| Faculty of Medicine University of Niš | Study program: INTEGRATED ACADEMIC STUDIES C MEDICINE ACCREDITATION 2018 | DF | |
|--|--|--|--|
| Course title: Molecular and human genetics | | | |
| Course head: Prof. dr Stevo Najman | | | |
| Course status: | required | required | |
| Semester : | Year of study: | Year of study: | |
| ECTS: 7 | Course code: M – I 1 | | |
| Course purpose: | | | |
| Acquisition of knowledge of theoretical principles and practical skills in Molecular and Human Genetics | | | |
| Course outcome: (knowledge | e, skills, attitudes) | | |
| good command of mic good command of PC s to describe and explain to understand the relat to understand the fund to understand and expl with phenotypic chang to apply the acquired k | skills, as a learning resource the transfer of information from DNA mole ionships of cytoplasmic and nuclear process lamental laws of heredity and to apply the k lain the mechanisms of genetic diversity and es. mowledge in Molecular and Human Genetic | ses nowledge in human situation. d to be able to associate genetic | |
| Nr. of classes of active teach | | | |
| Lectures: 45 | Practice: 24 | OFT: 6 | |
| Course content | | | |
| <u>Theory</u> Cell: Procarvotic and eucarve | otic organization. Genetic characteristics o | of viruses Basic characteristics of | |

Cell: Procaryotic and eucaryotic organization. Genetic characteristics of viruses. Basic characteristics hereditary material. Structure & function of DNA. Human genome. Genetic basis of reproduction: chromatin and chromosome. Cell cycle and mitosis. Meiosis. Gametogenesis. Cytogenetics: human karyotype. Chromosomal abberations. Numerical chromosomal abberations (polyploidy, aneuploidy, mechanisms of occurrence of chromosomal numerical abberations). Structural chromosomal abberations (causes, mechanisms of occurrence, classification. Deletions. Duplications. Isochromosomes. Inversions. translocations). Classical genetics: Mendelian laws of heredity. Autosomal dominant and recessive inheritance. Correlated inheritance. Inheritance of sex and sex-linked traits. Intra- and interlocus gene interactions. Polygenic inheritance. Molecular biology: transfers of hereditary informations in the cell. Genetic code. Transcription. Posttranscriptional RNA processing. Types & role of RNA. Translation. Regulation of gene expression in procaryotes. Mechanisms of regulation of gene expression in eucaryotes. Mutations. Mutagenic factors. Genetic recombinations. Crossing-over - mechanisms and significance. DNA molecule repair. Hereditary disorders caused by DNA repair errors. Human genetics: monogenic and polygenic human traits. Monogenic diseases. Multifactorial hereditary diseases. Gene families. Globin gene family. Immunogenetics. Oncogenetics (cell cycle control, protooncogenes, tumor-suppressor genes, genetic basis of malignant transformation, cancerogenesis). Chromosomopathies of autosomes and sex chromosomes. Genetics of development (molecular-genetic mechanisms in embryogenesis, human sex determination), genetic basis of sex differentiation and related disorders). Genetics of human populations (gene polymorphisms, population-genetic research in medicine). Methods in human genetics. Techniques and methods of DNA analysis. Use of DNA technologies and modern biotechnologies in medicine.

<u>Practice</u>

Use of ITs in biomedicine. Microscope and microscopy. Nucleus. Chromatin. Chromosome. Karyotype. Mitosis. Gametogenesis. Chromosomal abberations. Seminar in cytogenetics. Basic principles of heredity (tasks): mono-, di-, and test of crossbreeding, multiple alleles, polygenic inheritance, correlated inheritance, preparation of chromosomal maps, sex-linked inheritance, genealogic trees.lsolation of DNA (demonstration). Transfer of hereditary information in the cell (tasks). Regulation of gene expression, Mutations, Genetic recombinations (tasks). Seminar in molecular biology. Genetics of human populations (tasks). Technology of rDNA (tasks). Seminar in human genetics.

<u>Seminars</u>

Seminar in cytogenetics. Seminar in molecular biology. Seminar in human genetics.

Recommended literature:

| Recommended literature: | | |
|--|--|--|
| 1. Stanković Ž, Živanov-Čurlis J, Najman S: Biologija sa humanom genetikom (Osnovi citologije, | | |
| genetike i embriologije), Kodeks, Niš 2001. | | |
| 2. Najman S: Osnovi molekularne i humane genetike. Skripta. Savez studenata Medicinskog fakulteta u | | |
| Nišu, Niš 2002. | | |
| 3. Turnpenny P, Ellard S. Emerijevi osnovi medicinske genetike. Beograd. Datastatus; 2009. | | |
| 4. Matić G, Savić Pavićević D. Molekularna biologija 1. NNK Internatinal, Beograd, 2011. | | |
| 5. Najman S, Živanov-Čurlis J, Vukelić-Nikolić M. Molekularna i humana genetika – Praktikum sa | | |
| radnom sveskom za studente medicine i stomatologije. Medicinski fakultet u Nišu, Niš, 2017. | | |
| 6. Lecture handouts | | |
| Teaching methods: | | |
| Theory: lectures | | |
| Practice: microscopy, calculations and related tasks, interactive teaching (PCs) with practice log | | |
| notebooks. | | |
| seminars | | |
| consultations | | |
| Required previous passed exams: | | |
| None | | |
| Grade: (max. 100) | | |
| Preexam obligations | | |
| Credit: 0 – 10 points | | |
| Test: 0 – 20 points | | |
| Practice: 0 – 10 points | | |
| Final exam | | |

Final exam

• Written/oral exam: 0 – 60 points