



1. Identification data:	
Institution name:	Automous University of Nuevo León
Dependency name:	Medical School
Name of the educational program:	Clinical Chemist Biologist
Learning unit name:	Molecular Biology
Theory and/or practical classroom hours, total:	100
Frecuencies per week:	6
Overtime, total:	50
Modality type:	Schooled
Type of academic period:	Fifth Semester
Type of learning unit:	Mandatory
Curricular area:	ACFB
UANL Credits:	5
Production date:	21/06/2018
Last update date:	10/06/2024
Responsible (s) for design and updating:	Design:
	Celia Nohemí Sánchez Domínguez, PhD
	María Del Carmen Villalobos Torres, PhD
	Antonio Alí Pérez Maya, PhD
	Ana María G. Rivas Estilla, PhD
	Update:
	Ana María G. Rivas Estilla, PhD
	Ma. del Carmen Barboza Cerda, PhD
	Sonia A. Lozano Sepúlveda, PhD
	Celia N. Sánchez Domínguez, PhD





#### 2. Presentation:

The purpose of the Learning Unit (LU) of Molecular Biology (MB) is for the student to understand the importance of the main macromolecules of life: DNA, RNA, and proteins (Central Dogma of Molecular Biology), analyzing structural aspects, as well as the interactions and metabolic processes that occur between them and the application of molecular diagnostic tools.

In this learning unit, the contents are divided into 3 phases. In the first phase, the molecular bases of the Central Dogma (structure of nucleic acids; DNA replication, RNA transcription and protein translation, as well as gene regulation) are reviewed. In the second phase, the student knows the principles and foundations and identifies the main tools of molecular biology used for the manipulation and study of nucleic acids, which are crucial in real-world research and diagnostics. Finally, in the third phase, the applications of molecular biology are studied for the understanding of molecular medicine, molecular mechanisms of pathogenicity, the diagnosis of genetic and infectious diseases, gene therapy and biotechnological processes, all of which have direct applications in the field. During stages 2 and 3 the student will have the opportunity to execute basic molecular biology laboratory techniques, preparing them for their future careers in the field.

At the end of the LU, as an integrative product of learning, the student will develop a research project on the bases and techniques of Molecular Biology, as well as its applications in molecular medicine, the diagnosis of human diseases, and biotechnological processes, through exposure by the team before the group in a seminar and a final theoretical

#### 3. Purpose (s):

The purpose of the Learning Unit (LU) of Molecular Biology (MB) is for the student to analyze the structural aspects, interactions, and metabolic processes that occur between the main macromolecules of life: DNA, RNA, and proteins (Dogma Central of Molecular Biology). Likewise, you will learn and analyze the fundamentals and applications of Molecular Medicine, molecular diagnostic tools used for the interpretation of genetic and infectious diseases, and biotechnology.

This learning unit is located in the fifth semester. It is part of the basic instruction that the Clinical Chemistry Biologist student must acquire to base their professional practice within Molecular Medicine and Molecular Diagnostics. For this LU, use is made of the knowledge acquired in the LU of Biochemistry since it allows it to integrate the metabolism of biomolecules, including amino acids and nucleotides as fundamental elements of proteins and nucleic acids, respectively, of Morphological Sciences to recognize their aspects. Structural and functional of cells and tissues of





medical physiology by basing the homeostatic processes of the organism. In turn, the LU of MB contributes to obtaining the LU's competencies in both Clinical Biochemistry by providing basic knowledge of the molecular tests used in the diagnosis of molecular diseases and Clinical Pathology by implementing and interpreting laboratory tests for the diagnosis of molecular diseases.

Likewise, the LU of MB will be the basis of the optional UA of Molecular Diagnosis and Biotechnology since it will use the knowledge acquired in applying the tools of Molecular Biology to the molecular diagnosis of diseases (monogenic, multifactorial, and infectious), the studies of individual identification (forensic, paternity, and chimerism), and biotechnology.

The LU of MB serves as a platform for promoting logical, critical, and purposeful thinking in students. It encourages them to analyze the structural characteristics of the macromolecules of life in relation to their function and their impact on the organism's functioning. Moreover, it provides opportunities for students to discuss societal health challenges, fostering interventions with a critical attitude and professional commitment. This LU also contributes to the general well-being of students and their colleagues, promoting respect for working conditions in the classroom and laboratory. The LU of MB is not just about theoretical learning; it's also about skill development. Students are encouraged to apply their knowledge of the Central Dogma of Molecular Biology to solve problems. They also develop skills for executing chemical and/or biological procedures in the analysis of samples, which are crucial for the clinical diagnosis of genetic diseases. Furthermore, they learn to apply their knowledge to understand and interpret the production of recombinant proteins, their purification, and their use in solving health problems.

# 4. Graduation profile competencies:

#### General competencies to which this learning unit contributes:

Instrumentals

1. Use logical, critical, creative, and purposeful thinking to analyze natural and social phenomena that allow you to make relevant decisions with social responsibility in your sphere of influence.

#### Personal and social interaction

2. Intervene in the face of the challenges of contemporary society, locally and globally, with a critical attitude and human, academic, and professional commitment to contribute to consolidating general well-being and sustainable development.





#### Integrators

3. Resolve personal and social conflicts following specific techniques in the academic field and their profession for appropriate decision-making.

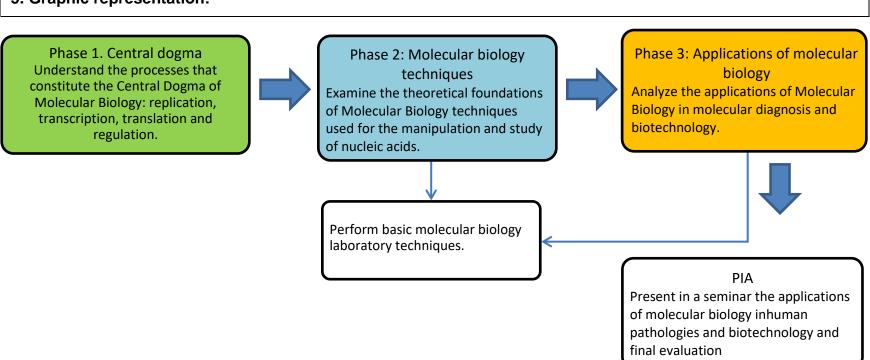
#### Specific competencies of the graduation profile to which the learning unit contributes:

- 1. Solve problems by applying knowledge of the chemical composition of matter, as well as its physicochemical properties to determine analytes in biological, environmental and food matrices.
- 2. Execute physical, chemical and/or biological procedures in obtaining, handling, storing and analyzing samples to contribute to a reliable clinical, toxicological, chemical, food, forensic and environmental diagnosis.





#### 5. Graphic representation:







# **PHASE I: The Central Dogma of Molecular Biology**

**Element of competence:** Understand the processes that constitute the central dogma of molecular biology, considering the normal bases of the molecular processes of the cell, to identify the consequences of its alterations

Evidence of learning	Performance criteria	Learning activities	Contents	Resources
Evidence 1 First partial evaluation on the processes that constitute the Central Dogma of Molecular Biology.	Individually solve a partial evaluation on the established date and time that includes the contents corresponding to phase 1.	<ul> <li>The teacher explains the topic and asks guidance and contextualization questions to consolidate concepts and resolve doubts.</li> <li>The student reads the corresponding chapters of the Biochemistry book (8th edition) by Emine E. Abalí according to the class calendar.</li> <li>The student actively participates in the sessions to review the contents</li> <li>The student accesses the Moodle and MS Teams platforms to retrieve teaching material published by the teacher and deliver the ponderable activities.</li> <li>The student analyzes the information corresponding to each topic after reading the corresponding chapter.</li> </ul>	DNA structure, replication and repair  - DNA structure  - Stages in the synthesis of DNA in prokaryotes.  - DNA replication of eukaryotic DNA.  - DNA repair.  - Characteristics and organization of viral genomes and plasmids  • Structure, synthesis and processing of RNA.  - Structure of RNA.  Transcription of prokaryotic genes.  - Transcription of eukaryotic genes.  - Post-transcriptional modification of RNA  • Protein synthesis.	Computer equipment or smart devices with Microsoft Office and internet connection  Electronic presentations  Biochemistry 8th Edition. Emine E Abali Chapters 31, 32, 33 and 34.  Current articles provided by the teacher  Moodle platform for delivery of ponderable activities  Microsoft Teams platform for organizing joint activities, delivering documents of ponderable activities, reviewing teaching materials and discussion between students and teacher.





• The student presents as a team during the session a topic from phase 1 assigned by the teacher.

#### Considerable activity 1.

Comparative tables of the Central Dogma of Molecular Biology. As a team, organize the information from chapters 31, 32 and 33 reviewed during phase 1 into tables, comparing the process between prokaryotes and eukaryotes. It is delivered on the established date and time through the MS Teams or Moodle platforms in accordance with the instructional guide and checklist.

work on the genetic code. Individually, prepare a written work about a disease caused by alteration of the genetic code.

It is delivered on the established date and time through the MS Teams or Moodle platforms in accordance with the instructional guide and checklist.

- The genetic code.
- Components necessary for translation.
- Codon recognition by tRNAs.
- Stages in protein synthesis
- Co-translational and posttranslational modification of polypeptide chains.
- Translation in prokaryotes and eukaryotes.
- Co-translational and post-translational modification.
- Regulation of gene expression in prokaryotes and Eukaryotes.
- Regulatory sequences and molecules. Regulation of gene expression in prokaryotes.
- Regulation of gene expression in eukaryotes.
- Role of MiRNAs, RNAi, IncRNA, IRES and Crispr/Cas9 in the processes of regulation of gene expression and editing

MS Forms platform for daily evaluations.

Electronic resources:

Google Scholar/google scholar National Center for Biotechnology Information (NCBI)

Web Pages:
YouTube:
DNA replication process
(Free Science), transcription
(process, intron removal and
post-transcriptional
modifications), translation
and regulation of the Lac
operon (Virtual Cell).





# Phase 2. Molecular Biology techniques in the manipulation and study of nucleic acids. Competition element(s):

Examine the fundamentals of molecular biology procedures used for the isolation, study, analysis, and manipulation of nucleic acids from different biological, environmental, and food matrices to justify their use in the various fields of application of Molecular Medicine, molecular biology, and biotechnology.

Evidence of learning	Performance criteria	Learning activities	Contents	Resources
Evidence 2 Second partial evaluation: molecular biology techniques used for the study of nucleic acids	Individually solve a partial evaluation on the established date and time that includes the contents corresponding to phase 2.	asks guidance and contextualization questions to	DNA Cloning Polymerase chain reaction and its variants: PCR-RFLP, real-time PCR, RT-PCR, hot	Electronic presentations  Biochemistry 8th Edition. Emine.E. Abali, Chapter 35  Web Pages: YouTube: Nucleic acid extraction, electrophoresis, cloning, PCR, sequencing and microarrays.  Molecular biology and genetic engineering 2nd. Edition. Angel Herráez. Chapter 14, 15 and 16  Molecular biology. Fundamentals and
			start PCR, long PCR, nested PCR, inverse PCR, PCR with	applications in health





		accordance with the instructional guide and checklist.  Considerable activity 4. Comparative table of Molecular Biology techniques.  The student individually prepares a comparative table of the techniques of PCR, cloning, library preparation, sequencing and microarrays where he describes the fundamentals, description of the technique, variants, applications, interpretation of results.  It is delivered on the established date and time through the MS Teams or Moodle platforms in	adapters, asymmetric PCR and digital PCR.  - Fundamentals of genomic sequencing and the human genome project (HUGO)  - Exome, transcriptome and proteome  - Analysis of gene expression: qPCR and Microarrays.	sciences, 2nd edition. Salazar Montes, AM.  Current articles provided by the teacher.  Moodle platform for delivery of ponderable activities  Microsoft Teams platform for and discussion between students.  MS Forms platform for daily evaluations.  Electronic resources:  Web Pages: YouTube: Nucleic acid extraction, cloning, electrophoresis, PCR, sequencing and microarrays
		accordance with the instructional guide and checklist		
Evidence 3				Teaching laboratory
Laboratory practices:	Review the theoretical foundations of the practice including the material to	<ul> <li>The student individually prepares the flow chart and the corresponding concept map,</li> </ul>	Biosafety and hygiene standards in the Molecular Biology laboratory.	Molecular Biology Laboratory Practice Manual.





	work in the laboratory and	delivering it at the beginning of	- Safety signs.	Digital, audiovisual or digital
3.1 1 Safety in the	the established safety	each session through the Moodle	- Waste disposal.	reading resources
Molecular Biology	standards.	platform.	<ul> <li>- Proper use of equipment</li> </ul>	
laboratory	Prepare the activities that	<ul> <li>The corresponding team explains</li> </ul>	and reagents.	Moodle platforms for delivery of
	support the corresponding	the theoretical foundations of the		ponderable activities
	practice.	practice, for which they prepare a	Quality parameters:	
3.2	It presents the necessary	Power Point presentation.	precision and accuracy.	
Quality control	material to work properly	<ul> <li>The teacher complements the</li> </ul>	<ul> <li>- Organization of a</li> </ul>	
	in the laboratory, taking	explanation of the practice	Molecular Biology Laboratory.	
	care of the established	presented by the designated team	- Proper use of	
	safety standards.	by asking questions and providing	micropipettes.	
	Prepare the	examples that reinforce the		
	corresponding activities,	theoretical knowledge of the	Extraction of human	
	work in an organized and	practice.	genomic DNA from a blood	
3.3	team manner on the	Students prepare a report	sample with the organic	
Genomic DNA	proposed experiments.	collaboratively using experimental	phenol-chloroform extraction	
extraction	The student will review	data, following the format	technique.	
	the requirements to carry	established in the practice manual.		
	out the practice cleanly, in		- DNA analysis by UV	
	a responsible and orderly		spectrophotometry.	
	manner, complying with		T (   DNA ( (; )	
0.4	the safety and hygiene		Total RNA extraction and	
3.4	standards established for		solid phase purification from	
Total RNA	the correct use of the		eukaryotic cell culture.	
extraction and	facilities. and correct		BNA analysis by UV	
solid phase	waste disposal.		- RNA analysis by UV	
purification	Appears to laboratory sessions on the		spectrophotometry.	
	established date and time.			
	•			
	time in accordance with			
	The report is delivered on the established date and time in accordance with			





3.5	the instructional guide and	•	• • Agarose gel	
Agarose gel	checklist.	e	electrophoresis for nucleic	
electrophoresis for		a	acids (DNA and RNA).	
nucleic acids.		•	<ul> <li>Determination of DNA</li> </ul>	
		a	and RNA integrity by analyzing	
			mages of agarose gels	
			stained with ethidium bromide.	
		•	<ul> <li>Biosafety and hygiene</li> </ul>	
			standards in the Molecular	
		B	Biology laboratory.	
		•	<ul> <li>Safety signs.</li> </ul>	
		•	<ul> <li>- Waste disposal.</li> </ul>	
		•	<ul> <li>- Proper use of</li> </ul>	
		e	equipment and reagents.	
		•	•	
		•	<ul><li>Quality parameters:</li></ul>	
		p	precision and accuracy.	
		•	<ul> <li>Organization of a</li> </ul>	
		N	Molecular Biology Laboratory.	
		•	<ul> <li>Proper use of</li> </ul>	
		n	micropipettes.	
		•	•	
		•	•	
		•	<ul> <li>Extraction of human</li> </ul>	
		•	genomic DNA from a blood	
			sample with the organic	
			phenol-chloroform extraction	
		te	technique.	
		•	•	
		•	<ul> <li>DNA analysis by UV</li> </ul>	
		S	spectrophotometry.	
		•	•	





	• Total RNA extraction and solid phase purification from eukaryotic cell culture.      • RNA analysis by UV spectrophotometry.      • Agarose gel electrophoresis for nucleic acids (DNA and RNA).      • Determination of DNA and RNA integrity by analyzing images of agarose gels etained with athidium bromide.
	stained with ethidium bromide.





# PHASE: 3 General applications of Molecular Biology in molecular diagnosis and biotechnology.

#### **Competition element:**

Analyze the general applications of Molecular Biology techniques in molecular diagnosis and biotechnology to substantiate their usefulness in Biomedicine and Industry.

Evidence of learning	Performance criteria	Learning activities	Contents	Resources
Evidence 4	Individually solve a	The teacher presents the key	Scope and applications of	Computer equipment or smart
Third partial	partial evaluation on		Molecular Medicine.	devices with Microsoft Office and
evaluation on	the established date	0		internet connection
molecular	and time that	contextualization questions to	Molecular Diagnosis:	FI
diagnosis and	includes the	consolidate concepts and	- Applications in hereditary	Electronic presentations
biotechnology.	contents	resolve doubts with the	diseases	Moodle platform for delivery of
	corresponding to phase 3.	support of infographics, videos and power point	<ul> <li>Applications in the molecular diagnosis of multifactorial</li> </ul>	Moodle platform for delivery of ponderable activities
	priase 5.	presentations.	diseases and cancer	portuerable activities
		procentations.	Applications in the	
		Prior to the theoretical	identification of pathogenic	- Books:
		session, the student reviews	agents: bacteria, fungi and	Biochemistry 8th Edition. Abali
		didactic material published by		Emine.E. Chapter 35
		the teacher.		
				Colleman W.B, Tsongalis G.J.,
		The student actively		Molecular Diagnostics For the
		participates in the online		Clinical Laboratory. Chapters 22, 23
		sessions to review the contents.		and 34 to 38.
		COINGINS.		Cox T. M., Molecular Biology in
		Considerable activity 5:		Medicine. Chapters 6 and 7.
		Table on different types of		Modicino. Chaptero o ana 7.
		molecular diagnoses.		Nussbaum R. L. Genetics in
		Analyzes the molecular tests		medicine. Chapter 11.





		used in the molecular diagnosis of hereditary, multifactorial (cancer) and infectious diseases.  Prepare as a team a table showing different diagnostic tests and deliver it on the date indicated by the teacher through the Moodle or Microsoft Teams platforms in accordance with the instructional guide and checklist.  • Considerable activity 6: Summary of the fundamentals and technical requirements to obtain a biotechnological product, using as an example a vaccine, transgenic organism or recombinant protein.		Antokoletz, A. F. G., Sarmiento, M. Á., Gaetan, R. A., Guzmán Rastelli, M. C., Carrera, M. F., Díaz, et al (2014). Biotechnology: Between cells, genes and human ingenuity. Ministry of Education of the Nation. Free book. Download from http://www.bnm.me.gov.ar/giga1/do cumentos/EL005063.pdf  Current articles provided by the teacher.  Electronic resources:  Google Scholar /google scholar ClinicalKey OMIM pages Web Pages: Induction, SDS-PAGE and biological activity.  You Tube Web Pages: Obtaining cDNA, Cloning
Evidence 5	Review the theoretical	The student individually creates the flow chart and	<ul> <li>PCR amplification of a fragment of the □-globin</li> </ul>	Computer equipment with Microsoft Office and internet connection.
Laboratory	foundations of the	the corresponding	gene.	
practice reports:	practice including	conceptual map.	<ul> <li>Verification of the</li> </ul>	Platforms:
	the material to	•	amplified product in	-Microsoft Teams
5.1	work in the laboratory and the	<ul> <li>A team previously designated by the teacher</li> </ul>	agarose gel.	-Microsoft Forms - Moodle





<b>Polymera</b>	se	cha	ain
reaction (	PC	R).	

5.2 DNA digestion with restriction enzymes

standards. Prepare the corresponding activities. Work virtually, organized and as a team on the proposed experiments. If the practice is carried out in person, the student will review the requirements to carry out the practice cleanly, in a responsible and orderly manner, complying with the safety and hygiene standards established for the correct use of the facilities and the correct disposal of waste. It is presented on the established date and time. The report is delivered on the established date

established safety

will make a Power Point presentation and explain the theoretical foundations of the practice,

- The teacher complements the explanation of the practice presented by the designated equipment and other audiovisual tools, asks questions and provides examples that reinforce the theoretical knowledge of the practice, indicates the important technical details for the adequate development of the experimental methodology to obtain results. optimal results.
- Students review the data, images and videos provided by the teacher as a team.
- The student prepares the practice report as a team following the established format, uploading it to the Moodle platform on the established date and time.

 Digestion of the amplified α-globin product with the restriction enzyme Bsu 36 I.

- Verification of the digestion of the amplified -globin product with the restriction enzyme Bsu 36 I.
- Analysis of results and usefulness of this test in the identification of the Glu6 →Val mutation.
- Molecular diagnosis of sickle cell anemia by identifying the Glu6 →Val mutation.

- Usefulness and importance of bioinformatics in biology and molecular diagnosis.
- Analysis of the α-globin gene sequences using bioinformatics tools.
- -In silico implementation of the molecular diagnosis of sickle cell anemia through the

**Power Point Presentations** 

Molecular Biology Laboratory Practice Manual. Digital, audiovisual or digital reading resources

5.3 Medical Bioinformatics.





	and time through the MS Teams or Moodle platforms in accordance with	identification of the Glu6  →Val mutation.  • • General review of the	
	the instructional	results obtained by the	
	guide and	group in all laboratory	
	checklist	practices. Guided tour of	
<i>5.4</i>		the department's	
Presentation,		molecular biology	
review and general		laboratories to learn	
discussion of the		about research	
results of the		applications of the	
laboratory		different techniques	
practices.		evaluated.	
Experience in a			
Molecular Biology			
laboratory			





# 7. Evaluation scheme of the Learning Unit broken down by Stages and Evidence of Learning:

Phase		Learning Challenge	Weighing
First phase (22%)	Ponderable activity 1	Comparative tables of the Central Dogma of Molecular Biology	2 %
,	Ponderable activity 2	Written work of the genetic code	2 %
	Evidencia 1	First partial evaluation (phase 1)	18 %
Second phase (31.5%)	Ponderable activity 3	Mental map of nucleic acid extraction techniques, electrophoresis, hybridization.	2 %
,	Ponderable activity 4	Comparative table of PCR techniques, sequencing, cloning, library preparation and hybridization techniques	2 %
	Evidencia 2	Second partial evaluation (phase 2).	19 %
	Evidencia 3 (3.1 a 3.5)	Laboratory (includes flow charts, concept maps, reports, presentations and class presentation of evidence 3.1 to 3.5)	9.4 %
Third phase	Ponderable activity 5	Table with examples of molecular diagnoses.	2 %
(26.5%)	Ponderable activity 6	Summary of the fundamentals and technical requirements to obtain a biotechnological product.	2 %
	Evidencia 4	Third partial evaluation (phase 3).	14 %
	Evidencia 5 (5.1 a 5.4)	Laboratory (includes flow charts, concept maps, reports, presentations and class presentation of evidence 5.1 to 5.4)	7.6 %
PIA	Integrative learning	Final Evaluation	10%
(20%)	product	Research Seminar on the molecular diagnosis of human diseases or any biotechnological application of biomedical interest	10%



10.

# UNIVERSIDAD AUTÓNOMA DE NUEVO LEÓN FACULTAD DE MEDICINA PROGRAMA ANALÍTICO



#### 8. Sources of support and consultation:

- 1. Abali Emine.E, *Bioquímica*. España: editorial Lippincott Williams &Wilkins. 8va. Edición.
- 2. AkademeiaUFM. (7 de enero de 2019). *Extracción y purificación de ADN*. [Archivo de video]. Recuperado de <a href="https://www.youtube.com/watch?v=a8d8ZNSX880">https://www.youtube.com/watch?v=a8d8ZNSX880</a>
- 3. <u>#BiotechReview #cDNA #mRNA</u>. cDNA Complementary DNA (2011). Recuperado de https://www.youtube.com/watch?v=rKPJpxCW2qw
- 4. Besme Allah. (12 de julio de 2017). *Pasos en la clonación de un gen*. [Archivo de video]. Recuperado de https://www.youtube.com/watch?v=lzBDO\_YFNW4
- 5. Brandon Ortiz Casas. (24 de agosto de 2018). *Electroforesis de ADN: Conceptos Básicos*. [Archivo de video]. Recuperado de https://www.youtube.com/watch?v=KGZBRfHQU Y
- 6. Brandon Ortiz Casas. (5 ene. 2018). Reacción en Cadena de la Polimerasa (PCR): Conceptos Básicos. [Archivo de video]. Recuperado de https://www.youtube.com/watch?v=msIMRqxbdOA
- 7. Brandon Ortiz Casas. (26 de febrero de 2018). Secuenciación Maxam-Gilbert: Conceptos Básicos. [Archivo de video]. Recuperado de <a href="https://www.youtube.com/watch?v=gP0uDYjA6jl">https://www.youtube.com/watch?v=gP0uDYjA6jl</a>
- 8. Brandon Ortiz Casas. (14 de abril de 2019). Secuenciación por Síntesis (Illumina): Conceptos Básicos. [Archivo de video]. Recuperado de https://www.youtube.com/watch?v=BimurK8vIYc
- 9. CanalDivulgación. (2 de enero de 2014). *PCR: Reacción en Cadena de la Polimerasa (divulgación científica IQOG-CSIC)*. [Archivo de video]. Recuperado de <a href="https://www.youtube.com/watch?v=TalHTjA5gKU">https://www.youtube.com/watch?v=TalHTjA5gKU</a>
- 11. Colleman W.B, Tsongalis G. J., Molecular Diagnostics For the Clinical Laboratorian.
- 12. DENISSE RODRIGUEZ ALVAREZ. (6 de junio de 2019). *DNA Microarrays*. [Archivo de video]. Recuperado de <a href="https://www.youtube.com/watch?v=hJMdso9Salo">https://www.youtube.com/watch?v=hJMdso9Salo</a>
- 13. Devlin, T. M. (2004). Bioquímica. España: editorial Reverté, S.A.
- 14. <u>EMAbiolog</u>. Clonación de un gen en un plásmido vector (2013). Recuperado de https://www.youtube.com/watch?v=mLd4WdQHeSM





- 15. GeneCards. Department of Molecular Genetics at the Weizmann Institute of Science. (2020). The Human Gene Database. Rehovot, Israel.: HBB Gene (Protein Coding) Hemoglobin Subunit Beta. <a href="https://www.genecards.org/cgi-bin/carddisp.pl?gene=HBB&keywords=sickle,cell#diseases">https://www.genecards.org/cgi-bin/carddisp.pl?gene=HBB&keywords=sickle,cell#diseases</a>
- 16.
- 17.
- 18. Martínez-Rodríguez, H.G. y cols. (2017). Manual de Laboratorio de Bioquímica. México: Facultad de Medicina, UANL.
- 19. Martin K. Gold Bio. A Deep Dive Into Induction with IPTG. Recuperado de <a href="https://www.goldbio.com/articles/article/a-deep-dive-into-iptg-induction">https://www.goldbio.com/articles/article/a-deep-dive-into-iptg-induction</a>
- 28. *MalaCards: The human disease database.* Department of Molecular Genetics at the Weizmann Institute of Science. (2020). Rehovot, Israel.: Sickle Cell Anemia (SKCA). Recuperado de <a href="https://www.malacards.org/card/sickle\_cell\_anemia?search=sickle%20cell%20anemia">https://www.malacards.org/card/sickle\_cell\_anemia?search=sickle%20cell%20anemia</a>
- 29. McKee, T. y McKee, J. (2014). Bioquímica las bases moleculares de la vida. 5ª edición. España: editorial McGraw-Hill.
- 30. Medicurio.(22 Dic 2016). <u>Duchenne Muscular Dystrophy and Dystrophin.Recuperado de https://www.youtube.com/watch?v=Ebu8W8Osuxk&t=32s</u>
- 31. Menor-Salvan C. <u>ChemEvol</u>. SDS-PAGE: ELECTROFORESIS EN GEL DE POLIACRILAMIDA (2019). Recuperado de <a href="http://www3.uah.es/chemevol/index.php/sds-page-electroforesis-en-gel-de-poliacrilamida/">http://www3.uah.es/chemevol/index.php/sds-page-electroforesis-en-gel-de-poliacrilamida/</a> 32.
- 33. NCBI. National Center for Biotechnology Information. http://www.ncbi.nlm.nih.gov/Omim/searchomim.html
- 34. Nussbaum R. L. Genetics in medicine. (2008). eBook Recuperado de http://www2.genoma.ib.usp.br/disciplinas/bio416/NGS-selecao\_thompson.pdf
- 35. OMIM. McKusick-Nathans Institute of Genetic Medicine, Johns Hopkins University School of Medicine. (2020). *OMIM- Online Mendelian Inheritance in Man.* Baltimore, MA, EU: #310200 MUSCULAR DYSTROPHY, DUCHENNE TYPE; DMD. Recuperado de: https://omim.org/entry/310200#molecularGenetics
- 36. OMIM. McKusick-Nathans Institute of Genetic Medicine, Johns Hopkins University School of Medicine. (2020). OMIM- Online Mendelian Inheritance in Man. Baltimore, MA, EU.# 603903 SICKLE CELL ANEMIA. Recuperado de <a href="https://omim.org/entry/603903?search=sickle%20cell%20anemia&highlight=%28anaemia%7Canemia%29%20cell%20sickle">https://omim.org/entry/603903?search=sickle%20cell%20anemia&highlight=%28anaemia%7Canemia%29%20cell%20sickle</a>
- 40.
- 42. Virtual Cell. Regulación: Operón Lac Recuperado de <a href="https://www.youtube.com/watch?v=oBwtxdl1zvk">https://www.youtube.com/watch?v=oBwtxdl1zvk</a> Virtual Cell (2008). Traducción. <a href="https://www.youtube.com/watch?v=5bLEDd-PSTQ&feature=youtu.be">https://www.youtube.com/watch?v=5bLEDd-PSTQ&feature=youtu.be</a>
- 43. Virtual Cell (2008).Transcripción. Proceso. Recuperado de <a href="https://www.youtube.com/watch?v=WsofH466lqk&feature=youtu.TRANSCRICPIAÓNbe">https://www.youtube.com/watch?v=WsofH466lqk&feature=youtu.TRANSCRICPIAÓNbe</a>





- 44. VirtualCell (2008).Transcripción. Adición de CAP y poli A. Recuperado de <a href="https://www.youtube.com/watch?v=eM\_8NUUg9YQ">https://www.youtube.com/watch?v=eM\_8NUUg9YQ</a>
- 45. VirtualCell (2008).Transcripción Remoción de intrones. Recuperado de <a href="https://www.youtube.com/watch?v=6ojQYMC7\_2A">https://www.youtube.com/watch?v=6ojQYMC7\_2A</a>
- 46. Watson, J.D., Baker, T.A., Bell, S.P., Gann, A., Levine, M. y Losick, R. (2006). Biología Molecular del Gen. España: editorial Médica Panamericana.
- 47.Watson, J. D., Myers, R. M., Caudy, A. A., Witkowski, J.A. (2006). Recombinant DNA: Genes and Genomes A Short Course. EUA: editorial Scientific American Books