Biology Department, College of Arts & Sciences, Valdosta State University SPRING 2011----COURSE SYLLABUS*

<u>BIOL 3100, Sections A & B. Microbiology</u> (CRN 21202 & 21203) - 4 credit hours <u>BIOL 5100, Sections A & B. Microbiology</u> (CRN 21231& 21232) - 4 credit hours**

Class: TR 8:00-9:15 am, 2022 Bailey Science Center

Laboratory: TR 3100/5100 Section A 10:00-11:25 am, 2068 Bailey Science Center

TR 3100/5100 Section B 2:00-3:25 pm, 2068 Bailey Science Center

<u>Instructor:</u> Dr. Jenifer Turco Email: <u>jturco@valdosta.edu</u>

Telephone: 229-249-4845 Office: 2091 Bailey Science Center

Office Hours: Tues. 4:30-5:30 pm & Thurs. 12:30-1:30 pm; or by appointment.

Course Description:

BIOL 3100 Microbiology 3-3-4 (4 credit hours)

Prerequisites: BIOL 2010, BIOL 2230, BIOL 2270, CHEM 1212/1212L. Recommended: CHEM 3402.

BIOL 5100 Microbiology 3-3-4 (4 credit hours)

Prerequisite: Admission into the graduate program or permission of the instructor. Survey of microbiology covering eubacteria, archaebacteria, protozoa, fungi, algae, and viruses. Includes fundamental techniques, microbial physiology and genetics, biotechnology, medical applications, and applied microbiology. Two 1.5 hour laboratory periods per week.

Required Textbook: BROCK BIOLOGY OF MICROORGANISMS, Twelfth Edition

by Michael T. Madigan, John M. Martinko, Paul V. Dunlap, and David P. Clark

Prentice Hall 2009

Required Lab Manual: BENSON'S MICROBIOLOGICAL APPLICATIONS, LABORATORY MANUAL

IN GENERAL MICROBIOLOGY (Complete Version), Eleventh Edition

by Alfred E. Brown McGraw-Hill, Inc. 2009

Other Required Items: A notebook for recording the results of laboratory work

Calculator that is not integrated with a cell phone

Permanent, fine-tip marking pen ("Sharpie") for labeling cultures in lab

One CD (or jump drive) for oral presentation

One thin, light-weight folder for handing in references & other assignments (Please do not

use a 3-ring binder to hand in assignments.)

Paper clips or stapler/staples for organizing references & assignments

Special notes to students:

- 1. In order to respect the privacy of each student, exam scores and grades will not be posted, given out by telephone, or sent to students by email.
- 2. Students should consult the VSU Student Handbook, Catalog, Semester Calendar, Schedule of Classes, & Registration Guide for information about VSU policies and procedures regarding registration, drop/add, and withdrawal. March 3 is midterm. Students are not permitted to withdraw after midterm except in cases of hardship.
- 3. Students requesting classroom accommodations or modifications because of a documented disability should discuss this need with the instructor at the beginning of the semester. These students must contact the Access Office for Students with Disabilities located in Farber Hall. The phone numbers are 245-2498 (V/VP) and 219-1348 (TTY).
- 4. Cell phones may not be used during examinations or at any time in class or lab.
- 5. Students must read and follow the Biology Department policy on plagiarism (available online through the departmental web site). The instructor may use a variety of methods for detecting plagiarism. Each student must be particularly careful to do his/her own writing on the oral presentations and on any assignments/reports that are to be completed individually. Plagiarism will result in a grade of "0" for the assignment. A student who plagiarizes on more than one assignment will receive a grade of "F" in the course.
- 6. No disruptive behavior will be tolerated during class or lab. A student who engages in disruptive behavior will be asked to leave.
- 7. Students who wish to use laptop computers as part of the class are required to sit in the first three rows of the classroom.

*This is a tentative syllabus. Changes to this syllabus will be announced during class or laboratory periods; alternatively, changes may be posted on BlazeView.

**Graduate students who are taking BIOL 5100 must meet with the instructor to discuss additional course requirements & grading.

Course Objectives:

(Page 3 shows how the objectives below are aligned with the VSU General Education Outcomes and the Biology Department Education Outcomes.)

After successful completion of this course, the student should be able to:

- A. List and describe the three domains of living organisms.
- B. List and describe the three types of noncellular infectious agents.
- C. List several activities of microorganisms that are beneficial to humans and the environment.
- D. List and briefly explain several current challenges in medical microbiology and infectious diseases.
- E. Compare and contrast the structure and function of the microorganisms in the domains Bacteria, Archaea, and Eukarya.
- F. List and describe the various strageties used by microorganisms to obtain carbon, energy, and electrons.
- G. Describe the growth of a pure culture of bacteria in a closed system, and perform mathematical calculations related to the exponential growth phase. Explain several ways in which bacterial growth can be measured.
- H. Compare and contrast the following processes as they occur in Bacteria, Archaea, and Eukarya: DNA replication, transcription, and translation.
- I. Describe several mechanisms through which gene expression is regulated in bacteria.
- J. Explain how viruses replicate, and explain in detail how HIV enters and replicates within host cells.
- K. Describe the causes and consequences of mutations.
- L. Describe the three mechanisms of horizontal gene transfer in bacteria, and explain their significance.
- M. Describe specific examples of the use of microorganisms in genetic engineering and biotechnology.
- N. Briefly explain the role of microorganisms in the evolutionary history of life on earth.
- O. List and describe a variety of methods and approaches that are used to detect and identify various microorganisms and noncellular infectious agents.
- P. Explain how physical methods and chemical agents (antiseptics and disinfectants) are used for controlling microbes.
- Q. State the mechanisms of action of various antibacterial, antifungal, and antiviral medications.
- R. Discuss the problem of antimicrobial drug resistance, and explain several ways in which the emergence of drug resistant bacteria can be minimized.
- S. Give examples of beneficial interactions between: (i) microorganisms and plants, (ii) microorganisms and animals, and (iii) different types of microorganisms.
- T. Describe the role of microorganisms in the cycling of nutrients, using examples from the carbon cycle, the nitrogen cycle, and the sulfur cycle.
- U. Describe in detail: (i) the innate defenses of humans and (ii) the adaptive immune response of a human to a foreign antigen.
- V. Explain how infectious diseases are transmitted, giving specific examples.
- W. List the major types of virulence factors observed in pathogenic bacteria, giving specific, detailed examples.
- X. List and describe several human diseases that are due to specific bacteria, viruses, and fungi.
- Y. Describe the general course of the disease caused by human immunodeficiency virus (HIV).
- Z. Properly handle microorganisms in a biosafety level 2 laboratory.
- ZA. Use a compound light microscope to examine various types of microorganisms.
- ZB. Keep accurate records of microscopic observations, as well as other laboratory and field work.
- ZC. Use culture media to grow bacteria and fungi in the laboratory, and maintain stock cultures.
- ZD. Use staining techniques, physiological tests, and rRNA sequences as aids in bacterial identification.
- ZE. Use dilutions to determine the colony-forming units per milliliter in a bacterial suspension and the plaque-forming units per milliliter in a viral suspension.
- ZF. Work with others to formulate an answerable question, develop a hypothesis, design and conduct an experiment, collect and organize data, and write a formal report in the format used in a scientific journal.
- ZG. Use library and electronic resources to obtain formal scientific articles related to a particular topic in microbiology.
- ZH. Read the articles mentioned in objective ZG and give an oral presentation based on them.

Alignment of Assignments with Course Objectives:

The course objective(s) aligned with each assignment are given on page 11.

$\underline{\textbf{Alignment of Course Objectives with VSU General Education Outcomes \& Biology Department Educational Outcomes:}\\$

The VSU General Education Outcomes (1-8) are available online at

 $\underline{http://www.valdosta.edu/academic/VSUGeneralEducationOutcomes.shtml}$

In this syllabus, the VSU General Educational Outcomes are designated as: VSU1-VSU8

The Biology Department Educational Outcomes (1-5) are in the Undergraduate Catalog on page 116. The catalog is available online through http://www.valdosta.edu. In this syllabus the Biology Department Educational Outcomes are designated as: B1-B5.

The course objectives that are aligned with the VSU General Education Outcomes and Biology Department Educational Outcomes are given below:

VSU or Biology Objective	Course Objective(s)
VSU1	D, R, V, X, Y
VSU2	D, R, V, X, Y
VSU3	ZF, ZG, ZH
VSU4	ZB, ZF, ZH
VSU5	all course objectives
VSU6	
VSU7	G, ZA, ZE, ZF, ZH
VSU8	
B1	Z, ZA, ZB, ZC, ZD, ZE, ZF, ZG, ZH
B2	A, B, D, H, J, K, L, N, O, R, X, Y
B3	A, D, E, F, G, H, I, O, P, Q, U, W, X, Y
B4	D, H, I, J, K, L, M, O, X, Y
B5	C, D, S, T, V

BIOLOGY 3100/5100. Microbiology - Class and Lab Schedule

Date		Topics/Lab Exercises (Additional notes for lab exercises)	Related material in text
Tues.	Jan. 11	General course information Microorganisms and microbiology	Chap. 1
Tues.	Jan. 11L	>Program #1, The Microbial Universe SUPPL. EX., HANDWASHING Wash your hands before leaving lab!	
Thurs.	Jan. 13	Microorganisms and microbiology An overview of microbial life Cell structure/function	Chap. 1 Chap. 2 Chap. 4
		Review the following topics on your own: Basics of chemistry and biochemistry; DNA structure & replication; transcription; translation	Chap. 3 Chap. 7
Thurs.	Jan. 13L	>EX. 19, CULTURE MEDIA PREPARATION (We will preparing nutrient broth and nutrient agar.) After able to explain how nutrient broth, nutrient agar Complete questions, p. 139-140. >PLEASE READ THE FOLLOWING BEFORE NEXT LABORATORY SAFETY (Read handout & p. ix-xv ex. 9, ASEPTIC TECHNIQUE SUPPL. EX., WINOGRADSKY COLUMN; EX. 54 WIN PAGES 654-657 IN THE TEXTBOOK.	er completing this lab, students should be plates, and nutrient agar slants are prepared. WEEK: in lab manual.)
Tues.	Jan. 18	Cell structure/function	Chap. 4
Tues.	Jan. 18L	>LAB ORIENTATION >LABORATORY SAFETY >EX. 9, ASEPTIC TECHNIQUE Wash your hands before leaving lab! PLEASE REMEMBER TO READ THE INFORMATION F	FOR EACH DAY'S LAB BEFORE COMING TO LAB

CONTINUED ON PAGE 4.....

Discuss the Winogradsky Column Project with your lab group. Decide on a question, formulate a hypothesis, and decide how you will conduct the experiment. Decide on your assignments for the Winogradsky Column Project, and bring any required materials to lab on Thursday, Jan. 27. Each group of 4 students will build <u>at least</u> two columns.

Date		Topics/Lab Exercises	Related material in text
Tues.	Jan. 18L	CONTINUED FROM PAGE 3	

- YOUR GROUP'S LAB REPORT ON THE WINOGRADSKY COLUMN PROJECT (DUE APRIL 7) must be written in the style of a scientific paper and must contain the following sections: <u>Title</u>, <u>Authors</u>, <u>Abstract</u>, <u>Introduction</u>, <u>Materials and Methods</u>, <u>Results</u>, <u>Discussion</u>, <u>Literature Cited</u>, <u>and an Appendix</u>. The Results section must include your group's organized data and observations on the Winogradsky columns, charts and/or graphs, selected drawings (or photographs), and a written description of the results.
- The Appendix must contain each lab group member's <u>original</u>, written notes and drawings (or photographs) for the project. <u>Each group member's work must be labeled with his or her name.</u>
- The overall format for the report must follow the "Instructions for Authors" for the Journal of Bacteriology (available online at http://jb.asm.org/misc/ifora.shtml)

	online at <u>http</u>	://jb.asm.org/misc/ifora.shtml)	
Thurs.	Jan. 20	Cell structure/function	Chap. 4
Thurs.	Jan. 20L	smear of Saccharomyces cerevisiae, and preparing smears from solid media [see F will use paper towels instead of bibulous >SUPPL. EX., EXAMINATION OF STAINED S	SIMPLE (POSITIVE) STAINING (On a single slide, prepare a a separate smear of <i>Escherichia coli</i> . Use the technique for Ex. 11, p. 96], & stain with crystal violet [see Ex. 12].) We spaper. Use this slide in the exercise below (SUPPL. Ex.). ELIDES AND WET MOUNTS OF THE YEAST <i>Saccharomyces</i> UM <i>Escherichia coli</i> (Hand in your drawings to the
Tues.	Jan. 25	Cell structure/function Eukaryotic microorganisms	Chap. 4 & 9 (pages 235-236) Chap. 18
Tues.	Jan. 25L	and teeth. Gently rub swab onto a DRY methylene blue, rinse, and blot dry. Exa bacteria on page 104 or 105. >EX. 13, NEGATIVE STAINING (We will us steps 1-7, but omit step #4. Draw a few Bacillus subtilis cells on page 104 or 10 on page 107. (You may need to consult 1) >If necessary, complete SUPPL. EX., EXA	emove a sterile swab from wrapping paper & swab your gum slide. Allow smear to air dry; then heat fix. Stain with amine with oil immersion objective. Draw epithelial cells and e nigrosin & the method in Fig. 13.1. On page 102, follow representative <i>Staphylococcus aureus</i> cells and 05. Answer questions 1-5, page 106; and answer questions Ex. 14 to answer the questions about the capsule stain.) AMINATION OF STAINED SLIDES AND WET MOUNTS OF THE IGUS) AND THE BACTERIUM <i>ESCHERICHIA COLI</i> tor at the end of lab, 10 points)
Thurs.	Jan. 27	Eukaryotic microorganisms	Chap. 18
Thurs.	Jan. 27L	next week. Work in groups of 4 and exponinutes. Expose one plate inside the build plates at room temperature until next week > SUPPL. EX., WINOGRADSKY COLUMN (WITHE PROCEDURE IN THE SUPPL. EX., BUT PLIN THE LAB MANUAL AS WELL.) Discuss > If necessary, complete SUPPL. EX., EXAYEAST Saccharomyces cerevisiae (A FUNDAL)	y. You will prepare the plates we will use ose 2 plates of Sabouraud dextrose agar to air for 45 lding and the other plate outside. Incubate the ek.) E WILL USE TEXT, P. 654-657

Date		Topics/Lab Exercises	Related material in text
Tues.	Feb. 1	Eukaryotic microorganisms Nutrition, culture, & metabolism of microorganisms	Chap. 18 Chap. 5 & 21
Tues.	Feb. 1L	>EX. 10, PURE CULTURE TECHNIQUES, STREAK-PLATE I You will use a loopful of water from one of your Wir microorganisms in this exercise. Use a prepared plate Eosin methylene blue agar for doing the quadrant strehis/her own streak plate. >CHECK WINOGRADSKY COLUMNS (Make macroscopin information. Observe biofilm slides. You may also plate detailed drawings of any microorganisms observed. & CYANOBACTERIA to aid you in recognizing different semester, be sure you see and draw examples of protogrelated to data collection & organization with your	nogradsky columns as the mixed sample of e of MacConkey agar, desoxycholate agar, or eak (method B on page 85). Each person will do to observations of columns, and record this prepare wet mounts, if desired. Make neat, Use the information in Ex. 6, PROTOZOA, ALGAE, at groups of organisms. At some point during the ozoa, algae, & cyanobacteria. Discuss issues
Thurs.	Feb. 3	Nutrition, culture, & metabolism of microorganisms	Chap. 5, 21, & 20
Thurs.	Feb. 3L	>FOR EX. 61, YOU WILL WORK IN GROUPS OF 4. PICE GROUP. OBTAIN A FRESHWATER SAMPLE AND BRIN >EX. 10, PURE CULTURE TECHNIQUES, STREAK-PLATE Examine plate from Tuesday. Pick a well-isolated comethod B on page 85) on the prepared plate of mediu well-isolated colony, take a VERY TINY sample from using method B on page 85. >FINISH EX. 7, THE BACTERIA (Complete table, p. 57. on page 58.) Use plates with fungal colonies in Ex. 3 >FINISH EX. 8, THE FUNGI (Fungi Study – Do NOT op them only in the biological safety cabinet. You will us more different molds. The instructor will describe the power (10x) objective and the high dry (40x) objective record a description of the appearance of the fungal colonies.	METHOD ONLY blony, and use it to do another streak plate (using am provided by the instructor. If you do not have m your plate and perform another streak plate, Answer short answer questions 1 & 2 8. ben fungal cultures in the lab. Open use transparent tape to prepare slides of two or its procedure. Examine the slides using the low ve. Draw the specimens on p.67, part A2. Also
Tues.	Feb. 8	Metabolic diversity of microorganisms	Chap.15, 16, & 17
Tues.	Feb. 8L	REMEMBER TO BRING 2 TUBES WITH FRESH WATER SEX. 61, BACTERIOLOGICAL EXAMINATION OF WATER water collected in 2 sterile, 50 ml tubes for this exerce >Ex. 10, PURE CULTURE TECHNIQUES, STREAK-PLATE Examine plates from Thursday. Hopefully, each grout isolate to use for their general unknown. If you are lead isolated colony, pick a well-isolated colony and transgroup's general unknown culture; please label it clear seat numbers. If your group has no plates that were a well-isolated colony and use it to do another streak prepared plate of medium provided by the instructor. colony from the new plate to transfer to a nutrient agas >MONITOR WINOGRADSKY COLUMNS Discuss plansgroup.	(You will work in groups of 4 and use the fresh rise.) METHOD ONLY up of 4 students will be able to decide today on a cooking at a streak plate prepared from a wellsfer it to a nutrient agar slant. This can be your rly with "UNKNOWN", your lab section, and prepared from a well-isolated coloony, then pick plate (using method B on page 85) on the During the next lab you will pick a well-isolated ar slant for use as your group's unknown. for the Winogradsky lab report with your
Thurs.	Feb. 10	EXAM 1 (Material covered through Feb. 8)	
Thurs.	Feb. 10L	>CONTINUE Ex. 61, BACTERIOLOGICAL EXAMINATION MacConkey agar instead of Endo agar. Record result CONTINUED ON PAGE 6	NOF WATER (MPN+EMB/MAC) We will use

Date		Topics/Lab Exercises	Related material in text
Thurs.	Feb. 10L	CONTINUED FROM PAGE 5	
		slant culture, please do this today. If you are colony, pick a well-isolated colony and transfe general unknown culture; please label it clearl	REAK-PLATE METHOD ONLY nasn't yet established a general unknown nutrient agar looking at a streak plate prepared <u>from</u> a well-isolated or it to a nutrient agar slant. This can be your group's y with " <u>UNKNOWN", your lab section, and seat</u> has no suitable colonies, please consult the instructor.
		TECHNIQUE) WORK IN GROUPS OF 2 FOR THIS	OCIATED WITH FRESH PRODUCE (SPREAD-PLATE EXERCISE. ALSO, PLEASE READ EX. 22 TO LEARN ING P. 155-157. HOWEVER, PLEASE NOTE THAT WE DURSE PACKET
Tues.	Feb. 15	Metabolic diversity of microorganisms	Chap.15, 16, & 17
Tues.	Feb. 15L	>FINISH EX. 61, BACTERIOLOGICAL EXAMINATI	ON OF WATER (Read results of EMB/MAC. We will omi C tests.) Answer questions 4-9 on p. 398-399. TERIA ASSOCIATED WITH FRESH PRODUCE KET
Thurs.	Feb. 17	Microbial growth	Chap. 6
Thurs.	Feb. 17L	work session on dilution problems; Ask >> Hand in 3 stapled articles in a folder (for scientific journals – 12 points). These articles	
Tues.	Feb. 22	Microbial molecular biology DNA structure & replication; transcription, tra	Chap. 3, 7, & 8
Tues.	Feb. 22L	manual. Complete drawings/questions, p. 119 >GENERAL UNKNOWN CULTURESREAD ABG (stock cultures) of the unknown and also gr your own unknown record sheet. Also record PREPARE A STREAK PLATE USING YOUR UNKNO >YOUR LAB REPORT ON THIS GENERAL UNKN a thin folder that contains the following: (i) a record sheets, including your drawings, (ii) a record sheets, including your drawings, (iii) a record sheets, includi	DUT STOCK CULTURES IN Ex. 20. <u>Prepare subcultures</u> <u>am stain it.</u> Record dates, work done, drawings, etc., or your results on the descriptive chart on page 263.

Date		Topics/Lab Exercises	Related material in text
Thurs.	Feb. 24	Microbial molecular biology DNA structure & replication; transcription, translation Regulation of gene expression	Chap. 7 & 8 Chap. 9
Thurs.	Feb. 24L	>CONTINUE WORK ON GRAM STAINING KNOWN AND UNKNOW >EXAMINE STREAK PLATE OF UNKNOWN. Measure diameter of colonies on your unknown record sheet and on the description	of colonies and record a description of the ve chart on p. 263. Consult p. 268 (Ex. 40)
Tues.	Mar. 1	Regulation of gene expression Viruses	Chap. 9 Chap. 10 & 19
Tues.	Mar. 1L	>SUPPL. EX., VARIOUS MEDIA (CULTURES FOR DESOXYCHOL AGAR: Escherichia coli, Staphylococcus aureus, Pseudomon (CULTURES FOR BLOOD AGAR: E. coli, S. aureus, Bacillus cer. >A THROAT CULTURE WILL ALSO BE PERFORMED ON A BLOOD SEX. 17, ACID-FAST STAINING (Ziehl-Neelsen method proced water for preparing the smears. On one slide prepare a smea smegmatis & Staphylococcus aureus, as well as a separate sto air dry, and then heat fix them. Put on gloves, and try to up any spills of carbol fuchsin.) Cover the smears with a culture over the edges of the slide. Hold the slide with a clothespin carbol fuchsin. Heat the slide intermittently over the flame of 5 minutes. Do NOT let the paper towel dry out—add more to cool and then remove the paper towel. Proceed with steps in Figure 17.1 on page 118. Complete drawings/questions, p. 119-122; omit questions 1 unknown culture on unknown record sheet, and on the description of the paper towel of the description of the paper towel of the pa	as aeruginosa, & unknown) reus, & unknown) D AGAR PLATE. lure) Use 0.1% albumin solution instead of a mixture of Mycobacterium mear of your unknown. Allow the smears be neat. (You are responsible for cleaning to piece of paper towel that does not extend or slide holder and soak the towel with of the bunsen burner so that it "steams" for carbol fuchsin as needed. Allow the slide is 2 through 7 as described in the lab manual & 2 on p. 122. Record results for riptive chart on p. 263. with your group.
Thurs.	Mar. 3	Viruses	Chap. 10 & 19
Thurs.	Mar. 3L	>EX. 33, ULTRAVIOLET LIGHT: LETHAL EFFECTS >FINISH SUPPL. EX., VARIOUS MEDIA Record results in the tallow, record results for your unknown on your unknown.	
		descriptive chart on p. 263. Consider the following question: Is the pattern of growth consistent with the results you obtained in the Gram stail	of your unknown on the selective media
		Consider the following question: Is the pattern of growth consistent with the results you obtained in the Gram stail >EX. 16, SPORE STAINING (Modified Schaeffer-Fulton Method Bacillus species provided as well as a separate smear of you then heat fix them. Put on gloves, and try to be neat. (You of malachite green.) Complete drawings/questions, p. 119-122; omit questions 1 culture on unknown record sheet, and on the descriptive characterists.	n of your unknown on the selective median? od) On one slide prepare a smear of the runknown. Allow smears to air dry, and are responsible for cleaning up any spills & 2 on p. 122. Record results for unknown on p. 263.
Tues.	Mar. 8	Consider the following question: Is the pattern of growth consistent with the results you obtained in the Gram stail >EX. 16, SPORE STAINING (Modified Schaeffer-Fulton Methon Bacillus species provided as well as a separate smear of you then heat fix them. Put on gloves, and try to be neat. (You of malachite green.) Complete drawings/questions, p. 119-122; omit questions 1 culture on unknown record sheet, and on the descriptive chamber Viruses Microbial genetics	n of your unknown on the selective median? od) On one slide prepare a smear of the runknown. Allow smears to air dry, and are responsible for cleaning up any spills & 2 on p. 122. Record results for unknown rt on p. 263. Chap. 10 & 19 Chap. 11
	Mar. 8 Mar. 8L	Consider the following question: Is the pattern of growth consistent with the results you obtained in the Gram stail >EX. 16, SPORE STAINING (Modified Schaeffer-Fulton Methon Bacillus species provided as well as a separate smear of you then heat fix them. Put on gloves, and try to be neat. (You of malachite green.) Complete drawings/questions, p. 119-122; omit questions 1 culture on unknown record sheet, and on the descriptive chamber of the provided statement of the provided stat	n of your unknown on the selective median? od) On one slide prepare a smear of the runknown. Allow smears to air dry, and are responsible for cleaning up any spills & 2 on p. 122. Record results for unknown rt on p. 263. Chap. 10 & 19 Chap. 11
		Consider the following question: Is the pattern of growth consistent with the results you obtained in the Gram stail >EX. 16, SPORE STAINING (Modified Schaeffer-Fulton Method Bacillus species provided as well as a separate smear of you then heat fix them. Put on gloves, and try to be neat. (You of malachite green.) Complete drawings/questions, p. 119-122; omit questions 1 culture on unknown record sheet, and on the descriptive chamber Viruses Microbial genetics >FINISH EX. 33, ULTRAVIOLET LIGHT (Observe demonstration)	n of your unknown on the selective median? od) On one slide prepare a smear of the runknown. Allow smears to air dry, and are responsible for cleaning up any spills & 2 on p. 122. Record results for unknown rt on p. 263. Chap. 10 & 19 Chap. 11

Date		Topics/Lab Exercises	Related material in text
Tues.	Mar. 8L	CONTINUED FROM PAGE 8	
		>Ex. 40, CULTURAL CHARACTERISTICS (You will inconutrient agar slant [use a straight inoculation line], a gelatin deep, & fluid thioglycollate medium.) >Ex. 18, MOTILITY DETERMINATION (TUBE METHOD of medium with <i>Staphylococcus aureus</i> , <i>Proteus vulga</i> >Ex. 21, CULTIVATION OF ANAEROBES – We will not particular attention to the information about oxygen which you will use for your unknown, as noted above	nutrient broth, motility medium [deep], nutrient ONLY) You will inoculate tubes of motility aris, (& your unknown, as noted above). do this exercise, but you should read it with requirements and fluid thioglycollate medium,
		>SUPPL. EX., PLAQUE ASSAY OF A PHAGE SUSPENSION READ pages 173-175 in the lab manual. Ask quest >MONITOR WINOGRADSKY COLUMNS (today &/or Th	tions on dilution problems.
Thurs.	Mar. 10	EXAM 2 (material covered through Mar. 8)	
Thurs.	Mar. 10L	>FINISH EX. 33, ULTRAVIOLET LIGHT (Observe demonants answer questions on p. 223-224.) >FINISH EX. 40. (Record results on unknown record Consult Ex. 21 for information about oxygen require. >FINISH EX. 18, MOTILITY (TUBE METHOD & WET MOTILITY (TUBE METHOD	sheet and on descriptive chart on p. 263. ements and fluid thioglycollate medium.) OUNT) (On pages 127-128, draw the motility tube et mount of the nutrient broth culture of your cope. Record the results of the motility tube test and in the descriptive chart on p. 263.) PENSION – Record results on board.
		SPRING BREAK	
Tues.	Mar. 22	Microbial genetics Genetic engineering & Biotechnology	Chap. 11 Chap. 12 & 26
Tues.	Mar. 22L	>PREPARE NEW STOCKS OF GENREAL UNKNOWNS >Ex. 41, OXIDATION AND FERMENTATION TESTS >Ex. 43, MULTIPLE TEST MEDIA (We will do ONLY SIM medium.) >EX. 42, HYDROLYTIC/DEGRADATIVE REACTIONS (Mespirit blue agar for the lipid hydrolysis test. On tributindicates a positive test for lipid hydrolysis.) INSTRUCTOR WILL MAKE BRIEF COMMENTS ABOUT PREAD EX. 44, USE OF BERGEY'S MANUAL, & EXAMIC SPERGEY'S MANUAL OF DETERMINATIVE BACTERIO You will need to consult it to find out which addition identify your unknown. You will need this information of the property of the p	Iodification: we will use tributyrin agar rather than utyrin agar, a clear zone around the bacterial grow T BERGEY'S MANUAL INE APPENDIX 2 (TEXT) bLOGY is on reserve in the library. Inal tests would allow you to more specifically ation for your lab report, which is due on April 7.
Thurs.	Mar. 24	Microbial genomics Microbial evolution & systematics Microbial identification & clinical microbiology	Chap. 13 Chap. 14 Chap. 32
Thurs.	Mar. 24L	>FINISH EX. 41(EXCEPT VP TEST, WHICH MUST BE INC >FINISH EX. 43 & 42. Record results on unknown record sheet, and on des Answer questions 4-9 in part B, p. 291-292. In addi >Work on lab reports (unknown & Winogradsky	CUBATED LONGER) scriptive chart on p. 263. ition, answer questions in part C, p. 293-294.

Date		Topics/Lab Exercises	Related material in text
Tues.	Mar. 29	Microbial genomics Microbial evolution & systematics Microbial identification & clinical microbiology Microbial growth control	Chap. 13 Chap. 14 Chap. 32 Chap. 27
Tues.	Mar. 29L	>Program #9, Microbial Control >EX 41, READ RESULTS OF VP TEST (See procedure on be record sheet, and on descriptive chart on p. 263. THIS IS THE LAST DAY FOR LAB WORK ON THE GENERAL PEX. 36, KIRBY-BAUER METHOD >EX. 37, EVALUATION OF ANTISEPTICS (PAPER DISK METHOD) >EX. 35, EFFECTIVENESS OF ALCOHOL >Do the following online exercise on your own: >SUPPL. EX., USING RIBOSOMAL RNA GENE SEQUENCES T	UNKNOWN. THOD- this exercise will be slightly modified
Thurs.	Mar. 31	>SUPPL. EX., Staphylococcus aureus EXPERIMENT (class Microbial growth control	Chap. 27
Thurs.	Mar. 31L	>VIDEO SEGMENTS >SUPPL. EX., Staphylococcus aureus EXPERIMENT >FINISH EX. 36, 37, & 35. Record data & answer questio >Work on lab reports with your group.	ons in lab manual. Brief class discussion.
Tues.	Apr. 5	Microbial ecology (selected topics)	Chap. 22-24, & 26
Tues.	Apr. 5L	>VIDEO SEGMENTS >CONTINUE SUPPL. EX., Staphylococcus aureus (Record antibiotic sensitivity tests that are described in this exer aureus for isolation on a plate of tryptic soy agar. This STUDENT ORAL PRESENTATIONS	cise. Remember to streak presumptive <i>S</i> .
Thurs.	Apr. 7	Innate immunity; adaptive immunity	Chap. 29, 30, & 31
Thurs.	Apr. 7L	>SUPPL. EX., BACTERIOLOGICAL ANALYSIS OF URINE (RE >FINISH SUPPL. EX., Staphylococcus aureus >Ex. 73, Latex agglutination test for S. aureus ide STUDENT ORAL PRESENTATIONS HAND IN WINOGRADSKY LAB REPORT	
Tues.	Apr. 12	Adaptive immunity Practical applications of immunology	Chap. 29, 30, & 31 Chap. 29-32
Tues.	Apr. 12L	>HAND IN SUPPL. EX., RIBOSOMAL RNA SEQUENCES (12 F) >CONTINUE SUPPL. EX., URINE (Inoculate Enterotube II) omit the Kirby Bauer antibiotic sensitivity tests. >SUPPL. EX., BACTERIAL CONJUGATION	See EX. 46, ENTEROTUBE II. We will
Thurs.	Apr. 14	EXAM 3 (Material covered through Apr. 12)	
Thurs.	Apr. 14L	>COMPLETE SUPPL. EX., BACTERIAL CONJUGATION — Ans be sure you understand what happened and why it hap spinish suppl. Ex., URINE & EX. 46 ENTEROTUBE II For Practical applications of immunology WORK ELISA AND IMMUNOFLUORESCENCE PROBLEMS (SI	swer the questions with this exercise & pened. HAND IN LAB REPORT Chap. 29-32

Date		Topics/Lab Exercises	Related material in text
Tues.	Apr. 19	Human-microbe interactions Epidemiology & public health	Chap. 28 Chap. 33
Tues.	Apr. 19L	HAND IN LAB REPORT ON GENERAL UNKNOWN STUDENT ORAL PRESENTATIONS	
Thurs.	Apr. 21	Human-microbe interactions Epidemiology & public health	Chap. 28 Chap. 33
Thurs.	Apr. 21L	STUDENT ORAL PRESENTATIONS	
Tues.	Apr. 26	Microbial diseases	Chap. 34-37
Tues.	Apr. 26L	STUDENT ORAL PRESENTATIONS	
Thurs.	Apr. 28	Microbial diseases	Chap. 34-37
Thurs.	Apr. 28L	STUDENT ORAL PRESENTATIONS	
Wed.	May 4	COMPREHENSIVE FINAL EXAM (EXAM 4) – 10):15 am – 12:15 pm

ADDITIONAL INFORMATION

<u>Course content:</u> We will not be covering all of the material in the textbook and lab manual. Please read the sections of the textbook and lab manual that pertain to the topics covered, and make use of the tables and illustrations. Study questions and online resources for the textbook may also be useful. **Specific assigned readings may be announced in class or lab.**

Laboratory:

- 1. Laboratory exercises are an integral part of microbiology. Students are expected to attend ALL laboratory sessions, to be on time at the beginning of the period, and to complete all assigned laboratory exercises. There will be no makeups for the laboratory exercises.
- 2. Microscopes will be assigned and spot checks will be made to ensure that they are clean and properly stored. Misuse or mishandling of the microscopes will result in the loss of points (20 points per occurrence). After you have finished using your microscope, please consult the "microscope checklist" to be certain that you have followed the proper procedures.
- 3. Each student must **read the laboratory exercises for the day, any additional required readings from the lab manual (noted in the syllabus), and any notes pertaining to the lab exercises (in the syllabus)** <u>before</u> coming to the laboratory. This will allow the student to complete the exercises in an efficient and informed manner.
- 4. Each student must record the results of the lab exercises and answer the related questions, as noted in the syllabus. In some cases, **lab reports** are due as indicated in the course schedule. If a student misses a portion of the lab work relating to a required lab report, the student's report will be worth a maximum of 85% of the points allotted for the report. Each student must turn in his/her own drawings (due Jan. 20) and rRNA report (due April 12). However, the Winogradsky Column Project report must be prepared with your lab group (due April 7). **For this report, each group member will evalulate the percentage of the work contributed by each of the group members, and individual scores will reflect the average percents.** For the remaining reports [general unknown (due April 19) and urine unknown (due April 14)], students may prepare their lab reports individually, or they may work with their lab groups and turn in joint reports.
- 5. Oral Presentations. During the laboratory portion of the course, each student will give an 8- to 10-minute oral report on a primary scientific article or case study selected from a list provided by the instructor. Students will draw numbers to indicate the order in which they will select articles and give their presentations. Once a topic is chosen it may not be changed. Students should search databases in GALILEO to find related, formal scientific articles. Some peer-reviewed, scientific journals are available in the Odum library and/or online. At least three formal articles (including the original article chosen) from PEER-REVIEWED, PROFESSIONAL JOURNALS must be used to prepare the presentation. Only one of these articles may be a review article. Complete, stapled (or paper-clipped) copies of these articles must be placed in a folder & handed in on Feb. 17. The folder with the articles must again be provided to the instructor on the day of the presentation. Articles must list references at the end, and references must be cited within the text. Informal articles, Web sites, Internet articles or fact sheets, newspaper articles, magazine articles, book reviews, and letters to the editor are NOT acceptable. Students should make every effort to ensure the accuracy of the information in their reports. Should a report contain inaccurate information, the presenter should expect to be questioned about it as well as about the source of the information.

For their presentations, students are required to use PowerPoint software. <u>Students must use a PowerPoint version that is compatible with the one available in the microbiology lab.</u> If you are in doubt, please bring your PowerPoint presentation to the

lab at least one week before the day of your presentation to verify that it will run. If you do not check your presentation ahead of time, you are responsible for having a backup method for showing your illustrations. Full-size print-outs of your PowerPoint slides are useful as backups, since they may be shown using the ELMO projector. Students may use visual aids in addition to PowerPoint. A projector for transparencies is available; handouts may also be used. There will be no makeups for the oral presentations. On the day of the presentation, the student must turn in: complete copies of the three references, a copy of the PowerPoint slides, and a copy of any other illustrations or notes used during the presentation.

ADDITIONAL EMPHASIS: IF YOU WANT A GOOD SCORE ON YOUR PRESENTATION, YOU MUST FOLLOW THE GUIDELINES ON THE PROVIDED EVALUATION FORM. A STUDENT WHOSE REPORT DOES NOT FOCUS ON THE PRIMARY SCIENTIFIC ARTICLE OR CASE STUDY WILL RECEIVE A SCORE OF ZERO.

Attendance, participation, and tardiness: In accordance with VSU policy, attendance and participation will be checked in the laboratory and in class. The remainder of this paragraph outlines the lab/oral presentation attendance policy. Attendance is required during ALL labs and oral presentations periods. A student who has perfect attendance during laboratory/oral presentation periods will receive 25 bonus points. A student who misses (or fails to complete) only one laboratory/oral presentation period will receive 15 bonus points. Missing (or failing to complete) additional laboratory/oral presentation periods will result in the loss of points as follows. Ten points will be deducted for the fourth missed (or incomplete) period; 20 additional points will be deducted for the fifth missed (or incomplete) period; 40 additional points will be deducted for the sixth missed/incomplete period, and 50 additional points will be deducted for each subsequent missed/incomplete period. Students who are habitually late for lab or oral presentation periods will be marked late. Coming late to lab or oral presentation periods three times will be counted as one absence. A student with more than 6 absences (or a student who fails to complete more than 6 laboratory or oral presentation periods) will not pass the course. There will be no makeups for the laboratory exercises or student presentations.

Examinations given during class periods:

- 1. Examinations 1-4 will cover material presented during both the class and laboratory portions of the course. The first three exams will be worth 185 points each. The final exam will be worth 225 points. Examinations will begin promptly at the times and dates indicated on the class schedule. The final examination will be comprehensive in that it will include material covered throughout the course. Exams 2 and 3 will be comprehensive in that up to 25% of the points on the exam may cover material presented before any earlier examination. Exams may include questions of the multiple-choice, matching, true-false, short-answer, and essay formats. A student who misses an examination should notify the instructor promptly. Arrangements for a make-up exam must be made within one week after the exam date; otherwise, a make-up exam will not be given. Make-up examinations may consist entirely of questions of the short answer and essay formats. Make-up examinations for exams 1, 2, and 3 will be worth 165 points rather than 185 points each.
- 2. STUDENTS ARE REQUIRED TO BRING TWO #2 PENCILS AND ERASERS TO ALL EXAMINATIONS. THE INSTRUCTOR WILL NOT PROVIDE PENCILS.

Late Assignments & Failure to Turn in Assignments:

Please make a calendar noting when assignments and lab reports are due. Turning in an assignment/report 1-4 days late will result in a deduction of 20% of the points for that assignment. Turning in an assignment 5-9 days late will result in a deduction of 50% of the points for that assignment. No points will be awarded for an assignment that is late by more than 9 days. Students will not be notified by the instructor for failing to turn in course assignments. Late assignments must be given DIRECTLY to the instructor. They may NOT be placed in the instructor's mailbox. It is also NOT ACCEPTABLE to slide late assignments under the instructor's office door.

Grading: Points for the course are allocated as follows:

EXAMS 1, 2, & 3 (Feb. 10, Mar. 10, & April 14) (185 points each x 3=555)	555	POINTS	
EXAM 4 (FINAL EXAM –May 4)	225	POINTS	
LAB REPORT (Drawings) (Course objective ZA) - (Jan. 20)	10	POINTS	
REFERENCES FOR ORAL PRESENTATION (Course objective ZG) - (Feb. 17)) 12	POINTS	
rRNA LAB REPORT (Course objective ZD) - (April 12)	12	POINTS	
LAB REPORT ON GENERAL UNKNOWN (Course objectives ZC, ZD) - (Apri	11 19) 33	POINTS	
LAB REPORT ON URINE UNKNOWN (Course objective ZD) - (April 14)	13	POINTS	
FORMAL (EXTENSIVE) LAB REPORT (Course objective ZF) - (April 7)	65	POINTS	
ORAL PRESENTATION (Course objective ZH) - (April 5-28)	75	POINTS	
TOTAL FOR COURSE	1000	POINTS	

There are FOUR REQUIREMENTS TO PASS the course:

- 1. Do not miss (or fail to complete) any more than 6 laboratories or oral report periods.
- 2. Complete and turn in all assignments and lab reports.
- 3. Obtain at least 40% of the points for **EACH** assignment and lab report.
- 4. Have a total of 600 or more points for the course.

Students should read the entire syllabus carefully so they understand the course policies & procedures.

The grade is "F" for a student who obtains less than 600 total points or fails to meet one of the other requirements for passing the course (see above list). GRADING SCALE: 900-1000, A; 800-899, B; 700-799, C; 600-699, D; < 600, F