## MED 1208: Biochemical basis of cardiorespiratory functions, alimentation and nutrition Module - 2016/2017 Batch

Year 1 Semester 2

Credits: 5

Responsible Department: Biochemistry

Module Coordinator: Dr. B.L. Goonapienuwala

Topic	Time	Objectives	T/L activity	Comments
1. Exchange and transport of respiratory gases				
1.1 Alveolar surfactant	1 hr	<ol> <li>Define surface tension and describe how it applies to lung mechanics, including the effects of alveolar size and the role of surfactants.</li> <li>Describe the principal components of pulmonary surfactant and explain the roles of each.</li> <li>Explain the biochemical basis of infant respiratory distress syndrome (IRDS), adult respiratory distress syndrome (ARDS) and chronic obstructive pulmonary diseases (COPD) based on the derangement of lung surfactant.</li> <li>List the common toxicants which affect function of surfactant</li> </ol>	Lecture (1hr)	
1.2 Transport of respiratory gases	7 hrs	<ol> <li>Describe the modes of O<sub>2</sub> transport in blood</li> <li>Draw and explain the O<sub>2</sub>-haemoglobin dissociation curve</li> <li>List the factors affecting O<sub>2</sub>-haemoglobin dissociation</li> </ol>	Lecture (2hrs) PD (3hrs) SGD (2hrs)	PD on respiratory gas transport  SGD based on the lectures on

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1.3 Acid base balance	5 hrs	<ol> <li>curve</li> <li>Explain how, temperature, pH, 2,3 DPG, variants of haemoglobin and CO affect the affinity of hemoglobin to O<sub>2</sub> based on O<sub>2</sub>-haemoglobin dissociation curve</li> <li>Describe the modes of CO<sub>2</sub> transport in blood</li> <li>Describe the importance of the chloride shift in the transport of CO<sub>2</sub> by blood</li> <li>Draw the carbon dioxide dissociation curves for oxyand deoxy-hemoglobin</li> <li>Define pH</li> <li>Describe the importance of acid-base regulation in human body</li> <li>List the important buffer systems of human body</li> <li>Explain the buffering actions of bicarbonate buffer system, phosphate buffer system, protein buffer system and ammonia buffer system</li> <li>Define acidaemia, acidosis, alkalaemia and alkalosis</li> <li>List common causes for respiratory acidosis, respiratory alkalosis, metabolic acidosis and metabolic alkalosis</li> <li>State the methods of assessing acid-base status</li> <li>Interpret the arterial blood gas analysis report up to the level of diagnosis of respiratory acidosis, respiratory alkalosis, metabolic acidosis and metabolic alkalosis</li> </ol>	Lecture (2hrs) PD (3hrs)	surfactant, respiratory gas transport & haemoglobin
2. Blood and				
circulation 2.1 Haemoglobin	1 hr	Describe the basic structure of haemoglobin	Lecture (1hr)	
2.1 naeinogiobin	T 111	<ol> <li>Describe the basic structure of haemoglobin</li> <li>Explain the importance of structure of haemoglobin for its function</li> <li>State the different physiological types of haemoglobin and their functional significance</li> </ol>	Lecture (IIII)	
2.2. Haem metabolism and	1 hr	<ol> <li>Outline the synthesis of haem and its regulation</li> <li>Describe the process of haem catabolism</li> </ol>	Lecture (1hr)	Jaundice will be done in detail under
metabonsin and		2. Describe the process of flacin catabolism		aone in actail anael

jaundice		3. Def	ne jaundice		Alimentation - Liver
		4. Exp	ain hyperbilirubinaemia and jaundice based on		
		hae	m metabolism		
2.3. Abnormal	5 hrs		the types of abnormal haemoglobin	Lecture (2hrs)	
haemoglobins			cribe the structural abnormalities of haemoglobin	PD (3hrs)	
			emoglobin C, methaemoglobin, alpha and beta		
			assemia		
			cribe the molecular basis of hemoglobin S,		
			noglobin C, alpha and beta thalassemia		
		•	ain the functional defects of haemoglobin S,		
			noglobin C, methaemoglobin, alpha and beta		
			assemia		
2.4. Plasma	6 hrs		important plasma proteins (including lipoproteins)	Lecture (3hrs)	PD on plasma
Proteins			cribe the functions of plasma proteins (overall)		proteins and
			e the tissues involved in synthesis of plasma		electrophoresis
		•	teins		
			cribe the role of serum albumin	PD (3hrs)	
			cribe the basic steps in metabolism of		
		=	omicrons, VLDL, TAG, HDL and LDL		
		· -	ain the basis of electrophoresis based on		
		•	aration of plasma proteins		
			ly the knowledge of electrophoresis in		
			sification of plasma proteins		
			cribe the importance of assessing plasma proteins		
O.E. N. 1212 and	2 1		isease diagnosis	Lastura (2bus)	
2.5. Nutritional	2 hrs		all the basic steps of erythropoiesis	Lecture (2hrs)	
factors affecting			the nutritional factors involved in erythropoiesis		
erythropoiesis		=	ain the role of iron, folate and vitamin B <sub>12</sub> in		
(Iron, Folate and		-	hropoiesis		
Vit B <sub>12</sub>			cribe the consequences of iron, folate and vitamin deficiency relating to erythropoiesis		
metabolism)					
			all common causes for iron, folate and vitamin $B_{12}$ ciencies		
			rpret the investigations of iron, folate and vitamin		
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	B <sub>12</sub> deficiencies (red cell morphology, serum iron,	
	ferritin, transferrin, TIBC, transferrin saturation,	
	serum vitamin B <sub>12</sub> , serum folate, intrinsic factor etc.)	
2 hrs	<ol> <li>Recall the functions of erythrocytes</li> </ol>	Lecture (2hrs)
	2. Describe the metabolic adaptations of erythrocytes	
	(glycolysis, HMP shunt, production of 2,3-BPG)	
	3. Explain the importance of above metabolic	
	adaptations for the functions of erythrocytes	
	4. Explain the importance of NADPH for erythrocytes	
	5. State the common enzyme defects of erythrocytes	
	metabolism (pyruvate kinase deficiency and glucose	
	6-phosphate dehydrogenase deficiency)	
	6. Describe the consequences of glucose 6-phosphate	
	dehydrogenase deficiency	
	7. Describe the structure of erythrocyte membrane and	
	cytoskeleton	
	8. State the diseases associated with alterations of	
	erythrocyte membrane and cytoskeleton	
	9. Describe the basis of hereditary spherocytosis,	
	hereditary elliptocytosis and hereditary ovalocytosis	
	based on the genetic defects of proteins associated	
	with erythrocyte membrane	
1 hr	1. Define hemolytic anemia	Lecture (1hr)
	2. Outline the causes of hemolytic anemia	
	(intracorpuscular and extracorpuscular)	
	3. Explain the mechanisms of haemolysis based on	
	above causes	
	4. Describe the fate of hemoglobin in excessive	
	intravascular haemolysis (haemoglobinaemia,	
	haemoglobinuria, jaundice etc)	
	5. Describe the general features of hemolytic anemia	
	based on the excessive haemolysis and abnormal	
	bilirubin metabolism (pallor, reduced haemoglobin,	
	reticulocytosis, enlargement of spleen, increased	
		ferritin, transferrin, TIBC, transferrin saturation, serum vitamin B <sub>12</sub> , serum folate, intrinsic factor etc.)  1. Recall the functions of erythrocytes 2. Describe the metabolic adaptations of erythrocytes (glycolysis, HMP shunt, production of 2,3-BPG) 3. Explain the importance of above metabolic adaptations for the functions of erythrocytes 4. Explain the importance of NADPH for erythrocytes 5. State the common enzyme defects of erythrocytes metabolism (pyruvate kinase deficiency and glucose 6-phosphate dehydrogenase deficiency) 6. Describe the consequences of glucose 6-phosphate dehydrogenase deficiency 7. Describe the structure of erythrocyte membrane and cytoskeleton 8. State the diseases associated with alterations of erythrocyte membrane and cytoskeleton 9. Describe the basis of hereditary spherocytosis, hereditary elliptocytosis and hereditary ovalocytosis based on the genetic defects of proteins associated with erythrocyte membrane 1 hr 1. Define hemolytic anemia 2. Outline the causes of hemolytic anemia (intracorpuscular and extracorpuscular) 3. Explain the mechanisms of haemolysis based on above causes 4. Describe the fate of hemoglobin in excessive intravascular haemolysis (haemoglobinaemia, haemoglobinuria, jaundice etc) 5. Describe the general features of hemolytic anemia based on the excessive haemolysis and abnormal bilirubin metabolism (pallor, reduced haemoglobin,

			MCV, elevated lactate dehydrogenase, etc.)	
2.8. Role of the	2 hrs	1.	State the functions of vascular endothelium	Lecture (2hrs)
vascular		2.	List vasodilators and vasoconstrictors produced by	
endothelium in			endothelial cells	
regulation of		3.	State how NO is synthesized and describe the	
blood			regulation	
pressure/blood		4.	State the role of NO, prostacyclins, endothelins etc.	
flow		5.	Outline the role of vascular endothelium in relations	
			to rennin-angiotensin-aldosterone system in	
			maintaining blood pressure	
		6.	Explain the rationale for the use of ACE inhibitors in	
			the regulation of blood pressure	
3. Muscle and				
exercise				
3.1. Biochemical	5 hrs	1.	Define oxygen debt	Lecture (2hrs)
Effects of		2.	Describe the metabolic pathways in relation to the	
exercise on			intensity of exercise	PD (3hrs)
muscle and other		3.	Describe the changes in the muscle following muscle	
tissues			fatigue after exercise and lactic acidosis	
		4.	Describe the biochemical basis of the recovery of	
			skeletal muscle after exercise	
			Describe energy cost of exercise in terms of BMR	
		6.	State the receptor changes in exercise (LDL, insulin	
			etc) and their impact	
		7.	Explain the changes in blood lipid profile with exercise	
3.2. identification	5 hrs	1.	Define "biomarker"	Lecture (2hrs)
of muscle			Describe the features of a good biomarker	
damage		3.	List the biomarkers that are useful in identifying	PD (3hrs)
			skeletal and cardiac muscle damage	
		4.	Explain how skeletal muscle damage could be	
			differentiated from cardiac muscle damage based on	
			change of serum parameters	
		5.	State the alteration in activity of the enzymes in	
			serum following skeletal muscle damage	

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		6. Define the term 'myoglobinuria'		
		7. State the effects of myoglobin on nephrons		
		8. Outline the basis for selection of biomarkers used in		
		the identification of muscle damage		
		9. List the enzymes used in diagnosis of myocardial		
		damage		
		10. Describe the place of troponin, LDH, CK levels in		
		relation to diagnosis of myocardial damage		
		11. Describe how myoglobinuria is detected in urine		
4. Digestion and				
absorption				
4.1. Liver and bile	11 hrs	<ol> <li>Describe the functions of the liver</li> </ol>		
		2. State the constituents of exocrine secretion and its		
		importance		
		3. Explain the metabolic functions of the liver		
		(Carbohydrate metabolism, synthesis and secretion of	Lecture (2hrs)	
		proteins, lipid metabolism, cholesterol catabolism,		
		haem metabolism)		
		4. State the constituents of bile and its importance	PD (2x3hrs)	PD 1 on objectives 4,
		5. Describe the factors that promote the formation of		8 & 9
		gall stones		
		6. State the composition of different types of gall stones	SGD (2hrs)	PD 2 on objectives
		commonly found		10 – 14
		7. Describe the role of liver in metabolism of xenobiotics		
		8. State the liver enzymes and other plasma parameters		SGD on all
		(plasma proteins) useful in diagnosis of liver		objectives
		dysfunction		
		9. Correlate the degree of cell damage with change in		
		the serum enzyme levels and plasma proteins		
		10. Describe the types of jaundice		
4.2. Jaundice		11. Differentiate pre-hepatic, hepatic and post-hepatic		
		jaundice based on etiology	Lecture (1hr)	
		12. Explain the biochemical changes that occur in each	, ,	
		type		
		7,70		

4.3. Digestion & absorption of carbohydrates	8 hrs	<ol> <li>Describe the biochemical basis of clinical features and laboratory investigations of jaundice</li> <li>Interpret the laboratory investigations of jaundice</li> <li>State the constituents that are important in the digestion of carbohydrate in saliva, pancreatic juice and brush border</li> </ol>	Lecture (1hr)	Lecture 1 from objectives 1 – 3
		<ol> <li>Explain carbohydrate digestion in the GI tract</li> <li>State the products of carbohydrate digestion and explain their absorption.</li> </ol>		
4.4. Digestion & absorption of proteins		<ol> <li>Name the important constituents of GI and pancreatic secretions involved in protein digestions and explain their functions in protein digestion</li> <li>Explain the importance of secreting proteolytic enzymes in their pro-forms and explain how the gut wall is protected from the activated proteolytic enzymes</li> <li>Outline the process of protein digestion</li> <li>State the products of protein digestion and explain how they enter the absorptive cell</li> </ol>	Lecture (1hr)	Lecture 2 from objectives 4 – 7
4.5. Digestion and absorption of lipids		<ol> <li>Recall the constituents of bile and explain their role in lipid digestion</li> <li>Explain the importance of emulsification process and micelle formation in the digestion process of lipids</li> <li>Name the enzymes involved in lipid digestion and explain their functions</li> <li>Describe the absorption of digestive products of lipids (including fat soluble vitamins)</li> <li>Explain the chylomicron formation in the enterocyte</li> <li>Describe the enterohepatic circulation and its importance in fat digestion</li> </ol>	Lecture (1hr) PD (3hrs) SGD (2 hrs)	Lecture 3 from objectives 8 – 13  PD from objectives 1,2,6,9 and 10

		14. State the changes that occur in the large intestine with respect to Fiber, Vitamin B <sub>12</sub> and Vitamin K		
4.6. Digestive disorders	4 hrs	<ol> <li>Explain how the following conditions affect digestion and absorption         <i>Digestive disorders</i> <ul> <li>Achlorhydria</li> <li>Intrinsic factor deficiency</li> <li>Biliary insufficiency</li> <li>Lactose intolerance</li> <li>Pancreatic exocrine insufficiency</li> <li>Coeliac disease</li> <li>Chronic diarrhoea</li> <li>Chronic alcoholism</li></ul></li></ol>	Lecture (2hrs) PD (2hrs)	PD on objective No. 4
5. Nutritional		assess the disorders stated in 1		
Biochemistry – 2			(4)	
5.1. Healthy diet	1 hr	<ol> <li>State what is healthy diet</li> <li>State what is "MyPlate"</li> <li>State the characteristic features of healthy diet and MyPlate</li> </ol>	Lecture (1hr)	
_		4. Describe Sri Lankan "healthy plate"		

5.2. Components of diet and food processing	10 hrs	products, de properties, pthe nutrition a. cereals b. pulses c. vegetabl d. oil seeds e. meat, fis	mportance, state the types and their escribe the nutritional value, antinutrient processing methods and their effects on hal value of the following in diet.  Ite, fruits and starchy foods and nuts including coconut is and egg	Lecture (1hr) Lecture (1hr) Lecture (1hr) Lecture (1hr) Lecture (1hr) Lecture (1hr)	
		colostrum ar	e nutritive value and the importance of nd breast milk	Lecture (1hr)	
			e methods used to minimize losses of Iring processing and increase the bio- of nutrients	PD (3hrs)	PD on objective No 3
5.3. Dietary fibre	1 hr	<ol> <li>State the typ</li> <li>State the foo</li> </ol>	erm dietary fibre bes of dietary fibre od items rich in dietary fibre health benefits of dietary fibre	Lecture (1hr)	
5.4. Energy requirement	6 hrs	<ol> <li>Recall why e</li> <li>Recall the so</li> <li>Define Resting Metabolic Ray</li> <li>(SDA)</li> <li>State factors</li> <li>Compare an requirement</li> <li>Calculate tot BMR factor</li> <li>Explain the Factors on the renergy in the</li> </ol>	energy is needed for the body burces of dietary energy ng Energy Expenditure (REE), Basal ate (BMR) and Specific Dynamic Action s influencing BMR d contrast BMR and the total energy t tal energy expenditure using BMR and FAO/WHO/UNU recommended intakes of e various phases of life (newborn, infants, I adolescents, adults and elderly,	Lecture (2hrs)	SGD on both energy and protein requirements

5.5. Protein		1	Explain why protein is essential in the diet	SGD (2hrs)
requirement			Explain what is nitrogen balance (zero, positive and	300 (21113)
requirement		2.	negative)	
		3.	State the parameters for evaluating quality of	
			proteins; Biological Value (BV), Net Protein Utilization	
			(NPU), digestibility and amino acid score, protein	
			digestibility corrected amino acid score (PDCAAS) etc.	
		4.	Define and compare Biological Value (BV), Net Protein	
			Utilization (NPU), digestibility and amino acid score	
			and protein digestibility corrected amino acid score	
			(PDCAAS)	
		5.	Compare the quality of protein sources commonly	Lecture (2hrs)
			used in Sri Lanka	
		6.	Discuss the FAO/WHO/UNU recommended intakes of	
			proteins in the various phases of life (newborn,	
			infants, children and adolescents, adults and elderly,	
			pregnancy, lactation)	
5.6. Protein	1 hr		Define wasting and stunting	Lecture (1hr)
energy			Explain kwashiorkor and marasmus	
deficiencies		3.	Explain the use of growth parameters on identifying	
			wasting and stunting	
		4.	Identify laboratory and clinical features associated	
			with malnutrition including kwashiorkor, marasmus,	
	6.1		wasting and stunting	(2)
5.7. Free radicals	6 hrs		State what are free radicals and antioxidants	Lecture (2hrs)
& antioxidants			State the types of free radicals	
		3.	State the sources of free radicals (exogenous and	
			endogenous) State the harmful effects of free radicals	
		3.	Explain the antioxidant systems in the body (enzymatic / non enzymatic and dietary)	
		6	Explain the relationship of oxidative stress in ageing	
		0.	and human diseases; atherosclerosis, diabetes,	
			haemolysis, neurodegenerative disorders, cancer etc	
	<u> </u>		macmoryolo, medioacychiciative alsoraers, cancer etc	

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5.8. Biochemical		<ol> <li>Describe the metabolism of alcohol in chronic</li> </ol>	Lecture (2hrs)	
effects of		alcoholism		
alcoholism		2. Describe the effects of alcohol on metabolism		
		3. State the possible organ failures in chronic alcoholism	SGD (2hrs)	
		4. Explain the consequences of chronic alcoholism		
		(chronic pancreatitis, fatty liver, liver cirrhosis)		
		5. Explain the nutritional problems arising in chronic		
		alcoholism (iron, vitamin B and protein)		
		6. Outline the management of nutritional deficiencies in		
		chronic alcoholism		
5.9. Dietary	1 hr	1. List the common diseases requiring special dietary	Lecture (1hr)	
management in		management (acute and chronic renal failures,	, ,	
diseases		chronic liver failure, dyslipidemia, diabetes, chronic		
		alcoholism, hepatitis, pancreatic insufficiencies,		
		malabsorption etc.)		
		2. Explain the biochemical basis of dietary management		
		of above diseases		
6. Student	3 hrs	Present and discuss the key areas that were learnt during the	Student	Holistic approach on
centered learning		module	presentations	the module.
activity			(3 hrs)	Revision of major
•				topics by presentation
				and discussion to
				improve the student-
				centered learning.

Lectures – 50 hours

SGD – 10 hours

Practical – 32 hours

Seminar – 3 hours