GENETIC					
1	Course Title:	GENETI	С		
2	Course Code:	VET1019			
3	Type of Course:	Compulsory			
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 f	face		
14	Course Coordinator:	Doç.Dr.	ÖZDEN ÇOBANOĞLU		
15	Course Lecturers:	Doç. Dr. Özden ÇOBANOĞLU Doç. Dr. Sena ARDIÇLI Araş. Gör. Dr. Deniz DİNÇEL			
16	Contact information of the Course Coordinator:	Doç. Dr. Özden ÇOBANOĞLU U.Ü. Veteriner Fakültesi Genetik Anabilim Dalı Nilüfer/BURSA E-mail:ocobanoglu@uludag.edu.tr Tel: 0 224 294 1241			
17	Website:	http://www.veteriner.uludag.edu.tr			
	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:	The course will provide the students with the necessary knowledge about the basic concepts related to the genetic structure of biological creatures during their further education.			
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.		
	Apply the principles of extensions to Mendelian inheritance, including codominance, gene interactions epistasis, multiple alleles, pleiotropy, lethal alleles, penetrance and sex-linked transmission.				
	Learn about cell division mechanisms in prokaryotic and eukaryotic organisms. Analyze basic genetic data using statistical procedures.				

			D er	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 basic aspects of the flow of genetic information from DNA to proteins by central dogma.			
6			sp ch	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.			
			10	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 basic and third generation DNA sequencing methodologies.			
		10		Obtain information about genetic markers and how to apply these techniques to animal breeding.			
21	Course Content:		,	1,7	<u> </u>		
		Co	ur	se Content:			
Week	Theoretical		Р	ractice			
1							
Activit	es			Number	Duration (hour)	Total Work Load (hour)	
Theore	ঞ্জিন্দ examples of non-Mendelian inhe	ritance;		14	2.00	28.00	
Practica	als/Labs		_	0	0.00	0.00	
Self stu	exceptions of the hotelian Genetics II	:	F	10	1.00	10.00	
Homew	vorks			0	0.00	0.00	
Project	multiple alleles, polygenic inheritance	510110py, − 9 ,		0	0.00	0.00	
Field S	tudies			0	0.00	0.00	
Midtern	hexerogeneity.			1	10.00	10.00	
Others				0	0.00	0.00	
Final E	Sex determination in different species linkage. X chromosome Inactivation.	s, sex dosage		1	12.00	12.00	
Total W	Vork Load					60.00	
Total w	genetic maternal effect, sex-influence ork load, 30 hr Icharacteristics and pedigree analysis	au 3.				2.00	
ECTS (Credit of the Course					2.00	
	Linkage and recombination, crossing chromosome theory, a genetic map or Drosophila melanogaster, linkage madihybrid and trihybrid cross by recomfrequencies between genes, interfere coefficient of coincidence.	of the apping in abination					
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, eriments, 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.	
8	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.	
12	Recombinant DNA Technology: Genetic Transfer in Bacteria: Transformation, transduction, and conjugation, plasmid structure in bacteria. 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			 Veteriner Genetik, Odabaşioglu F. İkinci Basim. Lazer Ofset MatbaaTesisleri San.Tic. Ltd. Şti. Ankara, 2012. Principles of Genetics. Sunstad D.P., Simmons M.J., and Jenkins J.B. John Wiley and Sons Inc. New York, USA, 1997. 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. Genetik. Yildirim A., Karadag Y., Kandemir N., Sakin M.A. 2. Baski. Nobel Yayin Dagitim, Ankara, 2010. Genetic Class Notes. Cobanoglu O. Bursa Uludag Univ., Faculty Veterinary-Medicine. Bursa, 2017. 		
23	Assesment				
TERM L	TERM LEARNING ACTIVITIES NUMBE R		WEIGHT		
Midterr	n Exam	1	30.00		
Quiz		1	10.00		
Home v	vork-project	0	0.00		
Final E	xam	1	60.00		
Total		3	100.00		
Contribution of Term (Year) Learning Activities to Success Grade		es to	40.00		
Contribution of Final Exam to Success Grade)	60.00		
Total			100.00		
1			The exams of the course will be in the form of multiple choice tests.		
24	ECTS / WORK LOAD TABLE				

CONTRIBUTION OF LEARNING OUTCOMES TO PROGRAMME **QUALIFICATIONS** PQ1 PQ2 PQ3 PQ4 PQ5 PQ6 PQ7 PQ8 PQ9 PQ1 PQ11 PQ12 PQ1 PQ14 PQ15 PQ16 ÖK1 ÖK2 ÖK3 ÖK4 ÖK5 ÖK6 ÖK7 ÖK8 ÖK9 ÖK10

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