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Total No. of printed pages = 4 BINA CHOWDHUR WELL LIBRARY (GIMT & GIPS) Azara, Hatkhowapara BP 505 T Guwahati - 781017 Roll No. of candidate 2021 B. Pharm 5th Semester End-Term Examination Pharmacy PHARMACEUTICAL JURISPRUDENCE (New regulation) Full Marks - 75 Time - Three hours The figures in the margin indicate full marks for the questions. 1. Multiple choice questions (MCQ)(Answer all Questions)  $(20 \times 1 = 20)$ Life period of drugs is covered under (a) Schedule H (b) Schedule P (c) Schedule M (d) None (ii) The form number required for the grant of license for wholesale of Schedule X drugs is (b) Form 20 B (a) Form 20 G (c) Form 21 B (d) None (iii) DTAB has -- nominated members. (a) 0 (b) 1 (c) 3 (d) 5 (iv) Special labelling requirement for Schedule H is (a) Symbol XRx in red left hand top corner (b) Symbol NRx prominently on left hand top corner (c) Symbol Rx prominently on left hand top corner (d) All of the above Examples of Schedule X drugs is (a) Amphetamine (b) Glutethimide

(d)

All of the above

(c)

Aqualone

-	(V1)	Nar	cotic drugs and Psychotropic	substa	nces act was passed in the year		
		(a)	1954	(b)	1919		
		(c)	1985	(d)	1948		
	(vii)	The first edition of Indian Pharmacopoeia was published in the year					
		(a)	1945	(b)	1955		
		(c)	1965	(d)	1975		
	(viii)	Pha	rmacy council of India is reco	onstitut	ed		
		(a)	Every 2 years	(b)	Every 3 years		
		(c)	Every 4 years	(d)	Every 5 years		
	(ix) Medicinal and toilet preparation act is also known as						
		(a)	Excise duty act	(b)	MTP Act		
		(c)	Narcotic Act	(d)	Poison Act		
	(x)	Non	a-bonded laboratory should eer.	be insp	ected — by the excise		
		(a)	Every day	(b)	Every week		
		(c)	Every month	(d)	Every year		
	(xi)	Dru	tuted in the year 1930 under the				
		(a)	RN Chopra	(b)	Mahadeva Lal Schroff		
		(c)	Bishnupada Mukherjee	(d)	None		
	(xii) The Indian Pharmacopoeia Committee was appointed in the year						
		(a)	1948	(b)	1949		
		(c)	1954	(d)	1962		
	(xiii)	Dru	g Inspector is appointed und	er			
39		(a)	42 of IPC	(b)	21 of IPC		
		(c)	46 of IPC	(d)	31 of IPC		
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(X1V)	) It a	drug contains a colour that is	not p	rescribed is called				
	(a)	Misbranded	(b)	Adulterated drug				
	(c)	Spurious	(d)	None				
(xv)	The of DPCO, 1995 contains forms required for approval fixation or revision of prices of bulk drugs and formulations.							
	(a)	Third Schedule	(b)	Second Schedule				
	(c)	First Schedule	(d)	None				
(xvi)	vi) Animal Welfare Board of India constituted under the Prevention of Cruelty to Animals act is situated in							
	(a)	New Delhi	(b)	Mumbai				
	(c)	Kolkata	(d)	Chennai				
(xvii	ii) Drugs and Magic remedies Act came into force in the year							
	(a)	1930	(b)	1955				
	(c)	1949	(d)	1954				
(xvi	ii) Gl	MP requirement of factory pres	nises	s, plants and equipments				
	(a)	Schedule M	(b)	Schedule P				
	(c)	Schedule Q	(d)	Schedule S				
(xix)	Who	Whole blood can be stored for ———						
	(a)	12 days	(b)	21 days				
	(c)	24 days	(d)	60 days				
(xx)	has formulated the code of ethics for the guidance pharmacist.							
	(a)	Pharmacy Council of India						
	(b)	Medical Council of India						
	(c)	UGC						
	(d)	AICTE						

2. Long answers (Answer two out of three)

- $(2 \times 10 = 20)$
- (a) Discuss in details about the qualifications and duties of a Drug inspector.
- (b) Write a detailed note on the Medical Termination of Pregnancy Act, 1971.
- (c) When did the Pharmacy Council of India constituted? Enlist the constitution and functions of PCI.
- 3. Short answers (Answer Seven out of nine)

 $(7 \times 5 = 35)$ 

- (a) Write down the offences and penalties for Narcotics and Psychotropic Substances Act.
- (b) Write a note on export of alcoholic preparations.
- (c) Define advertisements. Discuss in brief about the prohibited advertisements.
- (d) Write a note on prevention of Cruelty to Animals Act.
- (e) Write a note on the retail prices of formulations.
- (f) Discuss the ethics of a pharmacist in relation to his job.
- (g) Write a note on
  - (i) Drug Enquiry Committee
  - (ii) DTAB
- (h) Explain in brief about Right to Information Act.
- (i) Write down the details about Intellectual Property Rights (IPR).