

GPAT

2027-28



PHARMACEUTICAL JURISPRUDENCE

CLASS- 19

TIME- 8 : 00 P.M

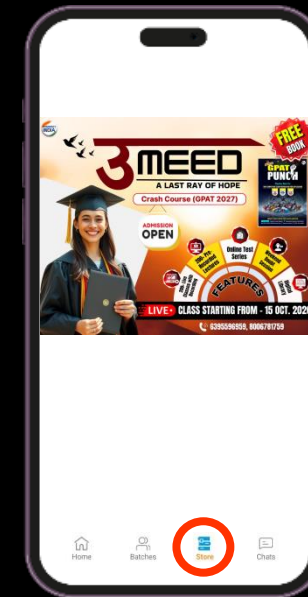
PART
2

DRUGS & COSMETICS ACT & RULES

UPDATED QUESTIONS TILL GPAT 2026

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01.

Schedule 'H' and Schedule 'S' as per the Drugs & Cosmetics Act deal with the following [GPAT-2025]

[P] Prescription drugs which are required to be sold by retail only on prescription of RMP

[Q] Standard for cosmetics

[R] Biological and special products

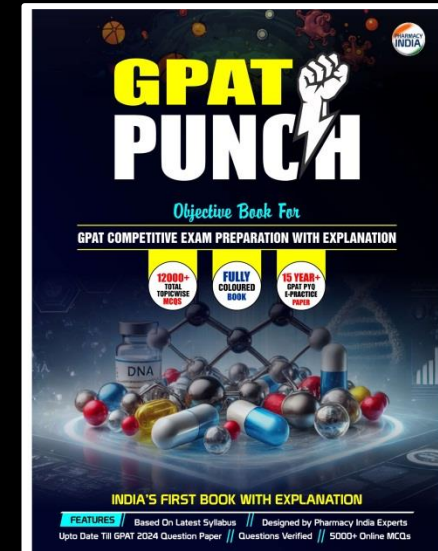
[S] List of coal tar colours permitted to be used in cosmetics and soaps

(a) [P], [Q]

(b) [P], [R]

(c) [Q], [S]

(d) [R], [S]



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01.

Schedule 'H' and Schedule 'S' as per the Drugs & Cosmetics Act deal with the following [GPAT-2025]

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[Q] Standard for cosmetics

[R] Biological and special products

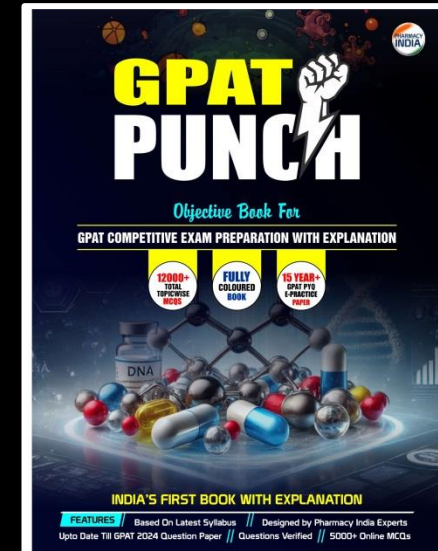
[S] List of coal tar colours permitted to be used in cosmetics and soaps

(a) [P], [Q]

(b) [P], [R]

(c) [Q], [S]

(d) [R], [S]



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Explanation:

Schedule H deals with prescription drugs that are sold by retail only on the prescription of a Registered Medical Practitioner. Schedule S specifies standards for cosmetics. Therefore Schedule H corresponds to [P] and Schedule S corresponds to [Q]. Biological and special products are dealt with under other schedules, while permitted coal-tar colours are covered separately.



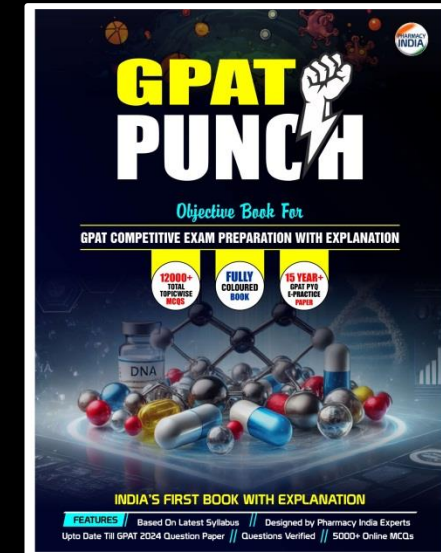
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02. **Schedule FF contains the list of the following [GPAT-2024]**

- (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



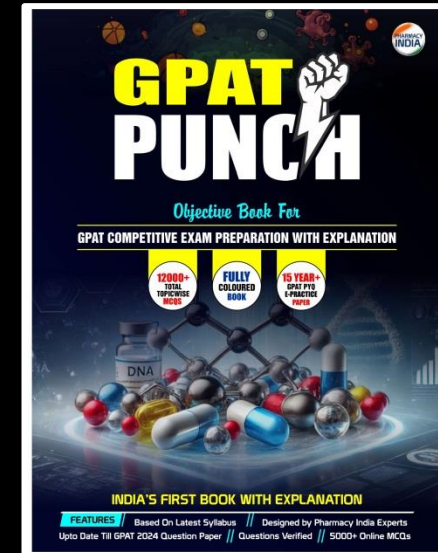
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- (b) Drug which are habit forming
- (c) Standards for ophthalmic preparation
- (d) Drug which are exempt from certain provisions applicable to manufacturing



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Explanation:

Schedule FF is specifically related to standards for ophthalmic preparations. Ophthalmic products require special standards because they are intended for application to the eye. Habit-forming drugs are not classified under Schedule FF. Exemptions from manufacturing provisions are also covered under other schedules.



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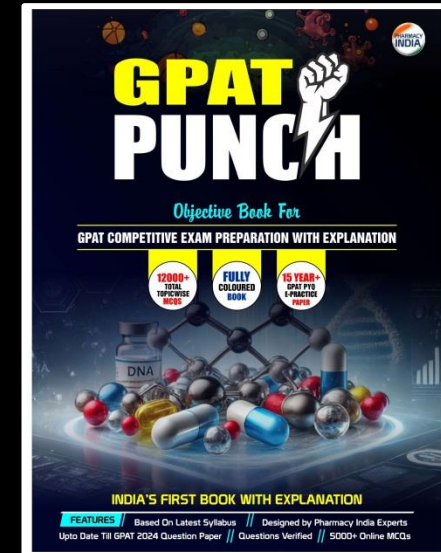
03. **Schedule D as per D & C Act is concerned with [GPAT-2023]**

(a) List of drugs exempted from the provision of import of drugs

(b) Diseases or ailments which a drug may not purport to prevent or cure

(c) Requirements of factory premises

(d) List of prescription drugs



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03.

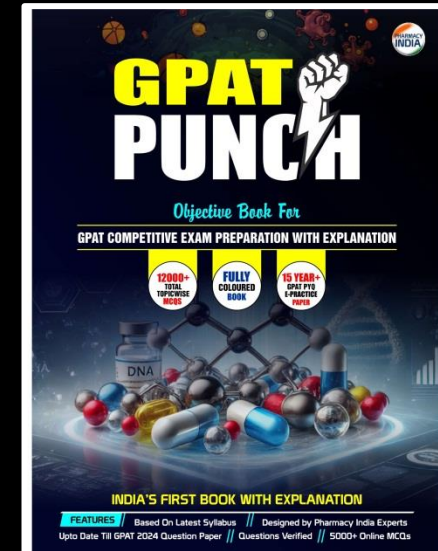
Schedule D as per D & C Act is concerned with [GPAT-2023]

(a) List of drugs exempted from the provision of import of drugs

(b) Diseases or ailments which a drug may not purport to prevent or cure

(c) Requirements of factory premises

(d) List of prescription drugs



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Explanation:

Schedule D deals with drugs that are exempted from certain provisions relating to import. Schedule J deals with diseases or ailments for which a drug cannot claim prevention or cure. Factory-premises requirements are associated with Schedule M. Prescription drugs are principally covered under Schedule H.



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04.

Listed below are the Schedules to the Drugs and Cosmetics Act. Match them [GATE-2000]

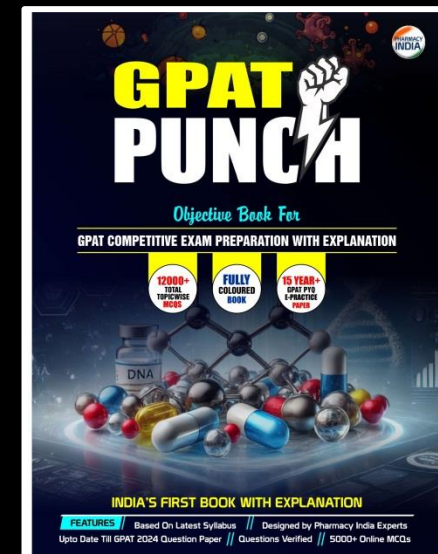
Schedule	Matching Description
1. Schedule M	[P] Standard for disinfectant fluids
2. Schedule O	[Q] Standard for ophthalmic preparation
	[R] Requirement of factory premises
	[S] Standard for cosmetics

(a) 1-[Q], 2-[S]

(c) 1-[R], 2-[P]

(b) 1-[R], 2-[S]

(d) 1-[P], 2-[S]



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04.

Listed below are the Schedules to the Drugs and Cosmetics Act. Match them [GATE-2000]

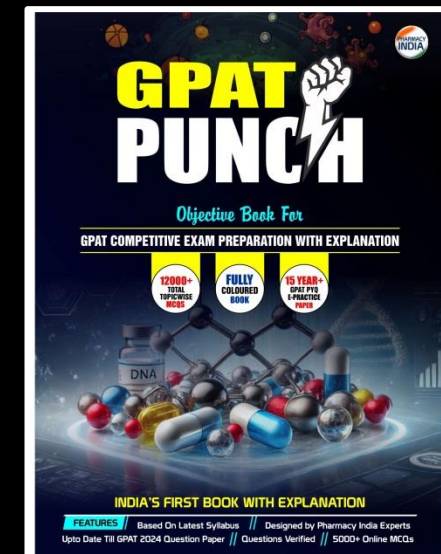
Schedule	Matching Description
1. Schedule M	[P] Standard for disinfectant fluids
2. Schedule O	[Q] Standard for ophthalmic preparation
	[R] Requirement of factory premises
	[S] Standard for cosmetics

(a) 1-[Q], 2-[S]

(b) 1-[R], 2-[S]

(c) 1-[R], 2-[P]

(d) 1-[P], 2-[S]



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Explanation:

Schedule M specifies requirements relating to factory premises and Good Manufacturing Practices. Therefore Schedule M corresponds to [R]. Schedule O contains standards for disinfectant fluids and therefore corresponds to [P]. Standards for ophthalmic preparations are associated with Schedule FF, not Schedule O.



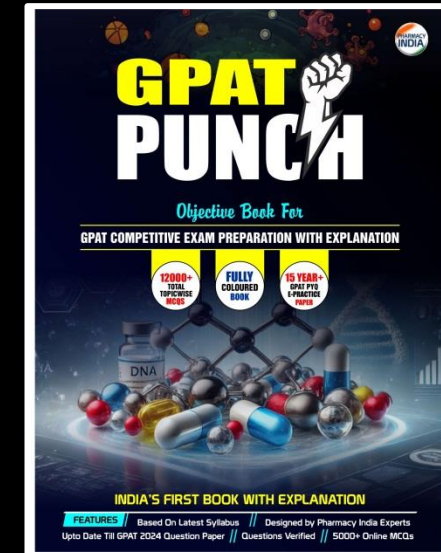
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05. An original licence or renewed licence to sell drugs remains valid up to [GATE-2000]

- (a) 31 March of the next year in which it is granted
- (b) 30 June of the following year in which it is granted or renewed
- (c) 31 January of the same year in which it is granted
- (d) 31 December of the following year in which it is granted or renewed



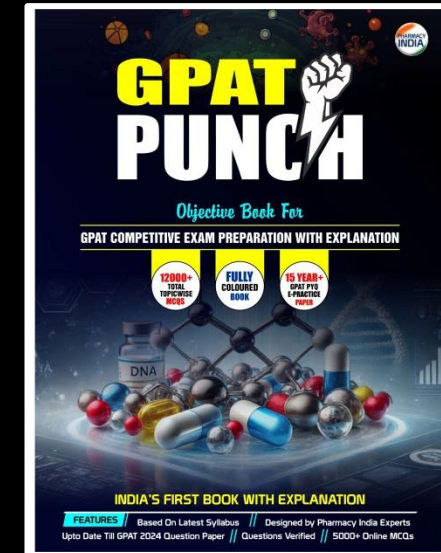
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- (a) 31 March of the next year in which it is granted
- (b) 30 June of the following year in which it is granted or renewed
- (c) 31 January of the same year in which it is granted
- (d) 31 December of the following year in which it is granted or renewed**



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Explanation:

An original or renewed licence for sale of drugs remained valid up to 31 December of the year following the year in which it was granted or renewed, unless suspended or cancelled earlier. Therefore, if a licence was granted or renewed during a particular year, its validity extended until 31 December of the succeeding year.



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06.

Given below are the Schedules as per D and C Act 1940.
Match them with information to be given in the label
[GATE-2000]

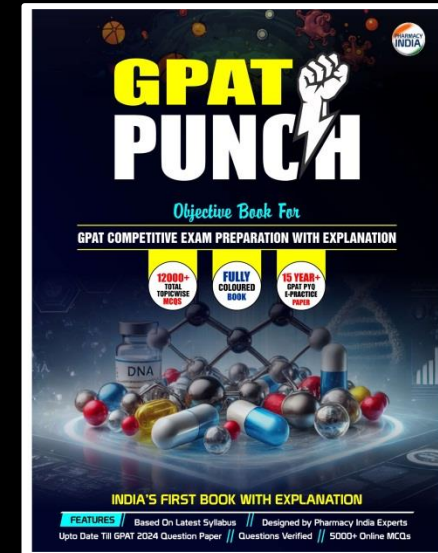
Schedule	Labelling Information
1. Schedule H	[P] For external use only
2. Schedule G	[Q] For therapeutic use only
	[R] Caution-It is dangerous to take this preparation except under medical Supervision
	[S] To be sold by retail on the prescription of a R.M.P. only

(a) 1-[Q], 2-[S]

(b) 1-[R], 2-[S]

(c) 1-[S], 2-[R]

(d) 1-[P], 2-[S]



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06.

Given below are the Schedules as per D and C Act 1940.
Match them with information to be given in the label
[GATE-2000]

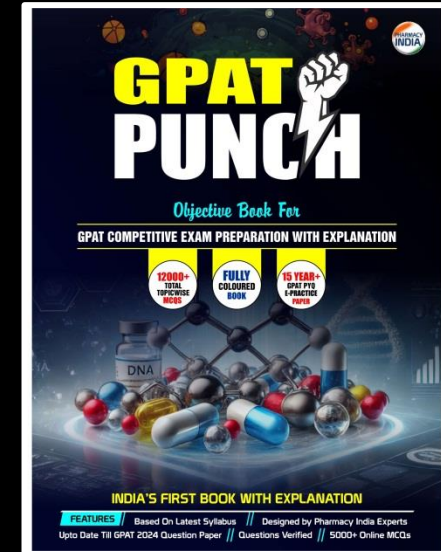
Schedule	Labelling Information
1. Schedule H	[P] For external use only
2. Schedule G	[Q] For therapeutic use only
	[R] Caution-It is dangerous to take this preparation except under medical Supervision
	[S] To be sold by retail on the prescription of a R.M.P. only

(a) 1-[Q], 2-[S]

(b) 1-[R], 2-[S]

(c) 1-[S], 2-[R]

(d) 1-[P], 2-[S]



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Explanation:

Schedule H drugs carry the warning that they are to be sold by retail only on the prescription of a Registered Medical Practitioner. Thus Schedule H matches [S]. Schedule G preparations carry the caution regarding use except under medical supervision. Therefore Schedule G matches [R].



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07.

Given below are different schedules as per the D and C Act.
Match them with items mentioned in [P] to [S] [GATE-1999]

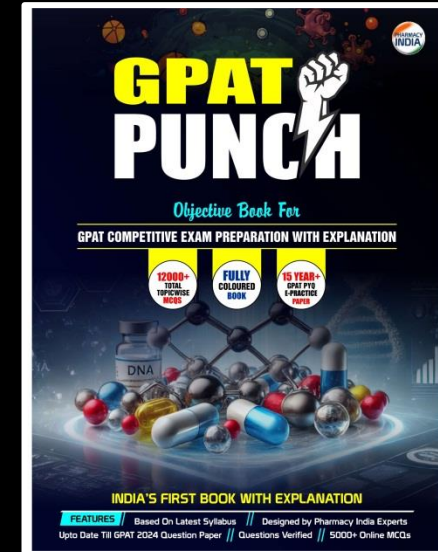
Schedule	Description
1. Schedule FF	[P] Standards for ophthalmic preparations
2. Schedule M	[Q] Diseases or ailments which a drug may not purport to prevent or cure
	[R] Life period of drugs
	[S] Requirements of factory premises

(a) 1-[Q], 2-[S]

(b) 1-[P], 2-[Q]

(c) 1-[R], 2-[P]

(d) 1-[P], 2-[S]



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07.

Given below are different schedules as per the D and C Act.
Match them with items mentioned in [P] to [S] [GATE-1999]

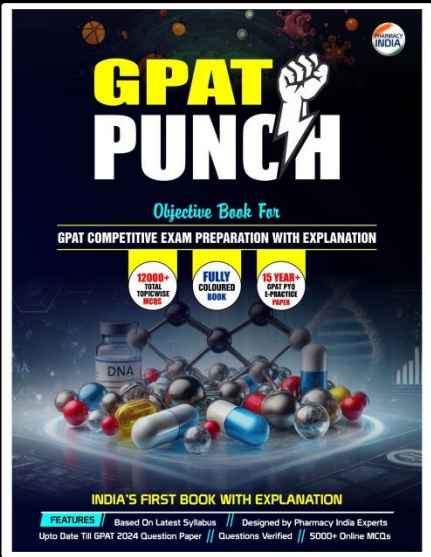
Schedule	Description
1. Schedule FF	[P] Standards for ophthalmic preparations
2. Schedule M	[Q] Diseases or ailments which a drug may not purport to prevent or cure
	[R] Life period of drugs
	[S] Requirements of factory premises

(a) 1-[Q], 2-[S]

(b) 1-[P], 2-[Q]

(c) 1-[R], 2-[P]

(d) 1-[P], 2-[S]



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Explanation:

Schedule FF specifies standards applicable to ophthalmic preparations and therefore matches [P]. Schedule M deals with requirements relating to factory premises and manufacturing practices and therefore matches [S]. Diseases for which drugs cannot claim prevention or cure are dealt with under Schedule J. The life period of drugs is covered under Schedule P.



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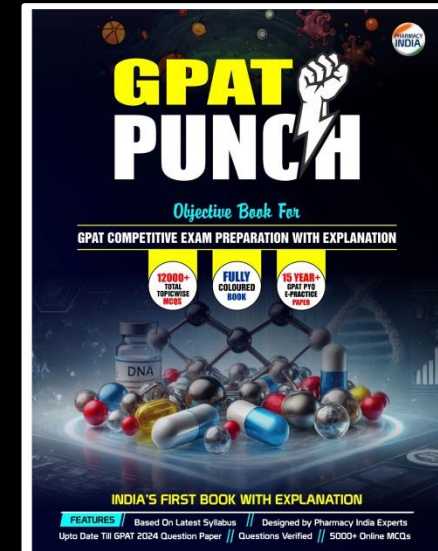
08. One of the forms mentioned below is used to issue license for wholesale of drugs other than specified in Schedule C, C1 and X. Choose the correct one [GATE-1999]

(a) 20 B

(b) 20 BB

(c) 21 B

(d) 20 A



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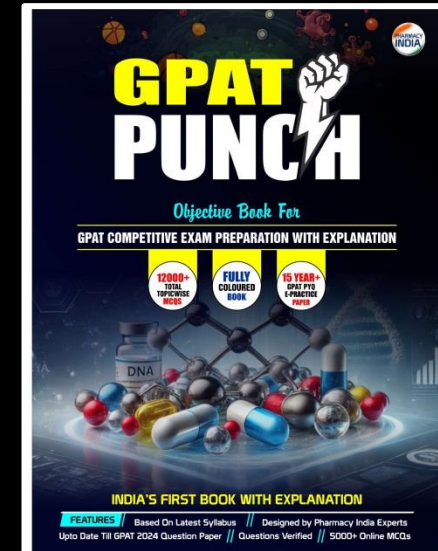
08. One of the forms mentioned below is used to issue license for wholesale of drugs other than specified in Schedule C, C1 and X. Choose the correct one [GATE-1999]

(a) 20 B

(b) 20 BB

(c) 21 B

(d) 20 A



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Explanation:

Form 20B is used for the wholesale sale of drugs other than those specified in Schedules C, C1 and X. Form 21B is associated with wholesale sale of drugs specified in Schedules C and C1. Form 20A is used for restricted retail sale under specified conditions. Form 20BB is not the appropriate form for the purpose asked.



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09. Given below from [P] to [S] are application forms for the specific purpose listed as per D and C Act. Match them
[GATE-1998]

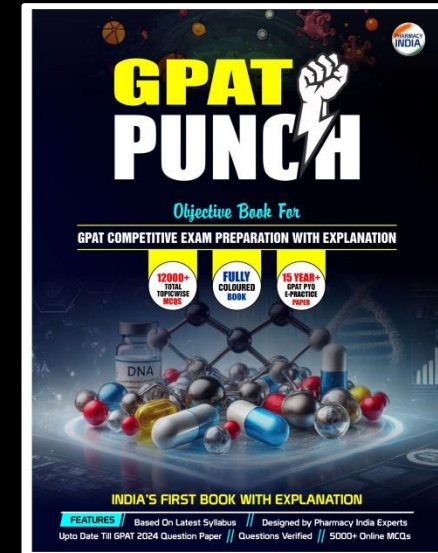
Purpose	Form
1. Manufacture of cosmetics	[P] Form No. 31
2. Retail sale of Schedule C and C1 drugs	[Q] Form No. 200
	[R] Form No. 20
	[S] Form No. 21

(a) 1-[P], 2-[R]

(b) 1-[Q], 2-[R]

(c) 1-[P], 2-[S]

(d) 1-[Q], 2-[S]



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09. Given below from [P] to [S] are application forms for the specific purpose listed as per D and C Act. Match them
[GATE-1998]

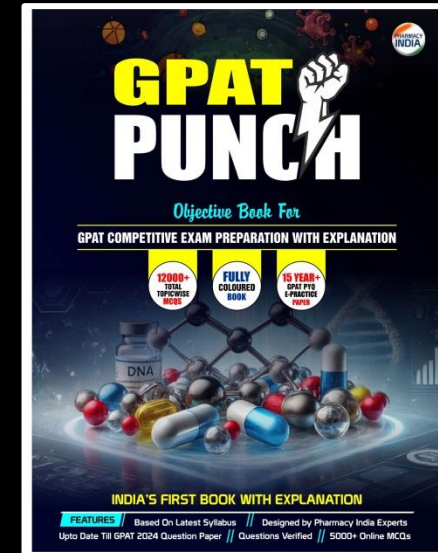
Purpose	Form
1. Manufacture of cosmetics	[P] Form No. 31
2. Retail sale of Schedule C and C1 drugs	[Q] Form No. 200
	[R] Form No. 20
	[S] Form No. 21

(a) 1-[P], 2-[R]

(b) 1-[Q], 2-[R]

(c) 1-[P], 2-[S]

(d) 1-[Q], 2-[S]



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Explanation:

Manufacture of cosmetics is associated with Form No. 31. Thus 1 corresponds to [P]. Retail sale of drugs specified in Schedules C and C1 is associated with Form No. 21 in the options. Therefore item 2 corresponds to [S].



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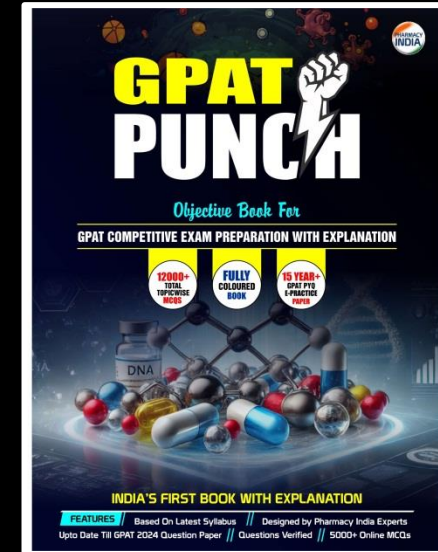
10. Licence for wholesale of drugs specified in Schedule C and C1 are issued in form [GATE-1995]

(a) 20 A

(b) 20 B

(c) 21 B

(d) 22



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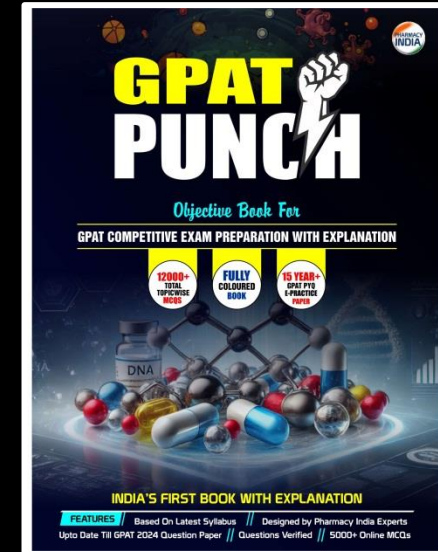
10. Licence for wholesale of drugs specified in Schedule C and C1 are issued in form [GATE-1995]

(a) 20 A

(b) 20 B

(c) 21 B

(d) 22



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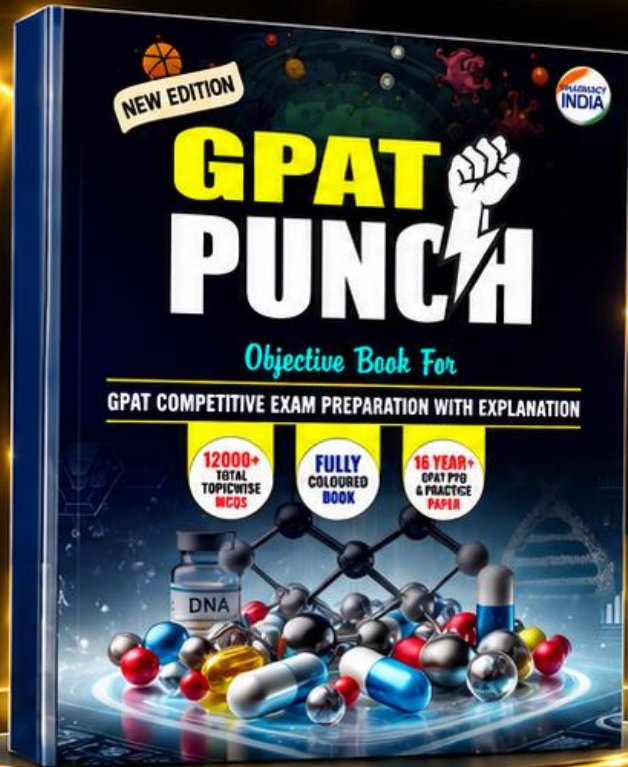
Explanation:

Form 21B is the licence form for wholesale sale of drugs specified in Schedules C and C1, excluding Schedule X drugs. Form 20B applies to wholesale drugs other than those in Schedules C, C1 and X. Form 20A is related to restricted retail sale.

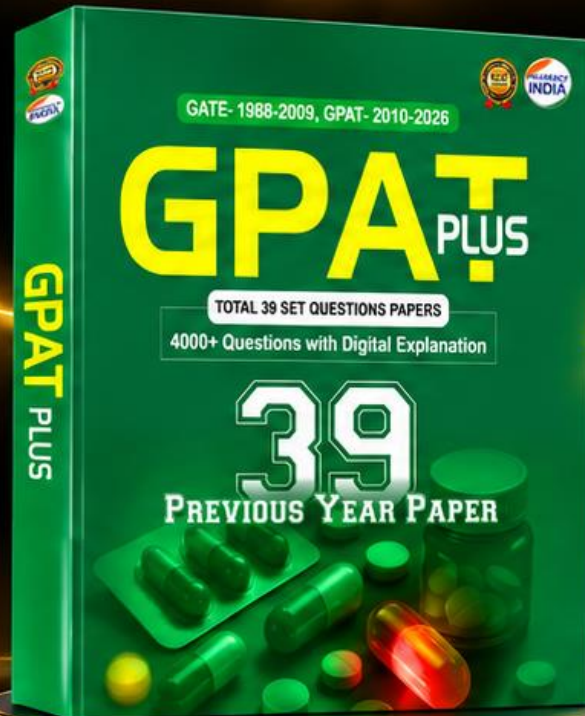


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11.

Listed below are some important drugs. Classify them as per the relevant Schedules of Drugs and Cosmetics Act [GATE-1994]

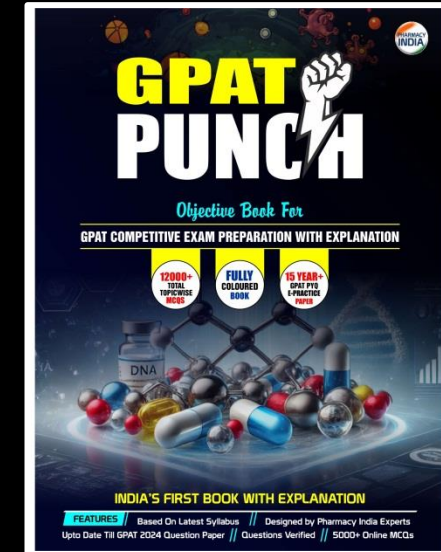
Drug	Schedule Code
1. Chlorpropamide	[P] Schedule G
2. Betamethasone benzoate	[Q] Schedule M
3. Amaranth	[R] Schedule H
4. Dexamphetamine	[S] Schedule Q
	[T] Schedule X

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

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

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

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



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11.

Listed below are some important drugs. Classify them as per the relevant Schedules of Drugs and Cosmetics Act [GATE-1994]

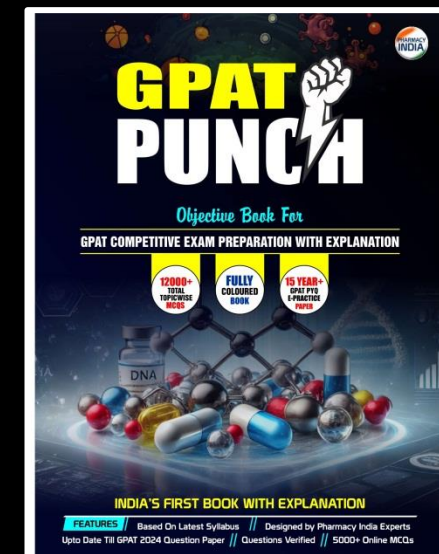
Drug	Schedule Code
1. Chlorpropamide	[P] Schedule G
2. Betamethasone benzoate	[Q] Schedule M
3. Amaranth	[R] Schedule H
4. Dexamphetamine	[S] Schedule Q
	[T] Schedule X

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

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

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

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



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Explanation:

Chlorpropamide is matched with Schedule G. Betamethasone benzoate is matched with Schedule H. Amaranth, being a permitted coal-tar colour, is associated with Schedule Q. Dexamphetamine is included under Schedule X.



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12.

A drug is deemed to be as indicated in 1 to 4 and the corresponding definitions are given in [P] to [T]. Match with the correct ones [GATE-1993]

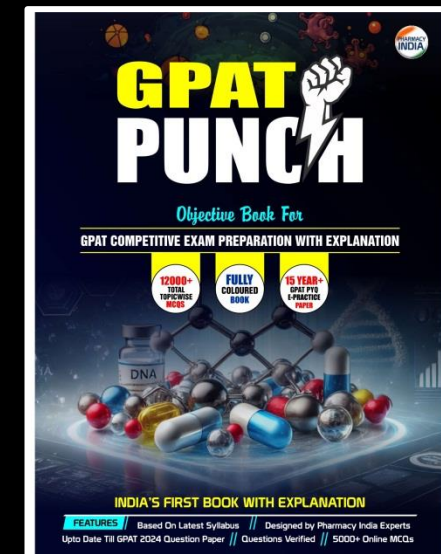
Type of Drug	Definition
1. Misbranded drug	[P] If it is marketed without prescription
2. Adulterated drug	[Q] If it is imported under a name which belongs to another drug
3. Spurious drug	[R] If it is not labelled in the prescribed manner
4. Drug of abuse	[S] If it contains any harmful or toxic substance
	[T] If it develops addiction

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

(b) 1-[P], 2-[Q], 3-[T], 4-[R]

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

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



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12.

A drug is deemed to be as indicated in 1 to 4 and the corresponding definitions are given in [P] to [T]. Match with the correct ones [GATE-1993]

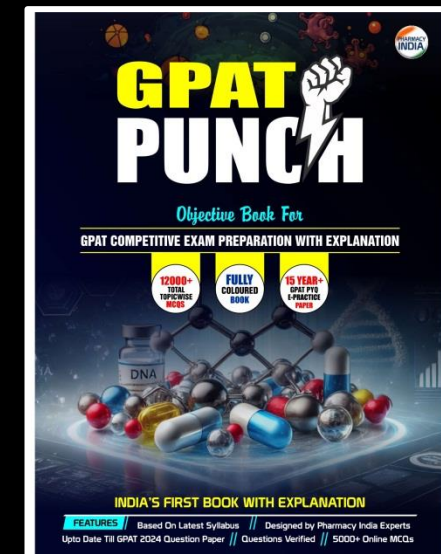
Type of Drug	Definition
1. Misbranded drug	[P] If it is marketed without prescription
2. Adulterated drug	[Q] If it is imported under a name which belongs to another drug
3. Spurious drug	[R] If it is not labelled in the prescribed manner
4. Drug of abuse	[S] If it contains any harmful or toxic substance
	[T] If it develops addiction

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

(b) 1-[P], 2-[Q], 3-[T], 4-[R]

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

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



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Explanation:

A misbranded drug includes one that is not labelled in the prescribed manner, so 1 matches [R]. An adulterated drug may contain harmful or toxic substances, so 2 matches [S]. A spurious drug may be imported under a name belonging to another drug, so 3 matches [Q]. A drug of abuse is associated with addiction, so 4 matches [T].



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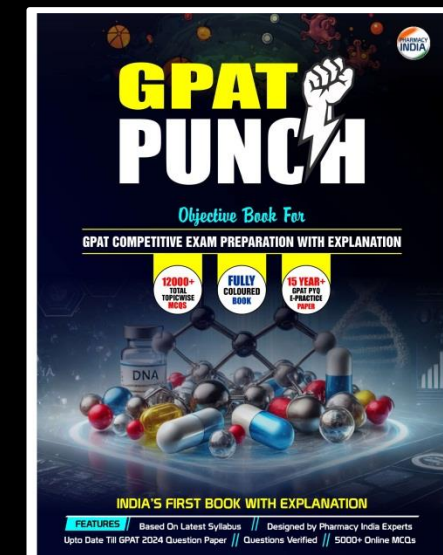


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13.

The following drugs are included under the schedules listed in [P] to [T]. Match them [GATE-1992]

Drug/Product	Schedule
1. Meprobamate	[P] Schedule E
2. Poisonous drugs	[Q] Schedule FF
3. Ophthalmic preparations	[R] Schedule C and C1
4. Biological and special products	[S] Schedule X
	[T] Schedule Q



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

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

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

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



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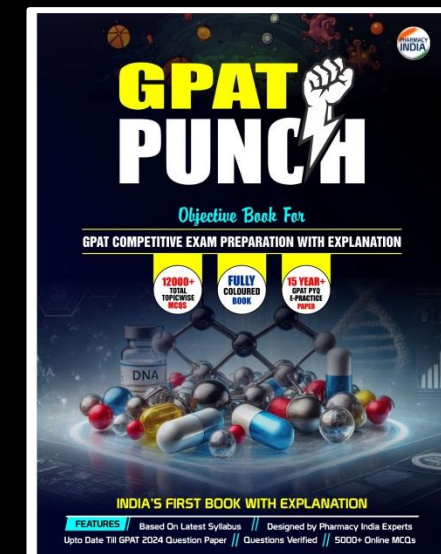


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13.

The following drugs are included under the schedules listed in [P] to [T]. Match them [GATE-1992]

Drug/Product	Schedule
1. Meprobamate	[P] Schedule E
2. Poisonous drugs	[Q] Schedule FF
3. Ophthalmic preparations	[R] Schedule C and C1
4. Biological and special products	[S] Schedule X
	[T] Schedule Q



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

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

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

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



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Explanation:

Meprobamate is placed under Schedule X in the matching given. Poisonous drugs correspond to Schedule E. Ophthalmic preparations are governed by Schedule FF. Biological and special products are included in Schedules C and C1.



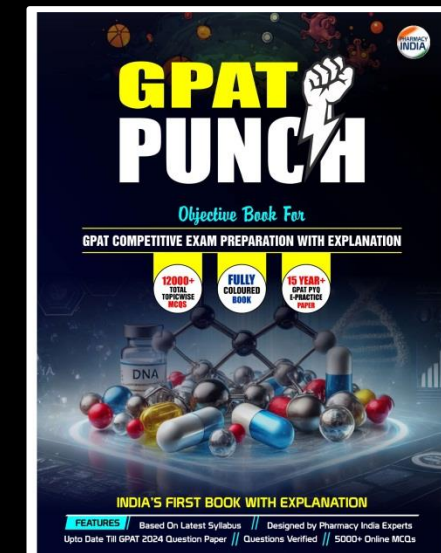
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14.

As per Drugs and Cosmetics Act, match correct schedule to their respective titles [GATE-1991]

Schedule	Title
1. Schedule P	[P] Standard for poison
2. Schedule Q	[Q] Standard for cosmetics
3. Schedule S	[R] Standard for ophthalmic preparations
4. Schedule FF	[S] Life period of the drug
	[T] Coal tar colour used in cosmetics



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

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

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

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



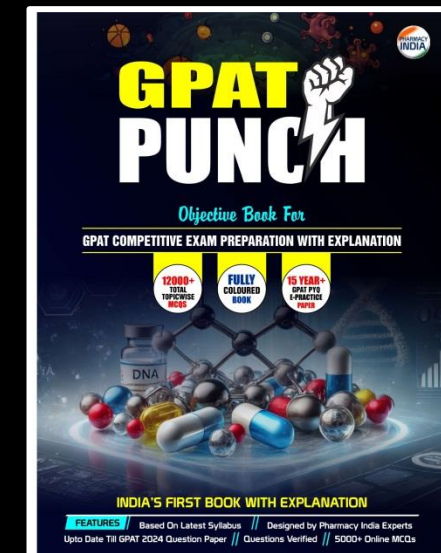
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14.

As per Drugs and Cosmetics Act, match correct schedule to their respective titles [GATE-1991]

Schedule	Title
1. Schedule P	[P] Standard for poison
2. Schedule Q	[Q] Standard for cosmetics
3. Schedule S	[R] Standard for ophthalmic preparations
4. Schedule FF	[S] Life period of the drug
	[T] Coal tar colour used in cosmetics



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

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

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

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



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Explanation:

Schedule P deals with the life period of drugs, so it matches [S]. Schedule Q deals with coal-tar colours permitted in cosmetics, so it matches [T]. Schedule S specifies standards for cosmetics and therefore matches [Q]. Schedule FF specifies standards for ophthalmic preparations and matches [R].



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15.

List below are some schedules 1 to 4 and the rules [P] to [T].
Match them correctly [GATE-1991]

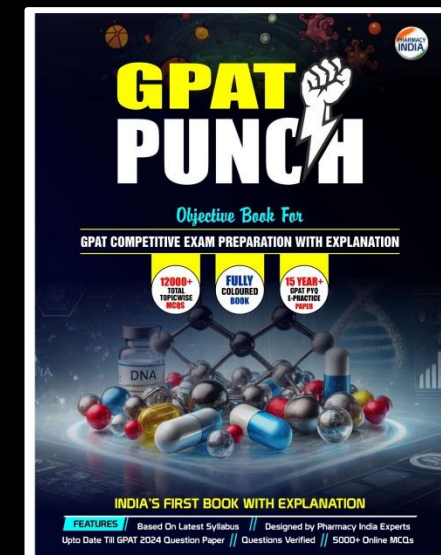
Schedule	Provision
1. C	[P] List of medicine required to be taken only under supervision of R.M.P
2. F	[Q] Biological and special products
3. G	[R] Provision applicable to vaccines, toxins, antigens and sera
4. M	[S] GMP requirement of factory premises, plants, equipment etc.
	[T] Standards for surgical dressing

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

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

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

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



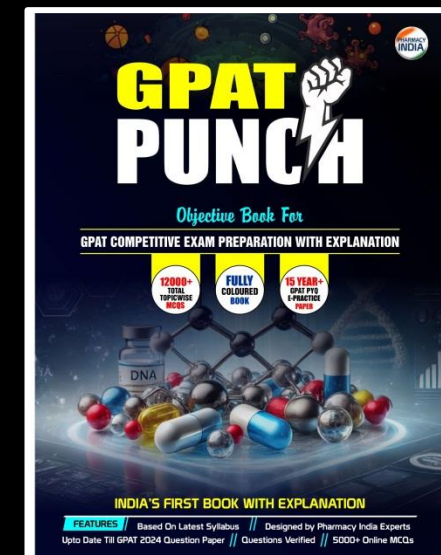
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15.

List below are some schedules 1 to 4 and the rules [P] to [T].
Match them correctly [GATE-1991]

Schedule	Provision
1. C	[P] List of medicine required to be taken only under supervision of R.M.P
2. F	[Q] Biological and special products
3. G	[R] Provision applicable to vaccines, toxins, antigens and sera
4. M	[S] GMP requirement of factory premises, plants, equipment etc.
	[T] Standards for surgical dressing



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

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

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

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



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Explanation:

Schedule C includes biological and special products and therefore matches [Q]. Schedule F contains provisions applicable to vaccines, toxins, antigens and sera and matches [R]. Schedule G covers medicines required to be taken under medical supervision and matches [P]. Schedule M concerns GMP requirements for factory premises, plant and equipment and matches [S].



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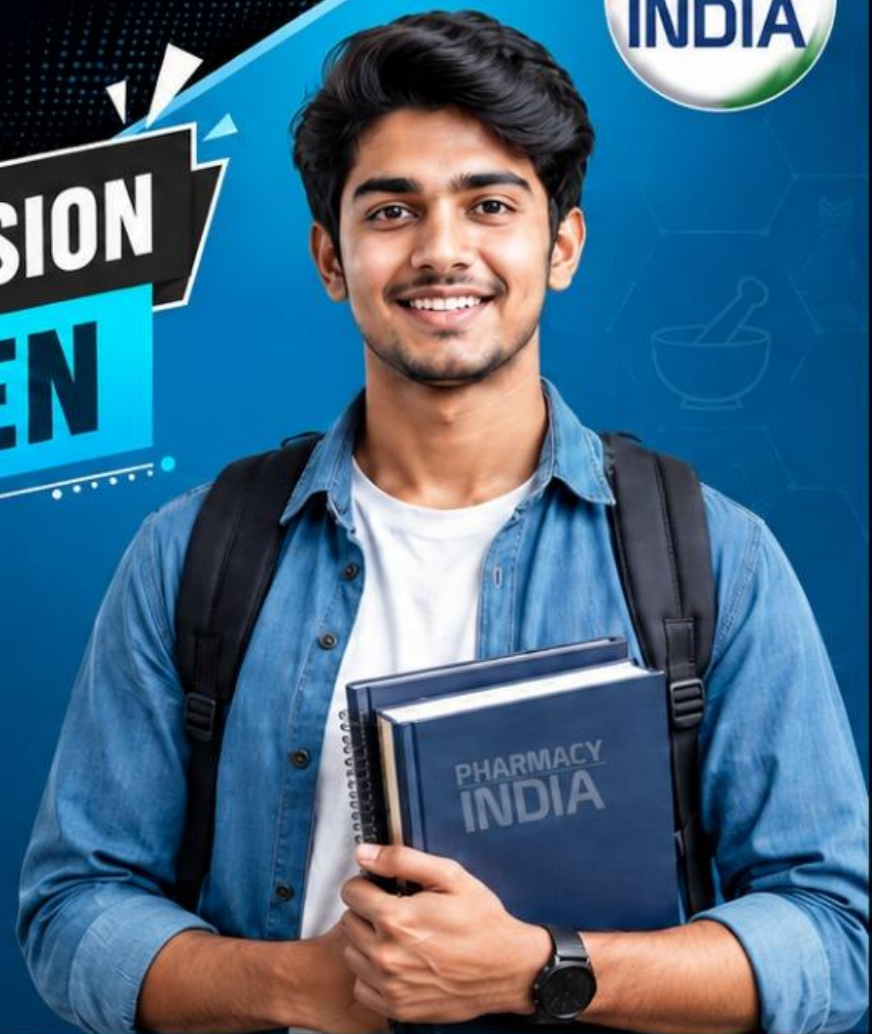
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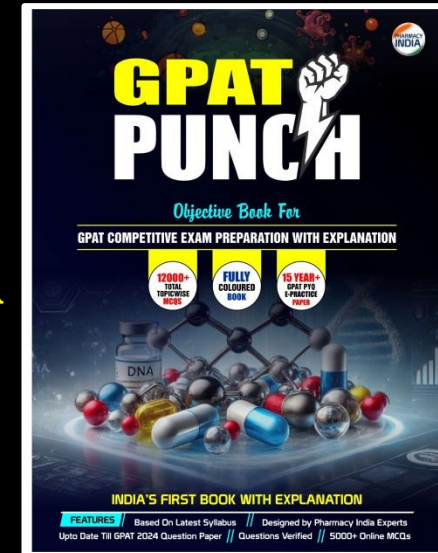


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16.

In Drugs and Cosmetics Act and Rules their under, list of substances that should be sold by retail only on prescription of registered medical practitioner is given in [GATE-1990]

- (a) Schedule H
- (b) Schedule V
- (c) Schedule X
- (d) Schedule Q



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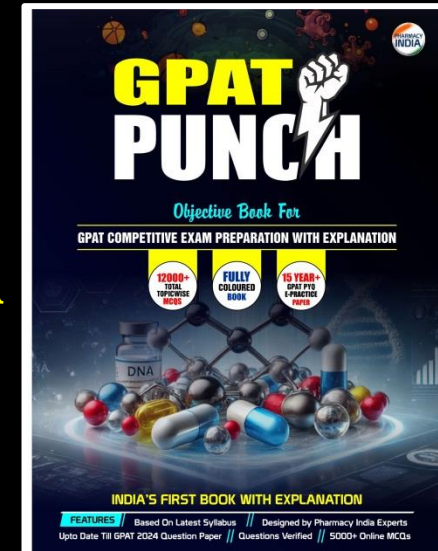


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16.

In Drugs and Cosmetics Act and Rules their under, list of substances that should be sold by retail only on prescription of registered medical practitioner is given in [GATE-1990]

- (a) Schedule H
- (b) Schedule V
- (c) Schedule X
- (d) Schedule Q



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Explanation:

Schedule H contains drugs that are required to be sold by retail only on the prescription of a Registered Medical Practitioner. Such drugs carry specific prescription-related labelling requirements. Schedule Q concerns permitted coal-tar colours for cosmetics. Schedule V is not the prescription-drug schedule asked here.



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17.

The following form under Schedule A of the Drugs and Cosmetics Act utilized for applying for licenses listed [P] to [T], match them [GATE-1990]

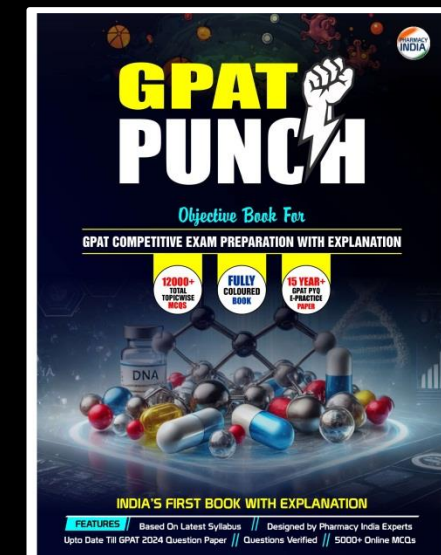
Form	Purpose
1. Form 8	[P] Application to import drugs for personal use
2. Form 12 A	[Q] Application for grant of license to sell, stock or distribute drug
3. Form 19	[R] Application to import biological products
4. Form 24 C	[S] Application to manufacture homeopathy drugs
	[T] Application to import drugs for research purposes

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

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

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

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



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17.

The following form under Schedule A of the Drugs and Cosmetics Act utilized for applying for licenses listed [P] to [T], match them [GATE-1990]

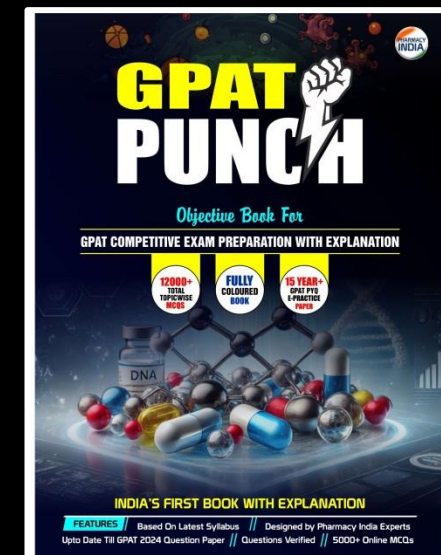
Form	Purpose
1. Form 8	[P] Application to import drugs for personal use
2. Form 12 A	[Q] Application for grant of license to sell, stock or distribute drug
3. Form 19	[R] Application to import biological products
4. Form 24 C	[S] Application to manufacture homeopathy drugs
	[T] Application to import drugs for research purposes

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

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

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

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



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Explanation:

Form 8 corresponds to application for import of the drugs indicated in the matching and therefore matches [R]. Form 12A relates to import for personal use and matches [P]. Form 19 is used for application to sell, stock or distribute drugs and matches [Q]. Form 24C is associated with manufacture of homoeopathic medicines and matches [S].



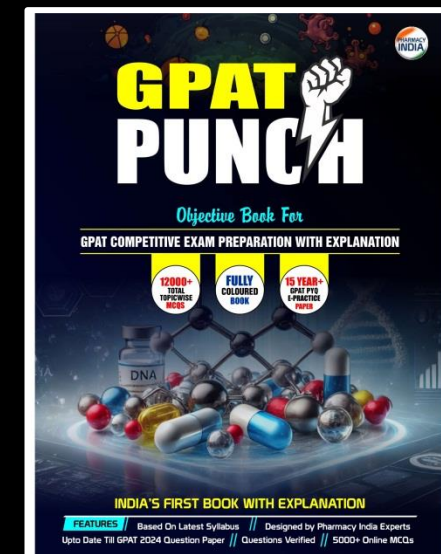
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18.

Match the following with the schedules listed in [P] to [T] correctly [GATE-1989]

List-I: Provision	List-II: Schedule
1. Requirements of factory premises	[P] Schedule P
2. Standards for disinfectant fluids	[Q] Schedule M
3. Life period of drugs	[R] Schedule N
4. List of minimum equipment for the efficient running of Pharmacy	[S] Schedule O



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

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

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

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



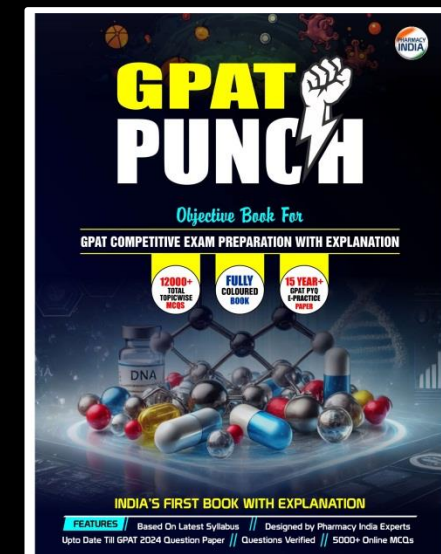
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18.

Match the following with the schedules listed in [P] to [T] correctly [GATE-1989]

List-I: Provision	List-II: Schedule
1. Requirements of factory premises	[P] Schedule P
2. Standards for disinfectant fluids	[Q] Schedule M
3. Life period of drugs	[R] Schedule N
4. List of minimum equipment for the efficient running of Pharmacy	[S] Schedule O



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

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

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

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



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Explanation:

Requirements of factory premises are covered under Schedule M, therefore 1 matches [Q]. Standards for disinfectant fluids are prescribed under Schedule O, therefore 2 matches [S]. The life period of drugs is specified under Schedule P, therefore 3 matches [P]. The list of minimum equipment required for efficient running of a pharmacy is given under Schedule N, therefore 4 matches [R].



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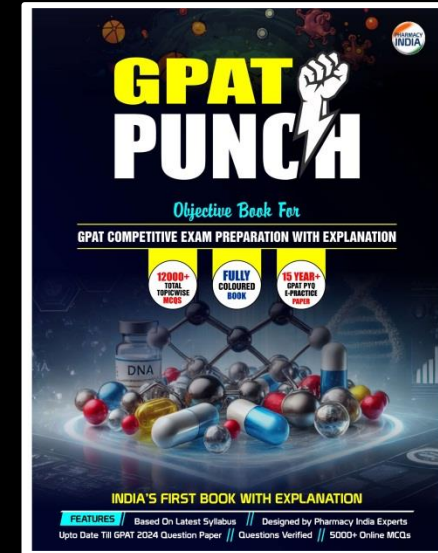
19. Licence to sell drug specified in Schedule C and C1 is given from number [GATE-1989]

(a) 19

(b) 18

(c) 21

(d) 24



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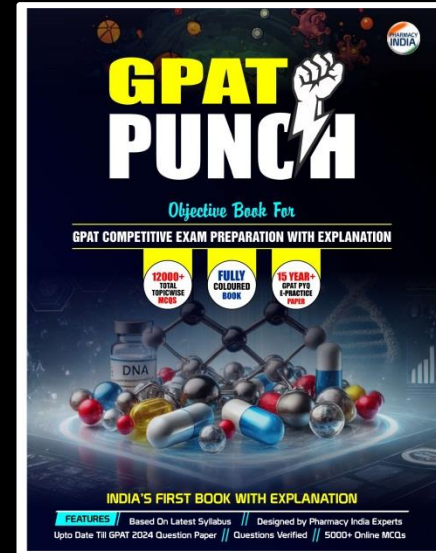
19. Licence to sell drug specified in Schedule C and C1 is given from number [GATE-1989]

(a) 19

(b) 18

(c) 21

(d) 24



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Explanation:

Form 21 is the licence form associated with retail sale of drugs specified in Schedules C and C1. Form 19 is generally the application form for obtaining sale licences. Forms 18 and 24 relate to different regulatory purposes.



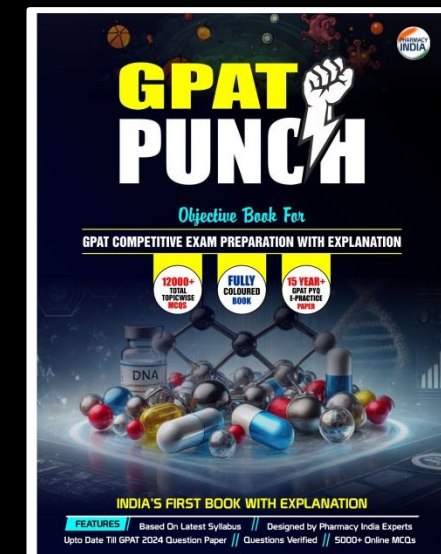
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20.

According to Drugs and Cosmetics Rules a list of schedule are as follows. Match the appropriate statement [P] to [S] with them [GATE-1988]

Schedule	Statement
1. Schedule G	[P] Drugs used under medical supervision
2. Schedule P	[Q] Drug used only under medical supervision
3. Schedule J	[R] Minimum equipment needed for a retail pharmacy
4. Schedule N	[S] Diseases that a drug should not claim to cure



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

(b) 1-[T], 2-[R], 3-[S], 4-[P]

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

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



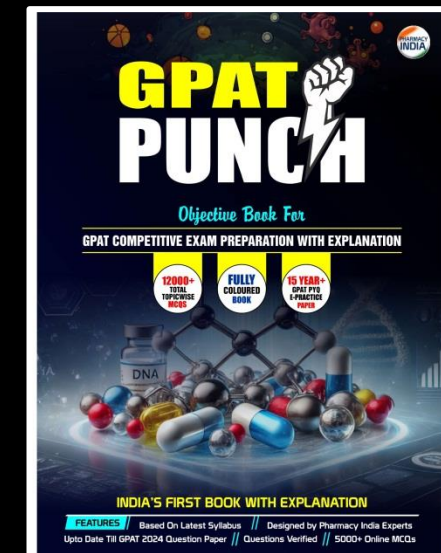
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Schedule	Statement
1. Schedule G	[P] Drugs used under medical supervision
2. Schedule P	[Q] Drug used only under medical supervision
3. Schedule J	[R] Minimum equipment needed for a retail pharmacy
4. Schedule N	[S] Diseases that a drug should not claim to cure



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

(b) 1-[T], 2-[R], 3-[S], 4-[P]

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

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



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Explanation:

Schedule G deals with drugs to be used under medical supervision and matches [P]. Schedule P specifies the life period of drugs and therefore matches [T]. Schedule J contains diseases for which drugs cannot claim prevention or cure and matches [S]. Schedule N gives the minimum equipment required for efficient running of a pharmacy and matches [R].



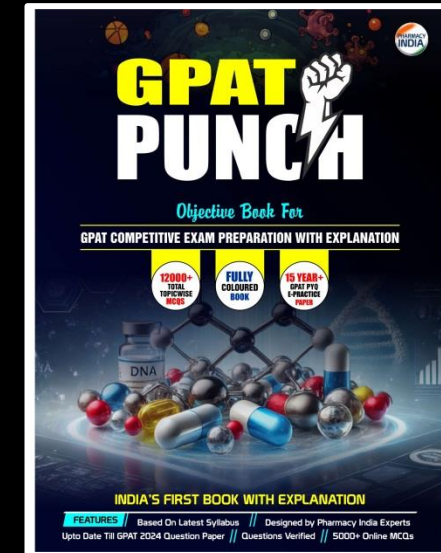
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21. Given the drug and their schedule [P] to [T]. Match them correctly [GATE-1988]

Drug	Schedule
1. B-Complex tablets	[P] Schedule C1
2. Calcium gluconate injection	[Q] Schedule F
3. Small pox vaccine	[R] Schedule H
4. Ampicillin capsule	[S] Schedule L



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

(b) 1-[P], 2-[Q], 3-[T], 4-[S]

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

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



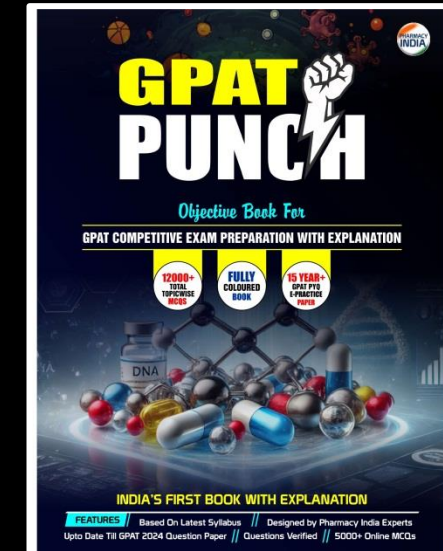
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21. Given the drug and their schedule [P] to [T]. Match them correctly [GATE-1988]

Drug	Schedule
1. B-Complex tablets	[P] Schedule C1
2. Calcium gluconate injection	[Q] Schedule F
3. Small pox vaccine	[R] Schedule H
4. Ampicillin capsule	[S] Schedule L



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

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

(b) 1-[P], 2-[Q], 3-[T], 4-[S]

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



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Explanation:

B-Complex tablets are matched with Schedule C1. Calcium gluconate injection is matched with Schedule C. Smallpox vaccine corresponds to Schedule F in the given matching. Ampicillin capsule is matched with Schedule H.



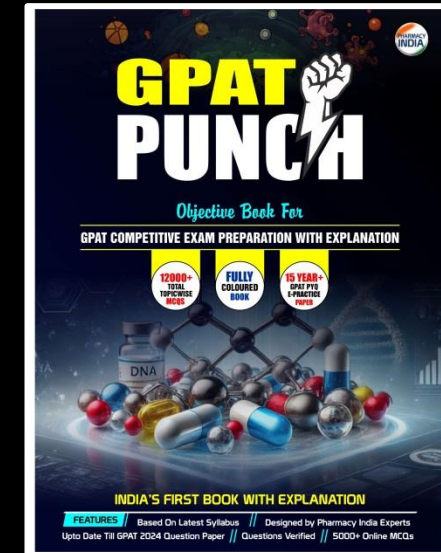
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22. In which year Drugs Rules were framed under the Drugs Act 1940?

- (a) 1940
- (b) 1945
- (c) 1960
- (d) 2000



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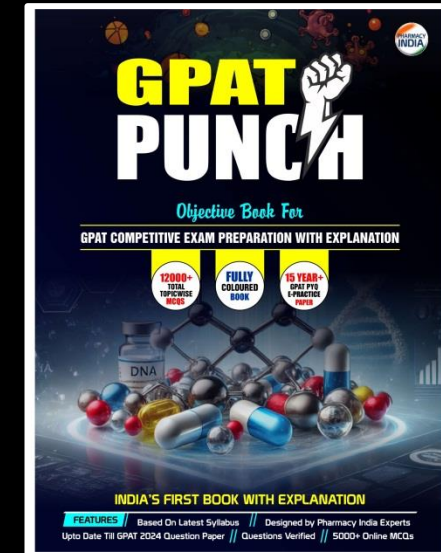
22. In which year Drugs Rules were framed under the Drugs Act 1940?

(a) 1940

(b) 1945

(c) 1960

(d) 2000



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Explanation:

The Drugs and Cosmetics Act was enacted in 1940. The detailed Drugs and Cosmetics Rules were subsequently framed in 1945. These Rules provide procedural and technical requirements for implementing the Act. The years 1960 and 2000 are not the year in which these Rules were originally framed.



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23.

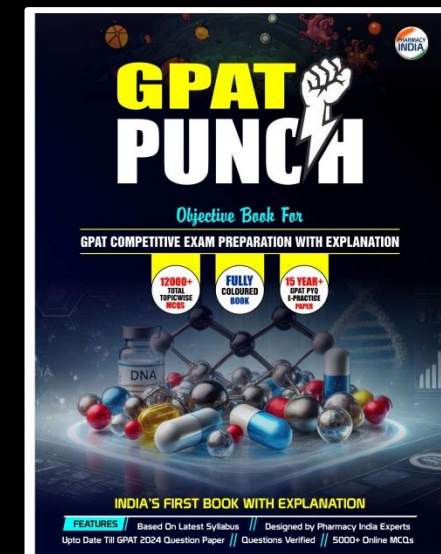
As per Schedule P Carbenicillin Sodium Powder should be stored in/at

(a) A cool place

(b) A cold place

(c) Temperature not exceeding 5°C

(d) Well closed container, protect from light, in cool place, protect from light



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23.

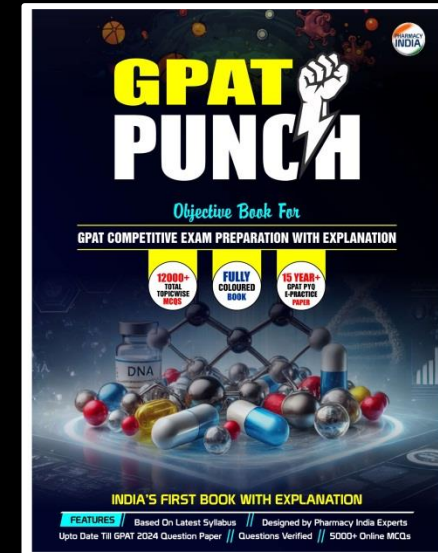
As per Schedule P Carbenicillin Sodium Powder should be stored in/at

(a) A cool place

(b) A cold place

(c) Temperature not exceeding 5°C

(d) Well closed container, protect from light, in cool place, protect from light



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Explanation:

Schedule P specifies storage conditions and life periods for various drugs. Carbenicillin Sodium Powder requires protection from environmental factors that may affect its stability. The option specifying a well-closed container, protection from light and storage in a cool place gives the complete condition. The other options mention only a single storage parameter and are incomplete.



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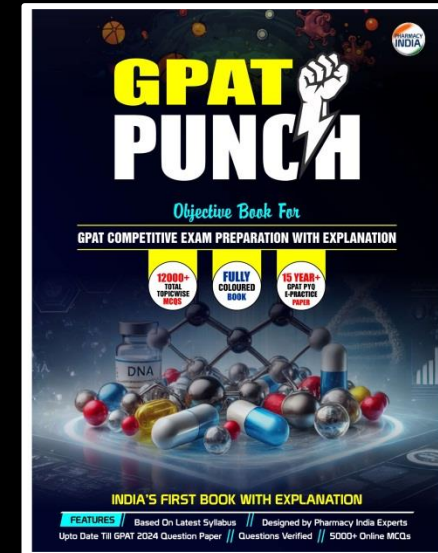


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24.

As per D and C Act “Schedule N” is related with

- (a) List of maximum equipments for efficiently running pharmacy**
- (b) Area for opening retail pharmacy**
- (c) List of minimum equipment for efficiently running pharmacy**
- (d) Area required to open wholesale drug store**



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24.

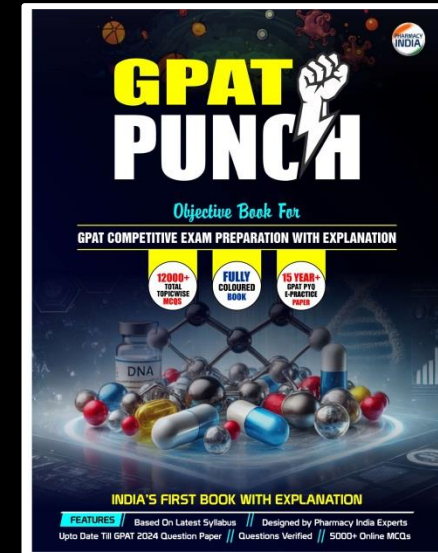
As per D and C Act "Schedule N" is related with

(a) List of maximum equipments for efficiently running pharmacy

(b) Area for opening retail pharmacy

(c) List of minimum equipment for efficiently running pharmacy

(d) Area required to open wholesale drug store



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Explanation:

Schedule N gives the list of minimum equipment required for the efficient running of a pharmacy. It does not prescribe a list of maximum equipment. It also does not primarily specify the floor area required for retail or wholesale drug stores. Its purpose is to ensure availability of essential equipment in a pharmacy.



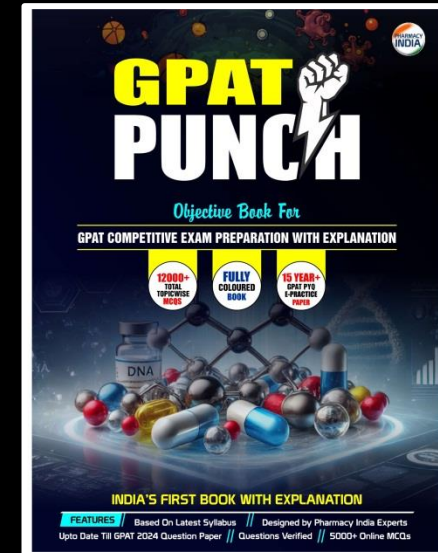
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25. For licence granted to sell by retail drugs specified in Schedule C and C1, drugs form no. required is

- (a) 20
- (b) 20B
- (c) 21B
- (d) 21



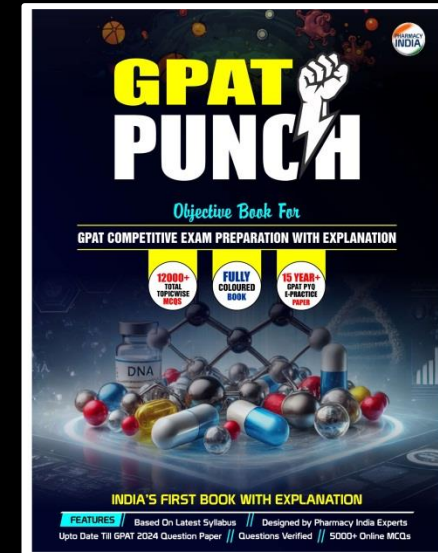
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25. For licence granted to sell by retail drugs specified in Schedule C and C1, drugs form no. required is

- (a) 20
- (b) 20B
- (c) 21B
- (d) 21**



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Explanation:

Form 21 is the licence granted for retail sale of drugs specified in Schedules C and C1. Form 21B is used for the corresponding wholesale licence. Form 20 relates to retail sale of drugs other than those specified in Schedules C, C1 and X. Form 20B is used for wholesale sale of those other drugs.



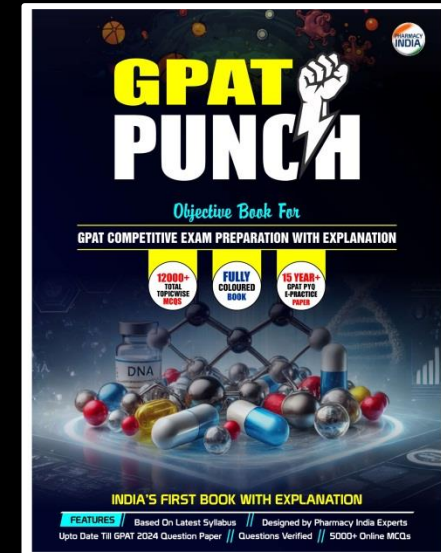
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26. For parenteral preparations in glass containers, the total minimum recommended area for Small Volume Injectables is:

- (a) 250 square metres
- (b) 400 square metres
- (c) 500 square metres
- (d) 350 square metres



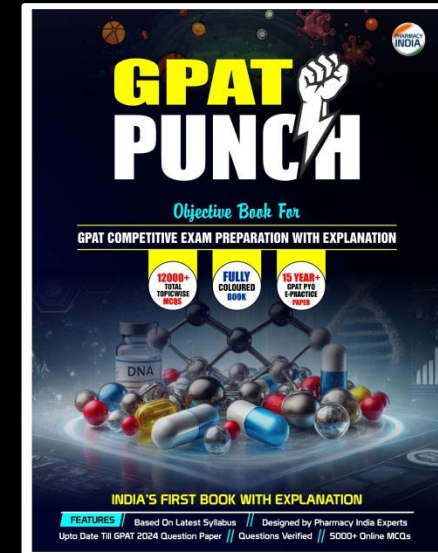
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26. For parenteral preparations in glass containers, the total minimum recommended area for Small Volume Injectables is:

- (a) 250 square metres
- (b) 400 square metres
- (c) 500 square metres
- (d) 350 square metres



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Explanation:

Under Schedule M, parenteral preparations in glass containers require separate manufacturing areas for different operations. For Small Volume Injectables (SVIs), a minimum of 150 square metres for basic installation and 100 square metres for ancillary area is recommended. Therefore, the total recommended area is $150 + 100 = 250$ square metres. For Large Volume Parenterals in glass containers, the requirement is different, with 150 square metres each for basic and ancillary areas.



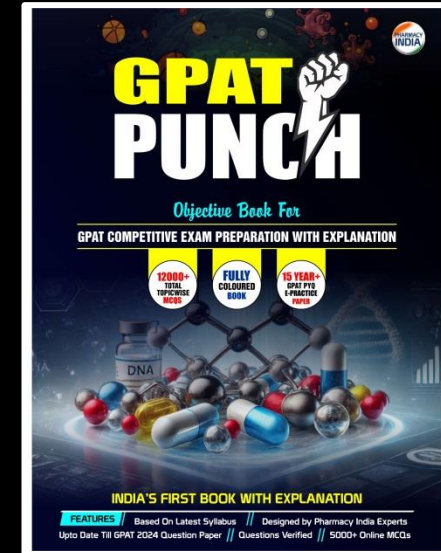
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27. Proforma for sending memorandum is included in which schedule?

- (a) D
- (b) A
- (c) FF
- (d) Q



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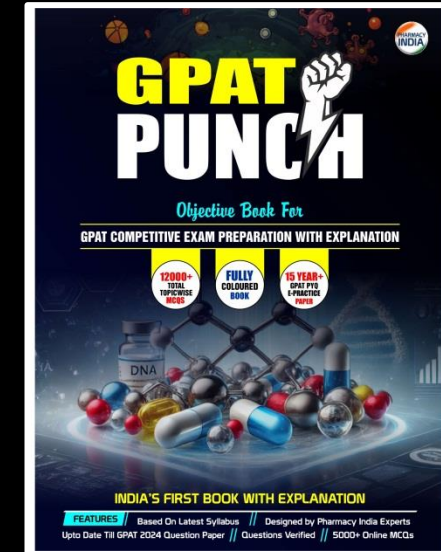
27. Proforma for sending memorandum is included in which schedule?

(a) D

(b) A

(c) FF

(d) Q



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Explanation:

Schedule A contains various prescribed forms and formats used under the Drugs and Cosmetics Rules. The proforma for sending a memorandum is therefore associated with Schedule A. Schedule FF deals with ophthalmic preparations. Schedule Q is associated with permitted coal-tar colours used in cosmetics.

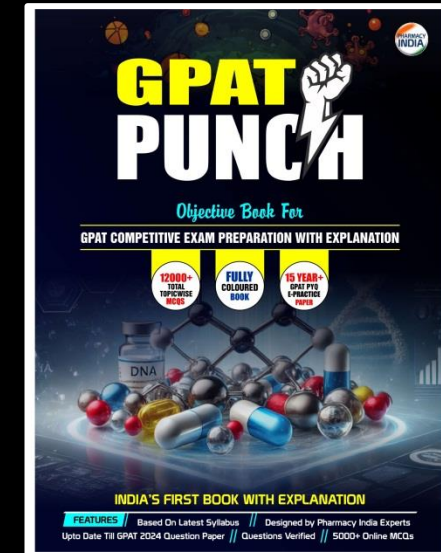


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- 28. As per D and C Act Schedule T states about**
- (a) Requirements of factory premises for the manufacture of medical devices**
 - (b) Requirements of factory premises for the manufacture of Homeopathy drugs**
 - (c) Requirements of factory premises for the manufacture of Ayurveda, Siddha and Unani drugs**
 - (d) Requirements of factory premises for the manufacture of Allopathy drugs**



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28.

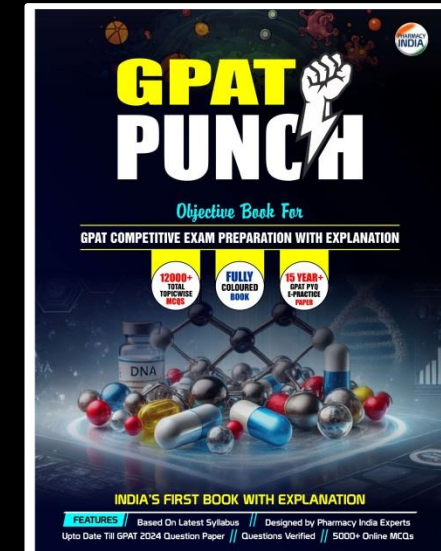
As per D and C Act Schedule T states about

(a) Requirements of factory premises for the manufacture of medical devices

(b) Requirements of factory premises for the manufacture of Homeopathy drugs

(c) Requirements of factory premises for the manufacture of Ayurveda, Siddha and Unani drugs

(d) Requirements of factory premises for the manufacture of Allopathy drugs



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Explanation:

Schedule T lays down Good Manufacturing Practices and requirements of premises for Ayurvedic, Siddha and Unani medicines. It specifies conditions intended to maintain appropriate manufacturing quality. Homeopathic manufacturing premises are dealt with under different provisions. Allopathic drug manufacturing requirements are principally associated with Schedule M.



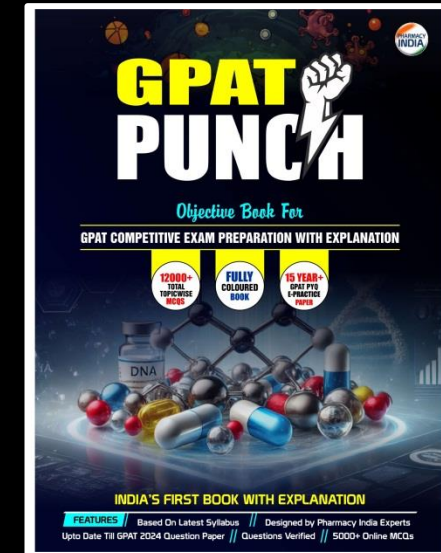
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29. Grant of licence to manufacture a drug requires

- (a) Form 24
- (b) Form 25
- (c) Form 26
- (d) Form 27



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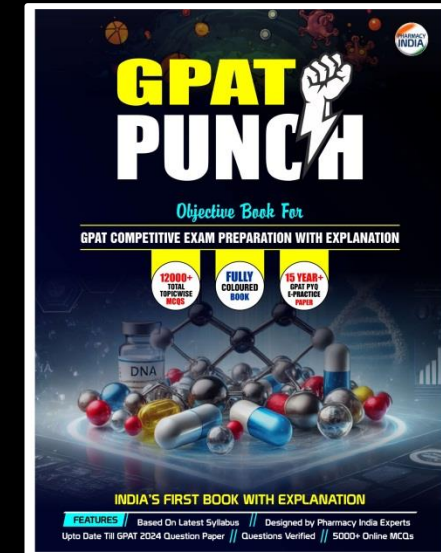
29. Grant of licence to manufacture a drug requires

(a) Form 24

(b) Form 25

(c) Form 26

(d) Form 27



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Explanation:

Form 24 is used as an application for obtaining a manufacturing licence for drugs other than those specified in Schedules C and C1. The licence granted on such an application is issued in Form 25. Form 27 is associated with an application concerning manufacture of Schedule C and C1 drugs. Form 26 serves a different licensing purpose.



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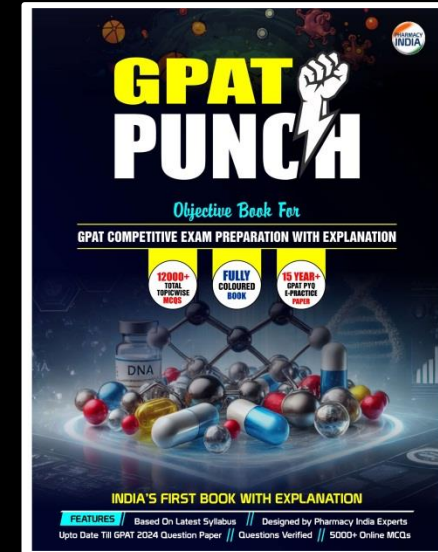


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30.

List of drugs which should be used by patient under medical supervision is covered under _____ as per D & C Act

- (a) Schedule G
- (b) Schedule H
- (c) Schedule I
- (d) Schedule J



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30.

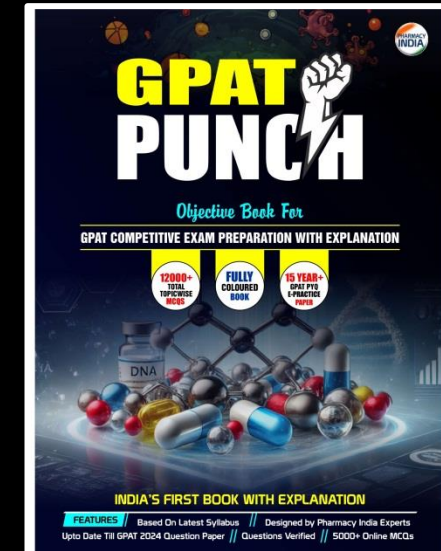
List of drugs which should be used by patient under medical supervision is covered under _____ as per D & C Act

(a) Schedule G

(b) Schedule H

(c) Schedule I

(d) Schedule J



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Explanation:

Schedule G contains drugs that are required to be taken under medical supervision. Their labels carry a caution regarding the danger of taking the preparation except under medical supervision. Schedule H is primarily associated with prescription-only retail sale. Schedule J deals with diseases for which drugs cannot claim prevention or cure.



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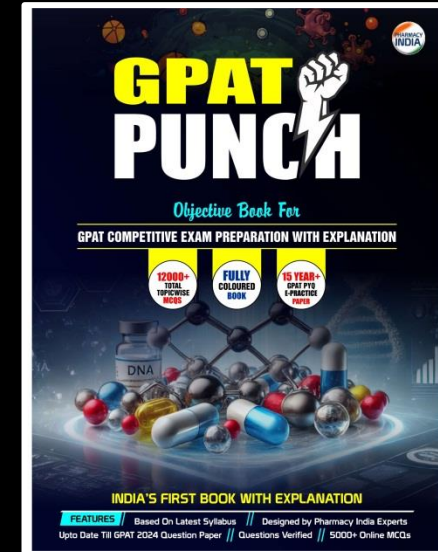
31. Schedule T of Ayurvedic Siddha and Unani medicines was introduced in

(a) 1982

(b) 1988

(c) 2000

(d) 2011



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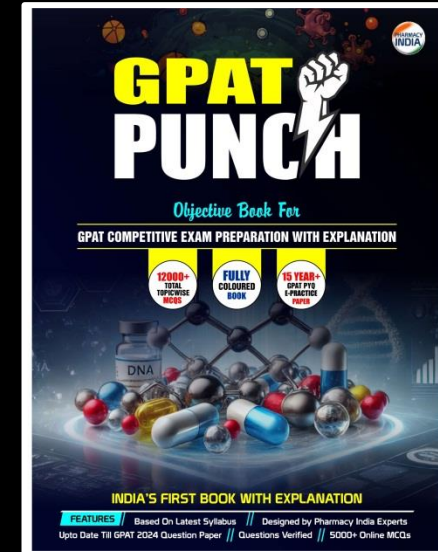
31. Schedule T of Ayurvedic Siddha and Unani medicines was introduced in

(a) 1982

(b) 1988

(c) 2000

(d) 2011



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Explanation:

Schedule T establishes requirements relating to Good Manufacturing Practices for Ayurvedic, Siddha and Unani medicines. In the examination material, its introduction is associated with the year 2000. The earlier years given in the options relate to other regulatory amendments and schedules.



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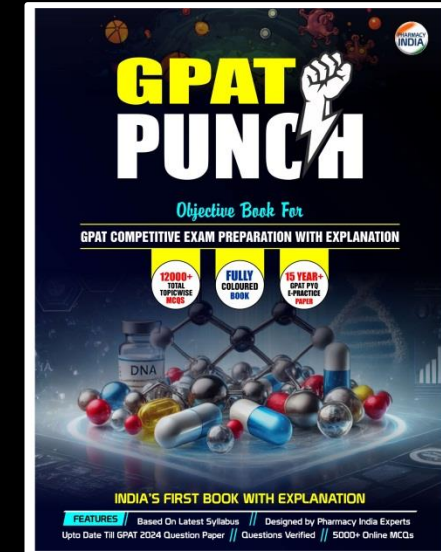


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32.

If blood group is “AB” then colour label used is

- (a) Red
- (b) White
- (c) Yellow
- (d) Blue



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32.

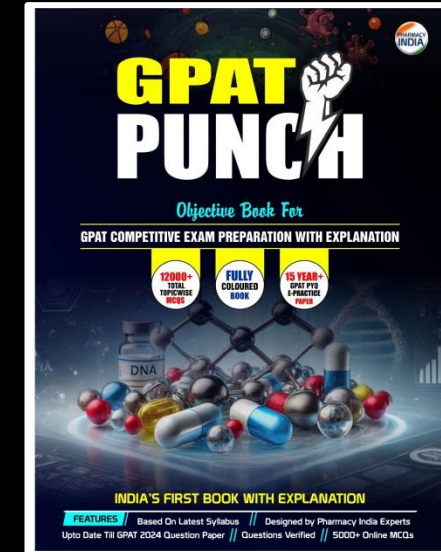
If blood group is “AB” then colour label used is

(a) Red

(b) White

(c) Yellow

(d) Blue



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Explanation:

Blood-group identification historically uses specified colour coding on blood labels. In the system, blood group AB is represented by a white label. The other colours listed correspond to other blood-group identification categories. Such colour coding helps rapid visual differentiation of blood groups.



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33.

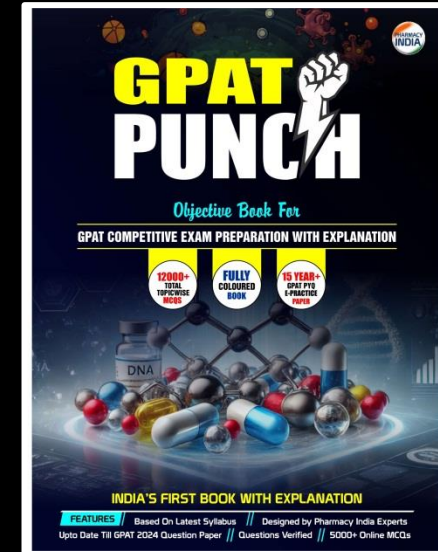
**Application for grant of a licence to manufacture
Ayurvedic, Siddha or Unani drugs requires**

(a) Form 24D

(b) Form 25E

(c) Form 20

(d) Form 21



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33.

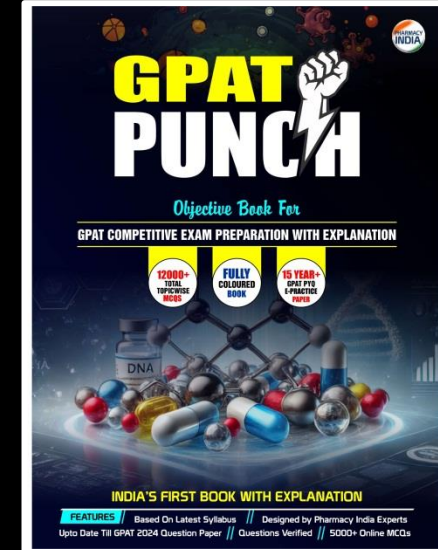
Application for grant of a licence to manufacture
Ayurvedic, Siddha or Unani drugs requires

(a) Form 24D

(b) Form 25E

(c) Form 20

(d) Form 21



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Explanation:

Form 24D is used for applying for a licence to manufacture Ayurvedic, Siddha or Unani drugs. Forms 20 and 21 are associated with sale licences for other categories of drugs. Form 25E is not the application form specified here.



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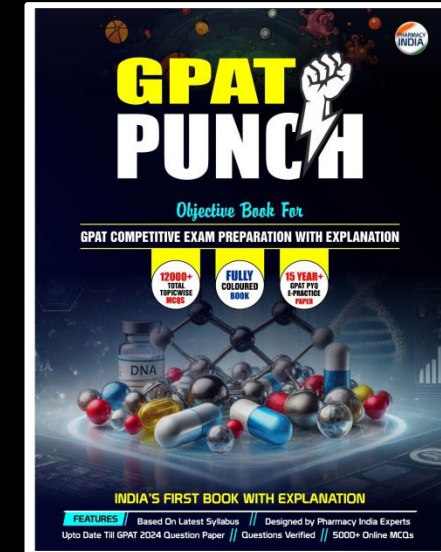


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34.

List of coal tar colours permitted to be used in cosmetics is covered under

- (a) Schedule O
- (b) Schedule P
- (c) Schedule Q
- (d) Schedule R



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34.

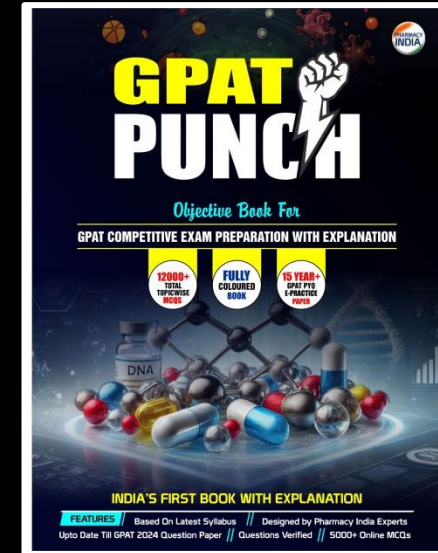
List of coal tar colours permitted to be used in cosmetics is covered under

(a) Schedule O

(b) Schedule P

(c) Schedule Q

(d) Schedule R



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Explanation:

Schedule Q contains the list of coal-tar colours permitted for use in cosmetics. Schedule O deals with standards for disinfectant fluids. Schedule P concerns the life period and related requirements of drugs. Schedule R is associated with standards for certain contraceptive products.



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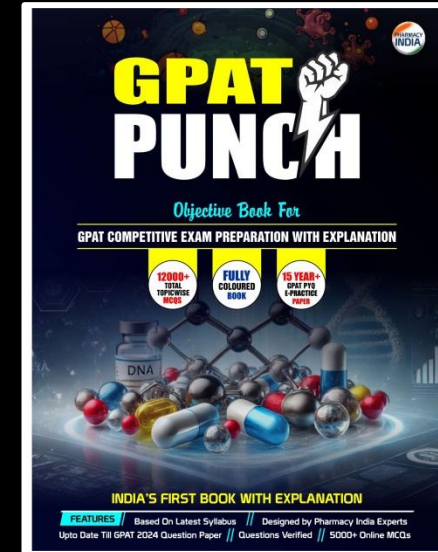
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35.

“Ampicillin capsule should be used within 24 months.”

This statement comes under

- (a) Schedule C
- (b) Schedule R
- (c) Schedule M
- (d) Schedule P



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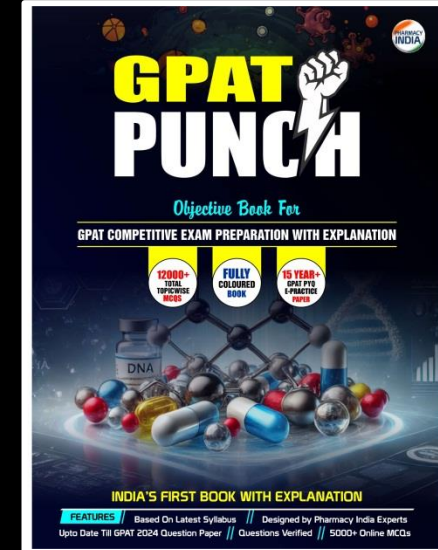
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35.

“Ampicillin capsule should be used within 24 months.”

This statement comes under

- (a) Schedule C
- (b) Schedule R
- (c) Schedule M
- (d) Schedule P**



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Explanation:

Schedule P specifies the life period of various drugs and pharmaceutical preparations. A statement that Ampicillin capsules should be used within 24 months refers to the product's specified life period. Schedule M primarily deals with manufacturing premises and GMP requirements. Schedules C and R have different regulatory purposes.



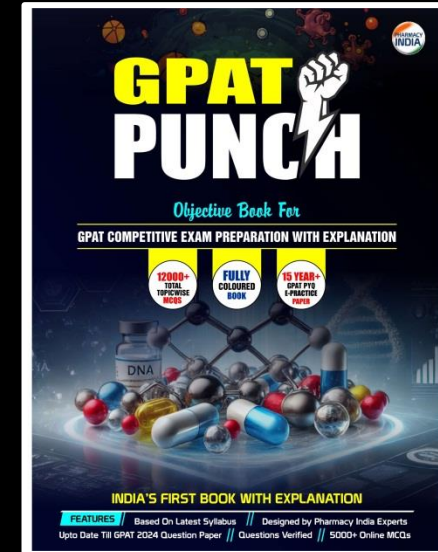
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36. One of the following drug is NOT in Schedule 'H'

- (a) Nalidixic acid
- (b) Naproxen
- (c) Phenometrazine
- (d) Nebivolol



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36.

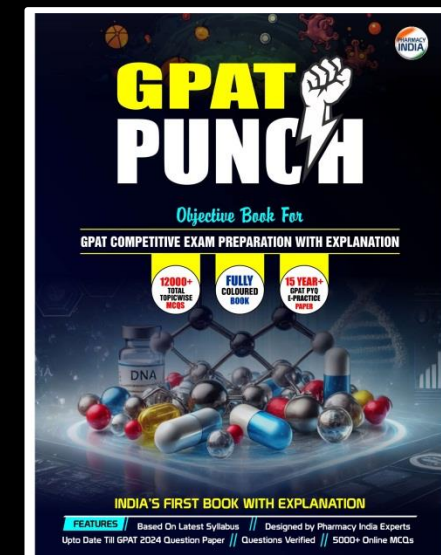
One of the following drug is NOT in Schedule 'H'

(a) Nalidixic acid

(b) Naproxen

(c) Phenometrazine

(d) Nebivolol



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Explanation:

Nalidixic acid, Naproxen and Nebivolol are treated as Schedule H drugs. Phenometrazine is not classified under Schedule H. Schedule H principally contains medicines requiring sale on prescription of a Registered Medical Practitioner. Certain habit-forming or specially controlled drugs are regulated differently.



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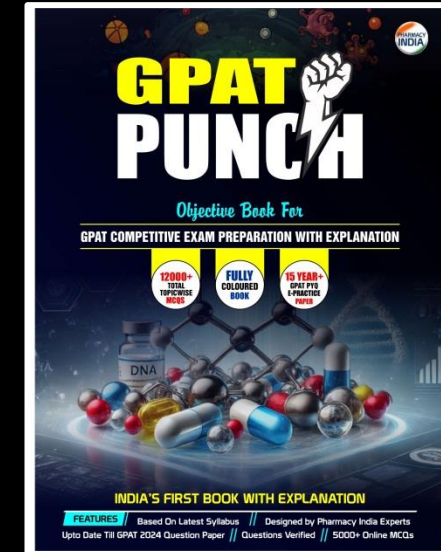


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37.

A drug is said to be Spurious drug when

- (a) It is marketed without prescription
- (b) It is imported under a name which belongs to another drug
- (c) It is not labelled in the prescribed manner
- (d) It contains any harmful or toxic substance



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37.

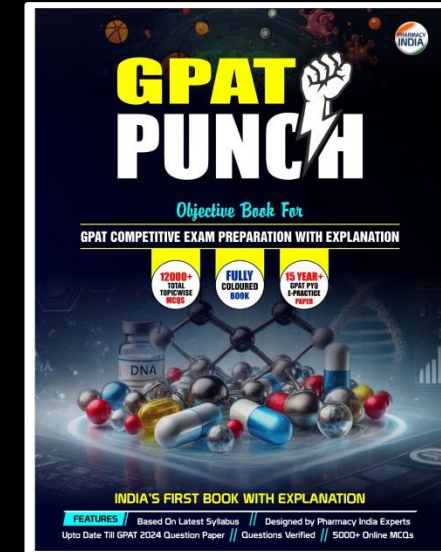
A drug is said to be Spurious drug when

(a) It is marketed without prescription

(b) It is imported under a name which belongs to another drug

(c) It is not labelled in the prescribed manner

(d) It contains any harmful or toxic substance



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Explanation:

A spurious drug includes a drug imported under a name that belongs to another drug. Improper labelling is associated with the concept of a misbranded drug. Presence of harmful or toxic substances is associated with adulteration. Marketing without prescription does not itself define a drug as spurious.



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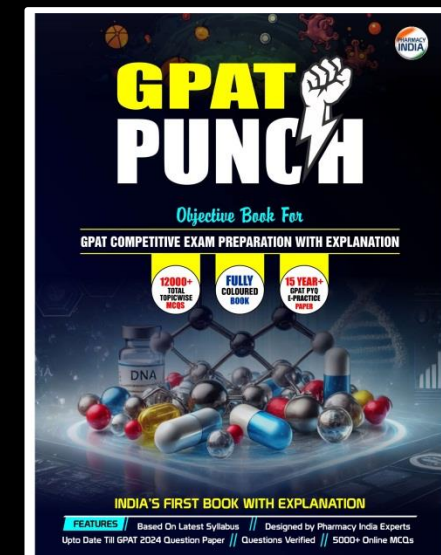


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38.

Government Analyst has to submit his report of drug analysis in _____ copies in Form _____

- (a) Two, 1
- (b) Three, 13
- (c) Four, 13
- (d) Five, 2



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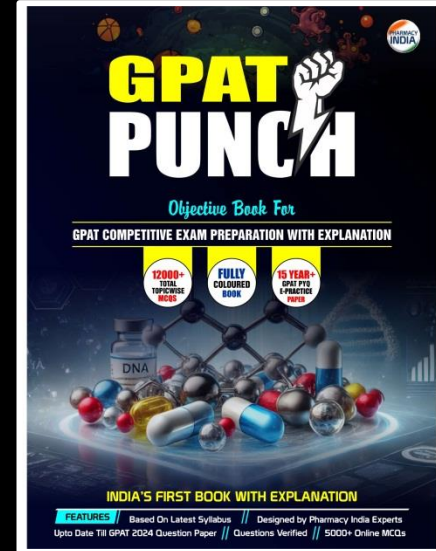
38. Government Analyst has to submit his report of drug analysis in _____ copies in Form _____

(a) Two, 1

(b) Three, 13

(c) Four, 13

(d) Five, 2



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Explanation:

The Government Analyst submits the prescribed analytical report in Form 13. The report is prepared in three copies. Form 13 records the result of analysis of the drug sample submitted for examination. The alternative combinations of copy number and form number are incorrect.



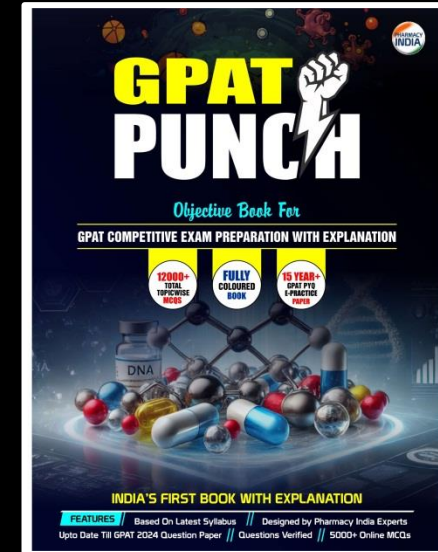
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39. The basic installation area for capsules according to Schedule M is

- (a) 25 sq. m
- (b) 30 sq. m
- (c) 20 sq. m
- (d) 15 sq. m



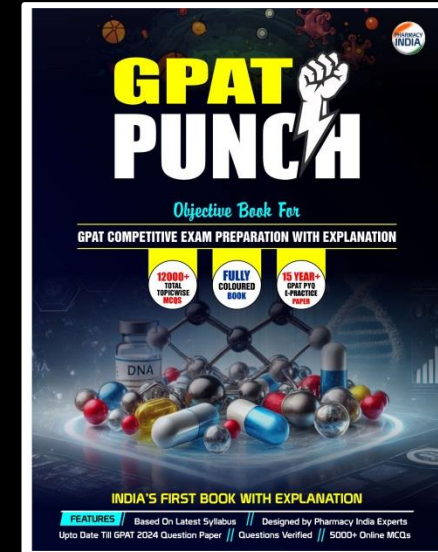
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39. The basic installation area for capsules according to Schedule M is

- (a) 25 sq. m
- (b) 30 sq. m
- (c) 20 sq. m
- (d) 15 sq. m



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Explanation:

Schedule M lays down requirements relating to manufacturing premises, plant and equipment. For capsule manufacturing, the basic installation area asked is 25 square metres. This space helps accommodate essential manufacturing equipment and operations.



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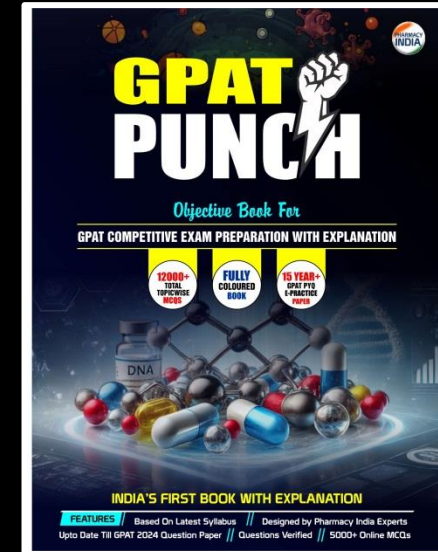


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40.

Application for a loan licence to manufacture cosmetics should be made in

- (a) Form 31B
- (b) Form 31A
- (c) Form 31L
- (d) Form 31E



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40.

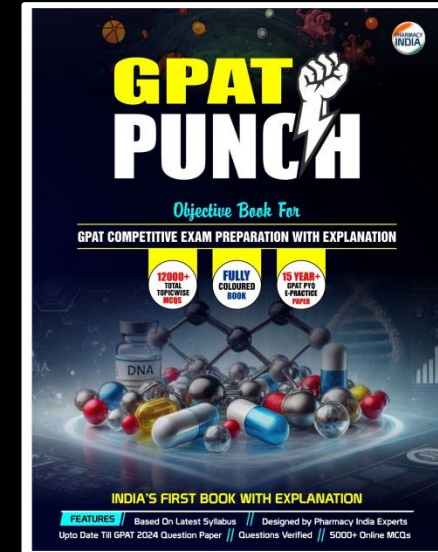
Application for a loan licence to manufacture cosmetics should be made in

(a) Form 31B

(b) Form 31A

(c) Form 31L

(d) Form 31E



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Explanation:

Form 31A is used to make an application for a loan licence for the manufacture of cosmetics. A loan licence allows manufacture using the facilities of another licensed manufacturer subject to prescribed conditions. Form 31 is associated with an ordinary application for manufacture of cosmetics.



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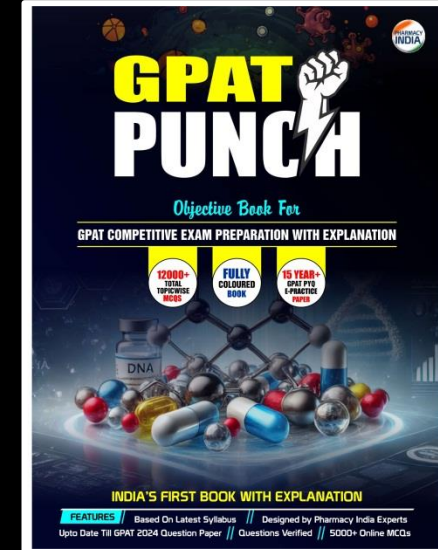


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41.

The chairman of the Drug Technical Advisory Board is

- (a) The Drug Controller of India
- (b) The Director, Central Drug Laboratory, Kolkata
- (c) The President, Pharmacy Council of India
- (d) The Director General of Health Service



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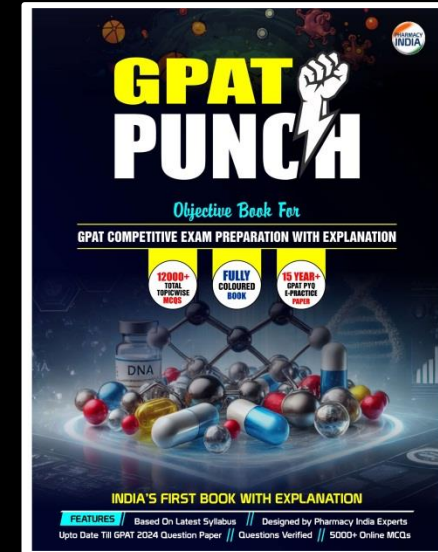


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41.

The chairman of the Drug Technical Advisory Board is

- (a) The Drug Controller of India
- (b) The Director, Central Drug Laboratory, Kolkata
- (c) The President, Pharmacy Council of India
- (d) The Director General of Health Service**



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Explanation:

The Director General of Health Services serves as the Chairman of the Drugs Technical Advisory Board. DTAB advises the Central and State Governments on technical matters arising from administration of the Drugs and Cosmetics Act. The Drugs Controller of India is an important member but is not the Chairman. The PCI President and CDL Director also do not hold the chairmanship.



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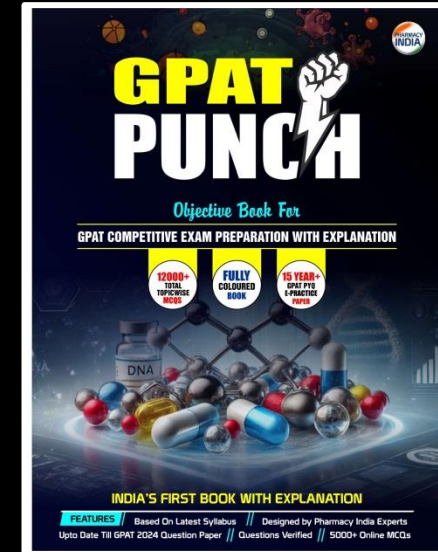
42. **Schedule M and Y were introduced in D & C Act in the year**

(a) 1982

(b) 1988

(c) 1985

(d) 1980



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42.

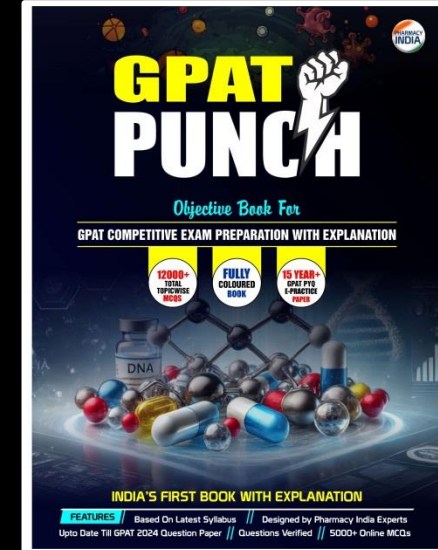
Schedule M and Y were introduced in D & C Act in the year

(a) 1982

(b) 1988

(c) 1985

(d) 1980



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Explanation:

Schedule M deals with Good Manufacturing Practices and requirements for pharmaceutical manufacturing premises. Schedule Y historically dealt with requirements relating to new drugs and clinical trials. Schedules M and Y were introduced in 1988.



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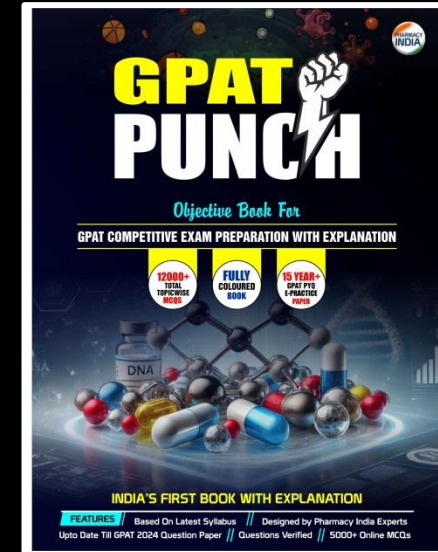


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43.

Schedule F1 deals with the provisions applicable to the production of

- (a) Ophthalmic preparations**
- (b) Surgical dressings**
- (c) Bacterial and viral vaccines**
- (d) Umbilical tape**



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43.

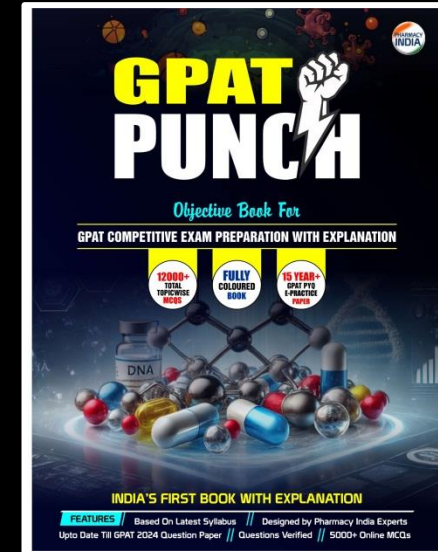
Schedule F1 deals with the provisions applicable to the production of

(a) Ophthalmic preparations

(b) Surgical dressings

(c) Bacterial and viral vaccines

(d) Umbilical tape



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Explanation:

Schedule F1 contains provisions applicable to the production of bacterial and viral vaccines. These biological products require specialized manufacturing and quality-control requirements. Ophthalmic preparations are principally covered under Schedule FF. Surgical dressings and umbilical tape are dealt with under other parts of the regulatory schedules.



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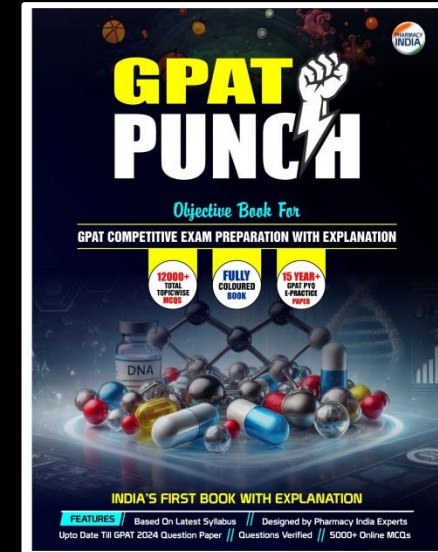


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44.

For the manufacture of drugs specified in Schedule C and C1 applicant has to make an application to the licensing authority in Form _____ and the licence is issued in Form _____

- (a) 24, 25
- (b) 27, 28
- (c) 24F, 25F
- (d) 25, 25



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44.

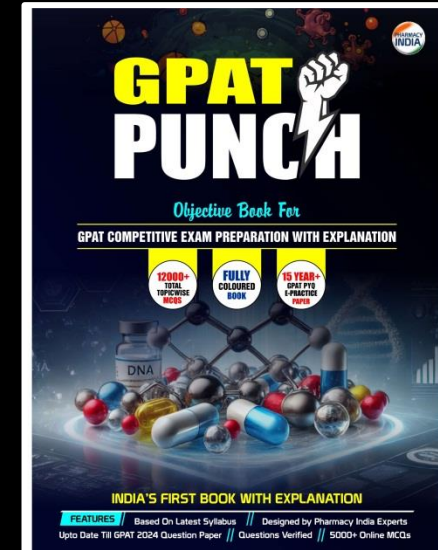
For the manufacture of drugs specified in Schedule C and C1 applicant has to make an application to the licensing authority in Form _____ and the licence is issued in Form _____

(a) 24, 25

(b) 27, 28

(c) 24F, 25F

(d) 25, 25



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Explanation:

An application to manufacture drugs specified in Schedules C and C1 is made in Form 27. After approval, the manufacturing licence is issued in Form 28. Forms 24 and 25 apply to manufacture of drugs outside the specified Schedule C and C1 category. The remaining combinations do not represent the required application-and-licence sequence.



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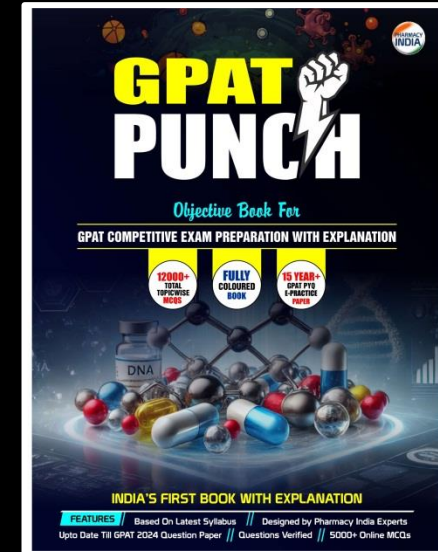


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45.

Application for permission to import or manufacture new drugs to undertake clinical trials should be made in

- (a) Form 21
- (b) Form 44
- (c) Form 11
- (d) Form 12A



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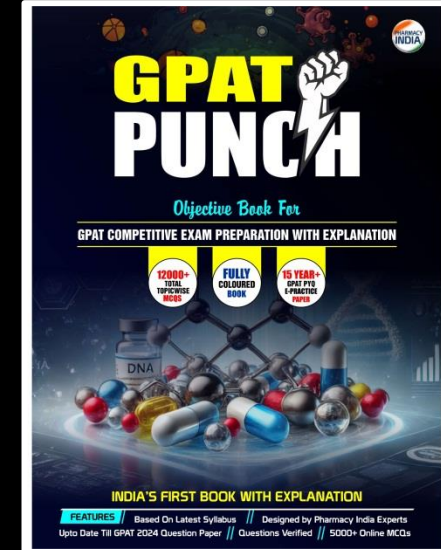


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45.

Application for permission to import or manufacture new drugs to undertake clinical trials should be made in

- (a) Form 21
- (b) Form 44**
- (c) Form 11
- (d) Form 12A



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Explanation:

In the regulatory framework referred to Form 44 was used for an application seeking permission to import or manufacture a new drug and undertake clinical trials. Form 21 is associated with retail sale of Schedule C and C1 drugs. Form 12A concerns import of small quantities for personal use. Form 11 has a different purpose under the Rules.



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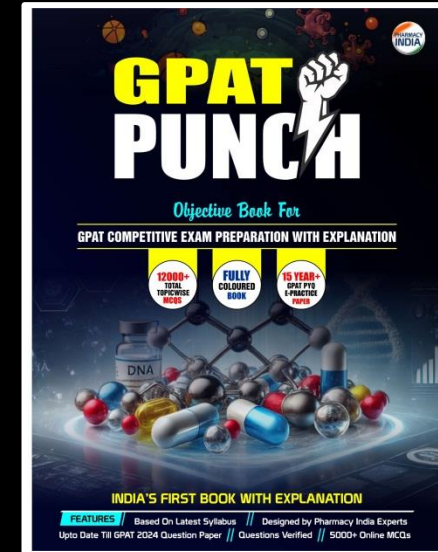


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46.

The warning “To be sold by retail on the prescription of a Registered Medical practitioner only” shall appear on the label of

- (a) Schedule H drugs and Schedule G drugs
- (b) Schedule X drug
- (c) Schedule H drugs and Schedule X drugs
- (d) Schedule G drug



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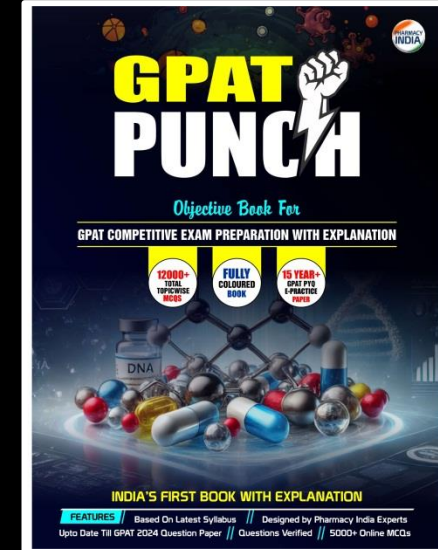


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46.

The warning “To be sold by retail on the prescription of a Registered Medical practitioner only” shall appear on the label of

- (a) Schedule H drugs and Schedule G drugs
- (b) Schedule X drug
- (c) Schedule H drugs and Schedule X drugs
- (d) Schedule G drug



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Explanation:

Schedule H medicines are prescription-only drugs and carry the prescribed retail-sale warning. Schedule X medicines are also subject to strict prescription controls and corresponding labelling requirements. Schedule G uses a different caution concerning administration under medical supervision. Therefore combining Schedules.



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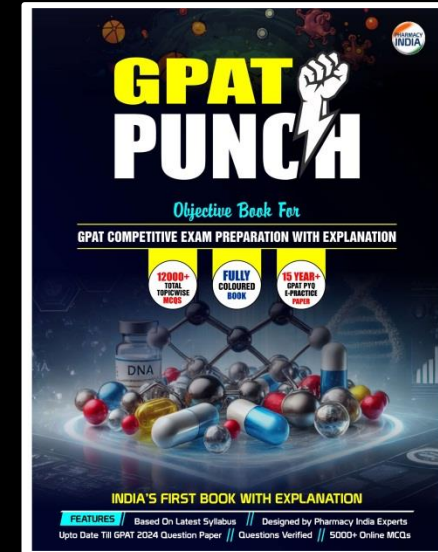


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47.

Which of the following diseases, for which a drug may not purport to prevent or cure, is NOT covered under Schedule J to the Drugs and Cosmetics Rules, 1945?

- (a) Diabetes
- (b) Obesity
- (c) Hypertension
- (d) Parkinsonism



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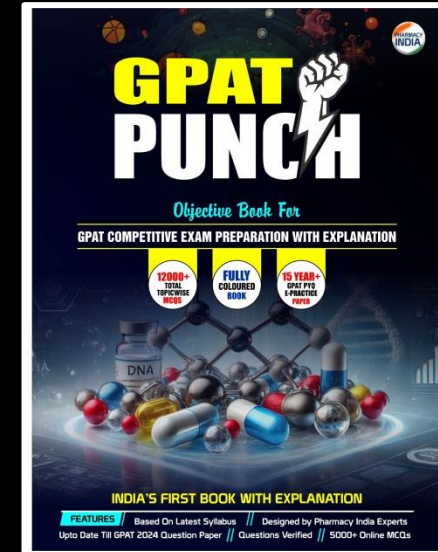


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47.

Which of the following diseases, for which a drug may not purport to prevent or cure, is NOT covered under Schedule J to the Drugs and Cosmetics Rules, 1945?

- (a) Diabetes
- (b) Obesity
- (c) Hypertension
- (d) Parkinsonism



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Explanation:

Schedule J contains diseases and ailments for which a drug is prohibited from claiming prevention or cure. In the Schedule J list relevant to Diabetes, Obesity, and High or Low Blood Pressure (Hypertension) are included.



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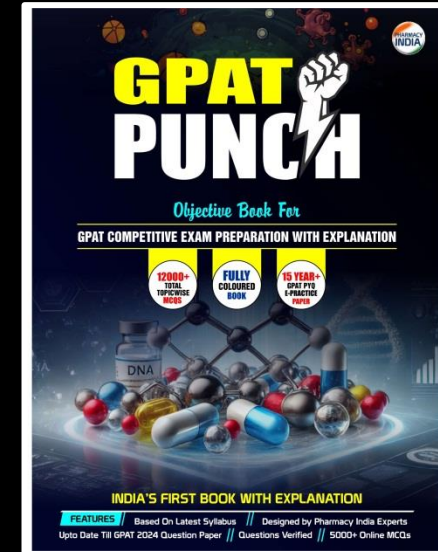


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48.

Schedule X of Drugs and Cosmetics Act comprises

- (a) List of incurable diseases
- (b) Guidelines for clinical Trials
- (c) List of generic drugs
- (d) None of the above



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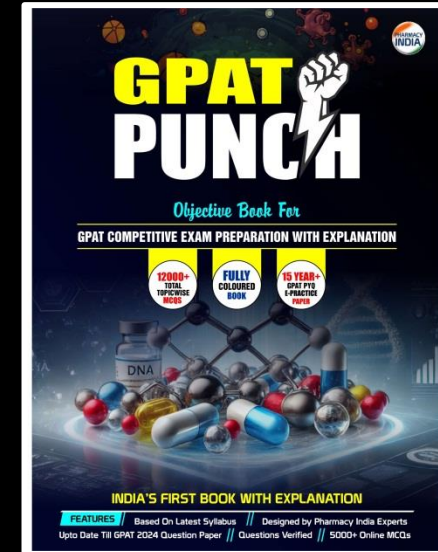


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48.

Schedule X of Drugs and Cosmetics Act comprises

- (a) List of incurable diseases
- (b) Guidelines for clinical Trials
- (c) List of generic drugs
- (d) None of the above**



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Explanation:

Schedule X contains specially controlled drugs subject to stringent sale, storage and record requirements. It does not contain the list of diseases that drugs cannot claim to cure. Clinical-trial requirements were historically associated with Schedule Y rather than Schedule X. Schedule X is also not simply a list of generic drugs.



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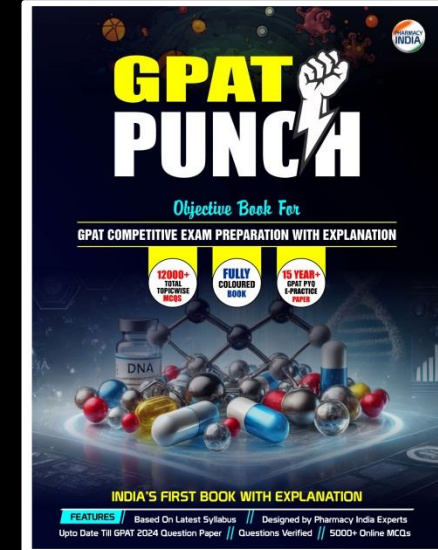


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49.

Chlorpropamide and Dexamphetamine drugs belong to

- (a) Schedule G and Schedule X respectively
- (b) Schedule X and Schedule H respectively
- (c) Schedule H and Schedule Q respectively
- (d) Schedule Q and Schedule M respectively



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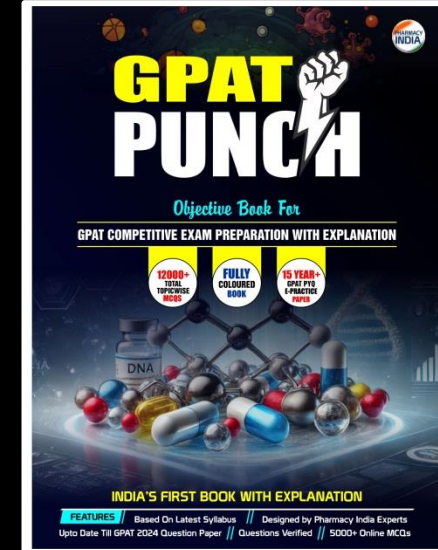


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49.

Chlorpropamide and Dexamphetamine drugs belong to

- (a) Schedule G and Schedule X respectively**
- (b) Schedule X and Schedule H respectively**
- (c) Schedule H and Schedule Q respectively**
- (d) Schedule Q and Schedule M respectively**



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Explanation:

Chlorpropamide is classified under Schedule G. Dexamphetamine is classified under Schedule X. Schedule G drugs require medical-supervision caution on their labels. Schedule X drugs are subject to stricter controls owing to their special regulatory status.



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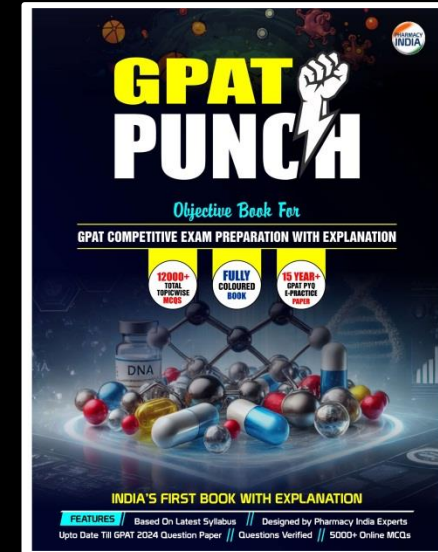


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50.

A product sample of paracetamol tablets claims to contain 500 mg paracetamol, but on analysis by Govt. Analyst it is found to contain 200 mg of paracetamol.

- (a) Adulterated drug
- (b) Misbranded drug
- (c) Spurious drug
- (d) Unethical drug



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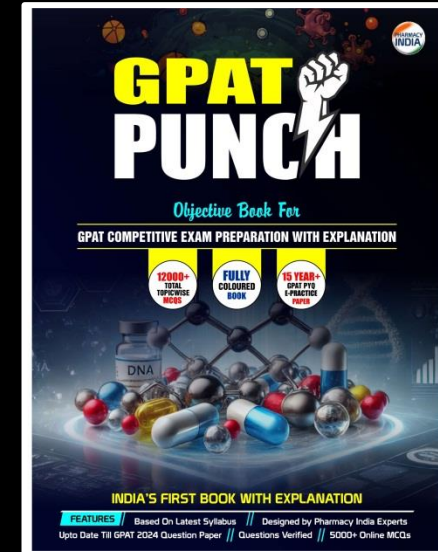


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50.

A product sample of paracetamol tablets claims to contain 500 mg paracetamol, but on analysis by Govt. Analyst it is found to contain 200 mg of paracetamol.

- (a) Adulterated drug
- (b) Misbranded drug**
- (c) Spurious drug
- (d) Unethical drug



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Explanation:

The product label claims that each tablet contains 500 mg of paracetamol whereas analysis shows only 200 mg. Thus the labelled representation regarding the drug's strength is false or misleading. It also does not describe substitution or imitation characteristic of a spurious drug.



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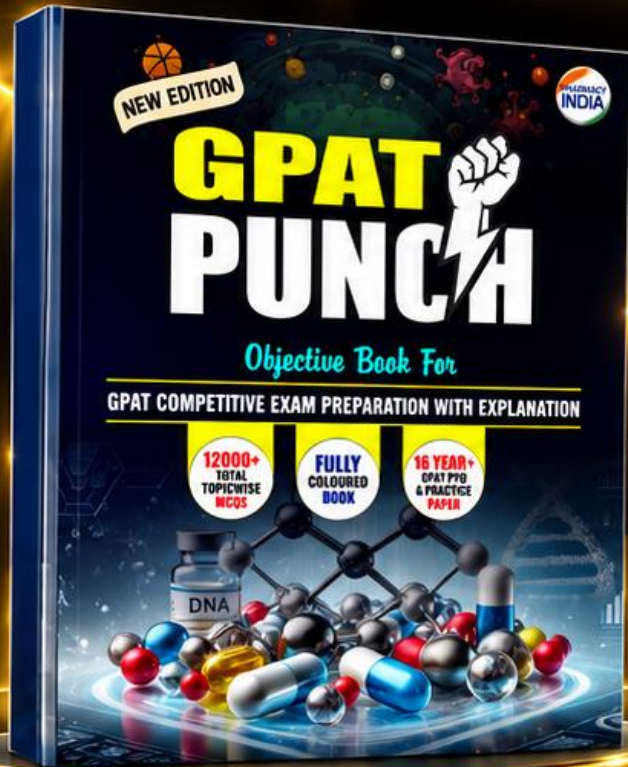


THANK
YOU!

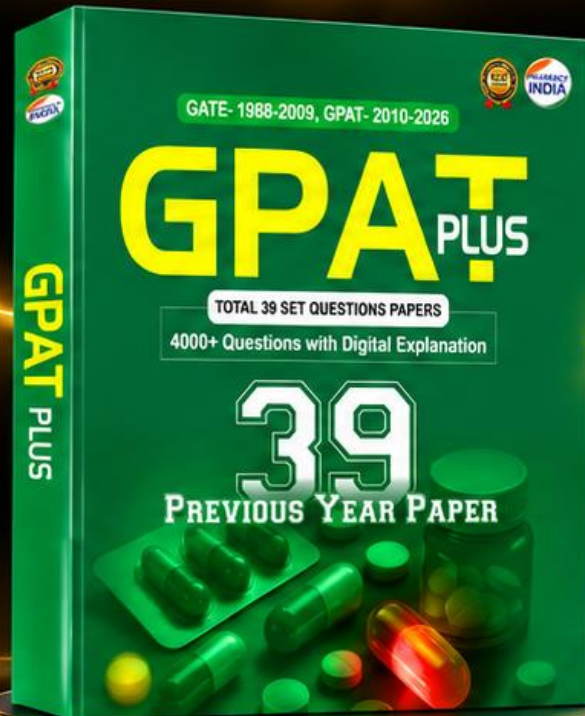


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