		GE	NETIC				
1	Course Title:	GENETI	C				
2	Course Code:	VET101	9				
3	Type of Course:	Compuls	sory				
4	Level of Course:	First Cycle					
5	Year of Study:	1					
6	Semester:	1					
7	ECTS Credits Allocated:	2.00					
8	Theoretical (hour/week):	2.00					
9	Practice (hour/week):	0.00					
10	Laboratory (hour/week):	0					
11	Prerequisites:	None					
12	Language:	English					
13	Mode of Delivery:	Face to t	face				
14	Course Coordinator:	Doç.Dr.	ÖZDEN ÇOBANOĞLU				
15	Course Lecturers:		Sena ARDIÇLI ör. Dr. Deniz DİNÇEL				
16	Contact information of the Course Coordinator:	Doç. Dr. Özden ÇOBANOĞLU e-mail: ocobanoglu@uludag.edu.tr U.Ü. Veteriner Fakültesi Genetik Anabilim Dalı Nilüfer/BURSA					
17	Website:						
18	Objective of the Course:	This course covers principles of prokaryotic and eukaryotic genetics. In this course, students will expand on the basic knowledge of genetics. This will involve learning new terminology and new core concepts about the principle of genetics which will be the basis for the other classes during their education. They will able to apply the general concept of genetics to veterinary science. The molecular basis of heredity, chromosome structure, patterns of Mendelian and non-Mendelian inheritance, and biotechnological applications will be covered in this course. Thus, the course provides the students with a review of analytical, molecular and cellular genetics along with new developments. Upon successful completion of this course, students should be able to recognize and describe genetic phenomena and demonstrate knowledge of important genetic principles.					
19	Contribution of the Course to Professional Development:						
20	Learning Outcomes:						
		1	History of Genetics, Major Events and Milestones in Genetics, Understand the principles of inheritance as formulated by Mendel; Mode of Inheritance.				
		2	Apply the principles of extensions to Mendelian inheritance, including codominance, gene interactions, epistasis, multiple alleles, pleiotropy, lethal alleles, penetrance and sex-linked transmission.				
		3	Learn about cell division mechanisms in prokaryotic and eukaryotic organisms. Analyze basic genetic data using statistical procedures.				
		4	Understand and relate the structure and function of the DNA and RNA molecules, realize their functional roles in encoding genetic material and obtain knowledge about gene expression.				

		5	Able to describe the bainformation from DNA					
		6	Distinguish the chromosomal number among different species and gain a cause and an effect of changes in chromosome number and structure. Learn how to identify and classify DNA mutations.					
		7	Understand gene transfer mechanisms in prokaryotic organisms and learn how to apply recombinant DNA technology to animal genomes theoretically.					
		8	Learn about gene regulation with emphasis on repressible vs. inducible operon systems.					
		9	Get information about sequencing methodolo		eration DNA			
		10	Obtain information abo apply these technique					
21	Course Content:							
		Co	urse Content:					
Week	Theoretical		Practice					
1	Introduction to the Course and Milest Genetics; Mendelian Genetics: The chromosomal basis of inheritanc Mendel's principles of segregation, a independent assortment, monohybric dihybrid and trihybrid crosses.	e, nd I,						
2	Variations on Mendelian Inheritance	l:						
Activit	es		Number	Duration (hour) Total Work Load (hour)			
Th g ore	Exceptions on Mendelian Genetics II	:	14	2.00	28.00			
Practica	als/Labs		0	0.00	0.00			
Self stu	dyuttipdepatiperatipelygenic inheritance	Э,	10	1.00	10.00			
Homew	vorks		0	0.00	0.00			
Project	heterogeneity.		0	0.00	0.00			
Field St			0	0.00	0.00			
Midtern	life Kage, X chromosome Inactivation,	dosage	1	10.00	10.00			
Others			0	0.00	0.00			
Final E	GAR Pacteristics and pedigree analysis	5. 5.	1	12.00	12.00			
	/ork Load				60.00			
Total w	Linkage and recombination, crossing ork load/ 30 hr. chromosome theory, a genetic map of	over, of the			2.00			
ECTS (Credit of the Course Joinybrid and trinybrid cross by recom	nination			2.00			
	frequencies between genes, interfere coefficient of coincidence.							
6	Identifying DNA and RNA as the Ger Material: Search for genetic material; the disco DNA by Griffith's Transformation Exp Avery, MacLeod and McCarty's expe Hershey-Chase bacteriophage exper and a discovery of RNA by Tobacco Virus (TMV) experiment.	overy of periment, priments, riment,						

7	The Structure and Analysis of DNA and RNA: Structure of nucleic acid, properties of pyrimidines and purines, the anatomy of DNA, a discovery of the structure of DNA, the DNA double helix as Watson and Crick model, polymorphism of DNA, structural features of DNA and a structure of RNA. DNA Packing in Prokaryotic and Eukaryotic Chromosomes:	
	DNA condensation, DNA supercoiling, nucleosomes, eukaryotic chromosomal organization, a structure of chromatin, chromosome folding, DNA packing. Gene Expression and Regulation: Repressible vs. inducible operon systems; Lac Operon and Tryptophan Operons in E. coli.	
9	DNA Replication in Prokaryotes and Eukaryotes: Models for DNA replication, Meselson-Stahl experiment, a mechanism of DNA replication in prokaryotes, replication of DNA in eukaryotes, enzymes required for replication, directionality of synthesis in DNA strands, DNA repair system, editing, and proofreading of DNA.	
10	The Central Dogma; Transcription, Translation and Protein Synthesis: Defining central dogma of molecular biology, transcription, RNA processing, genetic code, wobble base pairing, translation, protein synthesis, the structure of amino acid, principles of polarity in amino acid.	
11	The Genetic Mutations: Cause of mutation, types of mutations; spontaneous mutations, single base substitution and frameshift mutations, chromosomal disorders, nondisjunction in autosomal chromosomes, trisomies, nondisjunction of X chromosomes and induced mutations Genetic Transfer in Bacteria: Transformation, transduction, and conjugation, plasmid structure in bacteria.	
12	Recombinant DNA Technology: Type of vectors, techniques of recombinant DNA technology; electroporation, protoplast fusion, and injection: gene gun and microinjection.	
13	DNA Sequencing Techniques: Basic methods for sequencing; Maxam- Gilbert and Sanger methods, Whole genome sequencing and New DNA sequencing methods	
14	Basic Molecular Techniques: Polymerase Chain Reaction (PCR) and Its Steps and Application; Gel Electrophoresis System, Restriction Edonuclease; RFLP, AFLP, RAPD, Microsatellite and SNP Marker Analyzes, Microarray System and Marker Assisted Selection and Use of markers in Livestock.	

22	2 Textbooks, References and/or Other Materials:						1. Veteriner Genetik, Odabaşioglu F. İkinci Basim. Lazer Ofset MatbaaTesisleri San.Tic. Ltd. Şti. Ankara, 2012.									
							and	2. Principles of Genetics. Sunstad D.P., Simmons M.J., and Jenkins J.B. John Wiley and Sons Inc. New York, USA, 1997.								
								3. An Introduction to Genetic Analysis. Griffiths A.J.F., Miller J.H., Suzuki D.T., Lewontin R.C., Gelbart W.M. 5th Edition. W. H. Freeman and Company. New York, USA, 1993.								
							4. Genetik. Yildirim A., Karadag Y., Kandemir N., Sakin M.A. 2. Baski. Nobel Yayin Dagitim, Ankara, 2010.									
								5. Genetic Class Notes. Cobanoglu O. Bursa Uludag Univ., Faculty Veterinary-Medicine. Bursa, 2017.								
23	Assesm	ent														
								EWE	EIGHT							
Midterm	n Exam					1		30.	.00							
Quiz						1		10.	.00							
Home v	vork-pro	ject				C)	0.0	0							
Final Ex	xam					1		60.	.00							
Total						3	3	10	0.00							
	ution of s Grade		Year)	Learn	ing Act	tivities	s to	40.	40.00							
Contrib	ution of	Final E	xam to	o Suc	cess G	rade		60.	60.00							
Total								10	100.00							
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24	ECTS	/ WO	RK L	OAD) TAB	LE										
25			CON	TRIE	BUTIC	N O			-	OUTC ATIO	-	S TO I	PROG	GRAMI	ME	
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ÖK2	5	3	1	3	5	5	2	2 2	3 2	•	2 4	4 5	0 0	0	0	0
ÖK2 ÖK3	5	3 3	1	3 2	5 5					2			-	_		
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ÖK3	5	3	1	2	5	5 5	2	2 3	2	2 2 1	4	5 5	0	0	0	0
ÖK3 ÖK4 ÖK5 ÖK6	5 5 5 5 5	3 3 3 3	1 1 1 1	2 2 1 3	5 5 5 5	5 5 5 5 5	2 2 2 3 2	2 3 3 2 2	2 4 3 2	2 2 1 1	4 4 2 4	5 5 4 5	0 0 0	0 0 0 0 0	0 0 0 0 0	0 0 0 0 0
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ÖK3 ÖK4 ÖK5 ÖK6 ÖK7 ÖK8	5 5 5 5 5 5 5 5	3 3 3 3 3 3 3	1 1 1 1 1 1	2 2 1 3 2 2	5 5 5 5 5 5	5 5 5 5 5 5 5	2 2 2 3 2 2 2 2	2 3 2 2 3 3	2 4 3 2 4 4	2 2 1 1 2 2 2 1 1	4 4 2 4 4 4	5 5 4 5 5 5 5	0 0 0 0 0 0 0	0 0 0 0 0 0 0	0 0 0 0 0 0	0 0 0 0 0 0
ÖK3 ÖK4 ÖK5 ÖK6 ÖK7	5 5 5 5 5 5	3 3 3 3 3 3	1 1 1 1 1	2 2 1 3 2	5 5 5 5 5	5 5 5 5 5 5	2 2 2 3 2 2 2	2 3 3 2 2 3	2 4 3 2 4	2 2 1 1 2 2 2 1	4 4 2 4 4	5 5 5 4 5 5	0 0 0 0 0 0	0 0 0 0 0 0	0 0 0 0 0 0	0 0 0 0 0 0

LO: Learning Objectives PQ: Program Qualifications										
Contrib ution Level:	1 very low	2 low	3 Medium	4 High	5 Very High					