		St. Philomena's College (Autonomous), Mysore	
		PG Department of Biochemistry	
		Question Bank (Revised Curriculum 2018 onwards)	
		First Year- Second Semester (2019-20 Batch)	
	Cours	e Title (Paper Title): Pharmacology of Plant Products. QP Code -965	)3
Unit	Sl. No	Questions	Marks
1	1.	What is a crude drug? Give an example	2
1	2.	Define pharmacognasy.	2
1	3.	Name any two systems of classification of crude drug	2
1	4.	What is a pro drug? Give an example	2
1	5.	Define crude drug and pro drug	2
1	6.	Distinguish between pro drug and crude drug.	2
1	7.	Name any two systems of classification of crude drug	2
1	8.	Name any two seeds with the main active principle in it	2
1	9.	Name any two barks with the main active principle in it	2
1	10.	Define therapeutic index	2
1	11.	Mention any two sources of drugs with an example	2
1	12.	Name a drug with chemical, propioratory and brand name	2
2	13.	What is herbal tincture?	2
2	14.	Give examples for any two Ayurvedic plants with an application for each.	2
2	15.	Mention any two historical references for usage of plants as drugs.	2
3	16.	Write the resonating structure of ferulic acid radical.	2
3	17.	What are polyphenols? Give the natural sources of Polyphenols.	2
3	18.	Write the structure and sources of Gallic acid.	2
3	19.	List factors responsible for an efficient antioxidant activity of the Gallic	2
		acid molecule?	
3	20.	What are tannins? Give examples.	2
3	21.	What are the natural sources of Ellargic acid	2
3	22.	What are Alkaloids? Give examples.	2
3	23.	Report structure and sources of Ellargic Acid.	2
3	24.	Report structure and sources of Ferulic Acid.	2
3	25.	Report structure and sources of Nicotine.	2
3	26.	Report structure and sources of Caffeic Acid.	2
3	27.	What are Flavonoids? Give examples.	2
3	28.	Report structure and sources of cannabidiol.	2
3	29.	Illustrate Sesqiterpenes with examples.	2
3	30.	Illustrate diterpenes with examples.	2
4	31.	What are bioactive compounds? Give their biological significance.	2
4	32.	What are bioactive compounds? Give its classification	2

4	33.	What is chromatography? How can it be used in extraction or purification	2
		of bioactive compounds?	
4	34.	What is paper chromatography? How can it be used in extraction or purification of bioactive compounds?	2
4	35.	What is HPLC? How can it be used in extraction or purification of bioactive compounds?	2
4	36.	What is ion exchange chromatography? How can it be used in extraction or purification of bioactive compounds?	2
4	37.	What is thin layer chromatography? How can it be used in purification of bioactive compounds?	2
4	38.	What is gas chromatography? How can it be used in identification of bioactive compounds?	2
4	39.	What is column chromatography? How can it be used in extraction or purification of bioactive compounds?	2
4	40.	What is flash column chromatography? How can it be used in extraction or purification of bioactive compounds?	2
4	41.	What is molecular exclusion chromatography? How can it be used in extraction or purification of bioactive compounds?	2
4	42.	What is affinity chromatography? How can it be used in extraction or purification of bioactive compounds?	2
4	43.	What is spectroscopy? What are the different types? How can it be used in identification of bioactive compounds?	2
4	44.	What is UV-spectroscopy? What are the different types? How can it be used in identification of bioactive compounds?	2
4	45.	What is infrared spectroscopy? What are the different types? How can it be used in identification of bioactive compounds?	2
4	46.	What is NMR? How can it be used in identification of bioactive compounds?	2
4	47.	What is mass spectroscopy? How can it be used in identification of bioactive compounds?	2
4	48.	Which are the factors that affect extraction and purification protocols of bioactive compounds?	2
4	49.	How Phenolic Compounds are extracted Using Solvents?	2
4	50.	What is MAE? Give its significance	2
4	51.	What is Ultrasonic-Assisted Extraction? Give its significance	2
4	52.	How is Fractional Crystallization used in extraction of bioactive compounds? Give its significance	2
4	53.	What is Fractional Distillation ? Give its significance	2
4	54.	What is fractional liberation? Give its significance	2
4	55.	What is sublimation? Give its significance	2
4	56.	What is maceration? Give its significance	2
4	57.	What is percolation? Give its significance	2

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4	58.	What is reserve percolate method? Give its significance	2
4	59.	What is Soxhlet extraction? Give its significance	2
4	60.	What Is large scale extraction? Give its significance	2
4	61.	What is supercritical fluid extraction? Give its significance	2
4	62.	What are the different types of extracts?	2
4	63.	What is Quercetin? Give its structure	2
4	64.	What is Ellargic acid? Give its structure	2
4	65.	What is Curcumin? Give its structure	2
4	66.	What is diabetes mellitus? What are its types?	2
4	67.	What are the complications associated with diabetes mellitus?	2
4	68.	What are CVD's? Which are the different types?	2
4	69.	What are the complications associated with CVD's? give its symptoms	2
4	70.	What is Alzheimer's disease? What are the complications associated with	2
		it?	
4	71.	What is cancer? What are the different types of cancers?	2
4	72.	What are the complications associated with cancer?	2
5	73.	Mention the different drug delivery systems.	2
5	74.	Give the significance of sustain release dosage form.	2
5	75.	What is enteric coated drug delivery system? Give an example	2
5	76.	How does sustain release is different from that of enteric coated drug	2
		release?	
5	77.	Give the significance of enteric coated release dosage form.	2
5	78.	Define dosage form.	2
5	79.	Give the merits of sustain release dosage form.	2
5	80.	Give the demerits of sustain release dosage form.	2
5	81.	Mention the merits of enteric coated release dosage form.	2
5	82.	Mention the demerits of enteric coated release dosage form.	2
5	83.	What is sustained release drug delivery system? Give an example	2
5	84.	Differentiate between controlled release & sustain release.	2
5	85.	What is the purpose of an enteric coating on a tablet?	2
5	86.	What is liposome mediated drug delivery system?	2
5	87.	What is nanoparticle mediated drug delivery system?	2
6	88.	Define a. Pharmacokinetics b. Therapeutic Index	2
6	89.	Define Pharmacokinetics and Pharmacodynamics.	2
6	90.	Define Zero order process. Give example.	2
6	91.	Distinguish onset of action from onset time?	2
6	92.	What is PK/PD Modeling?	2
6	93.	Compare Pharmacokinetics with Pharmacodynamics.	2
6	94.	Formulate equation to determine <b>t1/2</b> of Zero order kinetics	2
6	95.	Formulate equation to determine t1/2 of First order kinetics	2
6	96.	Explain Peak Plasma Concentration.	2

6	97.	Explain Pharmacokinetic models.	2
6	98.	List the different approaches to pharmacokinetic analysis.	2
6	99.	Distinguish rate and order of reaction.	2
6		List the rate processes encountered in physiologic system.	2
6	101.	Explain Mixed order Kinetics.	2
6	102.	Report equation for elimination half-life	2
7		Prepare pictorial representation of One-Compartment Open Model IV Bolus	2
7	104.	Draw diagram representing One-Compartment Open Model -intravenous	2
		Infusion	
7	105.	What is partial agonist? Give example.	2
7	106.	Define receptors.	2
7	107.	What are agonists? Give example	2
7	108.	Define Pharmacodynamics.	2
7	109.	Mention the principles of drug action	2
7	110.	Describe Stimulation.	2
7	111.	What are depressants?	2
7	112.	Give examples of replacement drugs.	2
7	113.	Differentiate Affinity from Intrinsic Activity	2
7	114.	Differentiate Partial agonists from Agonist.	2
7	115.	What are the categories of body tissues in two compartment model?	2
7	116.	List the categories of two compartment models.	2
8	117.	What is a clinical trial? Give its significance	2
8	118.	What is a patent? How it is important?	2
9	119.	What Is AYUSH? What is its importance?	2
9	120.	What is Ayurveda? What is its importance?	2
9	121.	Enlist 4 important Ayurvedic drugs and its application	2
10	122.	What are Pre-clinical pharmacokinetic studies? Give its significance	2
10	123.	What are Pre-clinical pharmacodynamic studies? Give its significance	2
10	124.	What is Lipinski's rule? Give its significance	2
10	125.	What is high throughput screening? Give its significance	2
10	126.	What are the goals and applications of high throughput screening?	2
10	127.	Which is the biochemical techniques used in HTS? What is their	2
		significance?	
10	128.	What are a lead and a hit?	2
10	129.	What are the different approaches of HTS?	2
10	130.	What is vascular mapping? What is its significance?	2
10	131.	What is tissue perfusion? What is its significance?	2
10	132.	What is thermal shift analysis? What is its significance?	2
10	133.	How is combinatorial chemistry used in HTS?	2
1	1.	Discuss on therapeutic drug monitoring system.	5

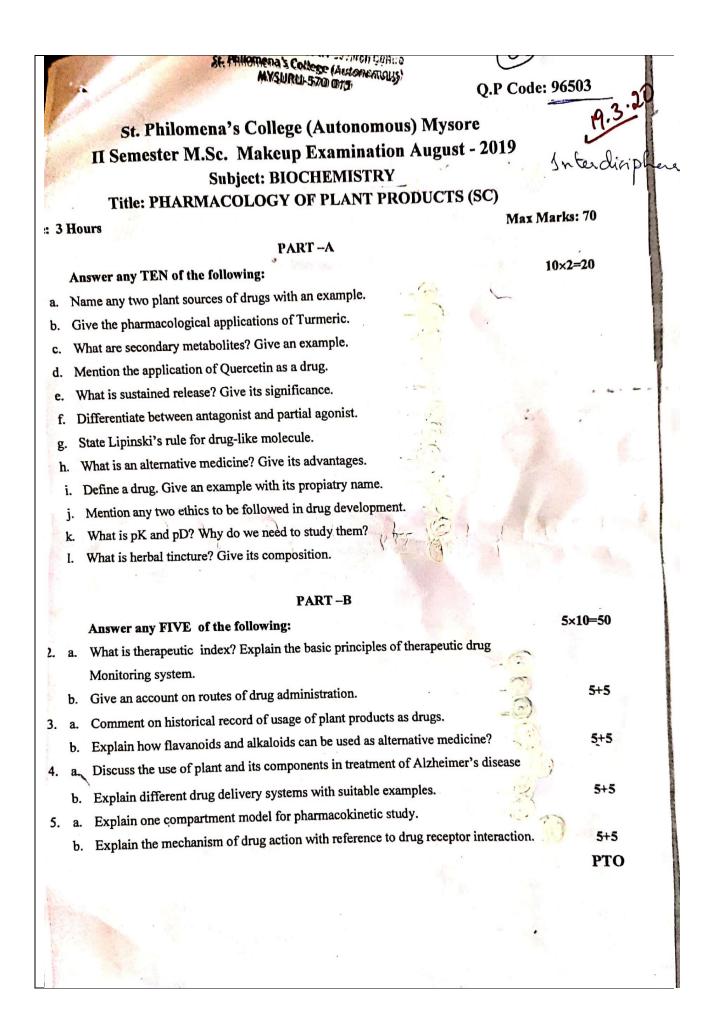
1	2.	Write a note on history and scope of pharmacognasy.	5
1	3.	Write a note on different sources of drugs.	5
1	4.	Explain the different routes of drug administration.	5
2	5.	Discuss briefly the history of plants usage as drugs.	5
2	6.	Write short note on history of usage of plants as drugs.	5
2	7.	Explain the historical aspect of usage of whole plants as drugs.	5
2	8.	Explain the historical aspect of usage of plant extracts as drugs.	5
2	9.	What are alternative medicines? Write a note on their pros and cons.	5
3	10.	Discuss about Ellargic acid as alternative Medicine.	5
3	11.	What is the role of Secondary metabolites in plant physiology?	5
3	12.	Explain the therapeutic importance of ferulic acid.	5
3	13.	What are Terpenes? Write the classification of terpenes with examples.	5
3	14.	Explain the role of qucertin in plant physiology and as alternative medicine.	5
3	15.	What is isoprene rule? Write the classification of terpenes with example	5
3	16.	Discuss the role of Quercetin as an alternative medicine.	5
3	17.	Discuss the therapeutic importance of Ellargic acid.	5
3	18.	Explain the role of CBD's in neural disorders.	5
3	19.	Explain the therapeutic importance of ferulic acid.	5
3	20.	Discuss the role of Nicotine as an alternative medicine.	5
3	21.	Discuss the role of Gallic acid as an alternative medicine.	5
3	22.	Summarize the role of secondary metabolites in plant physiology	5
3	23.	Discuss the structure and plant physiological functions of Gallic acid.	5
3	24.	Discuss the structure and plant physiological functions of ferulic acid.	5
3	25.	Discuss the structure and plant physiological functions of Ellargic acid.	5
3	26.	Explain the therapeutic importance of terpenes.	5
3	27.	Summarize the classification of alkaloids.	5
3	28.	Explain therapeutic importance of caffeine.	5
4	29.	Give a detailed account on the different types of biochemical techniques	5
		used in isolation purification and characterization of bioactive compounds	
4	30.	Give a detailed account on the different types of extraction protocols used	5
		to extract bioactive components from plant sources	
4	31.	Give the structure and biological significance of Quercetin	5
4	32.	Give the structure and biological significance of Ellargic acid	5
4	33.	Give the structure and biological significance of curcumin	5
4	34.	Give a detailed account on the medicinal plants used to treat diabetes mellitus	5
4	35.	Comment on the usage of medicinal plants in CVDS	5
4	36.	Discuss the usage of medicinal plants in treating Alzheimer's disease	5
4	37.	Briefly explain the MOA of podophyllum, taxanes and vinca.	5
4	38.	Add a lime light on the MOA of camptothecin and curcumin in cancer cells	5

4	39.	What is the pathophysiology of cancer? Explain the MOA of any one of the medicinal plant used to treat cancer	5
5	40.	Explain the characteristics of drug for formulation as sustained release dosage form	5
5	41.	Write a note on liposome mediated drug delivery system.	5
5	42.	What are sustained release and controlled release drug delivery systems? Explain on their pros & cons	5
5	43.	Write short note on nanoparticle mediated drug delivery system.	5
5	44.	Explain the characteristics of drug for formulation as enteric coated drug release.	5
6	45.	Explain the following terms: a. MEC, b. MSC, c. Therapeutic Index d.Peak Plasma Concentration. e. Onset time	5
6	46.	Write the advantages of Compartment modeling approach.	5
6	47.	What is compartment model? Explain one compartment open model.	5
6	48.	The half life for the 1 <sup>st</sup> order photolysis of cefotaxime solution containing 150 mg drug is 50 minutes. How long it will take for drug to decompose to 20% of its original amount?	5
6	49.	What is PK/PD Modeling? Elaborate on Linear model approach to explain PK/PD model.	5
6	50.	The half-life for first-order photolysis of cefotaxime solution containing 150 mg drug is 50 minutes. a. How long will it take for the drug to decompose to 20% of its original amount?	5
6	51.	Deduce mathematically rate constant for Zero Order Kinetics.	5
6	52.	Deduce mathematically rate constant for First Order Kinetics.	5
6	53.	A Penicillin solution containing 300 units/ml has Half-life of 8 days in Plasma. What will be the concentration in 7 days?	5
6	54.	A Penicillin solution has a Half-life of 6 days. How long it will take for the concentration to drop to 70% of the initial concentration?	5
6	55.	A Penicillin solution has an initial potentency of 90mg/ 6ml. After one month in a cold room, the concentration is found to be 80mg/6ml. What is the half-life of Penicillin solution under the storage conditions?	5
6	56.	Construct Plasma Drug concentration- Time profile and Label all terms involved.	5
6	57.	Summarize Clearance.	5
7	58.	Mathematically devise one compartment open model Intravenous infusion	5
7	59.	Mathematically devise one compartment open model Intravenous Bolus	5
7	60.	Describe briefly about principles of drug action.	5
7	61.	Explain action of Agonist with suitable example.	5
7	62.	Summarize principles of drug action.	5
7	63.	Illustrate Antagonist with suitable example.	5
7	64.	Explain partial agonist with suitable example.	5
7	65.	Describe the following Terms: I. Affinity ii. Intrinsic Activity iii. Partial	5

		Agonist	
8	66.	Explain in detail the evaluation of crude drug.	5
8	67.	Explain the various phases of clinical trial.	5
8	68.	Write a note on new drug development process.	5
8	69.	Briefly discuss on drug registration.	5
8	70.	Explain the process of clinical evaluation of drugs.	5
9	71.	What is AYUSH? Explain the applications of Ayurvedic drugs in treating	5
		diseases.	
10	72.	Explain the general steps involved in HTS.	5
10	73.	Explain the biochemical screen approach of HTS	5
10	74.	Explain the cell based and organism based screen approach of HTS	5

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## **Question Paper Pattern- Model Question Paper**



- 6. a. Give on account on phases of clinical trial.
  - 'b. Explain the process of new drug development.
- 7. a. Write short note on AYUSH.
  - b. Explain the process of preparation of phytochemical extract.
- 8. Write short notes on any two of the following:
  - a. Pre-clinical studies
  - b. Biological action of curcumin and Ellargic acid
  - c. Concept of prodrug

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