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(ISO/IEC - 27001 - 2013 Certified) MODEL ANSWER WINTER- 18 EXAMINATION

Subject Title: Pharmacology & Toxicology

Subject Code:

0813

Important Instructions to examiners:

- 1) The answers should be examined by key words and not as word-to-word as given in the model answer scheme.
- 2) The model answer and the answer written by candidate may vary but the examiner may try to assess the understanding level of the candidate.
- 3) The language errors such as grammatical, spelling errors should not be given more Importance (Not applicable for subject English and Communication Skills.
- 4) While assessing figures, examiner may give credit for principal components indicated in the figure. The figures drawn by candidate and model answer may vary. The examiner may give credit for anyequivalent figure drawn.
- 5) Credits may be given step wise for numerical problems. In some cases, the assumed constant values may vary and there may be some difference in the candidate's answers and model answer.
- 6) In case of some questions credit may be given by judgement on part of examiner of relevant answer based on candidate's understanding.
- 7) For programming language papers, credit may be given to any other program based on equivalent concept.



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Q.	Sub	Answer	Marking
No	Q. N.		Scheme
1		Define any EIGHT of the following terms with one example of each.	16M
1	a)	Haematinics:	1M def.
		Are the agents which raise the number of RBCs & the amount of haemoglobin to normal	Any one
		level when they are below normal.	example
		Examples: Iron, folic acid, Vit B ₁₂ .	1M.
	b)	Miotics:	
		These are the agents that produce miosis or constriction of pupil.	
		Eg. Parasympathomimetics like physostigmine, pilocarpine	
	c)	Anti-hypertensives:	
		These are the agents used in treatment of hypertension.	
		Eg. Clonidine, Atenolol, Losartan, Methyl dopa,	
		Hydralazine,Reserpine,Propranolol,Diazoxide,Thiazides etc	
	d)	Antiemetics:	
		These are the agents used in treatment of vomiting.	
		Eg: Phenothiazine derivative(Chlorpromazine), Hyoscine, Meclozine, Promethazine,	
		Diphenhydrazine	
	e)	Analgesics:	
		These are the pharmacological agents which relieve or suppress the pain sensation.	
		Examples: Narcotic analgesics like Morphine, Codeine etc., Non narcotics like Aspirin,	
		Paracetamol, Indomethacin, Ibuprofen, Piroxicam, Diclofenac etc.	
	f)	Nasal Decongestants:	
		These are the agents which help in relieving nasal congestion & decrease resistance to	
		airflow through the nose.	
		Eg.Ephedrine,Pseudoephedrine,Phenylephrine,Oxymetazoline,Xylometazoline etc.	
		Eg.Ephedrine,Pseudoephedrine,Phenylephrine,Oxymetazoline,Xylometazoline etc.	



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g)	Carminatives:
	These are the pharmacological agents which when administered expel gas from the
	stomach or intestine during the treatment of flatulence and colic.
	Examples: Peppermint oil, Dill oil, Cardamom tincture, Ginger tincture, Simethicone etc
h)	Antibiotics:
	These are the chemical substances produced by microorganisms having the property of
	inhibiting the growth of, or destroying other microorganisms in high dilution.
	E.g Penicillins,(Penicillin-G, Amoxicillin etc.) Cephalosporins, (Cefadroxil, Cefalor etc.),
	Aminoglyoside, antibiotics (Streptomyin, Kanamycin etc.) Erythromyin, Azitromycin etc.
i)	Local anesthetics:
	These are the pharmacological agents which when applied or injected block the
	conduction as well as generation of impulses in localized area & bring reversible loss of
	sensation without affecting degree of consciousness
	Examples: Cocaine, Procaine, Amethocaine, Xylocaine, Cinchocaine
j)	Antifungals:
	These are the agents which selectively eliminate fungal pathogen from a host and are used
	in treatment of fungal infections.
	Eg.Nystatin,AmphotericinB,Clotrimazole,Miconazole,Ketoconazole,Fluconazole,
	Terbinafin, Griseofulvin, Benzoic acid, Salicylic acid, Selenium sulphide etc.
k)	Disinfectants:
	These are the pharmacological agents having bactericidal properties that can be directly
	applied to inanimate objects like surgical instruments, O. T. area, wards etc., for making
	them free from microorganisms.
	Examples: Phenols, Formaldehyde.
1)	Anorexiants;
	These are the agents which suppress appetite & help in treatment of obesity.
	Eg. Amphetamine, Fenfluramine, Mazindol, Phenyl propanolamine.



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2		Attempt any FOUR of the followings	12M
2	a)	Write mechanism of action & uses of Penicillin.	1.5 M
		Mechanism of action- Penicillin is a bactericidal; it interferes with the synthesis of cell	MOA.
		wall, by inhibiting mucopolypeptide of gram positive bacteria. This makes the cell	1.5M any
		membrane of microorganisms susceptible to damage by solutes in surrounding medium,	three
		i.e. plasma. Penicillins are effective mainly against multiplying organisms.	uses.
		Uses:	
		In treatment of Respiratory Tract Infection like Diphtheria,	
		Urinary Tract Infections,	
		Tetanus,	
		Venereal diseases such as Syphilis, Gonorrhea.	
		Infections of Endocardium (Bacterial endocarditis)	
		It is used against Pneumococcal, Streptococcal, Staphylococcal, Meningococcal	
		Infections.	
2	b)	Give symptoms & treatment in barbiturate poisoning.	1M
		Symptoms:-	Sympto
		Shallow respiration, fall in B.P., cardiovascular collapse, decreased or no urinary output	ms
		(anuria) pulmonary complications.	2M For
		Treatment:-	Treatme
		Gastric lavage: - If patient is conscious and within 4 hrs of ingestion, patient can be	nt
		induced vomiting with concentrated salt solution or syrup of ipecac. If patient is	
		unconscious, simple stomach wash i.e. gastric lavage is performed.	
		Artificial respiration: Endotracheal intubation: to treat hypoventilation	
		Supportive measures: Intravenous fluids to prevent dehydration, to maintain blood	
		volume and use of vasopressor if needed.	
		Alkaline diuresis: - with sodium bicarbonate 1meq/kg iv. With or without mannitol (is	
		helpful only in the case of long acting barbiturates which are eliminated primarily by	
		renal excretion).	



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2	c)	Mention various routes under Parenteral administration.	
		Explain advantages & disadvantages of Intravenous route of administration of	1M
		drugs.	EACH
		Parenteral routes:	
		Injections: Intravascular, Intramuscular ,Intradermal, Subcutaneous ,	
		Intrathecal, Intraperitoneal, Intramedullary, Intraarticular	
		Inhalation.	
		Advantages of IV: (any three points)	
		Useful for Unconscious or uncooperative patients	
		Useful in case of Vomiting, diarrhoea	
		No irritation of stomach	
		Provides rapid onset of action	
		Useful in case of Emergencies (lifesaving route)	
		Accuracy of dosage is achieved.	
		Disadvantages of IV: (any three points)	
		Once injected drug can't be withdrawn.	
		Irritation of veins may cause thrombophlebitis	
		Only aqueous solutions can be given	
		Self-medication is difficult	
		Trained personnel required for administering drug	
		There is risk of embolism.	
2	d)	Define Drug tolerance. Mention different types of drug tolerance.	1M
		Definition: When an unusually large dose is required to elicit an effect produced by the	Def.
		normal therapeutic dose of drug, the phenomenon is called drug tolerance. OR It is the	2M
		insensitivity towards the use of the drug which earlier had shown the therapeutic effect.	Types
		Different types of drug tolerance:-	
		1. True tolerance	
		a) Natural: Species, Racial	
		b) Acquired: Tissue, Cross	



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	1		
		2. Pseudotolerance,	
		3. Acute Tolerance / Tachyphylaxis.	
2	e)	Classify antineoplastic agents with examples.	3M.
		I. Alkylating agents:	
		Nitrogen mustards:E.g.: Chlorambucil, Mechlorethamine , Chlorambucil	
		Ethylenimines:E.g.: Triethylenemelamine, Triethylene thiophosphamide	
		Alkylsulphones:E.g. : Busulphan	
		II. Antimetabolites:	
		Folic acid antagonists:E.g.: Methotrexate	
		Purine Antagonist:E.g.: 6-mercaptopurine	
		Pyrimidine Antagonist:E.g.: 5-Flurouracil, Cytosine	
		III. Radioactive Isotopes: E.g.: Radioiodine, Radiophosphorous	
		IV. Antibiotics: E.g.: Actinomycin-D, Mitomycin	
		V. Hormones: E.g.: Androgens, Estrogens, Corticosteroids	
		VI. Enzymes:E.g.: L-asparaginase	
		VII. Miscellaneous Agents:	
		Vinca alkaloids: E.g.: Vincristine, Vinblastin	
		Others:E.g.: Hydroxyurea, Cis- platin	
2	f)	Explain the terms Pharmacokinetics and Pharmacodynamics.	1.5M.
			EACH
		Pharmacokinetics : It is the study of movement or passage of drug across the body. It is	
		what body does to the drug.	
		It includes study of Absorption, Distribution, Metabolism & Excretion (ADME) of drug.	
		Pharmacodynamics:	
		It is the study of how drugs affect our body or Study of mechanism of action & pharmacological effects of drug. It is the study of what drug does to our body	



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3		Attempt any FOUR of the followings		12M
3	a)	Mention the drug of choice for the following	g condition	1M each
		i. Glaucoma : Pilocarpine, Timolol, Beta	xalol, Physostigmine, Acetazolamide,	
		Glycerine, Mannitol.		
		ii. Insomnia: Diazepam ,Alprazolam, Nit	razepam, Barbiturates	
		iii. Cardiac Arrhythmia: Quinidine, Ligr	nocaine, Propranolol, Practolol,	
		Procainamide		
3	b)	Mention the drug contraindicated in follow	ing conditions:	1M each
		i. Head injury: Morphine		
		ii. Peptic ulcer: Hydrocortisone, Heparin	and other anticoagulants, Aspirin and other	
		NSAIDs.		
		iii. Liver cirrhosis: Phenobarbitone sodium	n / Alcohol	
3	c)	Mention use and adverse effect of the follow	ving:	0.5M Us
		i) Streptomycin:		0.5M
		Use: Antibiotic, Antibacterial agent	, TB., Meningitis, UTI, RTI, Plague, Dysentry	ADE.
		Adverse effect:- Ototoxicity, skin	rash, dermatitis,	
		ii) Quinine sulphate:		
		Use: Antimalarial agent		
		Adverse Effect:-Cinchonism inclu	ide tinnitus, deafness, optic neuritis	
		iii) Reserpine:		
		Use: Antihypertensive, Tranquilizer.		
		Adverse Effect:-Nasal congestion, sa	divation, vasodilation, increased motility of	
		gut, weight gain, mental depression, r gastric ulceration, stomach cramps, h	nightmares, insomnia, suicidal tendency	



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3	d)	Give the route of administration of the following drugs.	1M each
		i. Diclofenac : Orally, IM,IV, topical	
		ii. Griseofulvin : Orally	
		iii. Mannitol: - Parenteral (IV)	
3	e)	e) Mention antidotes in case of poisoning due to:	
		i. Atropine : Physostigmine	
		ii. Morphine: Naloxone, Nalorphine	
		iii. Organophosphorous compounds: Atropine Sulphate, Pralidoxime (PAM),	
		Diacetyl monoxime (DAM),Obidoxime.	
3	f)	Name one drug which produces following effects.	1M
		i. Mydriatics: Atropine , Homatropine, Cyclopentolate, Tropicamide, Ephedrine etc	EACH
		ii. Hypolipidemic: Ciprofibrate,Fenofibrate,Atorvastatin and other Statins,Nicotinic	
		acid, Gemfibrozil,	
		iii. Brochodilator : Adrenaline , Salbutamol, Isoprenaline,Orciprenaline,Ephedrine,	
4		Attempt any FOUR of the followings	12M
4	a)	Classify hypnotics with examples.	3M
		Classification-	
		I) Barbiturates-	
		a) Long acting barbiturates e.g. Phenobarbitone	
		b) Intermediate acting barbiturates e.g. Cyclobarbitone	
		c) Short acting barbiturates e. g. Hexobarbitone	
		d) Ultra short acting barbiturates e. g. Thiopentone	
		II) Non barbiturates	
		a) Benzodiazepine e.g. Diazepam	



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		b) Alcohols e.g. Chloral hydrate	
		c) Aldehydes e. g. Paraldehyde	
		d) Miscellaneous e.g. Hyoscine.	
4	b)	Enlist factors modifying drug action and explain any two of them.	1M
		Body weight	Enlist
		• Age	2M Exp.
		• Sex	
		Presence of disease	
		Route of administration	
		Time of administration	
		Genetic factor	
		Emotional factor	
		Metabolic disturbances	
		• Cumulation	
		Drug-drug interaction	
		Additive effect	
		• Synergism	
		• Antagonism	
		Drug tolerance	
		Drug dependence	
		i) Age-	
		Metabolic& excretory systems are not well developed in infants & may be less functional	
		in geriatric patients. So dose in pediatric & geriatric patients has to be adjusted.	
		Children are more sensitive to CNS depressants.	
		Infants below 1 yr. do not have fully developed enzymes that metabolize drugs. This must	
		be considered while calculating optimum dosage for infants. In aged individuals, the	
		normal dose may prove	



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to be excessive due to their inability to metabolize drugs.

- ii) **Presence of disease** antibiotics like streptomycin and kanamycin are mainly excreted by the kidneys. So they may prove toxic on parenteral administration in patients with improper kidney function. It is also observed that drugs like barbiturates produce unusually long effects in cirrhotic patients.
- iii) **Synergism** synergism is a pharmacological cooperation which usually results in total effect greater than the sum of their individual effects.

E.g Codeine and aspirin as analgesics. Sometimes synergism results in prolongation of action of one of the drugs called time synergism. Eg. Procaine and adrenaline in combination increases the duration of action of the local anaesthetic procaine.

iv) Additive effect:

When the total pharmacological effect of two or more drugs administered together is equal to the sum of their individual pharmacological actions, the effect is called additive effect.

e.g. Ephedrine and Aminophylline show additive effect in the treatment of bronchial asthma.

v) Drug-drug interactions

Drug-drug interactions are the result of the use of two or more drugs, they may occur either outside the body or within the body. They may be useful or harmful to our body.

Eg: Interaction between Tetracycline antibiotics and antacids, calcium supplements, milk products etc

Interaction between Ciprofloxacin and antacids, calcium supplements, milk products etc

Antacids enhance ionization & so excretion of weakly acidic salicylates.

Administration of Dimercaprol in treatment of arsenic poisoning.

Synergistic effect of trimethoprim & sulphamethoxazole

vi) Antagonism

Phenomenon of opposing actions of two drugs on the same physiological system.



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		Types:	
		Chemical: . Eg: Between acid &alkali.	
		Competitive / Reversible Eg: Acetylcholine& atropine antagonize at muscarinic receptors.	
		Noncompetitive Eg: Acetylcholine & papaverine on smooth muscles	
		Physiological Antagonism Eg; Adrenaline in histamine reaction.	
		Functional: One drug opposes another drug by virtue of its function.eg acetylcholine and adrenaline.	
		(Explanation of any other factors can be considered)	
4	c)	Explain mechanism of action of sulphonamides.	3M
		Many microorganisms require Para amino benzoic acid (PABA) for the synthesis of folic acid. PABA & sulphonamides are similar in chemical structure such that bacteria are not able to differentiate them. There is also competition between these two substances for same receptor site. Bacteria take up sulphonamide instead of PABA & inhibit formation of folic acid which is required for the bacterial growth and have bacteriostatic action. Sulphonamides are useful against the microorganisms which synthesise their own folic acid.	
4	d)	Define Cardiotonics. Explain the action of digitalis in CCF.	1M def.
		Cardiotonics: These are the drugs which increases work performance of heart.	2M Expl.
		OR Drug which increases force of contraction of heart (positive inotropic action).	
		Digitalis improves cardiac functions in CCF by following ways	
		• It strengthens the heart muscles, increases the force of cardiac contraction,	



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		positive inotropic action	
		It decreases heart rate,	
		It increases cardiac output and it improves blood perfusion to all organs, decreases	
		heart size	
		Decreases venous congestion	
		Increase renal blood flow and perfusion.	
		Relieves edema by producing diuresis.	
4	e)	Give Pharmacological action of insulin and mention various types of Insulin preparations.	1.5M
		Insulin is the hormone secreted by beta cells islets of Langerhans of pancreas.	Types
			1.5M
		Simulates the uptake of glucose – Insulin decreases blood glucose concentration by indusing intelligence by the calls.	Action
		by inducing intake of glucose by the cells.	Any 3
		 Insulin also increases synthesis and storage of glycogen in peripheral tissue such as heart and skeletal muscles. 	THIS C
		Insulin increases fat storage in adipose tissue by stimulating lipogenesis and	
		inhibiting lipolysis.	
		Decreases gluconeogenesis and glycogenolysis.	
		Insulin facilitates amino acid uptake and protein synthesis and inhibits protein	
		breakdown.	
		Insulin preparation are as follows	
		• Insulin injection I.P (soluble insulin)	
		Insulin zinc suspension	
		Insulin zinc suspension I. P. (amorphous) (insulin semilente)	



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		o Insulin zinc suspension I. P. (Insulin lente)	
		• Insulin zinc suspension I. P. (Insulin ultra-lente) protamine Zinc insulin I.P	
		Isophane insulin, globin Zinc insulin	
4	f)	Enumerate types of epilepsy and mention one drug used in treatment of each type.	3M
		i) Grandmal epilepsy: Phenytoin, Phenobarbitone.	
		ii) Petitmal epilepsy: Ethosuximide, Trimethadione, Clonazepam, Valproic acid	
		iii) Myoclonic jerking : Phenobarbitone, Clonazepam	
		iv) Atonic/Akinetic epilepsy: Diazepam, Clonazepam	
		v) Infantile spasms: Phenobarbitone, Diazepam, Clonazepam	
		vi) Psychomotor : Carbamazepine, Phenobarbitone, Divalproex, Phenytoin etc	
		vii) Status Epilepticus :Diazepam	
5		Attempt any FOUR of the following:	12M
5	a)	Enlist properties of ideal general anaesthetics.	3M
		Properties of an ideal General Anaesthetic:	Any6
		1. It should be pleasant, nonirritant, should not cause nausea or vomiting which if occur	
		may disturb the stitches.	
		2. It should be potent so that oxygenation of patient does not suffer.	
		3. Induction and recovery should be fast with no unpleasant after effects.	
		4. It should provide adequate analgesia i.e. loss of pain sensation,	
		5. It should produce adequate muscle relaxation so that surgeon can perform surgery with	
		ease.	
		6. Its administration should be easy.	
		7. It should have wide margin of safety.	
5	b)	What is Angina pectoris? Discuss different drugs used in the treatment of angina	1M def.



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		pectoris.	2M.
		Angina pectoris: - It is a symptom resulting due to cardiac ischemia (localised anemia).	Different drugs
		It is described as a condition in which there is a compressing type of pain in the chest.	
		1. Nitrates:-	
		a) Short acting:- eg. Glyceryl trinitrate, nitroglycerine.	
		b) Long acting:- eg. Isosorbide dinitrate, isosorbide mononitrate, pentaerythritol	
		tetranitrate.	
		2. Beta blockers:- eg. propanalol, metaprolol, atenolol	
		3. Calcium channel blockers:-	
		eg. Verapamil, Diltiazem , Nifedifine, felodipine, amlodipine	
		4. Potassium channel blockers:-eg. Nicrorandil	
		5. Others:- Ranolazine, oxyphedrine.	
5	c)	What are cough suppressants and expectorants? Explain with examples.	1.5M
		1.Cough Suppressants:-	EACH
		These are the dugs that act in the central nervous system to raise the threshold of cough	
		centre or act peripherally in the respiratory tract to reduce tussal impulsesor both these	
		actions.	
		Ex. Codeine, Ethylmorphine, Noscapine, Dextromethorphan.	
		2. Expectorants: These are the drugs which causes production of demulcent respiratory	
		tract fluid that covers the irritant mucosa.	
		OR	
		These are the drugs which increases the secretion of the respiratory tract, thereby	
		reducing the viscosity of the mucus and help in removal of the content from the	
		respiratory tract.	
		Eg: Ammonium chloride, Potassium iodide, Ammonium bicarbonate, Ipecac etc.	
5	d)	What are diuretics? Give major clinical uses of diuretics.	Defn 1M
		Divertion. These are the pharmacological agents which when administered increases	Uses 2M
		Diuretics:- These are the pharmacological agents which when administered, increases	For Any



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		rate of formation of urine as well as excretion of urine.	4
		Examples: Mannitol, Theophylline, Acetazolamide, Furosemide, Spironolactone,	
		Chlorothiazide.	
		Clinical uses:-	
		1. Oedema	
		2. Cerebral oedema	
		3. Acute renal failure	
		4. Acute pulmonary oedema	
		5. Forced diuresis	
		6. Hypertension	
		7. Renal stones	
		8. Drug Poisoning	
_	. `		23.4
5	e)	Classify Parasympathomimetics with examples of each class.	3M
		Parasympathomimetics- These are the drugs which produce the actions similar to those	
		seen by the stimulation of parasympathetic nervous system.	
		Classification:	
		☐ Esters of choline- Methacoline, carbachol, Acetylcholine	
		☐ Cholinomimetic alkaloids- Piolcarpine, Muscarine	
		☐ Cholinestrase inhibitors-	
		a) Reversible :-Neostigmine, physostigmine, pyridostigmine.	
		b) Ireversible:- Organophosphorus compounds, (malathion, parathion)	
		☐ Synthetic compounds- Futrethonium	
5	f)	Explain the term bioavailability of drug and mention factor affecting the same.	1M Expl.
	- <i>)</i>	Bioavailability : It is the amount of drug which actually reaches systemic circulation or	2M



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		site of action from a given dosage forms after its administration. This amount of the drug	factors
		is responsible for its therapeutic effect. Depending on the dosage forms, bioavailability	140015
		differs. E.g. After Intravenous route bioavailability is 100%.	
		Factor affecting:	
		- woos wassing	
		o Physical state of drug	
		o Particle size	
		o Concentration	
		Absorbing surface	
		o Functional integrity of GIT.	
		o pH of drug and pH of GIT	
		o Formulation	
		o Routes of administration	
		0	
6		Give reasons for any <u>FOUR</u> of the following:	16M
6	a)	Chlorpromazine is called largactil.	4M
		Chlorpromazine is the major tranquilliser possessing large number of pharmacological	
		actions. Hence, called as largactil (large acting).	
		Its actions are as follows:	
		☐ In patients with major psychoses it produces psychomotor slowing, emotional	
		quietening and diminishes initiative and anxiety.	
		☐ It depresses cortical region and reduces spontaneous motor activity.	
		☐ Antiemetic- chlorpromazine depresses C.T.Z and acts as antiemetic	
		☐ Hypotensive- It decreases the blood pressure by decreasing sympathetic tone	
		☐ Antihistaminic action	
		□ Anticholinergic action	
		☐ It causes skeletal muscle relaxation	



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6	b)	Aspirin and other NSAIDS should be taken after food.	4M
		Aspirin if administered on empty stomach causes gastric irritation, gastritis, nausea,	
		vomiting.	
		It is known to decrease gastroprotective prostaglandin levels which leads to ulceration	
		To avoid all these gastric side effects, aspirin is advised on full stomach.	
6	c)	Eating of cheese is forbidden in patients with MAO inhibitor therapy.	4M
		Cheese contains tyramine which is metabolized in the liver by the enzyme Monoamine	
		Oxidase. If an individual is on MAO inhibitor therapy, overall amine metabolism is	
		decreased and so there are already increased levels of amines in the body. When patient	
		takes tyramine containing foods like cheese, then MAO inhibitors inhibit the metabolism	
		of tyramine. Thus, tyramine gets accumulated in the body. Tyramine causes release of	
		noradrenaline from its binding sites.	
		Increased level of noradrenaline causes hypertensive crisis.	
		Therefore, eating of cheese is forbidden while on MAO inhibitor therapy.	
6	d)	Ephedrine is preferred to atropine to produce mydriasis in elderly patients.	4M
		Ephedrine interact with alpha adrenergic receptors in eye and produces mydriasis.	
		☐ It does not cause paralysis of ciliary smooth muscles or tightening of suspensory	
		ligament.	
		☐ It does not result into cycloplegia or photophobia as in case of atropine.	
		☐ So to avoid these visual complications Ephedrine is preferred to produce mydriasis in	
		elder patients.	
6	e)	Chloramphenicol therapy is supplemented with haematinics.	4M
		The therapeutic dose of chloramphenicol also causes bone marrow depression and	
		inhibits erythropoiesis.	
		This results in aplastic aneamia.	
		• To overcome these side effects and to promote erythropoiesis process, haematinics	



(Autonomous)

(ISO/IEC - 27001 - 2013 Certified) MODEL ANSWER WINTER- 18 EXAMINATION

Subject Title: Pharmacology & Toxicology

Subject Code:

6	f)	It is important to complete the course of antibiotics.	4M
		Antibiotics are used to cure infections by killing bacteria or by inhibiting their growth.	
		If antibiotic is discontinued abruptly all bacteria may not get killed. They have the	
		potential to grow again or multiply & may further aggravate the symptoms. Patient	
		may become sick again, and the remaining bacteria may become resistant to that	
		antibiotic. Resistance once developed, patient fails to respond to that particular	
		antibiotic in future.	
		So it is essential to complete the course of antibiotics.	