudaliar committee was introduced?	1	CO2
Q. 11.	Name the Act that is applicable globally.	1	CO2
Q. 12.	When PCI got established?	1	CO2
Q. 13.	Define “Drug” as stated in Drugs and Cosmetics Act.	1	CO4
Q. 14.	Explain power of search and seizure.	1	CO4
Q. 15.	Schedule H is about .....	1	CO4
Q. 16.	Schedule X states that .....	1	CO4
Q. 17.	Name different types of IPR	1	CO5
Q. 18.	What is revocation of Patent?	1	CO5
Q. 19.	Expand SIC and CIC as Per RTI Act.	1	CO5
Q. 20.	Define Trademarks	1	CO5

<b>SECTION B (20 Marks)</b>			
<b>(2Qx10M=20 Marks)</b>			

<b>Attempt 2 Question out of 3</b>			
Q. 1.	Discuss in details requirements of Schedule D1 and D2 in import of drugs.	10	CO2
Q. 2.	Describe in detail about IPR.	10	CO5
Q. 3.	Write a note on NFCDA explaining its objectives and scope.	10	CO3

<b>SECTION-C (35 Marks)</b>			
<b>(7Qx5M=35 Marks)</b>			

<b>Attempt 7 Question out of 9</b>			
<b>Q. 1.</b>	Explain the constitution of Drug Technical Advisory Board.	<b>5</b>	<b>CO1</b>
<b>Q. 2.</b>	What do you mean by pharmaceutical code of ethics? What are the ethical responsibilities of a pharmacist in relation to job and medical profession?	<b>5</b>	<b>CO1</b>
<b>Q. 3.</b>	State conditions for grant of license for manufacturing of Drugs specified in Schedule C, C1 and X.	<b>5</b>	<b>CO2</b>
<b>Q. 4.</b>	Write down a note on Mudaliar committee and brief all Recommendations.	<b>5</b>	<b>CO2</b>
<b>Q. 5.</b>	Give a detail note on IAE.	<b>5</b>	<b>CO4</b>
<b>Q. 6.</b>	State objectives of NPPP 2012. Explain Scheduled and Non-Scheduled Formulations.	<b>5</b>	<b>CO4</b>
<b>Q. 7.</b>	When can pregnancies be terminated under the MTP act? Mention place requirement and offence and penalties related to the Act?	<b>5</b>	<b>CO3</b>
<b>Q. 8.</b>	When Drugs and Magic Remedies Act came into existence, State offence and Penalties related to Act.	<b>5</b>	<b>CO3</b>
<b>Q.9.</b>	Explain objectives of Medicinal and Toilet Preparations Act and Rules. Also State Levy and collection of Duties?	<b>5</b>	<b>CO5</b>