

	т	eaching Guide			
	Identifying Data			2022/23	
Subject (*)	Cellular Techniques		Code	610441001	
Study programme	Máster Universitario en Bioloxía Molecula	ar, Celular e Xenética			
		Descriptors			
Cycle	Period	Year	Туре	Credits	
Official Master's Degre	ee 1st four-month period	First	Obligatory	6	
Language	SpanishGalicianEnglish				
Teaching method	Face-to-face				
Prerequisites					
Department	BioloxíaDepartamento profesorado máste	er			
Coordinador	Yañez Sanchez, Julian E-mail julian.yanez@udc.es			c.es	
Lecturers	Bernal Pita da Veiga, María de los Ángele	es E-mail	angeles.bernal@	udc.es	
	Castro Castro, Antonio Manuel		antonio.castro@	udc.es	
	Insua Pombo, Ana Maria		ana.insua@udc.e	es	
	López Armada, María José		maria.jose.lopez	.armada@col.udc.es	
	Rioboo Blanco, Carmen		carmen.rioboo@	udc.es	
	Yañez Sanchez, Julian		julian.yanez@ud	c.es	
Web	https://campusvirtual.udc.gal		I		
General description	Experimental subject focused on microsc	opy (including image ana	alysis), plant and animal	cell culture, flow cytometry an	
	cytogenetic techniques.				
	1				

	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.
B3	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	Ability to express oneself correctly, both orally and in writing, in the official languages of the autonomous community
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	/ progra	mme
	cor	npetend	ces
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.		CC3
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 & Plant) 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).			

Planning			
Competencies	Ordinary class	Student?s personal	Total hours
	hours	work hours	
A2	14	28	42
A2 A1 A13 B3 B4	28	42	70
A2 B3 B4 C1 C3 C8	0	19	19
C9			
A2 B3	2	15	17
	2	0	2
	Competencies A2 A2 A1 A13 B3 B4 A2 B3 B4 C1 C3 C8 C9	A2 14 A2 A1 A13 B3 B4 28 A2 B3 B4 C1 C3 C8 0 C9 2 A2 B3 2	CompetenciesOrdinary class hoursStudent?s personal work hoursA21428A2 A1 A13 B3 B42842A2 B3 B4 C1 C3 C8019C9215

guidance only and does not take into account the	neterogeneity of the students.

	Methodologies				
Methodologies	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	Methodologies Description				
Supervised projects	pervised 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

Attendance at the practical lessons is a necessary condition to be evaluated. In case of not passing the evaluation in the first opportunity of the call, the grades obtained in the supervised works and practices will be kept for the second opportunity. Honors will be awarded preferably among students (face-to-face and non-face-to-face) presented in the evaluation corresponding to the first opportunity of the call.

Fraudulent performance of the tests or assessment activities, once verified, will directly imply the failing grade "0" in the call.

Sources of information



Basic

Complementary

CULTIVOS CELULARES Basra, 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 & amp; Francis, 2nd edition.Capes-Davis, A.
(2021). Freshney's culture of animalcells: a manual of basic technique and specialized applications. Wiley-Blackwell.
8th edition. Collin, H.A. e 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. Hammond, J., McGarvey, P., Yusibov, V. (1999). Plant Biotechnology. New products and
Applications. Springer Verlag.Loyola-Vargas, V.M. e Vázquez-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.& Freshney, R.I. (2006). Culture of cells
for tissue engineering. Wiley-Liss, Inc. TÉCNICAS DE MICROSCOPÍA E ANÁLISE DE IMAXE Watt, 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 FLUXO Ormerod, 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 CITOXENÉTICAS Czepulkowski, B. (2001).
Analyzing chromosomes. BIOS Scientific Publishers, Oxford.Gersen, S.L., Keagle, M.B. (2013). The principles of
clinical cytogenetics. Springer, New York.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.
- Artigos científicos sobre temas relacionados coa materia proporcionados a través da plataforma Moodle Páxinas
webYeralPubMed: http://www.nchi.nlm.nih.gov/pubmedCultivos

webXeralPubMed: http://www.ncbi.nlm.nih.gov/pubmedCultivos Celulares-https://inmunomundo.files.wordpress.com/2015/12/cultivo-celular.pdf-http://www.lgcstandards-atcc.org/Cito 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.htmhttp://rsbweb.nih.gov/ij/index.htmlhttp://www.invitrog en.com/site/us/en/home/support/Research-Tools/Fluorescence-SpectraViewer.html

Recommendations	
Subjects that it is recommended to have taken before	
Subjects that are recommended to be taken simultaneously	



 Subjects that continue the syllabus

 Other comments

 Program Green Campus

 Empower of SciencesTo help to achieve some sustainable immediate surroundings

 and fulfil with the point 6 of the Environmental Statement of the faculty of

 Sciences (2020), the documentary works that realise in this matter:to. They will request

 mostly in virtual format and computer supportb. To realise in

 paper:-they will not employ

 plastic-will realise

 impressions to double expensive-will employ paper

 recycled-will avoid the

 realisation of draftsTo Environmental Statement is available

 in:https://ciencias.udc.es/images/Facultade/Green_Campus/Regulamento_Comit%C3%A9_Green_Campus_FCiencias.pdf

(*)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.