Schedule

1. Schedule Tuf Drugs and Cosmeties Rules, 1945 deals with: [GPAT-2024]

(a) GMP for ASU drugs

(b) GMP for Pharmaceutical product

(c) GLP and requirement of premises and equipments

(d) GMP for Homeopathy medicine

2. Drugs covered under this schedule are not permitted for repacking license. Identify the correct schedule (GPAT-2023 Shift 1)

(a) Schedule H (b) Schedule G (c) Schedule O (d) Schedule C and CI

3. Requirements of factory premises for the manufacture of cosmetics are mentioned, in the Drugs and Cosmetics Rules, 1945 under (GPAT-2023 Shift I)

(a) Schedule M (b) Schedule M-1 (c) Schedule M-2 (d) Schedule M-3

4. Match List I with List 11 (GPAT-2023 Shift)

LIST I SCHEDULE	LIST II COVER THE
1. Schedule Y	[P] Shelf life of drugs
2. Schedule FF	[Q] Requirements for clinical trails
3. Schedule O	[R] Disinfectant liquids
4. Schedule P	[S] Ophthalmic ointments

Choose the correct answer from the options given below

(a) 1-[P], 2-[Q], 3-[R], 4-[S] (b) 1-[Q]. 2-[R], 3-[S].4-[P]

(c) 1-[S], 2-[R), 3-[Q]. 4-[P] (d) 1-(Q), 2-[S], 3-[R] 4-[P]

5. The Schedule in D & C Act that deals with the standards for disinfectant fluids is (GATE-2004, GPAT-2023 SHIFT-11)

(a) Schedule B (b) Schedule F (c) Schedule O (d) Schedule M

6. Match the following (GPAT-2021)

1. Schedule M [P] Regulations regarding Life period and storage of various drugs

2. Schedule FF [0] Regulations for manufacturing, Premises. Waste disposal requirements

3. Schedule G [R] Various drugs to be used under the medical supervision of plant and Equipments (Good Manufacturing Practices)

4. Schedule P [S] Standards for Ophthalmic preparations

Choose the correct answer from the options given below

(a) 1-[P], 2-[Q] 3-[R].4-[S]

(b) 1-[Q]2-[S] 3-[R].4-[P]

(c) 1-[S],2-[Q],3-[R],4-[P]

(d) 1-[R], 2-[P], 3-[Q].4-[S]

7. Schedule N is related with [GPAT-2015]

(a) Furniture and apparatus in pharmacy

(b) Water bath in pharmacy

(c) Indian Pharmacopoeia in pharmacy

(d) Good manufacturing practice

8. Secobarbital is drug [GPAT 2014]

(a) Schedule G

(b) Schedule H

(c) Schedule Q

(d) Schedule X

9. Which of the following Schedules include shelf life of drugs [GPAT-2011, 2013]

(a) Schedule F (b) Schedule M (c) Schedule G (d) Schedule P

10. The Schedule in Drugs and Cosmetics Act that deals with the requirements and guidelines for clinical trials, import and manufacture of new drugs is [GATE-2003, 2008]

(a) Schedule O (b) Schedule "M (c) Schedule F (d) Schedule Y

11. Schedule FF contains the list of the following [GATE-2002]

(a) Drug which can be marketed under generic names only

(b) Drug which are habit forming

(c) Standards for ophthalmic preparation

(d) Drug which are exempt from certain provisions applicable to manufacturing

Pharmaceutical Jurisprudence

Drugs and Cosmetics Act Rules are divided 18 parts.

Schedule	Applied to	Trick to Learn
A	Performa for application for licenses, issue & renewal, sending memoranda under the Act	A = Application
B	Rates of fee for test/analysis by Central or State Drug Labs	B = Bill (fees)
B1	Fee for test/analysis by Pharmacopeial Lab for Indian Medicine / Govt Analyst	B1 = Bill No.1 (special fees)
C	List of biological & special products (import, sale, distribution, manufacture under special provision)	C = Cell products
C1	List of other special products under special provision	C1 = Cell +1 (others)
D	Drugs exempted from provisions for import	D = Door open (Exempt)
E	List of poisonous substances (Ayurvedic, Siddha, Unani)	E = Evil = Poison
E_1	Poisonous drugs	
F	Requirements for blood bank & blood component preparation	F = Fluid (blood)
F1	Part I – Vaccines; Part II – Sera; Part III – Diagnostic agents (bacterial origin)	F_1 = First defense (vaccine/sera)
F2	Standards for surgical dressings	F2 = First Aid (dressings)
F3	Standards for sterilized umbilical tapes	F3 = Fresh cord tape
FF	Standards for ophthalmic preparations	FF = Focused on eyes
G	List of substances to be used only under medical supervision (labelled accordingly)	G = Guidance needed
H	List of prescription drugs	H = Handwritten Rx
J	List of diseases a drug may not claim to prevent/cure	J = Jail claims
K	Drugs exempted from certain manufacturing provisions	K = Kind exemption
L	Good Laboratory Practice (GLP) & premises/equipment requirements	L = Lab rules
M	GMP for factory premises, plant & equipment (drugs)	M = Manufacture GMP
M1	Requirements for homeopathic manufacture	M1 = Mother tincture
M2	Requirements for cosmetics manufacture	M2 = Mirror (cosmetics)
M3	Requirements for medical devices manufacture	M3 = Machine (devices)
N	Minimum equipment for efficient pharmacy running	N = Necessary items
O	Standards for disinfectant fluids	O = Odor kill (disinfect)
P	Life period of drugs	P = Period (shelf life)
P1	Pack sizes of drugs	P1 = Packet size
Q	Approved colours, dyes & pigments in cosmetics	Q = Queen's makeup (colors)
R	Standards for condoms & contraceptives	R = Reproductive safety
R1	Standards for medical devices	R1 = Replacement (devices)
S	Standards for cosmetics	S = Skin (cosmetics)
T	GMP for Ayurvedic, Siddha, Unani medicines	T = Traditional medicine
U	Manufacturing records (drugs: raw material & analysis)	U = Usage records
U1	Manufacturing records (cosmetics)	U1 = Usage record (cosmetics)
V	Standards for patent/proprietary medicines	V = Verified proprietary
W	Drugs to be marketed under generic names only	W = Without brand
X	Drugs with special provisions (import, manufacture, sale, labelling, packaging)	X = Extra control
Y	Clinical trial requirements & guidelines for new drugs (import/manufacture)	Y = Yes to trial