

School of Health Sciences

WRITTEN EXAMINATION

Course Genetics		
Examination Salster	ntame	en
Course code BM1360	3	Credits for written examination 4 hp
Date 20250609		Examination time 08.15-12:30
Examination respons	ible M a	aria Araceli Diaz
Teachers concerned N	/Iaria	Araceli Diaz, Johan Norden
Aid at the exam/appe	ndices	Calculator
Other		
Instructions		Take a new sheet of paper for each teacher.
mstructions		Take a new sheet of paper when starting a new question.
	\boxtimes	Write only on one side of the paper.
	\boxtimes	Write your name and personal ID No. on all pages you hand in.
	\boxtimes	Use page numbering.
	\boxtimes	Don't use a red pen.
	\boxtimes	Mark answered questions with a cross on the cover sheet.
Grade points		
For D: 50% correct of For C: 50% correct of For B: 50% correct of	n each n each n each	learning objective (6+6+4), 50% of total points, 16p. learning objective, 60% of total points, 19p. learning objective, 70% of total points, 22p. learning objective, 80% of total points, 25p. learning objective, 90% of total points, 28p.

Examination results should be made public within 18 working days

Good luck!

Code:		

Written exam: Genetics BM136G VT25, 4 hp, 20250609

This exam has 3 different parts, and you need to pass all of them to pass the exam (minimum 50% correct per part).

For E: 50% correct on each learning objective (6+6+4), 50% of total points, 16p.

For D: 50% correct on each learning objective, 60% of total points, 19p.

For C: 50% correct on each learning objective, 70% of total points, 22p.

For B: 50% correct on each learning objective, 80% of total points, 25p.

For A: 50% correct on each learning objective, 90% of total points, 28p.

You can answer multiple choice questions directly on this paper. Other questions on separate sheets of paper.

Le pe	earning objective 1: Describe principles for inheritance and edigrees, and make calculations on these (12 p)	JN
1-	What is the most likely mode of inheritance in these pedigrees?	1p
	 A. autosomal recessive B. autosomal dominant C. X-linked recessive D. X-linked dominant 	
2-	What is the most likely mode of inheritance in these pedigrees?	1p

	В. С.	autosomal recessive autosomal dominant X-linked Y-linked	
3-	disorder. T	has a history of cystic fibrosis, an autosomal recessive he parents are both carriers (heterozygous) for the disease. e probability that their child will be affected by cystic fibrosis?	1р
	A.	0%	
		25%	
		50%	
	D.	75%	
4-	Which of t	ne following is true about X-linked recessive disorders?	1p
	A.	They are more common in females	
	B.	They are passed from father to son	
	C.	They are more common in males	
	D.	They cannot be inherited	
5-	marries a	s a carrier for hemophilia, an X-linked recessive disorder, and man who is unaffected. What is the probability that their son ected by hemophilia?	1p
	Α.	0%	
		25%	
		50%	1
		100%	
6-	is caused in life. His family has	47, has just been diagnosed with Huntington's disease, which by a rare dominant allele. The disease is not shown until late daughter, age 25, has a 2-year-old son. No one else in the the disease. What is the probability that the daughter will e disease later in life?	1p
1			

netic	s BM136G ex	am, VT-25		Code:		
	A. B. C. D. E.	0 % 25 % 50 % 75 % 100 %				
7-	chromosor	nes in a chro ne would be Sequence A mix of s Sequence	closely examine the process of all for sequences of all for sequences.	other or from the fa mother and fathe irandparents	scle cells, that	1р
8-	What is the type O child A. B. C. D.	e likelihood od? It is impossik 25% if both p 50% if both p 25% if only t	nans exhibits cod of a type A father ble. parents are heteroz parents are heteroz the father is heteroz the mother is hetero	and a type B mo cygous cygous cygous		1p
9-	allele (G) cayellow-color monohybri and 1/2 are genotypes A. B. C. D.	auses green- ored pods. id cross in w	two alleles for the -colored pods, and You are prese hich approximate ch of the following	I the recessive al nted with the y 1/2 of the proge	lele (g) causes results of a eny are yellow,	1p

10-	A pea plant is heterozygous at the independent loci for flower color (Pp) and seed color (Yy). What types of gametes can it produce?		
	A. B. C. D. E.	two gamete types: pp and PP two gamete types: pY and Py four gamete types: pY , py , PY , and Py four gamete types: pP , Yy , pY , and Py one gamete type: $PpYy$	
11-	has galactos grandmother children, non	rtha are contemplating having children, but John's brother emia (a rare autosomal recessive disease) and Martha's also had galactosemia. Martha has a sister who has three e of whom have galactosemia. What is the probability that rtha's first child will have galactosemia?	1p
	A. B. C. D. E.	0 1/4 1/8 1/12 1/16	
12-	gives blue le You are mak heterozygous	dominant A allele inhibits leaf color. The dominant E allele saves, whereas the recessive e allele gives green leaves. ing a dihybrid cross of the maize plants by crossing two s plants with each other. ution of the different phenotypes will you see in the F1	1p
	A. B. C. D. E.	9 blue; 4 green; 3 no color 9 green; 4 blue; 3 no color 9 no color; 4 blue; 3 green 12 no color; 3 blue; 1 green 12 blue; 3 green; 1 no color	

~ I		
Code:		

lin	arning objective 2: Describe mitosis, meiosis, recombination and kage analysis, their produced effects on the next generation, and ake calculations on these (12 p)	MADC
13-	Which of the following statements correctly describes the difference between the leading and the lagging strands of DNA in DNA replication?	1p
	A. The leading strand is synthesized in the same direction as the movement of the replication fork, and the lagging strand is synthesized in the opposite direction.	9
	B. The leading strand is synthesized by adding nucleotides to the 3' end of the growing strand, and the lagging strand is synthesized by adding nucleotides to the 5' end.	
	C. The lagging strand is synthesized continuously, whereas the leading strand is synthesized in short fragments that are ultimately stitched together.	
	D. The leading strand is synthesized at twice the rate of the lagging strand.	
14-	The mitotic spindle plays a critical role in which of the following processes?	1p
	A. Separation of sister chromatids	
	B. Splitting of the cell (cytokinesis) following mitosis	
	C. Dissolving the nuclear membrane	
	D. Triggering condensation of chromosomes	
15-	Which of the following statements describes a characteristic feature of metaphase?	1p
	A. Separation of sister chromatids	
	B. Cytokinesis	
	C. Alignment of chromosomes on the equator of the cell	
	D. Separation of the centromeres	

Code:		

16-	Which of the following chromosome number		of a species that has a	1р
	B. The specie C. Each diplo	es is diploid and has 32 es has 16 sets of chromed d cell has eight homolo from this species has fo	osomes per cell. gous pairs of chromosomes.	
17-	mitosis? A. Chromoso B. Synapsis C. Alignment	g processes occurs in ome replication of chromosomes of chromosomes at the tion of chromosomes		1р
18-			to ab/ab, what percentage of are 10 map units apart?	1p
19-	chromosome as a heterozygous for k	a locus with alleles	nt is located on the same H and h. An organism ed with an organism with iles are produced:	2p
		Genotype	Number of progeny	
		GH/gh	12	
		Gh/gh	80	
		gH/gh	85	1
1				1
		gh/gh	20	
		notype of the heterozyg	Table 1	
	b) What is the ma	notype of the heterozyg	ous parent? (1p)	

Code:		

20-	Which of the following observations would most strongly suggest that two genes are linked?	1p
	 A. The genes exhibit independent assortment in a dihybrid cross. B. The recombination frequency between the genes is approximately 50%. C. The recombination frequency between the genes is significantly less than 50%. D. All possible phenotypic combinations appear with equal frequency in a testcross. 	
21-	Two different pure-breeding lines of tomatoes were crossed. The F1 progeny were phenotypically wild-type and heterozygous for three recessive traits: dwarf (d), yellow fruit (y), and narrow leaves (n). A testcross was performed by crossing the F1 (DdYyNn) with a homozygous triple-recessive plant (ddyynn). The resulting 10,000 progeny is shown below:	3p
	Phenotype Count DyN 3050 dYn 2980 dYN 970 DYn 1010 Dyn 700 dYN 690 DYN 80 dyn 60 Total 10,000	
	 a) What were the genotypes for the parental true-breeding lines? (1p) b) Determine the correct gene order. (1p) c) Draw a linkage map for the three genes, including map distances (in map units). (1p) 	

	earning objective 3: Explain the main concept as well as the use of mpler models in population genetics (8 p)	MADC
22-	If the frequency of A allele in a population is 0,2, the frequency of the homozygote genotype aa is:	1р
	A. 0.32	
	B. 0.16	
	C. 0.64	
	D. 0.8	

Code:		

23-	Allele frequencies for the ABO blood type among North Americans ar estimated at f(IA) = 0.3; f(IB) = 0.02; f(i) = 0.68. Assuming Hardy Weinberg equilibrium conditions for this locus, what is the expecte frequency of blood type AB?				
	A.	0.016			
	B.	0.012			
	C.	1.0			
	D.	0.006			
24-	Briefly	/ explain:	3р		
	a)	the conditions for assuming a Hardy-Weinberg Equilibrium (1p)			
	b)	Genetic variation and sources of genetic variation (1p)			
	c)	Natural selection (1p)			
25-	Given 150 G	a population of wildflowers in Oregon: 500 G1G1 (Purple flowers), 2G2 (Yellow flowers) and 50 G1G2 (Lavender flowers)	2p		
	Answer the following:				
	a)	Calculate the genotype frequencies for purple, yellow, and lavender flowers and the number of allele copies of G1 and G2. (1p)			
	b)	Using the allele frequencies, calculate the expected genotype frequencies under Hardy-Weinberg equilibrium and the expected number of individuals for each genotype. (1p)			
	c)	Is the population in Hardy-Weinberg equilibrium? Justify your answer with calculations. (1p)			
	d)	Pollen from a nearby plantation introduced 120 new yellow-flowered plants (genotype G2G2) into the population. What evolutionary mechanism does this represent, and how might it affect the population? (1p)			