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| In the event that teaching method. 1. Modifications: Contents would. 2. Methodologi. * Teaching method. * T | subject focused on micr | roscopy (includin | ng image analys | is), plant and animal o | cell culture, flow cytometry and | |
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| 1. Modifications Contents would 2. Methodologi * Teaching met The methodolo * Teaching met If necessary, sy face-to-face tea Practical session hybrid model) of analysis and in 3. Mechanisms Personalized at 4. Modifications Criteria and methodological | In the event that circumstances limit or prevent the access to the facilities of the Faculty, a hybrid or nonattendance | | | | | |
| Contents would 2. Methodologi * Teaching met The methodolo * Teaching met If necessary, sy face-to-face tea Practical session hybrid model) of analysis and in 3. Mechanisms Personalized at 4. Modifications Criteria and me | teaching method would be adopted, respectively, with the following specifications. | | | | | |
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| If necessary, so face-to-face tear Practical session hybrid model) of analysis and in 3. Mechanisms Personalized at 4. Modifications Criteria and me | ogies described will be | maintained in bo | th modalities. | | | |
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| Practical session hybrid model) of analysis and in 3. Mechanisms Personalized a 4. Modifications Criteria and me | ynchronous means of o | communication (| MSTeams,) w | vill be used in teaching | g activities that involve | |
| hybrid model) of analysis and in 3. Mechanisms Personalized a 4. Modifications Criteria and me | aching method. | | | | | |
| analysis and in 3. Mechanisms Personalized a 4. Modifications Criteria and me | ons in the laboratory w | vill be adapted to | the circumstand | ces; if necessary, they | will be replaced partial (in the | |
| 3. MechanismsPersonalized a4. ModificationsCriteria and me | or totally (in the non-att | tendance modali | ty) by non-atten | dance activities (video | o viewing, case studies, data | |
| Personalized a 4. Modifications Criteria and me | terpretation,) | | | | | |
| 4. Modification: Criteria and me | s for personalized atten | ntion to students. | | | | |
| Criteria and me | attention will be limited t | to telematic mea | ıns (email, Mood | dle, MSTeams,) | | |
| | s in the evaluation. | | | | | |
| * Frank a Cara ak | Criteria and methodologies for evaluation will be maintained in both modalities. | | | | | |
| " Evaluation of | * Evaluation observations: | | | | | |
| Assessments v | Assessments will be carried out electronically in both modalities | | | | | |
| 5. Modifications | 5. Modifications to the bibliography or webography. | | | | | |
| If necessary, co | If necessary, complementary sources or means of free access would be provided. | | | | | |

| | Study programme competences |
|------|------------------------------------------------------------------------------------------------------------------------------------------|
| Code | Study programme competences |
| A1 | Skills of working in a sure way in the laboratories knowing operation handbooks and actions to avoid incidents of risk. |
| A2 | Skills of using usual techniques and instruments in the cellular, biological and molecular research: that are able to use techniques and |
| | instruments as well as understanding potentials of their uses and applications. |

| A13 | Skills to become a professional in health, pharmacy, veterinary, animal production, biotechnology or food sectors. |
|-----|--------------------------------------------------------------------------------------------------------------------------------------------|
| В3 | Skills of management of the information: that are able to gather and to understand relevant information and results, obtaining conclusions |
| | and to prepare reasoned reports on scientific and biotechnological questions |
| B4 | Organization and work planning skills: that are able to manage the use of the time as well as available resources and to organize the work |
| | in the laboratory. |
| C1 | Adequate oral and written expression in the official languages. |
| C3 | Using ICT in working contexts and lifelong learning. |
| C8 | Valuing the importance of research, innovation and technological development for the socioeconomic and cultural progress of society. |
| C9 | Ability to manage times and resources: developing plans, prioritizing activities, identifying critical points, establishing goals and |
| | accomplishing them. |
| | |

| Learning outcomes | | | |
|---------------------------------------------------------------------------------------------------------------------------------------|------|-----------------|-----|
| Learning outcomes | | Study programme | |
| | | competences | |
| To understand the theoretical foundations on which microscopy (including image analysis), (plant and animal) cell culture, flow | AR1 | | |
| cytometry and cytogenetic techniques are based. | AR2 | | |
| To acquire basic skills in the management and use of instrumental and units required for the development of cellular | AR1 | | |
| techniques. | AR2 | | |
| | AR13 | | |
| To know the applications of the different cellular techniques. | AR2 | | |
| To design, plan and conduct experiments regarding the techniques learned. | AR1 | BR3 | |
| | AR2 | BR4 | |
| Adequate oral and written expression in the official languages | | | CC1 |
| Using ICT in working contexts and lifelong learning | | | ССЗ |
| Ability to manage times and resources: developing plans, prioritizing activities, identifying critical points, establishing goals and | | | CC8 |
| accomplishing them | | | CC9 |

| | Contents |
|-------------------------------|---------------------------------------------------------------------------------------|
| Topic | Sub-topic |
| Microscopy and image analysis | Fundamentals, techniques and applications of light and electron microscopy. |
| | Fluorescence and confocal scanning microscopy: advanced techniques and |
| | applications. Introduction to image processing and image analysis |
| (Animal & Dant) cell cultures | Introduction to cell cultures. Types of cell cultures. Cell culture requirements. |
| | Quantification of cellular parameters. Contamination. Cytotoxicity. |
| | In vitro cultures of plant tissues. Callus. Cultivation of plant cells in suspension. |
| Flow cytometry | General principles and methods of cytometry. Sample preparation and standardization |
| | of analysis procedures. Functional analysis of cells. |
| Cytogenetic techniques | Chromosome preparation and karyotype. Conventional in situ hybridization. Advanced |
| | techniques of fluorescence in situ hybridization (FISH). |

| class Student?s persor | nal Total hours |
|------------------------|----------------------------------|
| s work hours | |
| 28 | 42 |
| 42 | 70 |
| 19 | 19 |
| | |
| 15 | 17 |
| 0 | 2 |
| _ | 0 count the heterogeneity of the |

| | Methodologies | | |
|----------------------|----------------------------------------------------------------------------------------------------------------------------------|--|--|
| Methodologies | Description Description | | |
| Guest lecture / | Teacher will present the theoretical and practical contents of the subject (of the different techniques currently used in Cell | | |
| keynote speech | Biology). | | |
| Laboratory practice | Practical sessions represent an indispensable part of the course, in which practical aspects and applications of the different | | |
| | cellular techniques will be approached. Practical sessions will be developed in concrete laboratories and places: laboratories | | |
| | of the Faculty of Sciences, Laboratories of Biomedical Research Institute (INIBIC), Scientific Research Support Services (SAI) | | |
| | of UDC. | | |
| | Students will develop laboratory protocols and attend demonstrations about the use of research units. | | |
| Supervised projects | Students must carry out works, resolve problems and/or questions about specific aspects of the techniques used. | | |
| Mixed | It will consist of a written exam with questions-test and/or short answer questions about theoretical and practical contents and | | |
| objective/subjective | applications of the cellular techniques. | | |
| test | | | |

| Personalized attention | | |
|------------------------|--------------------------------------------------------------------------------------------------------------------------------|--|
| Methodologies | Description | |
| Supervised projects | Students (individually or in small groups) may consult their doubts about the contents and activities of the subject via phone | |
| | and/or electronic support. | |
| | | |

| | | Assessment | |
|----------------------|-------------------|---------------------------------------------------------------------------------------------------------------------|---------------|
| Methodologies | Competencies | Description | Qualification |
| Mixed | A2 B3 | The acquisition of knowledge about the theoretical foundations and applications, | 50 |
| objective/subjective | | clarity of explanations, ability to integrate and link information handled and the ability | |
| test | | to interpret data and solve problems will be taken into account. | |
| Laboratory practice | A2 A1 A13 B3 B4 | Attendance, skill in scheduled tasks and knowledge of the potential risks in laboratory practices will be assessed. | 20 |
| Supervised projects | A2 B3 B4 C1 C3 C8 | The ability to design (and plan) experiments, interpretate data and solve problems will | 30 |
| | C9 | be assessed. | |

Assessment comments

In order to be evaluated, students must attend to practical sessions.

In july there is the opportunity to retake only the tests. The january's score of supervised projects and practices are maintained.

Preferably, first class honors will be awarded in january.

Full-time and part-time students will be evaluated following this qualification guideline.

Sources of information

Basic

CULTIVOS CELULARESBasra, A.S. (2000). Plant growth regulators in agriculture and horticulture. Their role and commercial uses. Ed. Food Products Press. Benítez Burraco, A. (2005). Avances recientes en Biotecnología vegetal e ingeniería genética de plantas. Editorial Reverté.Boulton, A.A. e col. (1992). Practical cell culture techniques. Humana Press.Butler, M. (2008). Animal cell culture and technology. Taylor & press.Butler, M. (2008). Animal cell culture and technology. Taylor & press.Butler, M. (2008). Animal cell culture and technology. Edwars, S. (1998). Plant cell culture. Guilford Bios Scientific Publishers. Davis, J.M. (2011). Animal cell culture. Essential methods. Wiley-Blackwell.Doyle, A. e Griffiths, J.B. (2000). Cell and tissue culture for medical research. John Wiley and Sons. Fedoroff, S. e Richardson, A. (1992). Protocols for neural cell culture. Humana Press. Freshney, R.I. (2010). Culture of animal cells. A manual of basic technique and specialized applications. Wiley-Liss, Inc. 6a Edition. Hammond, J., McGarvey, P., Yusibov, V. (1999). Plant Biotechnology. New products and Applications. Springer Verlag, Loyola-Vargas, V.M. e Vázguez-Flota F. (2006). Plant cell culture protocols. Humana Press. 2nd Edition. Pollard, J.W. e Walker, J.M. (1997). Basic cell culture protocols. Humana Press. Shaw, A.J. (1996). Epithelial cell culture. A practical approach. Oxford University Press. Taji, A., Kumar, P., Lakshmanan, P. (2002). In vitro plant breeeding. Food Products Press.Trigiano, R.N. e Gray, D.J. (2004). Plant development and biotechnology. CRC Press.Tzfira, T. e Citovsky, V. (2006). Agrobacterium-mediated genetic transformation of plants: biology and biotechnology, Curr. Opin. Biotechnol. 17:147?154. Vunjak-Novakovic, G. & amp; nbsp; & amp; amp; Freshney, R.I. (2006). Culture of cells for tissue engineering. Wiley-Liss, Inc.TÉCNICAS DE MICROSCOPÍA E ANÁLISE DE IMAXEWatt, Ian M. (1996). The principles and practice of electron microscopy. Cambridge University PressHoppert, M. (1998). Electron microscopy in microbiology. Bios Scientific PublishersBozzola, John J. (1999). Electron microscopy : principles and techniques for biologists. Jones and Bartlett Publishers. Dykstra, Michael J. (2003). Biological electron microscopy theory, techniques, and troubleshooting. Kluwer Academic/Plenum PublishersRobin Harris. (1991). Electron microscopy in biology a practical approach. Oxford University Press. Hunter, Elaine Evelyn. (1984). Practical electron microscopy a beginner's illustrated guide. Praeger, cop.Slayter, Elizabeth M. (2000). Light and electron microscopy. Cambridge University PressHerman, B. (1998). Fluorescence microscopy. Bios Scientific PublishersDonat-P. Häder. (1992). Image analysis in biology. CRC Press, cop. Pertusa, JF. (2003). Técnicas de Análisis de imagen. Aplicaciones en Biología. Publicaciones de la Universidad de Valencia. CITOMETRÍA DE FLUXOOrmerod, M.G. (2009). Flow Cytometry: A Basic Introduction. 2a Ed. IRL Practical Approach series. Oxford University Press.Shapiro, H.M. (2004). Practical flow cytometry. Wiley-Liss. 4a ed. New York.TÉCNICAS CITOXENÉTICASCzepulkowski, B. (2001). Analyzing chromosomes. BIOS Scientific Publishers, Oxford.Gosden, J.R. (1994). Chromosome analysis protocols. Humana Press, Totowa (New Jersey). Kianian, S.F., Kianian P.M.A. (2016). Plant cytogenetics: methods and protocols. Spinger, New YorlLiehr, T. (2006). Multicolor FISH in human cytogenetics. Karger, Basel, Liehr, T. (2009). Fluorescence in situ hybridization (FISH)-application guide. Springer-Verlag, Berlin, Leitch, A.R., Schwarzacher, T., Jackson, D. (1994). In situ hybridization: a practical guide. Bios Scientific Publishers, Oxford. Verma, R.S. e Babu, A. (1989). Human chromosomes: manual of basic techniques. Pergamon Press. New York.

Complementary

- Artigos científicos sobre temas relacionados coa materia proporcionados a través da plataforma Moodle.- Páxinas webXeralPubMed: http://www.ncbi.nlm.nih.gov/pubmedCultivos

Celulares-http://www.cultek.com/aplicaciones.asp?P=Aplicacion_Cultivos_Celulares&opc=introduccionCito metría-Cytometry: http://www3.interscience.wiley.com/cgi-bin/jhome/33945Microscopía e Análise de imaxehttp://zeiss-campus.magnet.fsu.edu/index.htmlhttp://www.microscopyu.com/tutorials/http://www.olympusfluoview.com/index.htmlhttp://w3.uniroma1.it/MEDICFISIO/microscopy.htmhhttp://rsbweb.nih.gov/ij/index.htmlhttp://www.invitrog

| Recommendations |
|----------------------------------------------------------|
| Subjects that it is recommended to have taken before |
| |
| Subjects that are recommended to be taken simultaneously |
| |
| Subjects that continue the syllabus |

en.com/site/us/en/home/support/Research-Tools/Fluorescence-SpectraViewer.html



| Other comments |
|----------------|
| |

(*)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.