

Branning Conversity Barasat, Kelkata -70812\$

BRAINWARE UNIVERSITY



Term End Examination 2021 - 22 Programme – Diploma in Pharmacy Course Name – Pharmaceutical Jurisprudence Course Code - 2.4T (Year II)

Time: 1 Hr.35 Min.		Full Marks: 80
[The figure in the man	rgin indicates full marks.]	
Gr	roup-A	
(Multiple Choi	ice Type Question)	1 x 80=80
Choose the correct alternative from the following	ALCOHOL SELECT AND	
(1) Pharmacy act was established in		
a) 1948	b) 1940	
c) 1995	d) 1919	
(2) Pharmacy Act extends whole of the India exc	cept	
a) Delhi	b) West Bengal	
c) Both Delhi & West Bengal	d) Jhammu and Kashmir	
(3) In the "Joint State Pharmacy Council" electe cist is/are		pharma
a) 1 from each state	b) 3 to 5 from each state	
c) 2 from each state	d) 5 from each state	
(4) State Pharmacy Council should have the follow	owing number of elected members	:
a) 6	b) 5	
c) 9	d) 7	
(5) Which one of the following is/are ex-officio	member(s) of state pharmacy coun	cil?
a) Chief pharmacist of government hospital	 b) Chief administrative medicate 	cal officer of the st
c) .Assistant drug controller	d) All	
(6) TGA is regulatory agency of		
a) Denmark	b) Austria	
c) Australia	d) Zimbabwe	
(7) The first edition of Indian Pharmacopoeia wa	s published in the year of:	
a) 1940	b) 1050	

d) 1985

c) 1955

b) Every 3 years		
d) Every 6 y	ears	
	LIBRARY	
b) 6	Brathware University	
d) 8	Barasat, Kalkata -70012	
b) . 6th June	,1995	
d) 6th Marc	h,1995	
1		
b) DTAB		
d) All		
b) To furnis	h reports of results of tes	
d) All of the	ese	
Committee is		
	b) Maj. Gen. S.L.Bhati	
d) Dr.A.L.	d) Dr.A.L. Mudaliar	
b) 1931		
d) 1953		
Pharmacy Cour	ncil" hold office for a term	
b) 4 years		
d) 6 years		
ember(s) from ea	ach state.	
b) 1		
d) 4	700.	
and Developme	nt Committee is:	
b) Sir R.N.		
d) Sir J.S.	Hathi	
of		
b) 1953		
d) 1943	F1.0 CON ANN	
ct provides con	stitution of DTAB?	
b) Section		
d) Section	6	
DTAB is;		
b) 5		
d) 8		
b) 6		
60.87 P.S.		
	b) 6 d) 8 b) 6 d) 8 b) 6th June d) 6th Marc b) DTAB d) All b) To furnis d) All of the Committee is b) Maj. Ger d) Dr.A.L. b) 1931 d) 1953 Pharmacy Cour b) 4 years d) 6 years ember(s) from each b) 1 d) 4 and Developme b) Sir R.N. d) Sir J.S. of b) 1953 d) 1943 act provides con b) Section d) Section e DTAB is; b) 5 d) 8 the DTAB is:	

22) How many Nominated Representatives of the C C?	Central Government are there in the DC	
a) 2	b) 7	
c) 6	d) 3 Barasat, Kennan School Service Country	
23) The Chopra Committee was set up by the	Barasat, Kelkata -700128	
a) Drug Control General of India	b) Government of India	
c) Chairman, DTAB	d) Director, CDL	
24) The first Pharmaceutical industry in India is	Settle Committee	
a) The Bengal Chemical and Pharmaceutical W orks Ltd	b) The East India Pharmaceutical Works	
c) The Alembic Chemical Works	d) The Glaxo)
25) The register of sale of drugs is maintained and p	preserved for period of	
a) 1 year	b) 3 years	
c) 4 years	d) 5 years	
26) Central drug laboratory is located at		
a) Kolkata	b) Lucknow	
c) Hyderabad	d) Mumbai	
27) Names from the register can removed only by a	n order of the-	
a) PCI	b) Registration tribunal	
c) EC of the State PCI	d) Director of Health Services	
28) The Essential Commodity Act came into force i	in-	
a) 1945	b) 1950	
c) 1955	d) 1960	
29) Persons from teaching profession are nominated	d to PCI by	
a) State Government	b) Central Government	
c) Executive Committee	d) Election	
30) The Secretary of PCI may also acts as its		
a) Registrar	b) Treasurer	
c) Registrar and Treasurer	d) Registrar and President	
31) The period in hours of training to be under takes	n by a student pharmacist in a hospital i	
a) 500	b) 750	
c) 600	d) 800	
32) Standard for disinfectant fluids comes under		
a) Schedule O	b) Schedule R	
c) Schedule S	d) Schedule E	
33) Which pharmaceutical product is not included in	n Schedule C?	
a) Toxins	b) Sera	
c) Antigens	d) Capsules	
34) Blood bank comes under the schedule		
a) Blood bank comes under the schedule	b) Schedule D	
c) Schedule F	d) Schedule U	
35) If the drug contains filthy, putrid or decomposed	d substance then the drug is known as	
a) Misbranded drug	b) Adulterated drug	

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nufacture of Homeopathy drugs

nufacture of Allopathy drugs

b) 400 square meters

d) Requirements of factory premises for the ma

nufacture of medical devices

a) 250 square meters

c) Requirements of factory premises for the ma

nufacture of Ayurveda, Siddha, Unani drugs

(47) Minimum area required for parenteral preparation is....

c) 500 square meters	d) 150 square meters		
(48) Schedule X of Drugs and Cosmetics Act compr	rises		
a) List of incurable disease	b) Guidelines for clinical trials		
c) List of generic drugs	d) List of narcotic and psychotropic drugs		
(49) Which one of the following is prohibited to be	imported?		
a) Toilet preparation	b) Ayurvedic drugs		
c) Schedule C and G drugs	d) Misbranded drugs		
(50) Hatch Waxman Act is related to			
a) Banned drugs	b) Banned drugs	RV	
c) Dangerous drugs	d) Generic drugs Bratoware C Baraset, Kelka	Aiversity	
(51) ow many numbers of patients are required for p	phase II clinical trial?	p =100175	
a) 10-12 patients	b) 1-10 patient		
c) 100 patient	d) 500 patient		
(52) Purpose of phase-III clinical trial is	- 10 - 10		
To determine maximum tolerated dose in hu man, pharmacodynamic effects, adverse effects	b) Determine possible therapeutic uses, effectiv e doses range		
 c) Efficacy and safety of drug in larger number of patients (500 patients) 	 d) Long time adverse effects after marketing dr ug 		
(53) Standard for mechanical contraceptive comes u Act .	ander Schedule as per D and C		
a) R	b) A		
c) S	d) O		
(54) Pack size of drug is covered under Schedule			
a) P	b) R		
c) P1	d) O		
(55) Identify the special labelling requirement for So	chedule X drug		
 a) Symbol XRx is given in red at the left top co rner of the lable 	b) Symbol N in red displayed on left top corner of the labe		
c) Symbol N displayed on right top corner of the label	d) Symbol H displayed on right top corner of the label		
(56) In 1954 which one of the following act is passe	d?		
a) Narcotic and psychotropic substance Act	b) Drug and magic remedies Act		
c) The medical termination and pregnancy Act	d) Poisonous Act		
(57) Post marketing surveillance comes under clinic	al trial		
a) Phase I	b) Phase II		
c) Phase III	d) Phase IV		
(58) In 1970, which of the following act (s) was/wer	re enacted?		
a) Poisons Act	b) Drug and Cosmetics Act		
c) Both a & b	d) Patent act		
(59) The cosmetics that are prohibited to be importe	ed contains		
a) more than 10ppm lead	b) more than 20ppm lead		
c) more than 2ppm lead	d) more than 50ppm lead		
(60) Which one of the following form is needed for	the cosmetics manufacture?		

a) 6 months

c) 2 years

b) 12 months

d) 5 years

(72) The term of patent for ordinary invention fia) 7 years	rom the date of patent is-	Solda .
c) 15 years	b) 14 years	Bristowera Ghiversty Baraner, Kathate -70652
(73) Grant of patent, under Patent Act, may be o sement of acceptance within a period of	d) 10 years pposed by a person from the d	ate of all and
a) 4 months	the u	ate of adverti
c) 1 year	b) 8 months	
(74) VDRL antigen is to be tested and analyzed b	d) 2 years	
a) Drug Inspector	by the	
c) Serologist and Chemical Examiner	b) Excise Commissioner	
(75) Tests on oral policy vaccine	d) Drug Controller of Ind	lia
(75) Tests on oral polio vaccine are to be carried o	out at the	
a) National Institute of Communicable Disease, Delhi	 b) Indian Pharmacopoeial ad 	Lohani
c) Central Drug Laboratory, Kolkata		
(76) An example of artificial colour is	d) Central Drug Research	Institute Lucknow
a) Titanium dioxide		Juckilow
c) Cochineal	b) Caramel	
(77) Drugs and Chemists shops are	d) Curcumin	
(77) Drugs and Chemists shops are exempted froma) Opening time	the provision of	
c) Opening time&Closing time	b) Closing time	
(78) All the statements regarding CDV		
(78) All the statements regarding CDL are true exce	pt	
 a) To carry analysis of sample of drugs and cos metics sent by Court or Custom Collector c) To carry all duties suggested by Central or St ate Govt. 	b) Biological or microbiolog out by CDL, Kolkata	
	d) Analytical report with all plied by the Director	protocol of test sup
(79) atent and proprietary medicines can be imported a) Crude conditions	to some extent in	
c) Bulk form	b) Multi-dose vials	
(80) List of Ayurvedic and Unani poisons is given in s	schedule	
	b) F and F1	
	d) H	