| | | Teaching | Guide | | |
|---------------------|---|--------------------|---------------------|------------------------|---------------------------------|
| | Identifying E | Data | | | 2017/18 |
| Subject (*) | Molecular Genetics | Molecular Genetics | | Code | 610G02020 |
| Study programme | Grao en Bioloxía | | | | |
| | | Descrip | otors | | |
| Cycle | Period | Yea | ır | Туре | Credits |
| Graduate | 1st four-month period | Thir | d. | Obligatoria | 6 |
| Language | Galician | | | | |
| Teaching method | Face-to-face | | | | |
| Prerequisites | | | | | |
| Department | Bioloxía | | | | |
| Coordinador | Insua Pombo, Ana Maria | | E-mail | ana.insua@udo | .es |
| Lecturers | Insua Pombo, Ana Maria | | E-mail | ana.insua@udo | .es |
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| Web | | | | 1 | |
| General description | This course focuses on the conceptual and methodological bases necessary to understand the organization, expression, | | | | |
| | variation and manipulation of genetic material. It provides a molecular perspective to knowledge in "Genetics" (secon | | | | wledge in "Genetics" (second |
| | year) and knowledge necessary to a | ddress "Popu | ulation Genetics ar | nd Evolution", "Cytoge | enetics" and related courses of |
| | third and fourth year. | | | | |

| | Study programme competences | | |
|------|--|--|--|
| Code | Study programme competences | | |
| A5 | Analizar e caracterizar mostras de orixe humana. | | |
| A11 | Identificar e analizar material de orixe biolóxica e as súas anomalías. | | |
| A12 | Manipular material xenético, realizar análises xenéticas e levar a cabo asesoramento xenético. | | |
| A15 | Deseñar e aplicar procesos biotecnológicos. | | |
| A29 | Impartir coñecementos de Bioloxía. | | |
| A30 | Manexar adecuadamente instrumentación científica. | | |
| A31 | Desenvolverse con seguridade nun laboratorio. | | |
| B1 | Aprender a aprender. | | |
| B2 | Resolver problemas de forma efectiva. | | |
| В3 | Aplicar un pensamento crítico, lóxico e creativo. | | |
| B5 | Traballar en colaboración. | | |
| В7 | Comunicarse de maneira efectiva nunha contorna de traballo. | | |

| Learning outcomes | | | | |
|---|-----|-----------------|----|--|
| Learning outcomes | | Study programme | | |
| | cor | mpetend | es | |
| General knowledge and understanding of the molecular basis of the organization, expression, variation and manipulation of | A11 | B1 | | |
| genetic material | A12 | B2 | | |
| | A15 | В3 | | |
| | A29 | B5 | | |
| | | В7 | | |

| Knowledge of the basic methodologies used in Molecular Genetics. | A5 | B1 | |
|--|-----|----|--|
| | A11 | B2 | |
| | A12 | В3 | |
| | A15 | B5 | |
| | A29 | | |
| | A30 | | |
| | A31 | | |
| Ability to use sources of information of interest in Molecular Genetics. | A5 | B1 | |
| | A11 | B2 | |
| | A12 | В3 | |
| | A15 | | |
| | A29 | | |
| Ability to interpret and transmit information of Molecular Genetics | A29 | B1 | |
| | | B2 | |
| | | В3 | |
| | | B5 | |
| | | В7 | |

| | Contents |
|-----------------------------------|--|
| Topic | Sub-topic |
| 1 GENOME ORGANIZATION | Genome size. Prokaryotic and eukaryotic genomes. Single-copy and repetitive DNA |
| | sequences. Gene families. Centromeres. Telomeres. Organelle genomes. |
| 2 DNA REPLICATION | Semiconservative DNA replication: the Meselson and Stahl experiment. Modes of |
| | replication. Enzymology of the replication. DNA replication in Escherichia coli. DNA |
| | replication in eukaryotes. Telomere synthesis. Replication of mitochondrial and |
| | chloroplast DNA. |
| 3 SYNTHESIS AND PROCESSING OF RNA | Classes of RNA. RNA polymerases. Promoters and transcriptional apparatus. |
| | Transcription in prokaryotes and eukaryotes: initiation, elongation and termination. |
| | Interrupted genes: exons and introns. Processing of eukaryotic pre-mRNA. Synthesis |
| | and processing of pre-rRNA. Synthesis and processing of pre-tRNA. RNA edition. |
| | Revision of gene concept. |
| 4 TRANSLATION | The one gene-one enzyme hypothesis. The genetic code: characteristics and |
| | experiments to decipher the code. Initiation of translation. Elongation of the |
| | polypeptide chain. Termination of translation. Messenger RNA surveillance. |
| 5 MUTATION AND DNA REPAIR | Molecular basis of spontaneous mutations: replication errors, unequal crossing over, |
| | spontaneous chemical changes. Molecular basis of induced mutations: chemical and |
| | physical agents. Repair mechanisms: direct reversal of damaged DNA, excision |
| | repair, mismatch repair, repair of double-strand breaks, translesion synthesis. |
| 6 MOLECULAR MECHANISM OF GENETIC | The role of genetic recombination. Gene conversion. Models of homologous |
| RECOMBINATION | recombination: Holliday model and double-strand break model. Enzymes required for |
| | recombination. Site-specific recombination. Immunoglobulin gene rearrangements. |
| 7 TRANSPOSABLE GENETIC ELEMENTS | Transposable elements in prokaryotes: insertion sequences, composite transposons |
| | and noncomposite transposons. Replicative and non replicative transposition. |
| | Transposable elements in eukaryotes: transposons and retrotransposon. Evolutionary |
| | significance of transposable elements. |
| 8 RECOMBINANT DNA TECHNOLOGY | Restriction enzymes. Cloning vectors. DNA libraries: construction and screening. |
| | Southern and northern blotting. PCR. Restriction maps. DNA sequencing. |
| | Site-directed mutagenesis. |

| 9 APPLICATIONS OF RECOMBINANT DNA | Expression of eukaryotic genes in E. coli. DNA transfer to eukaryotic cells. Transgenic |
|--|---|
| TECHNOLOGY | animals. Transgenic plants. Gene therapy. Molecular markers. DNA fingerprinting. |
| | Genetic diagnosis. Synthetic genomes. Genome editing: CRISPR/Cas9 technology. |
| 10 GENOMICS | Physical and genetic mapping. Whole genome sequencing. Genome annotation. DNA |
| | microarrays. Reverse Genetics. Comparative genomics. Metagenomics. |
| 11 REGULATION OF GENE EXPRESSION IN BACTERIA | Jacob and Monod?s operon model for the regulation of lac genes in E. coli. Positive |
| | control of the lac operon. The arabinose operon of E. coli: positive and negative |
| | control. The triptophan operon of E. coli: negative control and attenuation. |
| | RNA-mediated regulation. |
| 12 REGULATION OF GENE EXPRESSION IN | Changes in chromatin structure. DNA methylation. Transcriptional control. |
| EUKARYOTES | RNA processing control. Control of mRNA stability. Control at the level of translation. |
| | RNA interference. Epigenetics. |
| 13 GENETIC CONTROL OF DEVELOPMENT | Basic events of development. Drosophila development stages. Maternal-effect, |
| | segmentation and homeotic genes in Drososphila. Homeobox genes in other |
| | organisms. General aspects of Caenorhabditis elegans development. Genetic control |
| | of flower development in Arabidopsis. |
| PRACTICE 1: DNA EXTRACTION | Genomic DNA extraction. Agarose gel electrophoresis for DNA. DNA quantification. |
| PRACTICE 2: PCR | PCR amplification of the CHD gene. Analysis of an intron polymorphism for bird |
| | sexing. |
| PRACTICE 3: DOT-BLOT | Nucleic acids hybridization: detection of microsatellite sequences by dot-blot |
| PRACTICE 4: BIOINFORMATICS. | Database search and comparison of nucleic acid sequences. Primer design. |
| | |

| | Planning | | | |
|---------------------------------|--------------------|----------------|--------------------|-------------|
| Methodologies / tests | Competencies | Ordinary class | Student?s personal | Total hours |
| | | hours | work hours | |
| Guest lecture / keynote speech | A5 A11 A12 A15 B2 | 28 | 42 | 70 |
| | B3 B7 | | | |
| Seminar | A5 A11 A12 A15 A29 | 8 | 12 | 20 |
| | B1 B2 B3 B5 B7 | | | |
| Laboratory practice | A5 A11 A12 A15 A30 | 15 | 7.5 | 22.5 |
| | A31 B1 B2 B3 B5 B7 | | | |
| Supervised projects | A5 A11 A12 A15 A29 | 0 | 29.5 | 29.5 |
| | B1 B2 B3 B5 B7 | | | |
| Mixed objective/subjective test | A5 A11 A12 A15 A29 | 6 | 0 | 6 |
| | B1 B2 B3 B7 | | | |
| Personalized attention | | 2 | 0 | 2 |

| | Methodologies | | |
|----------------------|--|--|--|
| Methodologies | Description | | |
| Guest lecture / | The teacher explains the main contents of each lesson. | | |
| keynote speech | | | |
| Seminar | Resolution/discussion of questions and problems realted to the subject. | | |
| Laboratory practice | The student conducts laboratory experiences following a protocol, under the supervision of the teacher. | | |
| Supervised projects | Solving of three questionnaires with exercises and questions related to some aspect of the subject as well as a practical case | | |
| | related to Bioinformatics. Questionnaires shall be delivered in writing and the practical case will be orally defended. Both | | |
| | activities are done in groups. | | |
| Mixed | Objective test, short answer questions and problem resolution. | | |
| objective/subjective | | | |
| test | | | |



| | Personalized attention | | |
|--|------------------------|--|--|
| Methodologies | Description | | |
| Supervised projects Individually or in group, students can attend tutorial sessions to consult any doubts that might arise from the different activities | | | |
| | | | |
| | | | |

| Assessment | | | |
|----------------------|--------------------|--|---------------|
| Methodologies | Competencies | Description | Qualification |
| Supervised projects | A5 A11 A12 A15 A29 | Assessment will be based on providing correct answers, clarity of explanations and | 25 |
| | B1 B2 B3 B5 B7 | documentary sources used. Questionnaires and practical case represent 15% and | |
| | | 10% of the final grade, respectively. The score depends on work carried out at | |
| | | personal (50%) and group level (50%). | |
| Mixed | A5 A11 A12 A15 A29 | The degree of general knowledge and understanding of the subject will be assessed. | 75 |
| objective/subjective | B1 B2 B3 B7 | Consists of two parts. | |
| test | | One is related to theoretical content and represents 70% of the score. | |
| | | The other is related to the laboraroty practices and represents 10% of the score. | |

Assessment comments

To pass the course, the score must be 5 or higher but with at least a 4 in each part of the test. If the sum of the score of all activities is higher than 5 but the score on one part of the text is lower than 4, the final score is 4.9 (failing score).

The grade of Non Attendance (NP) will be applied to the students that do no attend the official exam.

Preferably, first class honors will be awarded in January among students with a score of 9 or higher.

A mid-term exam will be held. A score of 5 or higher will be maintained until July.

In July there is the opportunity to retake only the test. The January's score of the supervised projects is maintained.

In the case of justified exceptional circumstances, additional measures may be taken, so that the student can pass the subject, such as flexibility in the delivery date of supervised projects, flexibility in practice schedules or a global assessment test of the learning results.

| | Sources of information |
|-------|--|
| Basic | - Griffiths, A.J.F., Wessler, S.R., Suzuki, Lewontin, R.C. Carroll, S.B. (2008). Genética. McGraw-Hill/Interamericana de |
| | España, Madrid |
| | - Klug, W.S., Cummings, M.R., Spencer, C.A (2013). Conceptos de Genética . Pearson/Prentice Hall, Madrid |
| | - Pierce, B.A. (2010). Genética: un enfoque conceptual. Médica Panamericana, Madrid |

Complementary

- Alberts, B., Johnson, A., Lewis, J., Raff, M., Roberts, K., Walter, P. (2010). Biología Molecular de la célula. Omega, Barcelona
- Benito, C., Espino, F.C. (2013). Genética: conceptos esenciales. Médica Panamericana, Madrid
- Brooker, R.J. (2015). Genetics: analysis and principles (5th ed). McGraw-Hill, New York
- Brown, T.A. (2008). Genomas (3ª ed.). Médica Panamericana, Buenos Aires
- Craig, N.L., Cohen-Fix, O., Green, R., Greider, C., Storz, G., Wolberger, C. (2014). Molecular Biology: principles of genome function. Oxford University Press, Oxford
- Hartwell, L.H., Goldberg, M.L., Fischer, J.A., Hood, L., Aquadro, C.F. (2015). Genetics: from genes to genomes (5th ed.) . McGraw-Hill, New York
- Herráez, A. (2012). Biología Molecular e Ingeniería genética. Elsevier, Ámsterdam
- Krebs, J.E., Goldstein, E.S., Kilpatrick, S.T. (2012). Lewin genes: fundamentos. Médica Panamericana, Madrid
- Lewin, B. (2008). Genes IX. McGraw-Hill. México
- Lodish, H., Berk, A., Kaiser, C.A., Krieger, M., Bretscher, A., Ploegh, H., Amon, A., Scott, M.P. (2016). Biología Celular y Molecular (7ª ed) . Médica Panamericana, Madrid
- Perera, J., Tormo, A., García, J.L. (2002). Ingeniería genética. Vol. I: Preparación, análisis, manipulación y clonaje de DNA. Síntesis, Madrid
- Perera, J., Tormo, A., García, J.L. (2002). Ingeniería genética. Vol. II. Expresión de DNA en sistemas heterólogos. Síntesis, Madrid
- Russell, P.J. (2010). iGenetics: a molecular approach (3º ed.) . Benjamin Cummings, San Francisco
- Snustad, D.P., Simmons, M.J. (2012). Genetics (6th ed). John Wiley and Sons, New York

Consultar a plataforma Moodle para fontes de información adicionais.

Recommendations

Subjects that it is recommended to have taken before

Biology: Basic Levels of Organisation of Life I (Cells)/610G02007

Biochemistry I/610G02011 Biochemistry II/610G02012 Microbiology/610G02015

Genetics/610G02019

Subjects that are recommended to be taken simultaneously

Subjects that continue the syllabus

Population Genetics and Evolution/610G02021

Cytogenetics/610G02022

Other comments

Recommendations: Attend class and follow the development of the course regularly. Check Moodle and email regularly to obtain the materials and know the schedule of activities. Attend tutorials to resolve any questions or difficulties that may arise. Consult the recommended bibliography. Keep up-to-date with course work.

(*)The teaching guide is the document in which the URV publishes the information about all its courses. It is a public document and cannot be modified. Only in exceptional cases can it be revised by the competent agent or duly revised so that it is in line with current legislation.