NWN-10061

## KNT/KW/16/6557

(Contd.)

## **B.Pharm Third Semester (C.B.S.) Examination**

## PHARMACEUTICAL JURISPRUDENCE AND ETHICS

## Paper—6

Time : Three Hours] [Max		hree Hours] [Maximum Marks : 8	30
<b>N.B.</b> :— (1)		- (1) Q. No. 1 is compulsory.	
		(2) Attempt any <b>four</b> questions from remaining.	
1.	Solv	we any <b>five</b> : $5\times4=2$	20
	(a)	Define:	
		(i) Adulterated drugs	
		(ii) Spuries drugs under Drugs and Cosmetic Act-1940.	
	(b)	Give formula for calculation of retail price of formulation.	
	(c)	Discuss various prohibited advertisement under Drugs and Magic Remedies Act.	
	(d)	Give offences and penalties under Pharmacy Act-1948.	
	(e)	Give background and objectives of Medicinal and Toilet Preparations Act.	
	(f)	Define "Opium derivatives" and "Hemp" under Narcotic Drug and Psychotropic Substances Ad	et.
	(g)	Central and State Consumer Protection Council.	
2.	(a)	Explain in depth qualifications, powers and duties of Drug Inspector under Drug and Cosmet Act-1940.	ic 8
	(b)	Give provisions for import of drug specified in schedule C and C <sub>1</sub> and for personal use.	7
3.	(a)	Give constitution and functions of Pharmacy Council of India.	8
	(b)	What is Central Register? Explain procedure for preparations of First Register.	7
4.	(a)	Give official definition of 'Advertisement', 'Magic Remedies'. Explain various advertisment which are exempted under Drug and Magic Remedies Act-1954 and Rules-1955.	ch 8
	(b)	Under which circumstances pregnancy can be terminated as per Medical Termination of Pregnancy.  Act-1970 and Rules-1975. Write about places used for termination of pregnancy.	су 7

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- 5. (a) Explain in depth bonded and nonbonded laboratory.
  - (b) Discuss role of pharmacist in relation to Job and Trade.

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- 6. (a) What sort of control the Central Government and State Government have over production and supply of dangerous drugs?
  - (b) Write about cultivation of opium. 7
- 7. (a) Explain in depth history of pharmaceutical legislation in India.
  - (b) What are the provisions for labelling and packing of drug as per Drug and Cosmetic Act-1940 and Rules-1945 for different formulations?

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