



1. Module identification code.		
Name of the institution:	Universidad Autónoma de Nuevo León	
Name of the school:	School of Medicine	
Name of the degree program:	Clinical Chemistry	
Name of the course (learning unit):	Basic Microbiology	
Total number of class hours-theory and practice:	80	
Class hours per week:	4 hours	
Independent study:	10	
Course modality:	Face-to-face instruction	
Module level:	Second semester	
Core/elective module:	Core	
Curriculum area:	ACFB	
UANL credit points:	3	
Create date:	August 28 th , 2017	
Date of last amendment made:	January 19 th , 2022	
Person(s) responsible for the design and amendment of the module:	Dr. C Miguel Ángel Becerril García	





2. Presentation:

The module (learning unit) of basic Microbiology consists of 4 phases.

In phase 1, the fundamentals of biochemistry will be revised through which the students will learn to differentiate biomolecules and to distinguish metabolic processes, this basis will let the student understand the structural and physiological characteristics of microorganisms as well as of infectious agents. This is distributed in the following phases:

During phase 2, microbiology will be examined from its historical point of view that will let the student provide the basis for the effects it has on current practice. He will classify microorganisms according to environmental factors and nutritional requirements that influence their growth.

He will differentiate genetic characteristics of microorganisms and will relate them to their virulence.

He will describe the characteristics of saprophyte, commensal and pathogen microorganisms in order to relate them to the human being as a host.

Finally, he will identify the use of physical and chemical agents in the control of microbial growth.

In phase 3, he will distinguish the morphology, growth, genetics and ecology of viruses, bacteria, fungi, protozoa and helminths. As a course integrative project/product (CIP), he will make a written report about a clinical case study assigned by the professor. In this report, he will provide an adequate methodological and conceptual strategy that will let him differentiate groups of microorganisms that are causing an illness.

3. Purpose:

The following module (LU) contributes to the graduate profile because it develops the necessary competences to select lab tests and provide the basis for such a choice that will let the student identify microorganisms through the analysis of the general concepts of their morphology, physiology, genetics and ecology, their impact on the environment, in human activity and their implication on the production of illnesses in the human being. All this, carried out under strict regulations of quality control that will let him make adequate and appropriate decisions to collaborate in the prevention, diagnosis, control and treatment of illnesses.

With regard to the general competences, the student will be able to apply autonomous learning strategies so that he can make adequate and appropriate decisions during sample analysis. This will also let him make a microbiological identification with accuracy and quality. Besides, he will show ethics, truth, honesty, and respect towards his classmates and professors in the classroom as well as in the laboratory.

He shows empathy when dealing with conflicts during team work in the classroom and in the laboratory, in such a way that he respects the ideas of his classmates and reaches agreements when facing different points of view during team work and in the practice laboratory.

In the LU of basic Microbiology, the student acquires competences that let him have the necessary basis for the collection and handling of samples for microbiological analysis.

Also, he handles chemical and biological materials according to national and international regulations in order to protect health and the environment.

The content of this module (learning unit) is interrelated with Cellular Biology because it describes the components and functions of cells both prokaryotes and eukaryotes; on the other hand, it provides the general aspects of the microorganisms that will be useful for their approach in depth in the modules (learning units) of Medical parasitology; Mycology and Virology; Immunology, Medical bacteriology, that contribute to the understanding of the infectious process and the behavior of microorganisms.





4. Competences of the graduate profile

General competences to which this module (learning unit) contributes:

Instrumental skills:

1. To apply autonomous learning strategies at different levels and fields of knowledge that allow them to make timely and relevant decisions in the personal, academic and professional spheres.

Personal and social interaction skills:

11. To practice the values promoted by the UANL: truth, equity, honesty, freedom, solidarity, respect for life and others, peace, respect for nature, integrity, ethical behavior and justice, in their personal and professional environment to contribute to building a sustainable society

Integrative skills:

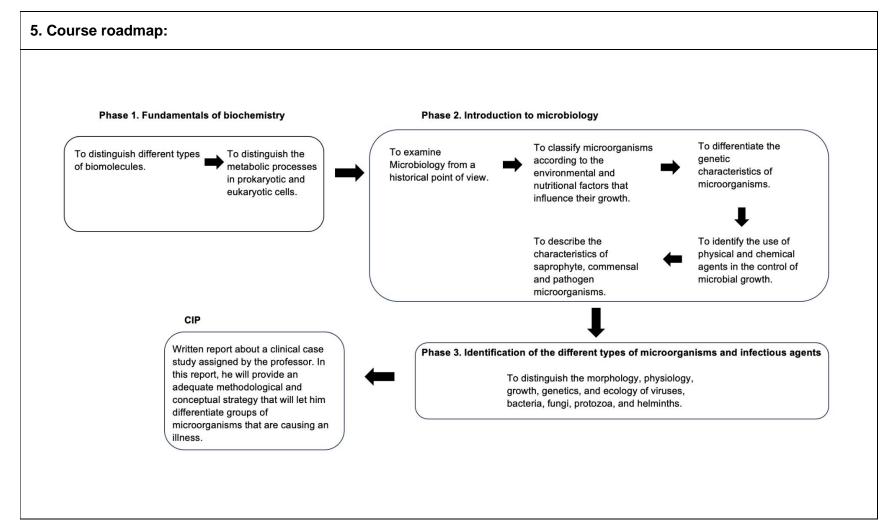
14. To resolve personal and social conflicts, in accordance with specific techniques in the academic field and in their profession for appropriate decision-making.

Specific competences of the graduate profile to which this module (learning unit) contributes:

- 2. To execute physical, chemical and/or biological procedures in the collection, handling, storage and analysis of samples to contribute to a reliable clinical, toxicological, chemical, food, forensic and environmental diagnosis.
- 3. To handle chemical and biological materials following official Mexican and/or international standards that guarantee their correct use and disposal to preserve health and the environment.











6. Structuring into stages or phases:

Stage 1: Fundamentals of Biochemistry.

Component(s) of the competence: To identify the biochemical basis as well as the metabolism of microorganisms and infectious agents that let the student enter into the study of microbiology.

Evidence of student learning	Performance criteria	Learning activities	Content	Resources
1.Diagram about biomolecules and cellular metabolism.	complete identification data. • Content: Includes the metabolism of carbohydrates, proteins and lipids. Highlights the critical aspects of the topic. • Synthesis: Uses diagrams only.	asking questions to the students. Accredited activity No. 1 Practice. Demonstration of the material used in the Microbiology laboratory. Cleaning and sterilization of material used in the Microbiology laboratory. (accredited activity).	1.2 pH and water 1.3 Carbohydrates 1.4 Lipids 1.5 Amino acids 1.6 Proteins 1.7 Enzymes 1.8 Nucleic acids 2. Cellular metabolism: 2.1 Metabolism, Anabolism, and Catabolism 2.2 Carbohydrate metabolism 2.3 Glycogenesis	Textbooks: Ferrier. 2017. Chapters: 1-19. Koolman. 2012. Chapters 2-3. Murray. 2013. Chapters 2-26. Alonso 2017. Chapter 1. Computer Projector Practice Laboratory Microscopes Laboratory glassware Reagents and lab disposables Diary Pen Pencil.





Stage 2: Introduction to Microbiology Component(s) of the competence:

To examine the microbiological agents from a genetic, nutritional, environmental and historical point of view; and their relation to the host; and their control using chemical and physical agents; in order to provide the theoretical basis about their effect in humans.

Evidence of student learning	Performance criteria	Learning activities	Content	Resources
2. Diagram about	Presentation:	-The student will read in	3. History of	Jawettz.2016.
historical events and	Turns in a diagram: six	advance the topics assigned.	Microbiology	Chapters 1-10.
factors that influence	sheets maximum,		3.1 Definition of	Prescott. 2009
the growth of	handwritten, with an	- The professor will present	Microbiology/Microorganis	Chapters 1-7, 11-13.
microorganisms,	additional cover page, and	the topics:	m.	Alonso. 2017.
•	complete identification	History of Microbiology,	3.2 Field of study.	Chapters 2-9.
genetics, microbiota	data.	environmental factors that	3.3 Discovery of	Computer
and antimicrobial	 Content: Includes 	influence microbial growth,	microorganisms.	Projector
agents.	the most relevant historical	classification of	3.4 Theory of spontaneous	Practice Laboratory
	processes	microorganisms, nutritional	generation.	Microscopes
	of microbiology;	factors for microorganism	3.5 Theories of Microbial	Laboratory glassware
	environmental and	growth and classification,	Evolution.	Reagents and lab
	nutritional factors that	microbial genetics. normal	3.6 Microbial Diversity.	disposables
	affect microbial growth;	microbiota, and antimicrobial	Prokaryotes/Eukaryotes.	Diary
	genetic characteristics of	agents.		Pen
	microorganisms;		4. Environmental factors	Pencil
	microbiota and its	- The professor will conduct a	that influence microbial	Web sites:
	relationship to the host;	discussion of the topics by	growth:	World Health Organization:
	and the impact of	asking questions to the	4.1 Microbial growth.	http://www.who.int/en
	physical and chemical	students.	4.2 Environmental factors	Centers for control and
	agents on microbial	Accredited activity No 2:	that influence microbial	disease prevention
	development.	Practice. Aseptic technique	growth: oxygen	http://www.cdc.gov/
	 Synthesis: Uses diagrams 	used to handle test tubes.	consumption, pH,	
	only.	Using the bacteriological	temperature and water.	Dirección general de
	 References: Uses three 	handle-rod.	4.3 Classification of	epidemiología,
	or more sources of		microorganisms in terms of	México:

Accredited activity No 3: oxygen consumption: https://www.gob.mx/s information. Planning: aerobious, anaerobious. alud/acciones-yprogramas/ Practice. Demonstration of the aerotolerant and direcciongeneral-Turns in the evidence deepidemiologia presence of microorganisms facultative. at the time and in the in nature. day programmed. 4.4 Classification in terms Accredited activity No. 4 of pH: acidophiles, Effect of molecular oxygen on neutrophiles and alkaliphiles. bacteria growth. Accredited activity No. 5 4.5 Classification of Practice. Preparation of microorganisms in terms of culture media. the temperature they require to grow: psychrophiles, mesophiles. Accredited activity No. 6 thermophiles and Practice. Culture of hyperthermophiles. microorganisms: the transfer of bacteria 5.- Nutritional factors for microorganism growth from one test tube to another. and classification: Accredited activity No. 7: 5.1 Nutritional factors 1a Written evaluation of the necessary to carry out content 1-5 metabolism: Macroelements (C, O, Accredited activity No. 8 Practice. Determination of the H, N, S and P) from carbohydatres, lipids, number of microorganisms proteins and nucleic acids. from a culture. ions (Mg2+, Ca2+, Fe2+ and Fe3+); and Accredited activity No. 9 Micronutrients (Mn, Zn, Co, Practice. Transmission of oral Mo, Ni and Cu). bacteria during verbal 5.2 Classification of communication microorganisms in terms of their carbon source: Accredited activity No. 10 autotroph, heterotrophs; in Practice. Sterilization by terms of energy source: means of physical agents. chemotroph and Sterilization by dry heat

(furnace) and humid heat phototroph; and of electron under pressure (autoclave). source: lithotroph and Effect of radiation on organotroph. microorganisms. 6. Microbial genetics: Accredited activity No. 11 6.1. Comparison of the Practice. Chemical agents: genetic structure of: Sterilization of medical viruses, bacteria, instruments. Skin antisepsis. yeasts, fungi, protozoa, and helminths. Accredited activity No. 12 6.2. Definition: Operon. 2ª Written evaluation about 6.3 Mechanisms of genetic content 6-8 variation: mutation and recombination. 6.4. Mechanisms of genetic recombination of the microorganisms: transformation, transduction. conjugation, and their relation to pathogenicity. 7.- Normal microbiota: 7.1Classification of microorganisms related to humans: saprophytes, commensals y pathogens. 7.2 Microbiota 7.3 Location of microbiota in the organism. 7.4 Definition of: Host, vector, saprophyte, commensal and pathogen. 8.- Antimicrobial agents:

8.1 Concept of microbial life and death. 8.2 Microbial control and its relation to microbial death. 8.3 Classification of microbial control based on its nature: physical agents and chemical agents. 8.4 Definition of: sepsis, asepsis, antisepsis, sanitation, disinfectant, antiseptic, germicide, bacteriostat, bactericidal, fungicidal, sterile, sterile, sterile, sterilezation, and antimicrobial. 8.5 Classification of the physical agents for	
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sterilization, and antimicrobial. 8.5 Classification of the physical agents for	ļ
antimicrobial. 8.5 Classification of the physical agents for	ļ
8.5 Classification of the physical agents for	
physical agents for	
microbial control: Dry heat,	
humid heat under	
pressure, humid heat	
without pressure and,	
ionizing radiation and non-	
ionizing radiation.	
8.6 Mechanisms of action	
of physical agents.	
8.7 Classification of	
chemical agents for	
microbial control:	
disinfectant, antiseptic, and	
antibiotic.	
8.8 Mechanisms and sites	
of action of disinfectant	
	li di
and/or antiseptic on the	İ
microbial cell.	

8.9 Tests to assess the effectiveness of a chemical agent.
8.10 Sites of action of antimicrobial agents within the microbial cell.
8.11 Definition of selective and nonselective toxicity. 8.12 Definition of susceptible and resistant.
8.13 Test to determine antimicrobial susceptibility.





Stage 3. Identification of the different types of microorganisms and infectious agents. Component(s) of the competence:

To identify the microorganisms by means of their morphology, physiology, growth, genetics and ecology of bacteria, viruses, fungi, yeast, protozoa, and helminths that have clinical importance in order to provide the basis for the selection of the tests that let the student recognize them in the laboratory.

Evidence of student	Performance criteria	Learning activities	Content	Resources
learning				
3 Comparison chart	Presentation:		9 Bacteria that have	Jawettz.2016
about helminths,	Turns in chart: four	-The student will read in	clinical importance:	Chapters 11-48.
protozoa, yeast, fungi,	sheets maximum,	advance the topics		Prescott. 2009.
viruses, and bacteria.	handwritten, with an	assigned.	9.1 Classification	Chapters 16-23.
	additional cover page,		according to growth,	Brock. 2009
	and complete	Accredited activity No.	physiological, and	Chapters 25-28.
	identification data.	13	morphological	Alonso. 2017
	Content:	Students will give an	characteristics; and of	Chapters 9-15.
	Includes the	oral presentation in	the mechanisms of	Computer
	comparison among	teams about the	transmission:	Projector
	helminths, protozoa,	following topics: viruses,	Gram-positive (aerobic	Practice laboratory
	yeast, fungi, viruses and	bacteria, fungi,	and anaerobic),	Microscopes
	bacteria.	protozoa, and helminths	Gram-negative (aerobic	Laboratory glassware
	Highlights the specific	that have clinical	and anaerobic),	Reagents and lab
	characteristics of each	importance.	Mycobacterias,	disposables
	group of		Actinomycetes,	Diary
	microorganisms.	The oral presentation	Spirochetes,	Pen
	Synthesis: Uses charts	should include the	Mycoplasm, obligate.	Pencil.
	only.	following sequence:	Intracelular bacteria:	Web sites:
	References:	structure. classification,	Rickettsia and	World Health
	Uses three or more	physiology,	Chlamydia.	Organization:
	sources of information.	pathogenicity factors,		http://www.who.int/en
	Planning:	lab tests used in the	9.2 Virulence factors	/
	Turns in the evidence at	diagnosis.	and bacterial	Centros de control y
	the time and in the day		pathogens.	prevención de
	programmed.	- The professor will		enfermedades
		conduct a discussion of	9.3 Diagnosis in the	infecciosas EEUU:
		the topics by asking	laboratory of the	http://www.cdc.gov/

questions to the illnesses caused by Dirección General de epidemiología, students. bacteria. México: Accredited activity No. 10.- Virus that have https://www.gob.mx/s clinical importance: alud/acciones-14 Practice. Separation of yprogramas/ two bacterial species. 10.1 Classification of direcciongeneralviruses in terms of the deepidemiologia Accredited activity No. nature of their genome: 15 DNA and RNA. Practice. Simple staining. Preparation of 10.2 Morphology and identification of the extensions on microscope slides. structural components Differential staining. of viruses. Gram's staining technique 1% 10.3 Classification of viruses in terms of their Accredited activity No. function of the presence 16 or absence of Practice. Bacterial packaging: naked and morphology. enveloped. Demonstration of the 10.4 Viral replication bacterial capsule. Bacterial mobility. cycle. Accredited activity No. 10.5 Pathogenicity 17 factors. Practice. General tests to differentiate bacterial 10.6 Cytopathological effects and the relation genders. to the tests used in their Accredited activity No. identification. 18. 3ª Written 11. Prions evaluation about the content 9-12 12.- Fungi that have Accredited activity No. clinical importance: 19. Practice. Microscopic 12.1Morphology observation of fungi and

	Protozoa and 12.2Pathogenicity	
helm	nths. factors.	
Accr	edited activity No. 12.3 Physiology of	
	Written growth.	
	nation about the	
	ent 13 and 12.4 Relationship of the	
14.	factors of pathogenicity	
	with the entryway into	
	the system; and the	
	mechanism of the	
	production of illnesses.	
	12.5 Tests used in their	
	identification.	
	13 Protozoa that	
	have clinical	
	importance:	
	13.1 Classification of	
	protozoa in terms of	
	their mobility and way of	
	phagocyting.	
	13.2 Morphology and	
	microscopic	
	identification	
	13.3 Mechanism of	
	transmission, location in	
	humans, parasitic	
	phase, diagnosing	
	phase.	
	μπασο.	
	13.4 Microscopio	
	13.4 Microscopic	
	identification.	
	40 F Tacksisses seed	
	13.5 Techniques used	
	in the identification of	
	protozoa.	

13.6. Species of clinical importance in each of their types: Sarcodina, Mastigophora, Ciliophora, Apicomplexa (Sporozoea)
14 Helminths that have clinical importance:
14.1Classification in terms of their adult form: nematodes, trematodes, and cestodes.
14.2 Species of clinical importance.
14.3 Mechanism of transmission.
14.4 Parasitic phase, Diagnosing phase.
14.5 Microscopic identification.
14.6 Test used in their diagnosis.





7. Summative evaluation:

Phase	Evidence	Weighing
	Evidence 1. Diagram about biomolecules and cellular metabolism	5%
Phase 1	Accredited activity 1. Practice. Demonstration of the material used in the Microbiology laboratory.	1%
	Evidence 2. Diagram about the historical events and factors that influence on the growth of microorganisms, genetics, microbiota and antimicrobial agents.	10%
	Accredited activity 2. Practice. Aseptic technique used to handle test tubes. Using the bacteriological handle-rod.	1%
	Accredited activity 3. Practice. Demonstration of the presence of microorganisms in nature.	1%
	Accredited activity No 3: Practice. Demonstration of the presence of microorganisms in nature.	1%
	Accredited activity No. 4 Effect of molecular oxygen on bacteria growth.	1%
	Accredited activity No. 5 Practice. Preparation of culture media.	1%
Phase 2	Accredited activity No. 6 Practice. Culture of microorganisms: the transfer of bacteria from one test tube to another.	1%
Phase 2	Accredited activity No. 7: 1 ^a Written evaluation about content 1-5	5%
	Accredited activity No. 8 Practice. Determination of the number of microorganisms from a culture.	1%
	Accredited activity No. 9 Practice. Transmission of oral bacteria during verbal communication.	1%
	Accredited activity No. 10 Practice. Sterilization by means of physical agents. Sterilization by dry heat (furnace) and humid heat under pressure (autoclave). Effect of radiation on microorganisms.	1%
	Accredited activity No. 11 Practice. Chemical agents: Sterilization of medical instruments. Skin antisepsis.	1%

	Accredited activity No. 12 2nd Written evaluation about content 6-8 5%	5%
	Evidence No 3 : Comparison chart among helminths, protozoa, yeast, fungi, viruses and bacteria.	10%
	Accredited activity No. 13 Students will give an oral presentation in teams about the following topics: Viruses, bacteria, fungi, protozoa, and helminths that have clinical importance.	5%
	Accredited activity No. 14 Practice. Separation of two bacterial species.	1%
	Accredited activity No. 15 Practice. Simple staining. Preparation of extensions on microscope slides. Differential staining. Gram's staining technique	1%
Phase 3	Accredited activity No. 16 Practice. Bacterial morphology. Demonstration of the bacterial capsule. Bacterial mobility.	1%
	Accredited activity No. 17 Practice. General tests to differentiate bacterial genders	1%
	Accredited activity No. 18. 3 ^a Written evaluation about content 9-12	5%
	Accredited activity No. 19. Practice. Microscopic observation of fungi and yeast. Protozoa and helminths.	1%
	Accredited activity No. 20. 4 ^a Written evaluation about content 13-14	5%
CIP	Course integrative project/product	35%
	Total	100

8. Course integrative project/product:

Written report about a clinical case study assigned by the professor. In this report, he will provide an adequate methodological and conceptual strategy that will let him differentiate groups of microorganisms that are causing an illness.



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