

BIMM 121 Laboratory in Microbiology Spring 2011

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Office hours: Mondays 10 am -11 am

Lecture: Tuesday/Thursday 8:00 am to 9:20 am, PCYH 122 (Pepper Canyon Hall, next to Gilman Parking structure)

Labs: York 2310 and 2332
Tuesday/Thursday: 9:30 am – 1:30 pm
Wednesday/Friday: 9:00 am – 1:00 pm

Course Structure:

This course will introduce you to the fundamentals of microbiology and allow you to explore the many ways in which microbes affect and are used in our lives. We begin the course with a foundation in basic techniques such as sterile techniques, microscopy, methods of quantitating microbes, and preparing and examining stained slides. The remaining duration of the course will comprise four main units: a comprehensive look at bacterial physiology, understanding the complex microbial community of soil, metagenomics as a tool in exploring complex communities, and the use of microbes in various aspects of our lives. Each of these units comprises several multi-day experiments and there will be considerable overlap in the execution, methodology, and analysis of data from each of these units. Throughout the course, you will also receive training in accurate data entry and analysis, scientific reasoning, and in clear and concise scientific writing.

Equipment:

For this lab you will need to purchase:

- A lab notebook (check with instructor to determine if notebook with carbons is required);
- A lab coat
- Eye protection (you may wear either safety glasses or goggles, but standard prescription eye glasses are not sufficient).
- A Sharpie permanent marker pen, preferably fine point (not extra fine or regular)

Attendance and Absences:

1. Your attendance is required at EVERY lab and through the entire lab period, until all the experimental work for the day is completed.
2. Absences will NOT be treated lightly. The labs are set up for groups of two or more and your absence will place an unnecessary burden on your partner. There are no make up labs and you will not be allowed in the lab on non-lab days or in the other Micro lab sections, although you may be asked to make up the work from the day you missed.
3. Documentation will be required for all unavoidable absences.
4. If you are likely to have interviews for graduate school, etc., please schedule them on non-lab days.
5. All absences without prior notification/permission and the appropriate paperwork will be considered unauthorized.
6. **50-point penalty** for the first unauthorized, unexplained absence from the lab. If there is a second such absence, you will be asked to drop the course or will be given an F.
7. If you are ill on a lab day or have an emergency, e-mail or call (instructor or lab partner) before the start of the lab. If you are ill enough to miss lab you must go to the student health center and provide documentation of your illness.

Assignment Deadlines and Submission:

1. A hard copy of each lab report is due in the first 10 minutes of the lab period or the first 10 minutes of the lecture period of the day on which your report is due. Check with your instructor as to where the report should be turned in. Reports turned in more than 10 minutes after the start of class will be considered late. Penalty for late reports will be 10% for each day late.
2. In addition to the hard copy of the report, you are required to submit an electronic copy to Turnitin.com. A link to the e-submission website will be provided on WebCT. Failure to submit on Turnitin.com will result in 0 (zero points) recorded for that report. Check the deadline of the Turnitin.com submission and make sure you adhere to it. Students agree that by taking this course all required papers would be subject to review for textual similarity by Turnitin.com for the detection of plagiarism. All submitted papers will be included as source documents in the Turnitin.com reference database solely for the purpose of detecting plagiarism of such papers. Use of the Turnitin.com service is subject to the terms of use agreement posted on the Turnitin.com site.
3. Additional points may be taken for late electronic submissions.

Regrade Requests:

All regrade requests should be submitted in writing within one week of receiving the graded material.

Grading Scheme

Quiz/Report/Midterm	Points
Daily quizzes at 5 points per week	40 points
TA assessment:	35 points
-Notebook checks	
-Techniques	
-Class participation	
2 surveys at 10 points each	20 points
Lab report 1	100 points
Lab report 3	100 points
Lab report 4	35 points
Assignments	20 points
3 Midterms	<u>250 points</u>
Total	600 points

Possible assignments.

- Dilution assignment
- Transposon mutagenesis assignment
- Metagenomics assignment
- Growth curve assignment

Most Likely Grade Distribution

A = 90% - 100%

B = 80% - 89.9%

C = 70% - 79.9%

D = 60% - 69.9%

F = below 60%

Notebook:

Spiral bound or composition notebook is OK. All notebooks should have a table of contents (handwritten OK) so on the first lab day leave several blank pages at the beginning of your notebook. Number your pages. Entries should be made in chronological order and EVERY day. Each day's entries on each experiment should begin with a brief (1 – 2 sentences) summary of work done on the same experiment the previous day.

How to use your notebook

Table of contents – update everyday – leave at least 4-5 pages for updating
Start a new page each day for each new experiment:

Purpose of experiment

Procedure

Outline or page from which protocol was taken

Note any changes

Note who did which part of the procedure – who inoculated controls, etc

Note which organisms you used – name and species of the controls, etc

Errors

Observations

Write

Draw

Questions and connections

Conclusion or summary

Answer any questions in the manual or that were raised in class.

Lab Performance and Participation

In addition to quizzes, midterms, lab reports and assignments, student evaluations will be based on the following criteria:

1. Lab techniques will be evaluated in class
2. Lab workshop participation

Subjective student evaluations will be based on the following criteria:

3. Pre-lab preparation
4. Careful management of lab procedures (e.g., sterile technique, proper waste disposal, experimental procedures, etc.)
5. Ability to adapt to unforeseen procedural changes

6. Caliber of thinking before asking questions
7. Scientific approach (e.g., proper use of notebooks, controls, experimental design)
8. Accuracy
9. Independence
10. Safety consciousness
11. General neatness in lab

Please note: **You will be expected to get into the habit of methodical, well-planned and organized work by the mid-term. This will help you with the experiments in the second half of the course.**

Course Website

This course is on WebCT (<https://webctweb.ucsd.edu>) and should automatically appear on your WebCT account as soon as you register for the class. We will use WebCT to post information on experiments, exams, schedules, readings and practice material, experimental data, report guidelines, etc. We strongly encourage you to use the Discussion board to post questions or answers to questions and to use it as a forum for exploring the material. The TAs and I will routinely check this website and answer any questions but feel free to respond as well. This website will also be used to post any announcements that pertain to the entire class. Please check the site regularly and update yourself on the information provided.

University Policy on Integrity of Scholarship

The principle of honesty must be upheld if the integrity of scholarship is to be maintained by an academic community. The University expects that both faculty and students will honor his principle and in so doing protect the validity of University grading. This means that all academic work will be done by the student to whom it is assigned, without unauthorized aid of any kind. Instructors, for their part, will exercise care in planning and supervising academic work, so that honest effort will be encouraged.

Student Responsibility:

Students are expected to complete the course in compliance with the instructor's standards. No student shall engage in any activity that involves attempting to receive a grade by means other than honest effort; for example:

- No student shall knowingly procure, provide, or accept any unauthorized material that contains questions or answers to any examination or assignment to be given at a subsequent time.

- No student shall complete, in part or in total, any examination, or assignment for another person.
- No student shall knowingly allow any examination or assignment to be completed, in part or in total, for himself or herself by another person.
- No student shall plagiarize or copy the work of another person and submit it as his or her own work.
- **If any work is plagiarized from that of another student, both students will be reported to the Office of Academic Integrity, even if one of the students has graduated already. Remember that most graduate schools check the undergraduate records for any indications of dishonesty before awarding a degree.**
- No student shall alter graded class assignments or examinations and then resubmit them for regrading.
- No student shall submit substantially the same material in more than one course without prior authorization.

Lab	Date	Experiment	Reports, Midterms, Reminders
Lab 1	Tues/Wed March 29/30	Registration, introductory remarks, safety lecture, etc. Sterile technique. Microbes in the environment <i>E.coli</i> and toilet paper experiment Aseptic technique, streak and spread plates Lollie, note that we will be asking TAs to do a 1:100 dilution of the mixed culture for the spread plates Use of pipettors: Demo	
Lab 2	Thurs/Fri March 31/April1	Sterile technique. Microbes in the environment: Observe results <i>E.coli</i> and toilet paper experiment: Observe results Streak and spread plates: Observe results. Microscopy: Learning to focus the light microscope Calibrating your microscope Observing stained slides Plant Pathogens: Set up <i>Agrobacterium</i> -kalanchoe infections	
Lab 3	Tues/Wed April 5/6	Microscopy: Making a wet mount and Phase Contrast Microscopy: Wet	

		<p>mounts and phase contrast:- view, identify, and measure</p> <p>Understanding dilutions: Understanding dilutions- theory</p> <p>Measuring microbial growth: Yeast Direct counts using a hemocytometer Using a spectrophotometer Counting viable cells using plating</p> <p>General microbiology: Introduction to selective and differential media (lecture only)</p>	
Lab 4	Thurs/Fri April 7/8	Workshop: Intro to microbes, notebook, data analysis, report formats, researching topics	
Lab 5	Tues/Wed April 12/13	<p>Microscopy: Staining Smear preparation and simple staining Gram stain: control organisms only</p> <p>Characterizing a Test Organism: Introduction: Receive test organism: wet mount and streak plate and slants</p> <p>Winogradsky column Understanding the set up</p> <p>Soil Enumeration and Enrichment: Lab Period 1 Simple Enumeration: Serial dilution, plating on TSA, SDA, GAA, and MacConkey</p>	Report 1 due
Lab 6	Thurs/Fri April	<p>Microscopy: Staining Complete staining of all controls</p>	

	14/15	<p>Characterization of a Test Organism</p> <p>Gram stain MacConkey – inoculate with known G+ and G- organisms Sticky test with known G+ and G- organisms Endospore test – inoculate NSM</p> <p>Macronutrient use – how organisms get energy to survive: Introduction: Hydrolysis and use of large extracellular materials Polysaccharides: Starch plates - inoculate Proteins: Skim milk plates and gelatin deeps - inoculate Lipids: Rhodamine plates - inoculate Inoculate with test organism</p> <p>Soil Enumeration and Enrichment: Lab Period 2</p> <ul style="list-style-type: none"> • Simple enumeration: colony counts • Extracellular degradation: Enumeration: Serial dilution and plating of soil sample on starch/rhodamine/skim milk/cellulose plates Enrichment of soil organisms: inoculate minimal media containing starch/olive oil/skim milk/cellulose with soil 	
Lab 7	Tues/Wed April 19/20	<p>Characterization of a Test Organism</p> <p>Macronutrient use – how organisms get energy to survive Polysaccharides: Starch plates - complete Proteins: Skim milk plates and gelatin deeps - complete Lipids: Rhodamine plates – complete</p> <p>How energy is produced – aerobic vs. anaerobic breakdown</p>	Midterm 1

		<p>of organic compounds</p> <p>Oxygen requirements – inoculate thioglycolate tube</p> <p>Acid and gas production from sugar fermentation – inoculate Methyl-Red and Voges-Proskauer – inoculate</p> <p>T-streak plate for fresh isolated colonies</p> <p>Soil Enumeration and Enrichment: Lab Period 3</p> <ul style="list-style-type: none"> • Extracellular degradation: <p>Enumeration: test differential media and count</p> <p>Enrichment: subculture</p>	
		<p>Non-lab day: You may check the growth in the thioglycolate tube to determine the oxygen requirement</p>	
Lab 8	Thurs/Fri April 21/22	<p>Characterization of a Test Organism:</p> <p>How energy is produced – aerobic vs. anaerobic breakdown of organic compounds</p> <p>Oxygen requirements –complete</p> <p>Acid and gas from sugar fermentation - complete</p> <p>Methyl-Red and Voges Proskauer – complete</p> <p>Cytochrome C test – complete</p> <p>Catalase test – complete</p> <p>Nitrate reduction - inoculate</p> <p>H₂S production – inoculate</p> <p>Soil Enumeration and Enrichment: Lab Period 4</p> <ul style="list-style-type: none"> • Extracellular degradation: 	

		<p>Enrichment:</p> <ul style="list-style-type: none"> • Serial dilutions and plating to enumerate enrichment • Centrifuge aliquot of enrichment and freeze pellet <p>Winogradsky column</p> <p>Examine for evidence of anaerobiosis</p>	
Lab 9	Tues/Wed April 26/27	<p>Characterization of a Test Organism:</p> <p>How energy is produced – aerobic vs. anaerobic breakdown of organic compounds</p> <ul style="list-style-type: none"> • Nitrate reduction – complete test • H₂S production – complete test <p>Special metabolic functions: Standards only</p> <ul style="list-style-type: none"> • Indole production from tryptophan, catabolite repression – inoculate • Urease test – inoculate • Differential utilization of citrate by enterics - inoculate <p>Motility – inoculate plate and deep with test organism</p> <p>Survival in extreme conditions:</p> <ul style="list-style-type: none"> • Low pH • High pH • Low temp • High temp • High salt • Control 	

		<ul style="list-style-type: none"> inoculate appropriate broth with test organism <p>Soil Enumeration and Enrichment: Lab Period 5</p> <ul style="list-style-type: none"> Extracellular degradation: Enrichment: Complete colony counts and calculations Extreme conditions Enrichment: Inoculate medium as assigned <p>Winogradsky column</p> <p>Examine for evidence of anaerobiosis and H₂S production</p>	
Lab 10	Thurs/Fri April 28/29	<p>Characterization of a Test Organism:</p> <p>Special metabolic functions: Standards only</p> <ul style="list-style-type: none"> Indole production from tryptophan, catabolite repression – complete Urease test - complete Differential utilization of citrate by enterics – complete <p>Motility – complete</p> <p>Survival in extreme conditions:</p> <ul style="list-style-type: none"> Score growth/no growth in each tube <p>Soil Enumeration and Enrichment: Lab Period 6</p> <ul style="list-style-type: none"> Extreme conditions Enrichment: Serial dilution and plating of enriched sample Centrifuge aliquot and freeze pellet 	
		<p>TAs set up serial dilution and plating of soil sample for “Screening for Antibiotic Producers”</p>	

Lab 11	Tues/Wed May 3/4	<p>Soil Enumeration and Enrichment: Lab Period 6</p> <ul style="list-style-type: none"> • Extreme conditions <p>Enrichment: Colony counts and calculations</p> <p>Nitrogen fixation: Free-living - Anabaena</p> <ul style="list-style-type: none"> • Inoculate BG11 and BG11-0 with <i>Anabaena</i> <p>Metagenomics: Lab Period 1</p> <ul style="list-style-type: none"> • Step 1: Chromosomal DNA preps from frozen cell pellets from various soil enrichments • Step 2: Set up 16S rRNA PCR 	Report 2 due
Non lab day		TAs run gel of PCR reactions	
Lab 12	Thurs/Fri May 5/6	<p>Metagenomics: Lab Period 2</p> <ul style="list-style-type: none"> • Step 3: Evaluate PCR results • Step 4: Purify PCR product • Step 5: Set up ligations in pGEM-T <p>Screening for Antibiotic Producers: grid plates</p>	
Lab 13	Tues/Wed May 10/11	<p>Metagenomics: Lab Period 3</p> <ul style="list-style-type: none"> • Step 6: Transform ligations and plate on selective media <p>Screening for Antibiotic Producers: Identify antibiotic producers, measure ZOI</p> <p>Evaluation of antibiotics by the Kirby Bauer method Spread plates with standards and test efficiency of antibiotics</p>	

		<p>Nitrogen Fixation – Free-living - Anabaena Subculture in BG11 and BG11-0 – check materials list</p>	
Lab 14	Thurs/Fri May 12/13	<p>Metagenomics: Lab Period 3</p> <ul style="list-style-type: none"> • Step 7: Select white colonies and streak out for sequencing <p>Evaluation of Antibiotics by the Kirby Bauer Method Measure ZOI, identify any resistant colonies</p> <p>Growth curve experiment Growth and graphing of <i>Vibrio natriegens</i></p> <p>Identification of an Unknown Organism: Lab Period 1</p> <ul style="list-style-type: none"> • Receive unknown • Check morphology by microscopy – wet mount • Gram stain 	Midterm 2?
Lab 15	Tues/Wed May 17/18	<p>Identification of an Unknown Organism: Lab Period 2</p> <ul style="list-style-type: none"> • Streak TSA and MacConkey plates • Streak TSS slants <p>Metagenomics: Lab Period 4</p> <ul style="list-style-type: none"> • Analyze of sample sequence data – computer lab <p>Characterization of a Test Organism:</p> <ul style="list-style-type: none"> • Each group or set of groups outlines and explains characteristics of their assigned test organism • Create elimination flow chart for identification of genera <p>Transposon mutagenesis: Lab Period 1</p> <ul style="list-style-type: none"> • Step 1: Set up conjugation of <i>E.coli</i> and <i>Salmonella</i> 	<p>Midterm 2? Computer lab day</p>

Lab 16	Thurs/Fri May 19/20	<p>Transposon mutagenesis: Lab Period 2</p> <ul style="list-style-type: none"> • Step 2: Plate exconjugants for selection and counterselection • Save LB DAP recipient control plates for later use <p>Identification of an Unknown Organism: Lab Period 3</p> <ul style="list-style-type: none"> • Check TSA plates for temperature preference • Check morphology and motility by microscopy – wet mount • Gram stain and sticky test • Examine MacConkey • Complete Cytochrome C and catalase tests • Inoculate all media (except thioglycolate) provided with your unknown organism <p>Yogurt: Inoculate milk with starter yogurt</p>	
Lab 17	Tues/Wed May 24/25	<p>Transposon mutagenesis: Lab Period 2</p> <ul style="list-style-type: none"> • Count colonies and calculate transposition efficiency • Save all plates <p>Yogurt: measure pH, gram stain</p> <p>Identification of an Unknown Organism: Lab Period 4</p> <ul style="list-style-type: none"> • Complete all tests • Save all tests for reexamination • Inoculate thioglycolate tubes <p>Metagenomics: Step 8: Begin/complete analysis of all sequences</p>	Computer lab
		Non-lab day: Examine thioglycolate tubes	
Lab 18	Thurs/Fri	Identification of an Unknown Organism: Lab Period 5	Computer lab

	May 26/27	<ul style="list-style-type: none"> • Complete thioglycolate and any other tests <p>Transposon mutagenesis: Lab Period 3</p> <ul style="list-style-type: none"> • Step 3: Screen transposants (mutants) for loss of function mutations <p>Nitrogen Fixation Free-living: <i>Anabaena</i>: check for heterocysts Symbiotic: <i>Rhizobium</i>: Observe nodules</p> <p>Plant Pathogen Observe <i>Agrobacterium</i>-kalanchoe interaction</p> <p>Metagenomics: Complete analysis/discussion of all sequences</p>	
Lab 19	Tues/Wed May 31/June 1	<p>Identification of an Unknown Organism:</p> <ul style="list-style-type: none"> • Report due today- check with instructor <p>Transposon mutagenesis: Lab Period 4</p> <ul style="list-style-type: none"> • Check screens • Streak out mutants on LB/TSA plate for future use <p>Winogradsky column Observation and sampling Check out Review Potluck</p>	Unknown Report due?
Lab 20	Thurs/Fri June 2/3	Midterm 3 will be held during normal lecture or lab hours.	Midterm 3