# M.TECH/BT/3<sup>RD</sup> SEM /BIOT 6153/2015 2015

# Biopharmaceuticals (BIOT 6153)

Time Allotted: 3 hrs

Full Marks: 70

Figures out of the right margin indicate full marks.

Candidates are required to answer Group A and any 5 (five) from Group B to E, taking at least one from each group.

Candidates are required to give answer in their own words as far as practicable.

Gro	up – A	
(Multiple Choice	Type Questions)	
1. Choose the correct alternatives for the following:		10 x 1=10
(i) Organisms suitable for use in modified li-	ve vaccines are produced b	y
(a) inactivation	(b) genetic recombination	
(c) attenuation	(d) complement fix	
(ii) A suitable organism for use in recombin	ant vaccines is	
(a) influenza virus	(b) smallpox virus	
(b) polionyelitis virus	(d) vaccinia virus.	
(iii) One problem why vaccines fail to work i	n very young infants is the	presence of
(a) maternal antibodies	(b) glycoproteins	
(b) endotoxin	(d) serum.	
(iv) IL-10 acts to		on the there easi
(a) enhance T-cell response	(b) activate macrophages	
(c) suppress cytokine production	(d) suppress antibody production.	
(v) Hemophilia A is due to deficiency of clott	ing factor :	
(a) X (b) XIII	(c) XIII	(d) V.
(vi) In JAK-STAT pathway phosphorylation of	ccurs at	
(a) Tyrosine residue	(b) Alanine residue	
(c) Tryptophan residue	(d) None of these.	
(vii) DNAse is used to control:		
(a) Cystic fibrosis	(b) Gaout,	
(c) Cancer	(d) none.	
(viii) Cytokine receptor is composed of		
(a) two separate subunit	(b) one subunit	
(c) both a and b	(d) none of these.	

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- ix) Urate oxidase is used to control
  - (a) milk intolerance

(b) cystic fibrosis

(c) gout

- (d) None of these.
- k) In JAK-STAT pathway phosphorylation occurs at
  - (a) tyrosine residue

(b) alanine residue

(c) tryptophan residue

(d)none of these.

#### Group - B

- a) Discuss about the oral and pulmonary route for delivery of biopharmaceuticals.
- b) What are the roles of Bioavailability and Bioequivalence studies during drug testing? [3+3]+[3+3]=12
- (a) What are the principles behind rational drug design?
- (b) Comment on the impact of genomics on drug discovery.
- (c) Describe the different aspects of phase-I, phase-II and phase-III clinical trials for drug testing.

4+4+4=12

### Group - C

- a) Define cytokine. Give two examples.
- b) Schematically describe JAK-STAT pathway.

$$(2+1)+9=12$$

- a) What are different types of interferons? Write down the specific mode of action of interferon.
- b) Write notes on cytokine receptor.

$$(2+5)+5=12$$

# Group - D

- a) What are polyclonal antibodies?
- (b) Discuss the role of anti-D immunoglobulins in human therapeutics.
- (c) How are snake and spider antivenoms used as therapeutic agents?

2+5+5=12

- 7.(a) Mention the uses of HepatitisB and Tetenus immunoglobulins as biopharmaceuticals.
  - (b) How can you design a cancer vaccine?
- (c) Discuss the benefits of bone marrow stem cells in human therapeutics.

4+4+4=12

#### Group - E

- 8.(a) What is CHO cell line?
  - (b) Why it is preferred host for cloning mammalian protein pharmaceuticals?
  - (c) Write down different steps for expressing human protein gene in CHO cell line?

2 + 3 + 7 = 12

- 9. Write short notes on (any three):
  - i) Human blood substitute
  - ii) Hematopoeitic stem cell
  - iii) Alpha galactosidase as pharmaceuticals
  - iv) Erythropoietin
  - v) Platelet function in blood clotting

 $(3 \times 4) = 12$