

Question	Answer	Mark
1 (a)	A → C → D → B → F → E	1

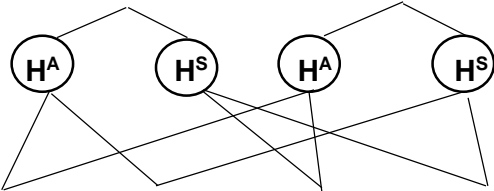
(b)	Structure X : Centriole	1
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	Function : Organize the formation of spindle fibre	1
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(c)

Stage of process	Mitosis	Meiosis I	
Prophase	No synapsis occur// no crossing over between chromosomes// no formation of chiasmata	Homologous chromosomes pair up during synapsis// crossing over occurs//formation of chiasmata// chiasmata hold homologous chromosomes together	1
Metaphase	Chromosome/ sister chromatids align at metaphase plate// sister chromatids of chromosome face each pole// both side of centromere attaches to spindle fiber// no chiasmata	(Pairs of) homologous chromosomes/ bivalent/ tetrad align/ line up/ arranged at metaphase / cell equator/ metaphase plate/ plane// one chromosome of each pair face each pole// only one side of centromere attached to spindle fibre// chia	1
Anaphase	Sister chromatids separated and move to the opposite poles	(Pairs of) homologous chromosomes separated and move to the opposite poles	1

(d)	DNA replication occur once// cytokinesis occurs at the end of the process	1
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Question	Answer	Mark
2 (a) (i)	Allele is the alternative forms of a gene	1
(ii)	<p>Parental phenotype : Normal male Normal female</p> <p>Parental genotype : $H^A H^S$ $H^A H^S$ x $H^A H^S$</p> <p>Gamete : H^A H^S H^A H^S</p>  <p>F₁ genotype : $H^A H^A$ $H^A H^S$ $H^A H^S$ $H^S H^S$</p> <p>F₁ phenotype : Normal Normal Normal Sickle cell Anaemia</p>	1 1 1
	<p>Explanation:</p> <ul style="list-style-type: none"> - Sickle cell anaemia is controlled by recessive allele// Sickle cell anaemia are express by homozygous recessive allele// Sickle cell anaemia condition is recessive phenotype. - A mating between heterozygous parent/ carrier genotype parents may formed sickle cell anaemia condition in child. 	1 1
(iii)	$\frac{1}{4}$	
(b) (i)	Sex-linked gene	1
(ii)	Haemophiliac male: $X^h Y$	1
	A woman with normal blood clotting but have a son with haemophilia: $X^H X^h$	1
(iii)	<ul style="list-style-type: none"> — His mother is a carrier/ heterozygous — Haemophilia is X-linked gene — Son always inherit X-chromosome from mother — Male only have one X-chromosome 	1 1 1 1
	Any 3	
3 (a)	(Extremely) large population size	1
	Random mating	1
	No (net) mutation	1
	No migration/No gene flow	1
	No natural selection	1
	Any 2	

Question	Answer	Mark
(b)	From, $p^2 + 2pq + q^2 = 1$ Frequency of homozygous recessive genotype, q^2 = 20/100 =0.20 So, frequency of recessive allele, q = $\sqrt{0.20}$ = 0.45 Since, $p + q = 1$ Frequency of dominant allele, p = $1 - q$ = $1 - 0.45$ = 0.55 Frequency of heterozygous genotype = $2pq$ = $2(0.55)(0.45)$ = 0.50 Therefore, the number of heterozygous squirrels in the population = $2pq \times 100$ = 0.50×100 = <u>50 (squirrels)</u>	1 <

Any 2

Question	Answer	Mark
(v)	The Okazaki fragments cannot be joined (by forming phosphodiester bonds).	1
(vi)	<i>because DNA polymerase III cannot form a new strand in the 3' to 5' direction// because DNA polymerase III only can synthesizes new strand in the direction 5' to 3'.</i>	1
(b)	<u>Stage: initiation</u>	
	i. A small ribosomal subunit binds to an mRNA and moves along the mRNA until it reaches the start codon, AUG	1
	ii. An initiator tRNA with anticodon (UAG) base-pairs with the start codon, AUG	1
	iii. This tRNA carries the amino acid methionine (Met)	1
	iv. The arrival of a large ribosomal subunit completes the initiation complex	1
	v. Proteins called initiation factors are required to bring all the translation components together	1
	vi. Hydrolysis of GTP provides the energy to the assembly	1
	vii. The initiator tRNA is in the P site. The A site is available to the tRNA bearing the next amino acids.	1
Max 6 marks		
5 (a) (i)	Base substitution	1
(ii)	Sickle cell anaemia	1
	• In DNA, base T is replaced with base A // the base sequence in DNA change from CCT to CAT	1
	• Codon in mRNA change from GAA to GUA // Glutamic acid/ Glu is replaced with Valine/ Val	1
Any 1		
(b) (i)	Allopolyploidy	1
(ii)	18 // $n + n = 18$	1
(iii)	Q is sterile because the chromosomes are not homologous/ from different species // Q chromosomes do not pair during meiosis.	1

Question	Answer	Mark														
(c)	<table><tr><th>Klinefelter syndrome</th><th>Turner syndrome</th></tr><tr><td>Aneuploidy condition is Trisomy / $2n+1$</td><td>Aneuploidy condition Monosomy / $2n-1$</td></tr><tr><td>Male individual</td><td>Female individual</td></tr><tr><td>Genotype XXY</td><td>Genotype XO</td></tr><tr><td>Individual has 47 chromosomes/ $44 + XXY$</td><td>Individual has 45 chromosomes/ $44 + XO$</td></tr><tr><td>Underdeveloped testes</td><td>Underdeveloped ovary</td></tr><tr><td>Tall / enlarge breast / feminized voice / knocked knees / long limbs</td><td>Short stature / webbed neck / poorly develop breasts</td></tr></table>	Klinefelter syndrome	Turner syndrome	Aneuploidy condition is Trisomy / $2n+1$	Aneuploidy condition Monosomy / $2n-1$	Male individual	Female individual	Genotype XXY	Genotype XO	Individual has 47 chromosomes/ $44 + XXY$	Individual has 45 chromosomes/ $44 + XO$	Underdeveloped testes	Underdeveloped ovary	Tall / enlarge breast / feminized voice / knocked knees / long limbs	Short stature / webbed neck / poorly develop breasts	<div>1/0</div> <div>1/0</div> <div>1/0</div> <div></div> <div>1/0</div> <div>1/0</div> <div></div> <div>1/0</div>
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Any 5																
6 (a) (i)	Structure Y : Plasmid	1														
	Enzyme : Restriction enzyme/ endonuclease	1														
(ii)	Cut DNA at the same restriction site/specific base sequence	1														
	Produce sticky ends (of the gene of interest and the plasmid vector) that are complementary to each other	1														
(iii)	DNA ligase	1														
(iv)	Able to accept foreign DNA in multiple cloning site/ MCS	1														
	Able to replicate freely in host cell // present of origin of replication	1														
	Has selectable marker gene/ <i>amp^R</i> / <i>amp^R</i> / <i>lacZ</i>	1														
Any two																
(b)	i. mRNA only consist exon/ coding sequence of gene//no introns.	1														
	ii. mRNA is isolated from the target cell	1														
	iii. The reverse transcriptase enzyme catalysed the synthesis of first DNA strand/ single cDNA using mRNA as a template (and short poly-dT as a DNA primer).	1														
	iv. mRNA is degraded by another enzyme/ mRNA degrading enzyme	1														
	v. DNA polymerase is used to catalyse the synthesizes of the second DNA strand/ double stranded cDNA.	1														
	v. Produce cDNA which carries the complete coding sequence of gene /no intron.	1														

Any 5

vi.

Question	Answer	Marks
7 (a)	<u>First trimester</u>	
	i. trophoblast cells of blastocyst secrete human chorionic gonadotropin (hCG).	1
	ii. hCG maintains the corpus luteum.	1
	iii. corpus luteum continue to secrete estrogen and progesterone.	1
	iv. estrogen and progesterone maintain the thickening of endometrium for embryonic development.	1
	v. Progesterone level is high to prevent the miscarriage / maintain the pregnancy.	1
	vi. hCG level increases dramatically during the first trimester.	1
	vii. End of first trimester, placenta takes over the roles of corpus luteum to secrete estrogen and progesterone.	1
	<i>Any 2 points</i>	
	<u>Second trimester</u>	
	i. hCG secretion declines.	1
	ii. Corpus luteum degenerates.	1
	iii. Placenta completely takes over the production of estrogen and progesterone.	1
	iv. Levels of estrogen and progesterone continue to increase to maintain the thickening of endometrium	1
	v. High level of progesterone inhibits contraction of uterine muscle	1
	<i>Any 2 points</i>	
	<u>Third trimester</u>	
	i. Estrogen reach the highest level during last week of pregnancy	1
	ii. This trigger formation of oxytocin receptor on uterus for preparation of birth process	1
	iii. Progesterone level decreases allow the contraction of uterine muscle.	1
	<i>Any 2 points</i>	

Parturition

- | | | |
|------|---|---|
| i. | Final week of pregnancy, oestrogen reaches its highest level, which stimulates the formation of oxytocin receptors in the uterus. | 1 |
| ii. | Progesterone level drops off, which stimulates the uterus contraction that will lead to birth. | 1 |
| iii. | Oxytocin released by the fetus and mother's posterior pituitary gland stimulates powerful contraction on uterus. | 1 |
| iv. | Oxytocin also stimulates placenta to secrete prostaglandins, which further enhance uterine contraction. | 1 |
| v. | The contractions stimulate further release of oxytocin & prostaglandins. | 1 |
| vi. | which in turn stimulates further contractions // contraction become stronger and more frequent. | 1 |
| vii. | Action of oxytocin and prostaglandins form a positive-feedback mechanism. | 1 |

*Any 6 points***Maximum = 12 marks**

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|----------------|---|--|---|
| (b) | i. | Secondary oocyte consists of granulosa cells / follicular cells of corona radiata, zona pellucida, plasma membrane, cytoplasm, cortical granule, nucleus and first polar body. | 1 |
| | ii. | <u>Granulosa cell</u> is secretory cells in Graafian follicle that surrounds the oocyte | 1 |
| | iii. | It supplies nutrients to the developing oocyte | 1 |
| | <i>Any one</i> | | |
| | iv. | <u>Zona pellucida</u> is a layer of glycoprotein that surround plasma membrane of a mammalian egg cell. | 1 |
| | v. | It develops as a jelly coat | 1 |
| <i>Any one</i> | | | |
| vi. | <u>First polar body</u> is a small cell produced in meiosis I during development of oocyte (and finally degenerates). | 1 | |

*Maximum: 4 marks***TOTAL MARK****80**