

School: School of Science
Program/s: Biomedical Science
Year: 2nd Semester: 1X

Examination: End Semester Examination

Examination year: December - 2022

Course Code: BM502 Course Name: Regenerative Biomedicine and Tissue Engineering

 Date:
 02/12/2022
 Total Marks:
 40

 Time:
 11:30 AM to 01:30 PM
 Total Pages:
 2

Instructions:

→ Write each answer on a new page

→ Draw neat and well-labelled diagrams wherever required

→ * COs=Course Outcome mapping. # BTL=Bloom's Taxonomy Level mapping

Q. No.	Det	Mark	s COs	BTL
2.1	Choose the correct option	12	COI	BTLI
		1.0	CO2	BTL2
	Which of the following are not myeloid of		CO3	BTL3
	a. Macrophages	b. Monocytes	CO4	
	c. Neutrophils	d. T cells	CO5	
	Which one of the following genes was NOT part of transcription factors used to generate induced pluripotent stem (iPS) cells from mouse skin fibroblasts? a. Oct 4			
	3. What is the major concern of using retrov pluripotent genes?	al vectors to deliver or activate		
	a. Not very efficient and needs repeated transfections	b. Slow and unstable		
	c. Vector and transgenes remain in the ge			
	and can be reactivated in differentiated ce	s excised by transposase		
	4. What are stem cell scientists investigating	oday?		
	a. When and how embryonic stem cells m			
	decisions to produce more specialised cel			
	c. How stem cells might be used to treat of			
	5. Neural stem cells from the brain can diffe	entiate into which types of cell?		
	a. Only specialized brain cells b. Specialized brain and skin			
	a. o.n., specialized orani cons	cells		
	c. All types of specialized cells	d. Only specialized blood cells		
	6 Biometerial belonging to cate	on, possess high tensile strength		
	Biomaterial belonging to category, possess high tensile strength. Bioceramics			
	c. Metal alloys	d. All of the above		

	7	which leads to	time of cell . It is			
	the tra	nsformation process during anit	type of cell-cell interaction, chiefly drive			
	the transformation process during epithelial to mesenchymal transition a. Cadherins, heterodimeric h. Integrins, homodimeric					
	c. Selec	tins, homodimeric	b. Integrins, homodimeric			
			d. None of the above			
	8 Number of hematopoetic stem cells increase with response to the increase in the					
	activity of cells					
		eoclasts	b. Osteoblasts			
	c. Mes	senchymal cells	d. None of the above			
	Stem cells for hair follicle niche are found in					
	a. Dermal papilla b. Basal layer c. Dermal sheath					
	c. Der	mal sheath	d. None of the above			
	10cells generate inner root sheath and hair shaft during the growth phase of hair follicles					
	a Dermal parille					
	c Der	mal sheath	b. Matrix cells			
	c. Dei	mar sheath	d. None of the above			
	H. Wnt (ligand) is a				
	a. Pro		b. Lipoprotein			
	c. Gly	coprotein	d. None of the above			
		•	30.0000			
	12. The Wnt signalling in HSC niche is described as					100
	a. an autocrine induction b. an endocrine industrion					
	c. a para	crine induction	d. None of the above			
Q.2	Ancwontho	fallowing to all				
Q.2	Answer the following in short. Any six					
	1. What are the differences between adult stem cells and embryonic stem cells?					
	2. What are induced pluripotent stem cells?					
	3. What is Somatic Cell Nuclear Transfer?				COI	
					CO2	BTLI
	the stell cells come from:			12	CO3	BTL2
	the ney entitleteristics of biolitaterials to be used for tissue engineering?				CO4	BTL3
	 Explain types of epithelial to mesenchymal transitions and their specific role development and survival of humans. 		chymal transitions and their specific roles in the		CO5	
	7. Define the roles of cells found in "rosette structure".					
	Define the role of Slug/Snail during epithelial to mesenchymal transition.					
0.2	Anguar the fellowing in 1 a 19					
Q.3	Answer the following in detail. Any four					
	1. What are the types of mesenchymal stem cells? What can mesenchymal stem cells					
	differentiate into?					
	Provide detailed insight into the gut-epithelial stem cell niche and explain the					
	roles of various cell types.					
				COI		
	y an analysis and stem cens being used for neurodegenerative			CO2	BTL1	
	diseases. Citing an example of any one such disease; explain at length the		16	CO3 CO4	BTL2 BTL3	
	application of stem cells in clinical application of treating neurodegenerative diseases.					1.00000
				CO5		
	4. For type I diabetes patients the only hope is beta cell regeneration and					
	replacement. Give your views and discuss in detail the possibility of developing					
	beta cells from stem cells for such application. Describe the underlined					
		echanism.				
			echnology for drug discovery and			
	de	velopment.				
		******	*****End of Question Paper********		-	