Engineering Biology C3 - Exam

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Oranierone Marchall A Danck A AUG COS

1 Antisanse strand: 3'-TACGIAACCAATT-5'
mRNA encoding: 3'-AUGICAAUGIC-5'

- DNA polymerase can proceed only 5' to 3'.

 Thus short fragments are produced as the replication fork expands. Otherwise DNA polymerase used to use the replication fork expands. Otherwise DNA replication fork expands to use the polymerase would need to use until the fork polymerase would need to use to replicate the lagging strand.
- (3) longth of coding region = 1800 base pairs.

 each amine acid corresponds to 3 base pairs

 no of codons = 1800 = 600 codons.

ule will get 599 amino acids because the last one is stop codon. molecular weight = 599×110

The P-site (for keptidyl) his the second binding site for tRNA in the ribosome. The other two sites are the A-site (Aminoayl), which is the first binding site in the ribosome, and the E-site (exit), the third. During protein translation, the P-site holds the tRNA which is linked to the growing polypeptic chain.

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(5) to 3' polarity refers to the carbon numbers of the sugars ribose and deoxyribose ring. It refers to the orientation of a single strand of 9 trefers to the orientation of a single strand of DNA or RNA. The phosphate group attached to DNA or RNA. The phosphate and the hydrory's the 5' end of one nucleotide and the hydrory's group at the 3' end of another nucleotide have group at the 3' end of another nucleotide have the polarical to form phosphatester bonds, and the polarical to form phosphate backbone that provides the sugar-phosphate backbone that gives DNA its structural rigidity. Any single gives DNA | RNA will always have an unbound strad of DNA | RNA will always have an unbound 5' phosphate at one end and an unbound 3' hydrory group at the opp and

6) when a protein is denatured, secondary and tertiary structures are altered but the peptide bonds of the primary structure between the amino acido are left intart. Since all structural levels of the protein determine its function, the prolein can no longer perform its function once it has been donatured. F) Number of ontigen binding sites

ig G = 2, ig M = 10, ig A = 2

(8) Defined Media

Total cells =

Composed of pure

dilution factor = 8

cell count = 800

no of square 2 5

tio chamicals off the as blood or milk or years extracted for which Shelf. Exact chamical accurate chamical composition composition is known is not determined. dil factor x 1000
no of squares alls/ml total alla X counted (9) total cells/ml=

800 X 8 x 1000

complex media

Contains complex material

of biologically origin such

cells/mL = 1280000

10) After RNA is transcribed, it is often modified to create a mature RNA that is ready to be transplated translated. Processing mRNA involves the removal of introns that do not code for protein. Splice osome remove the introno and ligate the exono together, often in different sequences than their original order of newly transcribed mRNA. A GTP cap is added to 5'- end and a poly-A tail at 3' and - This mature mRNA than leaves the nucleus and enters the cytoplasm. Once in the cytoplasm, the length of teme the mRNA resides there can be altered to control the amount of protein that is synthesized

(1) Snoculation is the process of introducing micro-organisms or suspension of microorganisms into a culture medium. Incubation is the process of allowing inoculated microorganisms to grow under required around 1-1-1-

growing conditions.

The principle of Pour Plate method: In this method, serial dillutions of the inoculum (serially dilluting the primary specimen) are added within Sterile petri plates to which is poured melted and cooled again medium and completely mixed by revolving the plates unlich are then left to solidify

(13) In order to be capable of engaging the key elamates of adaptive immunity (specifically, momony, diversity) of adaptive immunity (specifically, momony, diversity) antigens have to be processed and presented to antigen presentation is mediated immune cells. Antigen presentation is mediated immune cells. I molecules, and the class II by MHC class I molecules, and the class II molecules found on the surface of antigen-presety molecules found on the surface of antigen-presety aclos and certain other cells.

MHC class I and II molecules are similar in function:

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they deliver short peptides to the cell surface, allowing they deliver short peptides to the cell surface, allowing they deliver short peptides to the cell surface, allowing they deliver short peptides to the cell surface, allowing they deliver short peptides to the cell surface, allowing they deliver short peptides to the cell surface, allowing they deliver short peptides to the cell surface, allowing they deliver short peptides to the cell surface, allowing they deliver short peptides to the cell surface, allowing they deliver short peptides to the cell surface, allowing they deliver short peptides to the cell surface.

these keptides to be recognised by CD8+ and CD4+
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Tells. The diff is that the keptides originate from
Tells. The diff is that the keptides originate from
different sources; endogenous and errogenous. There
different sources; endogenous and errogenous in which
is also so called cross- presentation in which
errogenous antiques can be presented by MMC class I
errogenous antiques can also be presented
errogenous antiques can also be presented
molecules. Endogenous antiques can also be presented
by MMC class II when they are degraded through
autophagy.

19 methionine and tryptophon are the only two amino acids that are coded for by just and amino acids that and UCG respectively) a single codon (AUG and UCG respectively)

(5) Fats and oils are called triglycerides because they are esters composed of three fatty acids units joined to glyceral, a trihydroxy alcohol.

A triglyceride is called a fat if it is a solid at 25°C: it is called an oil if it is liquid at that temperature. These differences in melting points reflect differences in the degree of unsaturation and number of carbon atoms in the constituent fatty acids.

(6) Mentralizing antibodies can inhibit the infertivity by binding to the pathogen and block the by binding to the pathogen and block the molecules needed for cell entry. This can be molecules needed for cell entry interfering due to the antibodies statically interfering due to the antibodies statically interfering uith the pathogens or tosins attaching to host cell receptors.

(in pul)

No. of cells/ml = No. of colonies × 1000

dilution × V deeme of culture

(in pul)

= 400 × 10 × 10 3 cells/l = 4× 10 cells/letr

(18) The fundamental underlying mechanisms of autoimmunity is défertine climination or control of self-reactive lymphocytes. The genetic and environmental factors also contribute to autoimmunity.

HECMOH HDOH CHZOH (H₂OH H CHLOH CHLOIT