



UNIVERSITÀ
DEGLI STUDI DI MILANO-BICOCCA

SYLLABUS DEL CORSO

Fundamentals of Cell Biology and Genetics

2627-1-H4104D002

Aims

The course will provide the essential theoretical knowledge of biology and genetics, also focusing on the possible future application in the medical field. The subjects of the course will provide the necessary knowledge to understand the vital processes, both on the cellular and molecular level, as well as the laws of heredity and the processes involved in the generation of phenotypic diversity. The acquired knowledge will contribute to better understand the processes involved in normal and pathological situations.

Knowledge and understanding - at the end of the course the student will be able to use the provided information to understand the molecular mechanisms of cell biology and the basic mechanisms of disease genetics.

Ability to apply knowledge and understanding - at the end of the course the student must be able to apply the acquired knowledge to understand the molecular basis of human pathology and inheritance.

Independent judgment - at the end of the course, the student will be able to put together information from different fields (biology, genetics) to understand and interpret the mechanisms of human disease.

Communication skills - at the end of the course the student will have acquired adequate scientific terminology and will be able to explain the topics covered in the course with correct language.

Learning ability - at the end of the course the student will be able to understand and critically evaluate the scientific literature regarding the genetics and biology of human pathologies.

Contents

Structure and function of the most important cellular macromolecules; DNA duplication and repair mechanisms; transcription and RNA processing; translation and protein sorting; molecular and cellular mechanisms responsible for gene expression and its regulation, analyzing epigenetic mechanisms, transcriptional and post-transcriptional

regulation; signal transduction pathways; molecular and cellular mechanisms which control the cell cycle, cellular growth and differentiation as well as cell-to-cell interactions; basic concepts of heredity and the transmission patterns of inherited traits; mechanisms which can generate phenotypic variants in men; methodologies used in genetic analysis; most important biotechnological applications in medicine (gene-based and cell-based therapy).

Detailed program

GENERAL BIOLOGY – Classification of living organisms – Structure of prokaryotic and eukaryotic cells – Viruses, classification, lytic and lysogenic cycle.

MOLECULAR BIOLOGY. Chemical composition and molecular organization of the cell – water, carbohydrates, lipids, proteins and nucleic acids. Identification of the chemical compound carrying the genetic information – Molecular basis of inheritance – DNA replication. Telomerases – Mechanisms of DNA repair. Correlation with human diseases, aging and cancer. - RNA, structure and function – Transcription and RNA maturation – The genetic code, and its biological implication (redundancy, frameshift) – Translation – Protein sorting – Gene expression regulation in prokaryotes and eukaryotes – Molecular genetic tools (restriction enzymes, vectors, Southern/Northern/Western blotting, PCR, Sanger sequencing and Next Generation Sequencing, microarrays). Molecular cloning.

CELL BIOLOGY – Structure and function of the cytoskeleton – Cell adhesion mechanisms – Endocytosis and Exocytosis – Cell-to-cell communication in complex organisms – Signal transduction and the role of protein kinases – Cell cycle and its regulatory mechanisms - Mitosis and Meiosis – Apoptosis – Cell differentiation processes: embryonal and adult stem cells - Tumors.

GENETICS – Human reproduction – Genetic variability – Inheritance – Genes: genotype and phenotype – Diploidy and reproduction. Homologous chromosomes, alleles and loci, homozygosity and heterozygosity – Mendel's laws – Alleles: wild-type, mutated and multiple ones, dominance and recessivity – Mendel's law's implementation: epistasis, penetrance and expressivity – Sex chromosomes. Sex determination – How to build and analyze a family tree – Chromosome X inactivation. Its implication in the phenotypic manifestations of genetic diseases – Test cross and inheritance of genes localized on different chromosomes – Crossing over and its genetic consequences – Recombination frequencies calculation and genetic maps – Principles and consequences of mitochondrial inheritance and genomic imprinting – Examples of monogenic inheritance: blood groups (ABO, Rh), color blindness – Multigenic inheritance and quantitative genetics – Characters showing a threshold effect – Multifactorial disorders – Population genetics and Hardy-Weinberg equilibrium.

CYTOGENETICS – Methods for chromosome analysis – Normal human karyotype – Chromosomal and genomic mutations and their effect on meiosis and phenotype – Deletions, inversions, duplications, translocations and non-disjunctions – Down's, Turner's and Klinefelter's syndrome – Chromosomal mutations and leukemia: Philadelphia chromosome and Burkitt's lymphoma – Germinal and somatic mutations, mosaicism.

MOLECULAR GENETICS: Relationship between DNA content and organism complexity – DNA assembly in the nucleus of eukaryotic cells – Structural differences between prokaryotic and eukaryotic genes – Genome organization in prokaryotic and eukaryotic cells. Characteristics of human genome – Gene mutations: development mechanisms – Mutation consequences on gene products – Examples of autosomal dominant and recessive mutations, as well as X-linked ones – Mitochondrial gene mutations – Genomic instability - DNA polymorphisms and their use as genetic markers – Elements of developmental biology – Immunogenetics. Generation of antibody diversity - Cancer genetics. Oncogenes and tumor suppressor genes (Rb1, WT1 and p53) – Strategies for the diagnosis of genetic diseases (direct and indirect) – The human genome project: future implications – Gene therapy: general concepts and applications.

Prerequisites

Basic sciences (chemistry, physics)

Teaching form

54 two-hour classes delivered in mixed modality: in presence (at least 70%) and recorded (up to 30%).

3 two-hour classes in presence, with an initial part delivered by the professor aiming to generate an interactive work by the students.

4 six-hour interactive practical training sessions delivered in presence.

Textbook and teaching resource

Alberts and Johnson. Molecular biology of the cell. Seventh edition. Garland Science, 2022; H. Lodish, A. Berk, S.L. Zipursky, P. Matsudaira, D. Baltimore, J. Darnell. Molecular cell biology, Freeman, 2021; Thompson & Thompson Genetics and Genomics in Medicine, 9e, 2023; Peter Russell iGenetics: A Molecular Approach. Pearson, 2016

Semester

first and second semester

Assessment method

The final assessment will be an oral exam. It will consist of an interview on the topics covered in class and in the exam texts and will allow for an in-depth evaluation of the acquired knowledge, as well as the critical ability and the ability to place individual topics within a broader context (for example, by evaluating possible interactions).

Office hours

by appointment

Sustainable Development Goals

GOOD HEALTH AND WELL-BEING | QUALITY EDUCATION | GENDER EQUALITY
