Faculty:	Science and Technology
Program No. & Name of the Examination:	1P00145 / / T.Y. B. Pharmacy (SEM-V) (Choice Based) (R-2019)
Subject (Paper Code):	66115 / / Pharmaceutical Jurisprudence

MCQ's

1	What is adulterated drug
a	**Whole or in part of any filthy, putrid or decomposed substance**
b	Miss branded drug
c	Drug in Phase I trial
d	Drug in Phase II trial
2	As per Drug and Cosmetic Act, which of the following is not defined as medicine?
a	Glucometer
b	paracetamol
c	Glimepiride
d	**Coding program**
3	For import of drugs specified in schedule X .....form is required.
a	**Form 10 A**
b	Form For registration
c	Certification
d	Form 25 B
4	CLAA is appointed by .....
a	State government
b	**Central government**
c	PCI
d	Government analyst

5	Which indication must be prescribed on the label of ophthalmic ointments?

a	**Use within 1 month of opening**
b	It is dangerous to take this preparation except under medical supervision
c	Consult to Physician if irritation persist, discontinue the use
d	Not for human use
6	The Schedule H on the label denotes
a	**To be sold by retail on the prescription of registered medical practitioner only**
b	Biologicals
c	Opthalmic
d	Good manufacturing Practices
7	Which of the following is not the duty of a Drug Analyst?
a	Analysis of Drug sample
b	Preparation of Drug analysis report
c	send the analysis report
d	**Market the drug**
8	Schedule ---------- of Drugs and Cosmetics Act includes requirements and guidelines of factory premises, plants, and equipment
a	Schedule P
b	Schedule Y
c	**Schedule M**
d	Schedule H

9		The name of Registered Pharmacist can be removed from the register, if
	a	**His name was entered by error in the register**
	b	His name is recommended to enter by another person
	c	He was not aware about it
	d	Anytime it can be removed without specifying the reason
10		Regulation that prescribe minimum standard of education required for qualification as a pharmacist is called as
	a	Pharmacy regulation

	b	Teaching regulation
	c	**Education regulation**
	d	Central pharmaceutical regulation
11		Displaced person" means any person who, on account of the setting up of the Dominions of India and Pakistan,has on or before _____, left or been displaced from his place of residence in such area and who has since then been residing in India.
	a	15th August 1947
	b	**First April of March 1947**
	c	26th January 1947
	d	first day of March, 1947
12		A person can get registered in___state(s) & become a Registered pharmacist.
	a	five
	b	Two
	c	**One**
	d	Three
13		Who among the following is an ex-officio member of the pharmacy Council of India?
	a	The Director of state FDA
	b	The Drugs Controller each state
	c	The Director of Medical council
	d	**The Director General Health Service**
14		What is the minimum education qualification required to register as a Pharmacist?
	a	Bachelor of computer science
	b	Masters in Pharmacy
	c	Management of Business Administration
	d	**Diploma in Pharmacy**
15		Name the medical preparations which are considered as capable of being misused as ordinary alcoholic beverages.
	a	Unrestricted preparation
	b	**Restricted preparation**
	c	Bonded manufactory
	d	Non-bonded manufactory

16		Which of the following establishments may be exempted from duty on medical preparations containing alcohol manufactured in India.
	a	**Charitable hospital**
	b	Retail Medical stores
	c	Supermarket
	d	Commercial establishment
17		The official and non-official preparations, capable of being consumed as ordinary alcoholic preparations are called as
	a	**Restricted preparations**
	b	Unrestricted preparations
	c	Asavas
	d	Aristas
18		The medicinal and toilet preparation act was passed with the main objective
	a	To regulate the production of medicines
	b	**Collect duties of excise on medicinal toilet preparations containing alcohol, opium, Indian hemp, etc.**
	c	To conduct clinical trial
	d	To sale the products
19		Opium means
	a	Coagulated juice of opium poppy and its mixture including preparation containing less than 0.2% cocaine
	b	**Coagulated juice of opium poppy and its mixture excluding preparation containing less than 0.2% morphine**
	c	Coagulated juice of opium poppy and its mixture excluding preparation containing more than 2% cocaine
	d	Coagulated juice of opium poppy and its mixture including preparation containing more than 1 % morphine
20		Diacetylmorphine is also known as
	a	Hemp

	b	Cannabis
	c	Charas
	d	**Heroin**
21		Medicinal cannabis is also known as
	a	Opium
	b	**Hemp**
	c	Heroin
	d	Charas
22		Opium poppy includes
	a	**Plant of papaver somniferum**
	b	Leaf of Erythroxylon
	c	Ecgonine
	d	Cocaine

23		Which of the following examples is the example of a prohibited advertisement?
	a	**Advertisements of magic remedies for the treatment of certain diseases and disorders.**
	b	Advertisements by Government
	c	Leaflets or literature along with packings of drugs
	d	Therapeutic index published by a licensed manufacturer
24		Which of the following is other than a magic remedies
	a	Mantras
	b	Talismans
	c	Kavachas
	d	**Pharmacy Education Regulations**
25		The Drugs and magic remedy (OA) Act was passed in _____.
	a	**1954**
	b	1948
	c	1985
	d	1972

26		Prevention of cruelty to animals act was passed in
	a	**1960**
	b	1985
	c	1945
	d	1956
27		Any research undertaken by an individual , company, firm, corporation or institutionon behalf of a foreign individual , company, firm, corporation or institution for any consideration is called
	a	**Contract research**
	b	Collaborative research
	c	Experiment
	d	Research methodology
28		Who has the power to fix selling price of scheduled formulations
	a	State Government
	b	**Central Government**
	c	Lok Sabha
	d	Rajya Sabha
29		Ceiling price means:
	a	Price fixed by the Government for Non-Scheduled formulations in accordance with the provisions of this Order
	b	Price fixed by the Government for Marketed formulations in accordance with the provisions of this Order
	c	Price fixed by the Government for formulations under clinical trial in accordance with the provisions of this Order
	d	**Price fixed by the Government for Scheduled formulations in accordance with the provisions of this Order**
30		Which act's prime objective is to make sure that the essential drugs are available to all at a reasonable price.
	a	DMR (OA) act
	b	**DPCO 2013**
	c	D and C act
	d	MTP(ED) act
31		DPEA stands for

	a	**Drugs Prices Equalization Account**
	b	Drugs Pricing Equity Account
	c	Drugs Policy Equalization Account
	d	Drugs price equalization administration.

32		When the Drug Bill was introduced?
	a	1938
	b	**1940**
	c	1942
	d	1944
33		Who was the Chairman of Drug Enquiry Committee?
	a	**RN Chopra**
	b	RL Chopra
	c	ML Shroff
	d	B Mukharjee
34		Which of the following is recommended of Drug Enquiry Committee?
	a	Control Sale of Spurious Drug
	b	Control Price of Drug
	c	**Establishment of CDL**
	d	Law for Illegal Traffic of Drug
35		Bengal chemical and pharmaceutical works was started by?
	a	**Acharya PC Ray**
	b	Mr. Bathgate
	c	TK Gajjar
	d	B Mukharjee
36		Pharmaceutical ethics are the ethics in relation to
	a	Architectural profession
	b	Civil profession
	c	Engineering profession

	d	**Pharmacy profession**
37		Standard of conduct is called
	a	Behaviour
	b	Law
	c	**Ethics**
	d	Etiquettes
38		The code of pharmaceutical ethics helps Pharmacists conduct in relation to
	a	**His job**
	b	His life
	c	His relatives
	d	His family
39		The legislative intent of the Medicinal and Toilet Preparation Act was to
	a	**Levy and collection of duties on preparation of alcohol & narcotics**
	b	Levy and collection of duties on preparation of alcohol only
	c	Levy and collection of duties on preparation of narcotics only
	d	Levy and collection of duties on preparation of methanol
40		Under MTP act termination is permitted maximum up to ---- weeks
	a	**20**
	b	21
	c	22
	d	23
41		Right to Information act 2005 is enacted on
	a	13 Jun 2005
	b	**15 Jun 2005**
	c	16 Jun 2005
	d	17 Jun 2005
42		Right to Information act 2005 is commenced on

	a	**12 Oct 2005**
	b	13 Oct 2005
	c	14 Oct 2005
	d	15 Oct 2005
43		RTI means right to
	a	Information which would prejudicially affect
	b	Information which might incite an offense
	c	**Information which can be assessed by public authority**
	d	Information which is forbidden by court
44		Under RTI fees for application is Rs
	a	100
	b	25
	c	**10**
	d	150
45		Term and product process of patent is
	a	**20 years for both**
	b	20 years for process patent and 10 years for product patent
	c	10 years for process patent and 10 years for product patent
	d	10 years for process patent and 20 years for product patent
46		Compulsory license can be granted after how many years of the expiration of a granted patent?
	a	2 Years
	b	**3 Years**
	c	4 Years
	d	5 Years
47		A complete specification should be filed within how many months of filing the provisional specification as per Indian Patent Act 2005?
	a	**12**
	b	18
	c	24
	d	6
48		Prior art generally does not include
	a	Knowledge disclosed only in Patents

	b	Knowledge disclosed in publications
	c	**Knowledge disclosed only to closed group of members bound by non-disclosure agreement**
	d	Knowledge available in public domain
49		Patent is ---------- property
	a	Negotiable
	b	Non negotiable
	c	**Transferable**
	d	Non transferable

4. The full fledged pharmacopeia committee was formed in the year *

Mark only one oval.

- 1948
- 1955
- 1971
- 1965

5. Joint State Council is constituted under *

Mark only one oval.

- The Pharmacy Act
- Drugs and Cosmetics Act
- Drugs and Magic Remedies Act
- Bombay Shop Act

6. Which of the following is a recommendation of Drug Enquiry committee? *

Mark only one oval.

- Establishment of CDL
- Control price of drug
- Control sale of spurious drug
- Law for Illegal traffic of drug

7. Following is the duty of Government analyst *

Mark only one oval.

- To forward to the government the reports of analytical and research work
- To inspect any premises
- To take samples of any drug or cosmetics
- To Enter and search manufacturing premises

8. Cosmetics means any article intended to - *

Mark only one oval.

- Alter the appearance of human body
- Affect the structure of human body
- Destroy vermin
- Mitigate a disorder

9. Which of the following term is used for drugs that is imported under a name which belongs to another drug, purports to be the product of a manufacturer of whom it is not a truly a product? *

Mark only one oval.

- Spurious drug
- Misbranded drug
- Adulterated drug
- New drug

10. Which of the following is advisory body? *

Mark only one oval.

- Drugs Consultative Committee
- Central Drug Laboratory
- Government Analyst
- Drug Inspectors

11. Choose the correct option *

Mark only one oval.

- Life period of drugs: Schedule P
- Pack sizes of drugs: Schedule D
- Biological and special products: Schedule D
- Standards for surgical dressings: Schedule H

12. GMP requirements for manufacture of Ayurvedic (including Siddha) and Unani drugs is given in *

Mark only one oval.

- Schedule T
- Schedule M-II
- Schedule M
- Schedule U

13. XRx on the label of the product indicate: *

Mark only one oval.

- The product contains a substance specified in Schedule X
- The product contains a substance specified in Schedule N
- The product contains a substance specified in Schedule G
- The product contains a substance specified in Schedule J

14. Application forms and licenses types are available under *

Mark only one oval.

- Schedule A
- Schedule B
- Schedule X
- Schedule Y

15. The objective of the Drug and Magic Remedies(OA) Act 1954 is *

Mark only one oval.

- To prohibit certain types of advertisements relating to magic remedies which falsely claim and mislead public
- To control sale of drugs
- To market the drugs
- To analyse the drugs

16. Hashish under NDPS act is *

Mark only one oval.

- Hemp
- Opium
- Coca
- Cocaine

17. Which is incorrect (under the provisions of the DPCO)? *

Mark only one oval.

- Ceiling price is fixed by manufacturer and retailer
- Ceiling price is fixed by the Central Government
- Ceiling price is only for Scheduled formulations
- Ceiling price does not include local taxes

18. The legislative intent of the Medicinal and Toilet Preparations Act was to *

Mark only one oval.

- levy excise duty on preparations containing alcohol and narcotics
- levy excise duty on preparations containing ethanol
- collect excise duty on preparations containing methanol
- collect customs duty on preparations containing alcohol and narcotics

19. As per Food safety Act, following is excluded from the definition of "Food" *

Mark only one oval.

- Animal feed
- Genetically modified food
- Infant food
- Alcoholic drinks

20. Prior art does not include *

Mark only one oval.

- Knowledge disclosed to only closed group of members bound by non disclosure agreement
- Knowledge disclosed in publications
- Knowledge disclosed only in patents
- Knowledge available in public domain

21. Patent is a ----------property. *

Mark only one oval.

- Transferable
- Non transferable
- Negotiable
- Non negotiable

22. As per Indian Penal Code, a person can be arrested with a warrant if *

Mark only one oval.

- The offence is non-cognizable
- The offence is committed in front of a police officer
- The offence is cognizable
- The offence is committed outside the country

23. CTD is divided into --------- modules *

Mark only one oval.

- 5
- 6
- 7
- 4

4. Who was the first chairman of Drug Enquiry committee? *

Mark only one oval.

- RN Chopra
- RL Chopra
- ML Shroff
- B Mukherjee

5. The institution running pharmacy course is approved by *

Mark only one oval.

- Pharmacy Council of India
- Medical Council of India
- Indian Pharmaceutical Association
- State pharmacy Council

6. First Schedule to D & C act gives *

Mark only one oval.

- list of Ayurvedic, Siddha Unani books
- records to be maintained for raw materials
- colours to be used in medicines
- drugs exempted from certain provisions

7. The administrative bodies established under the drugs and cosmetics act having advisory role include *

Mark only one oval.

- Drug Technical Advisory Board
- Drug Inspector
- Government Analyst
- Customs collectors

8. Which of the following committee suggested to expand the scope of schedule K to include OTC drugs *

Mark only one oval.

- Mashelkar Committee
- Chopra Committee
- Hathi Committee
- Goel Committee

9. Cosmetics means any article intended to - *

Mark only one oval.

- Alter the appearance of human body
- Affect the structure of human body
- Destruct vermis of insects
- Clean the human body

10. Schedule E stands For- *

Mark only one oval.

- List of poisonous substances (omitted)
- Standards for surgical dressing
- Standards for medical devices
- List of drugs to be sold on Prescription of RMP

11. Choose the correct option *

Mark only one oval.

- Habit forming drugs : Schedule X
- Pack sizes of drugs: Schedule D
- Biological and special products: Schedule D
- Standards for surgical dressings: Schedule H

12. Fees for test or analysis by the Central Drugs Laboratories are available under *

Mark only one oval.

- Schedule B
- Schedule M
- Schedule X
- Schedule Y

13. Requirements and guidelines to undertake clinical trials are available under *

Mark only one oval.

- Schedule Y
- Schedule C
- Schedule G
- Schedule X

14. Schedule G is concerned with *

Mark only one oval.

- List of drugs required to be taken only under medical supervision
- List of habit forming, psychotropic and other such drugs
- Standards of cosmetics
- Drugs required to be sold by retail only on prescription of a RMP

15. Advertisement on 'Cure for cancer' falls under which type of advertisement from below? *

Mark only one oval.

- Prohibited
- Exempted
- Bonafide
- Permitted

16. Preparations containing more than 0.2% morphine are called *

Mark only one oval.

- Opium derivatives
- Opium poppy
- Poppy straw
- Poppy straw concentrates

17. For the manufacture of a medicinal preparation containing alcohol in bond a licence is required *

Mark only one oval.

- Excise Commissioner
- State Licensing Authority
- Central Licensing authority
- Narcotic Commissioner

18. The formula to calculate ceiling price includes *

Mark only one oval.

- 16 % to retailer
- Margin to wholesaler
- All applicable taxes
- Excise duty

19. As per Food standards and safety Act, 'which of the following doesnot fall under category of 'Food'? *

Mark only one oval.

- Plants prior to harvesting
- Genetically modified food
- Infant food
- Chewing gum

20. Compulsory licenses for patents are granted *

Mark only one oval.

- Statutory
- Voluntary
- Virtual
- Implied

21. A complete specifications should be filed within ------- months of filing the provisional specification as per Indian Patent Act 2005. *

Mark only one oval.

- 12
- 18
- 24
- 6

22. Proceedings carried out by a Police officer for collection of evidence is called *

Mark only one oval.

- Investigation
- Inquiry
- Trial
- Custody

23. CFR stands for *

Mark only one oval.

- Code of Federal Regulations
- Center of Federal Regulations
- Center of Federal Register
- Code of Federal Register

4. Bengal Chemical and Pharmaceutical works was started by *

Mark only one oval.

- Acharya PC Ray
- Mr Bathgate
- TK Gajjar
- B Mukherjee

5. The main object of the pharmacy Act, 1948 is to *

Mark only one oval.

- Regulate the profession of pharmacy
- Control the advertisement
- Prevent the unnecessary pain or suffering on animals
- Prevention of food adulterants

6. Central drug laboratory is situated at *

Mark only one oval.

- Kolkata
- Lucknow
- Gaziabad
- Mumbai

7. _____is advisory to the central government in matters concerning administration of the D and C act. *

Mark only one oval.

- Drugs Consultative committee
- Central Drugs Laboratory
- Customs collectors
- Government Analyst

8. _____ is cosmetic, if it is an imitation of, or is a substitute for another cosmetic or resembles another cosmetic in a manner likely to deceive or bears upon it or upon *

Mark only one oval.

- ◯ Spurious Cosmetic
- ◯ Adulterated Cosmetic
- ◯ Substandard Cosmetic
- ◯ Misbranded Cosmetic

9. The drug enquiry committee is also known as *

Mark only one oval.

- ◯ Chopra Committee
- ◯ Hathi Committee
- ◯ Mashelkar Committee
- ◯ Goel Committee

10. What do you mean by DTAB? *

Mark only one oval.

- ◯ Drug Technical Advisory Board
- ◯ Drug Testing Advisory Body
- ◯ Drug Technical Appellant Body
- ◯ Drug Technical Admission Board

11. Choose the correct option *

Mark only one oval.

- ◯ Standards for disinfectants : Schedule O
- ◯ Pack sizes of drugs: Schedule D
- ◯ Biological and special products: Schedule D
- ◯ Standards for surgical dressings: Schedule H

12. Standards for condoms made of rubber latex are available under *

Mark only one oval.

- Schedule R
- Schedule C
- Schedule G
- Schedule X

13. The drugs which can be sold out in retail against prescription of registered medical practitioner only are available under *

Mark only one oval.

- Schedule H
- Schedule C
- Schedule G
- Schedule X

14. Standards for ophthalmic preparations are available under *

Mark only one oval.

- Schedule FF
- Schedule A
- Schedule G
- Schedule Y

15. As per DMR Act' 1954, Classes of Exempted Advertisements include *

Mark only one oval.

- Any advertisement relating to a drug which is sent confidentially in the prescribed manner to RMP
- Advertisement may Directly or indirectly give a false impression regarding the true character of the drug
- Advertisement of the certain drugs used in the procurement of miscarriage in women or prevention of conception in women
- Advertisement of the certain drugs used in the correction of menstrual disorder in women

16. Cocaine is obtained from a plant of the genus *

Mark only one oval.

- Erythroxylum
- Papaver
- Cannabis
- Cacao

17. As per the Drugs Prices Control Order, market share is based on *

Mark only one oval.

- Moving annual turnover
- Maximum retail price
- Ceiling price
- Margin to Wholesaler and retailer

18. Finished preparations issued from a bonded manufactory are *

Mark only one oval.

- not less than 57ml
- not less than 2273 ml
- more than 60 ml
- 100ml

19. The term of office of Chairperson of Food Safety and Standards Authority of India is *

Mark only one oval.

- Three years
- Five years
- Seven years
- Ten years

20. Term of process and product patent is *

Mark only one oval.

- 20 years for both
- 20 years for process patent and 10 years for product patent
- 10 years for product patent and 10 years for process patent
- 20 years for product patent and 10 years for process patent

21. Only GMO's which do not fall under section _____ are patentable. *

Mark only one oval.

- 3(b)
- 3(a)
- 3(c)
- 3(d)

22. Restriction/surveillance on the movement of a person is called *

Mark only one oval.

- Custody
- Arrest
- Inquiry
- Summons

23. Animal studies, clinical studies and bioavailability are part of *

Mark only one oval.

- NDA
- ANDA
- BLA
- IND

Faculty:	Science and Technology
Program No. & Name of the Examination:	1P00145 / / T.Y. B. Pharmacy (SEM-V) (Choice Based) (R-2019)
Subject (Paper Code):	66115 / / Pharmaceutical Jurisprudence

DESCRIPTIVE QUESTION'S BANK

Elaborate on the procedure for the import of drugs
Ellaborate on constitution of DTAB committe.
Explain the procedure of dispatch of sample for test
explain labelling and packing of requirements of opthalmic ointment

1.Write note on Education Regulations under pharmacy act 1948
2.Give objectives of Pharmacy Act 1948. Define 1. Registered Pharmacist, Displaced Person as per Pharmacy Act.
3.Elaborate the objectives of Pharmacy Act 1948. Differentiate between State Pharmacy Council and Joint State Pharmacy Council.
4.What is the objective of MTP Act 1955 and give an account of bonded manufactory?
5.What is the objective of MTP Act 1955 and give account of non-bonded manufactory?
6.Define Cannabis, Opium and opium derivatives and elaborate on measures taken by the central government to control illicit traffic also enlist the functions of the Narcotic Commissioner.
7.Define Magic Remedy. What are the steps taken by Government to have control over magic remedies?
8.Define Formulation as per DPCO 2013. Comment on legislative intention of DPCO 2013. 9.Explain significance of pharmaceutical policy 2017.
10.What are the objectives of DPCO2013? Define 'Bulk Drug' and 'Ceiling Price'
11. Define Drug and misbranded drug as per D and C Act 1940 explain the form and manner of application for import license. 12.Enlist the qualification requirements for Drug Inspectors and give account of duties and powers of Drug Inspector. 13. Define Dutiable goods, Alcohol as per MTP(ED) Act. Differentiate between Restricted and Unrestricted preparations. Explain the procedure to be followed for issuing license for manufacture of medicinal and toilet preparations as per MTP(ED) Act. 14. What is illicit traffic and describe measures to prevent illicit traffic as per NDPS Act 1985?

15. Elaborate on duties of pharmacist as per pharmacy practice regulations 2015.

16. Describe the process and control cultivation of opium by central government.

17.What is DEC? Discuss its recommendations.

18. Comment on the calculation of ceiling price for scheduled formulations as per DPCO 2013.

19.Elaborate on labelling of homeopathy medicines as per D & C Act 1940.

20.Define “Invention” as per Indian Patent Act. Elaborate on the criteria to be satisfied by an invention to be patentable in India.

21.List the inventions which are not patentable as per the provisions of Indian Patent Act,

22. Describe in detail offences and penalties as per NDPS Act 1985

23.Write a note on loan licensing

24. Write a note on DCC committee from D and C act 1940

25. Give the constitution, working and functions of PCI.

26.Describe the constitution and functions of the Institutional Animals Ethics Committee.

27. Define Ethics, Pharmaceutical ethics, morality and law

28.Elaborate qualification of Drug analyst and duties of drug analyst as per 1940 as per D and C Act 1940

29.Write a note on central license approving authority

30. Define Ethics, Pharmaceutical ethics, morality and law. Add a note on code of pharmaceutical ethics.

31. Give account of different types of licenses for manufacture of drugs for sale.

32. Write short note on Central drug laboratory.

33.What is meant by 'manufacture in bond'? Outline the procedure that should be followed for obtaining license for manufacture in bond including the conditions that are to be fulfilled?

34. Write a note on central license approving authority

35.Elaborate qualification of Drug analyst and duties of drug analyst as per 1940 as per D and C Act 1940

36.What is DEC? Discuss its recommendations.

37. Define cannabis and describe in detail offences and penalties as per NDPS Act 1985.

38.Define Drug and misbranded drug as per D and C Act 1940 explain the form and manner of application for import license.

39. Define Advertisement and magic remedy. List classes of prohibited advertisements and exempted advertisements as stated by DMR(OA) Act 1954.

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41.Elaborate on labelling of homeopathy medicines as per D & C Act 1940.

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44. Write a note on loan licensing

45.Write a note on central license approving authority

46. As per D and C act, elaborate on labelling of homeopathy medicines

47. Define Drug and misbranded drug as per D and C Act 1940 explain the form and manner of application for import license.

48. Enlist the qualification requirements for Drug Inspectors and give account of duties and powers of Drug Inspector.

49. Write short note on Central drug laboratory.

50. What is meant by 'manufacture in bond'? Outline the procedure that should be followed for obtaining license for manufacture in bond including the conditions that are to be fulfilled?

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