

		MONDAY September 4	TUESDAY September 5	WEDNESDAY September 6 Day 1	THURSDAY September 7 Day 2	FRIDAY September 8 Day 3
M O L E B I O				Introductions; Syllabus; Textbooks	Chemistry Skills QUIZ How to make various molar solutions Video Tutorials: safety measurement making solutions	Making Solutions Practice;
				Bring a calculator!	Read DNA Science (pg. 321-330); Read Lab #1	Read DNA Science (pg. 321-330); Read Lab #1; Be ready for Practical
H W						

	MONDAY September 11 Day 4	TUESDAY September 12 Day 5	WEDNESDAY September 13 Day 6	THURSDAY September 14 Day 1	FRIDAY September 15 Day 2
M O L E B I O	Making Dilutions; Making Solutions Practice; Pipetting Techniques	<u>Lab #1:</u> Measurements, Micropipetting, and Sterile Techniques Solutions Practical	<u>Lab #1:</u> Measurements, Micropipetting, and Sterile Techniques Solutions Practical	Lecture: Basics of Chemistry	Lecture: Water, Buffers, and pH
H W	Read DNA Science (pg. 321-330); Read Lab #1; Be ready for Practical	Read DNA Science (pg. 321-330); Lab #1 due Thursday; Be ready for Practical	Read DNA Science (pg. 321-330); Lab #1 due Thursday; Be ready for Practical	Khan Academy/ Bozeman Biology/ other video links	Read through "The Nature of Amino Acids" handout Read "Gel Electrophoresis MANUAL"

	MONDAY September 18 Day 3	TUESDAY September 19 Day 4	WEDNESDAY September 20 Day 5	THURSDAY September 21 Day 6	FRIDAY September 22 Day 1
M O L E B I O	Henderson Hasselbach Equation	H-H questions Lecture: Basics of Gel Electrophoresis (Part I)	<u>Lab #2:</u> Gel Electrophoresis: Practicing Loading Gels prepping for RUN 1: using 50X TAE buffer; dilution to 1X TAE; making 0.8% agarose in 1X TAE	<u>Lab #2:</u> Gel Electrophoresis: Practicing Loading Gels prepping for RUN 1: finish making solutions; cast gels	<u>Lab #2:</u> Gel Electrophoresis RUN 1 - control
H W	Complete #1, 2, 4 on "The Nature of Amino Acids" handout; Read "Gel Electrophoresis MANUAL"	Read in text: 113-115; 357-359; Read Lab #2	Read Lab #2	Read Lab #2	Lab #2 due Wednesday;

	MONDAY September 25 Day 2	TUESDAY September 26 Day 3	WEDNESDAY September 27 Day 4	THURSDAY September 28 Day 5	FRIDAY September 29
M O L E B I O	<p><u>Lab #2:</u></p> <p>Gel Electrophoresis prepping for RUN 2: different [agarose] or voltage condition</p>	<p><u>Lab #2:</u></p> <p>Gel Electrophoresis RUN 2 - variation</p>	<p>EXAM #1</p> <p>GEL ELECTROPHORESIS</p> <p>HENDERSON-HASSELBACH EQUATION</p> <p>SOLUTION STUFF</p>	<p>Lecture:</p> <p>Carbon Skeletons and Amino Acids</p>	<p>Lecture:</p> <p>Protein Structure</p>
H W	<p>Lab #2 due Thursday;</p> <p>EXAM #1 WEDNESDAY</p>	<p>Lab #2 due tomorrow;</p> <p>EXAM #1 TOMORROW</p>	<p>Read pages 48 – 53;</p> <p>Starting Lectures about proteins tomorrow</p>	<p>Read pages 48 – 53;</p> <p>Read SciAm: Proteins</p>	<p>Read pages 48-53;</p> <p>Read SciAm: Proteins;</p> <p>Read “Understanding the Nature of Amino Acids”</p>

	MONDAY October 2 Day 6	TUESDAY October 3 Day 1	WEDNESDAY October 4 Day 2	THURSDAY October 5 Day 3	FRIDAY October 6
M O L E B I O	Learning all 20 Amino Acids!	Learning all 20 Amino Acids! Lecture: Ionization of Amino Acids; Isoelectric Point	Lecture: Ionization of Amino Acids; Isoelectric Point	Lab #3: Electrophoretic Separation of Proteins myoglobin hemoglobin cytochrome c serum albumin How to make our buffers from scratch!	STAFF DAY
H W	Read pages 48-53; Read SciAm: Proteins; Read "Understanding the Nature of Amino Acids"	Read pages 48-53; Read SciAm: Proteins; Read "Understanding the Nature of Amino Acids"	Read Lab #3; Complete questions #3, 5 - 7 in "Understanding the Nature of Amino Acids"	Read Lab #3; BE READY FOR PRACTICAL APPLICATION OF CHEMISTRY!	

		MONDAY October 9	TUESDAY October 10 Day 4	WEDNESDAY October 11 Day 5	THURSDAY October 12 Day 6	FRIDAY October 13 Day 1
M O L E B I O	NO SCHOOL		<u>Lab #3:</u> Electrophoretic Separation of Proteins Prepping RUN 1: preparing 5X tris-glycine stock buffer (1 per group) diluting to 1X tris-glycine running buffer @ pH 8.6 (1 per group) preparing 3.5% agarose in 1X tris-glycine running buffer	<u>Lab #3:</u> Electrophoretic Separation of Proteins Prepping RUN 2: preparing 1X sodium acetate running buffer @ pH 5.5 (1 per group) preparing 3.5% agarose in 1X sodium acetate running buffer	<u>Lab #3:</u> Electrophoretic Separation of Proteins cast gel for RUN 1 diluting proteins learning staining/destaining procedure	<u>Lab #3:</u> Electrophoretic Separation of Proteins RUN 1: Tris-Glycine at pH 8.6 cast gel for RUN 2
	H W		Read Lab #3; Read protein article from Scientific American Read "Protein Gel Electrophoresis MANUAL"	Read Lab #3; Read protein article from Scientific American Read "Protein Gel Electrophoresis MANUAL"	Read Lab #3;	Read Lab #3; EXAM #2 WEDNESDAY

		MONDAY October 16 Day 2	TUESDAY October 17 Day 3	WEDNESDAY October 18 Day 4	THURSDAY October 19 Day 5	FRIDAY October 20 Day 6
M O L E B I O		<u>Lab #3:</u> Electrophoretic Separation of Proteins RUN 2: Acetate at pH 5.5	<u>Lab #3:</u> Electrophoretic Separation of Proteins RUN 3: Tris-Glycine-SDS SDS-PAGE	EXAM #2 AMINO ACIDS PROTEINS GEL ELECTROPHORESIS	<u>Lab #4:</u> Introduction to Spectrophotometry	<u>Lab #4:</u> Introduction to Spectrophotometry
	H W	Lab #3 due Friday; EXAM #2 WEDNESDAY	Lab #3 due tomorrow; EXAM #2 TOMORROW	Read Lab #4	Be ready for Lab #4!	Lab #4 due tomorrow

	MONDAY October 23 Day 1	TUESDAY October 24 Day 2	WEDNESDAY October 25 Day 3	THURSDAY October 26 Day 4	FRIDAY October 27 Day 5
M O L E B I O	Lecture: Enzymes General Properties	Lecture: Enzymes Factors that Affect Reaction Rates Michaelis-Menton equation Lineweaver-Burk	Lecture: Enzymes Kinetics Questions	Lecture: Enzyme Regulation	<u>Lab #5:</u> Enzyme Kinetics (preview of lab)
H W	Read over NOTES; Read over Lab #5 (yes, I know it's long!)	Read over NOTES; Read Lab #5	Read over NOTES; Read Lab #5	Read over NOTES; Read Lab #5	Read Lab #5; Complete Pre-Lab Questions on pages 5 and 6 Be ready to work on Monday!

	MONDAY October 30 Day 6	TUESDAY October 31 Day 1	WEDNESDAY November 1 Day 2	THURSDAY November 2 Day 3	FRIDAY November 3 Day 4
M O L E B I O	<u>Lab #5:</u> Enzyme Kinetics establishment of standard Curve; cellobiase reaction with and without enzyme	<u>Lab #5:</u> Enzyme Kinetics effect of temperature on cellobiase reaction	<u>Lab #5:</u> Enzyme Kinetics effect of pH on cellobiase reaction	<u>Lab #5:</u> Enzyme Kinetics effect of enzyme concentration on cellobiase reaction	<u>Lab #5:</u> Enzyme Kinetics effect of substrate concentration on cellobiase reaction
H W	Finish Activity 1 questions and graphs; Lab #5 due next Monday	Finish Activity 2 questions and graphs; Lab #5 due next Monday	Finish Activity 3 questions and graphs; Lab #5 due next Monday	Finish Activity 4 questions and graphs; Lab #5 due next Monday EXAM #3 MONDAY	Finish Activity 5 questions and graphs; Lab #5 due Monday EXAM #3 MONDAY

	MONDAY November 6 Day 5	TUESDAY November 7 Day 6	WEDNESDAY November 8 Day 1	THURSDAY November 9 Day 2	FRIDAY November 10
M O L E B I O	<p>EXAM #3</p> <p>ALL ABOUT ENZYMES!</p>	<p>Lecture: DNA</p> <p>History of Discovery</p>	<p>Lecture: DNA</p> <p>DNA Structure & Replication</p>	<p>Lecture: DNA</p> <p>DNA Structure & Replication</p>	
H W	<p>Read pg. 3 - 17; 24 - 34; 36 - 43</p>	<p>Read pg. 3 - 17; 24 - 34; 36 - 43</p>	<p>Read pg. 3 - 17; 24 - 34; 36 - 43</p>	<p>Read pg. 3 - 17; 24 - 34; 36 - 43</p> <p>Read 192 - 198</p>	

	MONDAY November 13 Day 3	TUESDAY November 14 Day 4	WEDNESDAY November 15 Day 5	THURSDAY November 16 Day 6	FRIDAY November 17 Day 1
M O L E B I O	Lecture: DNA Amplification via PCR	Lecture: DNA DNA Sequencing	Lecture: DNA Human Genome Project	Lecture: Basics of Gel Electrophoresis (Part II)	Lab #6: PV92 PCR Bioinformatics (protocol review)
H W	Read pg. 3 - 17; 24 - 34; 36 - 43 Read 192 - 198	Read pg. 3 - 17; 24 - 34; 36 - 43 Read Lab #6	Read pg. 3 - 17; 24 - 34; 36 - 43 Read Lab #6	Read pg: 192 - 198 Read Lab #6	Be ready for Part I of Lab #6 tomorrow!

	MONDAY November 20 Day 2	TUESDAY November 21 Day 3	WEDNESDAY November 22	THURSDAY November 23	FRIDAY November 24
M O L E B I O	<p><u>Lab #6:</u></p> <p>PV92 PCR Bioinformatics</p> <p>Part I: cheek cell DNA template preparation</p>	<p><u>Lab #6:</u></p> <p>PV92 PCR Bioinformatics</p> <p>Part II: amplification of DNA via polymerase chain reaction (PCR)</p>			
H W	<p>Read Lab #6</p> <p>Answer Questions #1 - 3 on page 27</p>	<p>Read Lab #6</p> <p>Lab #6 due next Wednesday</p> <p>Answer Questions #1 - 5 on page 28</p>			

	MONDAY November 27 Day 4	TUESDAY November 28 Day 5	WEDNESDAY November 29 Day 6	THURSDAY November 30 Day 1	FRIDAY December 1 Day 2
M O L E B I O	<p><u>Lab #6:</u> PV92 PCR Bioinformatics</p> <p>Part III: gel electrophoresis of amplified PCR samples</p>	<p><u>Lab #6:</u> PV92 PCR Bioinformatics</p> <p>Part IV: analysis and interpretation of results</p>	<p>Lecture: Protein Synthesis</p> <p>Defining a Gene; RNA</p>	<p>Lecture: Protein Synthesis</p> <p>Transcription</p>	<p>Lecture: Protein Synthesis</p> <p>Translation</p>
H W	<p>Read Lab #6 Lab #6 due Wednesday Answer Questions #1 - 4 on page 29</p>	<p>Read pg. 53 - 58; 65 - 67 Lab #6 due tomorrow Answer Questions #1 - 2 and finish table on page 30</p>	<p>Read pg. 53 - 58; 65 - 67</p>	<p>Read pg. 53 - 58; 65 - 67</p>	<p>Read pg. 53 - 58; 65 - 67 EXAM #4 TUESDAY</p>

	MONDAY December 4 Day 3	TUESDAY December 5 Day 4	WEDNESDAY December 6 Day 5	THURSDAY December 7 Day 6	FRIDAY December 8 Day 1
M O L E B I O	Lecture: Protein Synthesis Mutations	EXAM #4 DNA AND PROTEIN SYNTHESIS	Lecture: Restriction Enzymes	<u>Lab #7:</u> Restriction Enzyme Simulation	<u>Lab #8:</u> Restriction Enzyme Simulation using NEB Cutter
H W	Read pg. 53 - 58; 65 - 67 EXAM #4 TOMORROW	Read pg. 107 - 115 in DNA Science Read through Labs #7, 8, and 9	Read pg. 107 - 115 in DNA Science Read through Labs #7, 8, and 9	Lab #7 due tomorrow; Read Lab #8	Lab #8 due tomorrow; Read Lab #9 (textbook pages 351 - 374) Lab #9: Do Questions 1 - 6

	MONDAY December 11 Day 2	TUESDAY December 12 Day 3	WEDNESDAY December 13 Day 4	THURSDAY December 14 Day 5	FRIDAY December 15 Day 6
M O L E B I O	<p><u>Lab #9:</u> DNA Restriction Analysis</p> <p>run restriction digest prepare: 10X TBE stock buffer dilute 1X running buffer 25 mL of 0.8% agarose</p>	<p><u>Lab #9:</u> DNA Restriction Analysis</p> <p>prepare gel before class load and run gel after class* visualization with SYBR stain</p>	<p><u>Lab #9:</u> DNA Restriction Analysis</p> <p>analyze results</p>	<p><u>Lab #9:</u> DNA Restriction Analysis</p> <p>re-analyze results using Logger Pro</p>	<p><u>Lab #10:</u> Effects of DNA Methylation on Restriction</p> <p>discuss concept of lab review protocol</p>
H W	<p>Read Lab #9 (textbook pages 351 - 374)</p> <p>Come in to prepare gel before class! Get Logger Pro!</p>	<p>Read Lab #9 (textbook pages 351 - 374)</p> <p>Lab #9 due Friday Do Questions 7 - 10</p>	<p>Read Lab #9 (textbook pages 351 - 374)</p> <p>Lab #9 due Friday Do Questions 11 - 14</p>	<p>Read Lab #10 (textbook pages 375 - 384)</p> <p>Lab #9 with Logger Pro extension due tomorrow</p>	<p>Read Lab #10 (textbook pages 375 - 384)</p>

	MONDAY December 18 Day 1	TUESDAY December 19 Day 2	WEDNESDAY December 20 Day 3	THURSDAY December 21 Day 4	FRIDAY December 22 Day 5
M O L E B I O	<p>Lab #10:</p> <p>Effects of DNA Methylation on Restriction</p> <p>run methylase reaction; prepare 0.8% agarose</p>	<p>Lab #10:</p> <p>Effects of DNA Methylation on Restriction</p> <p>run restriction reaction</p>	<p>Lab #10:</p> <p>Effects of DNA Methylation on Restriction</p> <p>cast gels in boxes; load and run gel; visualize with SYBR-stain</p>	<p>Lab #10:</p> <p>Effects of DNA Methylation on Restriction</p> <p>analyze fragments; clean up stations</p>	(catch up day)
H W	<p>Lab #10 due Friday: Questions #1 - 6 on page 383 - 384</p>	<p>Lab #10 due Friday: Questions #1 - 6 on page 383 - 384</p>	<p>Lab #10 due Friday: Questions #1 - 6 on page 383 - 384</p>	<p>Lab #10 due tomorrow: Questions #1 - 6 on page 383 - 384</p>	<p>Read pages: 130 - 131; 116 - 129 in DNA Science</p>

	MONDAY December 26	TUESDAY December 27	WEDNESDAY December 28	THURSDAY December 29	FRIDAY December 30
M O L E B I O					
H W					

	MONDAY January 1	TUESDAY January 2 Day 6	WEDNESDAY January 3 Day 1	THURSDAY January 4 Day 2	FRIDAY January 5 Day 3
M O L E B I O		Lecture: Bacterial Genetics	Lecture: Bacterial Genetics	Lecture: Bacterial Genetics	<u>Lab #11:</u> Engineering a Plasmid
H W		Read pages: 130 - 131; 116 - 129 in DNA Science	Read pages: 130 - 131; 116 - 129 in DNA Science	Read Lab #11	Lab #11 due Monday

	MONDAY January 8 Day 4	TUESDAY January 9 Day 5	WEDNESDAY January 10 Day 6	THURSDAY January 11 Day 1	FRIDAY January 12 Day 2
M O L E B I O	<p>Plasmid Mapping Activity 1</p>	<p>Plasmid Mapping Activity 2</p>	<p>Prepping for Bacterial Culture Techniques Labs using an autoclave preparing LB plates preparing LB/amp plates</p>	<p>Prepping for Bacterial Culture Techniques Labs using an autoclave preparing group bottles of LB broth setting up the group stations reviewing the tools of microbiology</p>	<p>Prepping for Bacterial Culture Techniques Labs relearning learning serial dilutions reviewing the lab protocols</p>
H W	<p>Plasmid Mapping Activity 1 due tomorrow;</p>	<p>Plasmid Mapping Activity 2 due next Tuesday; Read pages 331 - 350 (Lab #2*) in DNA Science</p>	<p>Plasmid Mapping Activity 2 due Tuesday; Read Labs #12 - 14; Read pages 116-119, 331-350</p>	<p>Plasmid Mapping Activity 2 due Tuesday; Read Labs #12 - 14; Read pages 116-119, 331-350</p>	<p>Plasmid Mapping Activity 2 due Tuesday; Read Labs #12 - 14; Read pages 116-119, 331-350</p>

	MONDAY January 15	TUESDAY January 16 Day 3	WEDNESDAY January 17 Day 4	THURSDAY January 18 Day 5	FRIDAY January 19 Day 6
M O L E B I O		<p><u>Lab #12:</u> Bacterial Culture Techniques isolation of individual <i>E. coli</i> MM294 colonies understanding antibiotic resistance</p>	<p><u>Lab #12 - 13:</u> Bacterial Culture Techniques observing effects of antibiotic resistance preparation of an overnight suspension culture</p>	<p><u>Lab #13 - 14:</u> Bacterial Culture Techniques determining the number of individual cells in an overnight culture serial dilutions in practice</p>	<p><u>Lab #14:</u> Bacterial Culture Techniques calculation of cells in suspension</p>
H W		<p>Read Labs #12 - 14; Read pages 116-119, 331-350</p>	<p>Lab #12 Questions; Read Lab #13 - 14</p>	<p>Lab #13 Questions; Read Lab # 14</p>	<p>Lab #14 Questions; Read Lab #15</p>

	MONDAY January 22	TUESDAY January 23	WEDNESDAY January 24	THURSDAY January 25	FRIDAY January 26
M O L E B I O	REGENTS WEEK – NO CLASSES				
H W					

	MONDAY January 29 Day 1	TUESDAY January 30 Day 2	WEDNESDAY January 31 Day 3	THURSDAY February 1 Day 4	FRIDAY February 2 Day 5
M O L E B I O	<p><u>Lab #15:</u> Bacterial Culture Techniques (media making day) reviewing the Part A and Part B protocols; setting up the schedules for sample readings preparation of tubes and plates for tomorrow</p>	<p><u>Lab #15:</u> Bacterial Culture Techniques <u>Part A</u> - determination of <i>E. coli</i> growth pattern via spectrophotometry</p>	<p><u>Lab #15:</u> Bacterial Culture Techniques analysis of Part A setting up the schedules for sample readings preparation of tubes and plates for tomorrow</p>	<p><u>Lab #15:</u> Bacterial Culture Techniques <u>Part B</u> - determination of <i>E. coli</i> cell count via spectrophotometry and serial dilutions</p>	<p><u>Lab #15:</u> Bacterial Culture Techniques analysis of part B</p>
H W	Be ready for Lab #15 Part A tomorrow;	Be ready for Lab #15 analysis	Be ready for Lab #15 Part B tomorrow	Be ready for Lab #15 Part B tomorrow	Lab #15 due Monday; Read pages: 122 - 125 385 - 398 (DNA Science Lab #5*)

		MONDAY February 5 Day 6	TUESDAY February 6 Day 1	WEDNESDAY February 7 Day 2	THURSDAY February 8 Day 3	FRIDAY February 9 Day 4
M O L E B I O		Making Media LB plates LB/amp plates discuss theory of bacterial transformation	Making Media LB plates LB/amp plates review transformation procedure	<u>Lab #16:</u> Rapid Colony Transformation of <i>E. coli</i> with Plasmid DNA streaking starter plates	<u>Lab #16:</u> Rapid Colony Transformation of <i>E. coli</i> with Plasmid DNA transformation using pAMP	<u>Lab #16:</u> Rapid Colony Transformation of <i>E. coli</i> with Plasmid DNA analysis of results using pAMP streaking starter plates [pGREEN] protocol
	H W	Read pages: 385 – 398 (DNA Science Lab #5*) Read “Transformations” handout	Read pages: 385 – 398 (DNA Science Lab #5*) Read “Transformations” handout	Read pages: 385 – 398 (DNA Science Lab #5*)	Read pages: 385 – 398 (DNA Science Lab #5*)	Answer questions #1 – 4 on pages 395 – 396

	MONDAY February 12 Day 5	TUESDAY February 13 Day 6	WEDNESDAY February 14 Day 1	THURSDAY February 15 Day 2	FRIDAY February 16 Day 3
M O L E B I O	<p>Lab #16: Rapid Colony Transformation of <i>E. coli</i> with Plasmid DNA</p> <p>transformation with pGREEN using different heat shock times</p>	<p>Lab #16: Rapid Colony Transformation of <i>E. coli</i> with Plasmid DNA</p> <p>analysis of results using pGREEN and different heat shock times streaking starter plates</p> <p>Bonnie Bassler Video</p>	<p>Lab #16: Rapid Colony Transformation of <i>E. coli</i> with Plasmid DNA</p> <p>transformation using pVIB</p>	<p>Lab #16: Rapid Colony Transformation of <i>E. coli</i> with Plasmid DNA</p> <p>analysis of results using pVIB</p>	<p>EXAM #5 DNA RESTRICTION ANALYSIS & BACTERIAL TRANSFORMATION</p> <p>make media for Labs #17 - 18 streak starters</p>
H W	<p>Answer questions #5 - 6 on pages 395 - 396</p>	<p>Read pVIB handout; EXAM #5 FRIDAY</p>	<p>Read pVIB handout; EXAM #5 FRIDAY</p>	<p>Read pVIB handout; EXAM #5 TOMORROW</p>	<p>Read pages: 399 - top of 407 (DNA Science Lab #6*)</p>

	MONDAY February 19	TUESDAY February 20	WEDNESDAY February 21	THURSDAY February 22	FRIDAY February 23
M O L E B I O					
H W					

		MONDAY February 26 Day 4	TUESDAY February 27 Day 5	WEDNESDAY February 28 Day 6	THURSDAY March 1 Day 1	FRIDAY March 2 Day 2
M O L E B I O	<p>Making Media</p> <p>make media for Labs #17 - 18 streak starters</p>	<p>Lab #17:</p> <p>Assay for an Antibiotic Resistance Enzyme</p> <p>transformation of MM294 with pAMP review lab protocol during incubation</p>	<p>Lab #17:</p> <p>Assay for an Antibiotic Resistance Enzyme</p> <p>preparation of an overnight suspension culture of transformed <i>E.coli</i>+pAMP review tomorrow's protocol</p>	<p>Lab #17:</p> <p>Assay for an Antibiotic Resistance Enzyme</p> <p>preparation of control and pAMP 'sup' via centrifugation review tomorrow's protocol</p>	<p>Lab #17:</p> <p>Assay for an Antibiotic Resistance Enzyme</p> <p>assay of β-lactamase via spectrophotometry streak starter plates review Monday's protocol</p>	
	H W	<p>Read pages: 399 - top of 407 (DNA Science Lab #6*)</p>	<p>Read pages: 399 - top of 407 (DNA Science Lab #6*)</p>	<p>Read pages: 399 - top of 407 (DNA Science Lab #6*)</p>	<p>Read pages: 399 - top of 407 (DNA Science Lab #6*)</p>	<p>Answer questions #1 - 6 on pages 406 - 407</p> <p>Read pages: 411 - 422 (DNA Science Lab #7*)</p>

	MONDAY March 5 Day 3	TUESDAY March 6 Day 4	WEDNESDAY March 7 Day 5	THURSDAY March 8 Day 6	FRIDAY March 9 Day 1
M O L E B I O	<p>Lab #18:</p> <p>Purification and Identification of Recombinant GFP</p> <p>transformation of MM294 with pGREEN</p> <p>review lab concept during incubation</p>	<p>Lab #18:</p> <p>Purification and Identification of Recombinant GFP</p> <p>preparation of an overnight suspension culture of transformed <i>E.coli</i>+pGREEN</p> <p>review tomorrow's protocol</p>	<p>Lab #18:</p> <p>Purification and Identification of Recombinant GFP</p> <p>lysozyme used to lysate cells to release proteins</p>	<p>Lab #18:</p> <p>Purification and Identification of Recombinant GFP</p> <p>purification of GFP by HIC (Hydrophobic Interaction Chromatography)</p>	<p>Lab #18:</p> <p>Purification and Identification of Recombinant GFP</p> <p>PAGE analysis of purified GFP</p> <p>stain/destain with Coomassie Blue</p>
H W	<p>Read pages: 411 - 422 (DNA Science Lab #7*)</p> <p>Review pages: 340 - 343 (DNA Science Lab #2B*)</p>	<p>Review Procedure on pages 414 - 416</p>	<p>Review Procedure on pages 414 - 416</p>	<p>Review Procedure on pages 419 - 420</p>	<p>Answer questions 1 - 4 on page 416</p> <p>Read pages: 125 - 127 423 - 441 (DNA Science Lab #8*)</p>

MONDAY March 12 Day 2		TUESDAY March 13 Day 3		WEDNESDAY March 14 Day 4		THURSDAY March 15 Day 5		FRIDAY March 16 Day 6		
M O L E B I O	Making Media LB plates; LB/amp plates; LB/kan plates; LB/amp/kan plates LB broth ; LB/amp broth toothpicks; Q-tips GTE (glucose/tris/EDTA); TE (tris/EDTA); SDS/NaOH; KOAc; isopropanol; ethanol Streak Starter Plates				<u>Lab #19:</u> Purification and Identification of Plasmid DNA rapid colony transformation of <i>E.coli</i> with pAMP		<u>Lab #19:</u> Purification and Identification of Plasmid DNA calculating transformation efficiency preparation of an overnight suspension culture of our <i>E.coli</i> MM294/pAMP transformants		<u>Lab #19:</u> Purification and Identification of Plasmid DNA preparation of duplicate minipreps (PART I)	
	Read pages: 423 - 441		Read pages: 423 - 441 Review pages: (391 - 395)		Review page: 423 - 441 (342)		Review pages: 423 - 441 (427 - 428)		Review pages: 423 - 441 (428 - 429)	
H W										

	MONDAY March 19 Day 1	TUESDAY March 20 Day 2	WEDNESDAY March 21 Day 3	THURSDAY March 22 Day 4	FRIDAY March 23
M O L E B I O	<p>Lab #19: Purification and Identification of Plasmid DNA</p> <p>preparation of duplicate minipreps (PART II)</p>	<p>Lab #19: Purification and Identification of Plasmid DNA</p> <p>set up and run restriction digest of purified pAMP with <i>Bam</i>HI/ <i>Hind</i>III</p> <p>prepare 1X TBE buffer prepare 0.8% agarose</p>	<p>Lab #19: Purification and Identification of Plasmid DNA</p> <p>separate DNA fragments by electrophoresis</p>	<p>Labs #20 - 23: Discussion of the next Lab Stream</p> <ul style="list-style-type: none"> • Ligation • Transformation • Replica Plating • Miniprep 	
H W	<p>Review pages: 423 - 441 (431 - 435)</p> <p>Answer questions 1 - 3 on page 430</p>	<p>Review pages: 423 - 441 (435 - 436)</p>	<p>Review pages: 437 - 439</p> <p>Answer questions 1 - 3a, b, c on page 439 (use Logger Pro for #1)</p>	<p>Review pages: 443 - 455 (447 - 448) (DNA Science Lab #9*)</p>	

	MONDAY March 26 Day 5	TUESDAY March 27 Day 6	WEDNESDAY March 28 Day 1	THURSDAY March 29 Day 2	FRIDAY March 30
M O L E B I O	<p>Lab #20:</p> <p>Recombination of Antibiotic Resistance Genes</p> <p>restriction digest of the plasmids pAMP and pKAN</p> <p>prepare 0.8% agarose</p>	<p>Lab #20:</p> <p>Recombination of Antibiotic Resistance Genes</p> <p>separate digested plasmid fragments by electrophoresis ("The Prudent Control")</p>	<p>Lab #20:</p> <p>Recombination of Antibiotic Resistance Genes</p> <p>ligation of digested plasmid DNA</p> <p>prepare 0.8% agarose</p>	<p>Lab #20:</p> <p>Recombination of Antibiotic Resistance Genes</p> <p>separate ligated DNA fragments by electrophoresis</p> <p>streak starters</p>	
H W	<p>Review pages: 443 - 455 (448 - 451) (DNA Science Lab #9*)</p>	<p>Review pages: 443 - 455 (453 - 454)</p>	<p>Review pages: 443 - 455 (454*)</p> <p>Read pages: 457 - 470 (DNA Science Lab #10*)</p>	<p>Answer questions 1 - 3, 5 - 7 on page 455</p> <p>Read pages: 457 - 470 (DNA Science Lab #10*)</p>	

	MONDAY April 2	TUESDAY April 3	WEDNESDAY April 4	THURSDAY April 5	FRIDAY April 6
M O L E B I O					
H W					

	MONDAY April 9 Day 3	TUESDAY April 10 Day 4	WEDNESDAY April 11 Day 5	THURSDAY April 12 Day 6	FRIDAY April 13 Day 1
M O L E B I O	<p>Lab #21: Transformation of <i>E. coli</i> MM294 with Recombinant DNA</p> <p>streak starter plates of <i>E.coli</i> MM294</p>	<p>Lab #21: Transformation of <i>E. coli</i> MM294 with Recombinant DNA</p> <p>prepare overnight culture of <i>E.coli</i> MM294</p>	<p>Lab #21: Transformation of <i>E. coli</i> MM294 with Recombinant DNA</p> <p>preparation of a 250 mL mid-log suspension culture (pg. 347-348) preparation of competent cells (pg. 460-462)</p>	<p>Lab #21: Transformation of <i>E. coli</i> MM294 with Recombinant DNA</p> <p>perform <i>E. coli</i> transformation with recombinant DNA</p>	<p>Lab #21: Transformation of <i>E. coli</i> MM294 with Recombinant DNA</p> <p>analysis of transformation</p> <p>Answer Questions #1-4 on page 469-470</p>
H W	<p>Review pages: 457 - 470 (340 - 342)</p>	<p>Review pages: 457 - 470 (347 - 348) (460 - 462)</p>	<p>Review pages: 457 - 470 (463 - 468)</p>	<p>Review pages: 457 - 470 (468 - 470)</p>	<p>Complete Questions #1, 3, & 4 on pg. 469 - 470</p> <p>Read pages: 473 - 477 (DNA Science Lab #11*)</p>

		MONDAY April 16 Day 2	TUESDAY April 17 Day 3	WEDNESDAY April 18 Day 4	THURSDAY April 19 Day 5	FRIDAY April 20 Day 6
M O L E B I O		<p>Lab #22:</p> <p>Replica Plating to Identify Mixed <i>E.coli</i> Populations</p>	<p>Lab #22:</p> <p>Replica Plating to Identify Mixed <i>E.coli</i> Populations</p> <p>analysis of replica plates start prepping the Miniprep Answer Questions #1-3 on page 477</p>	<p>Lab #23:</p> <p>Purification and Identification of Recombinant Plasmid DNA</p> <p>preparation of an overnight suspension culture of our transformed <i>E.coli</i></p>	<p>Lab #23:</p> <p>Purification and Identification of Recombinant Plasmid DNA</p> <p>preparation of duplicate minipreps (PART I)</p>	<p>Lab #23:</p> <p>Purification and Identification of Recombinant Plasmid DNA</p> <p>preparation of duplicate minipreps (PART II)</p>
	H W	<p>Review pages: 473 - 477</p>	<p>Complete Questions #1 - 3 on pg. 477</p> <p>Read pages: 481 - 499 (DNA Science Lab #12*)</p> <p>Review pages: (340 - 342)</p>	<p>Review pages: 481 - 499 (484 - 485)</p>	<p>Review pages: 481 - 499 (486)</p>	<p>Review pages: 481 - 496 (487 - 490)</p>

	MONDAY April 23 Day 1	TUESDAY April 24 Day 2	WEDNESDAY April 25 Day 3	THURSDAY April 26 Day 4	FRIDAY April 27 Day 5
M O L E B I O	<p><u>Lab #23:</u> Purification and Identification of Recombinant Plasmid DNA</p> <p>set up and run restriction digest of purified ligation products with <i>Bam</i>HI / <i>Hind</i>III</p>	<p><u>Lab #23:</u> Purification and Identification of Recombinant Plasmid DNA</p> <p>separate DNA fragments by electrophoresis</p>	<p><u>Lab #23:</u> Purification and Identification of Recombinant Plasmid DNA</p> <p>recap of our 'simple recombinant'</p>	<p>EXAM #6 LIGATION OF, TRANSFORMATION WITH, AND PURIFICATION/ IDENTIFICATION OF RECOMBINANT DNA</p>	<p><u>Lab #24:</u> Creating a Library of Bacteriophage λ and pBLU</p> <p>Making Media: LB plates LB/amp/X-gal plates CaCl₂</p>
H W	<p>Review pages: 481 - 496 (490 - 491)</p>	<p>Review pages: 481 - 496</p>	<p>Based on your analysis, make scale restriction maps of your M1 and M2 plasmids. #9 on page 496</p>	<p>Read the bottom of page 455!</p>	<p>Review pages: (447 - 448)</p>




	MONDAY April 30 Day 6	TUESDAY May 1 Day 1	WEDNESDAY May 2 Day 2	THURSDAY May 3 Day 3	FRIDAY May 4 Day 4
M O L E B I O	<p>Lab #24: Creating a Library of Bacteriophage λ and pBLU</p> <p>restriction digest of the plasmid pBLU and λ DNA with <i>Bam</i>HI and <i>Hind</i>III; predict fragments of pBLU and λ DNA prepare 0.8% agarose</p>	<p>Lab #24: Creating a Library of Bacteriophage λ and pBLU</p> <p>separate digested plasmid fragments by electrophoresis ("The Prudent Control")</p>	<p>Lab #24: Creating a Library of Bacteriophage λ and pBLU</p> <p>ligation of digested plasmid DNA prepare 0.8% agarose</p>	<p>Lab #24: Creating a Library of Bacteriophage λ and pBLU</p> <p>separate ligated DNA fragments by electrophoresis</p>	<p>Lab #24: Creating a Library of Bacteriophage λ and pBLU</p> <p>reviewing classic protocol for preparing competent cells</p>
H W	Review pages: (448 - 450)	Review pages: (454)	Review pages: (454)	Review pages: (336 - 338)	Review pages: (336 - 338)

	MONDAY May 7 Day 5	TUESDAY May 8 Day 6	WEDNESDAY May 9 Day 1	THURSDAY May 10 Day 2	FRIDAY May 11 Day 3
M O L E B I O	<p>Lab #24: Creating a Library of Bacteriophage λ and pBLU</p> <p>streak starter plates of <i>E.coli</i> MM294</p>	<p>Lab #24: Creating a Library of Bacteriophage λ and pBLU</p> <p>prepare overnight culture of <i>E.coli</i> MM294</p>	<p>Lab #24: Creating a Library of Bacteriophage λ and pBLU</p> <p>preparation of a 250 mL mid-log suspension culture (pg. 347-348)</p> <p>preparation of competent cells (pg. 460-462)</p>	<p>Lab #24: Creating a Library of Bacteriophage λ and pBLU</p> <p>perform <i>E. coli</i> transformation with recombinant DNA</p>	<p>Lab #24: Creating a Library of Bacteriophage λ and pBLU</p> <p>analysis of transformation</p> <p>selection of colonies for miniprep and re-streak on new LB/amp/X-gal plates</p>
H W	<p>Review pages: (342)</p>	<p>Review pages: (347 - 348) (460 - 462)</p>	<p>Review pages: (463 - 468)</p>	<p>Review pages: (468 - 470) (336 - 338)</p>	<p>Review pages: (464 - 485)</p>
	<p>Period 1: -----</p> <p>Period 6: -----</p>	<p>Period 1: -----</p> <p>Period 6: -----</p>	<p>Period 1: -----</p> <p>Period 6: -----</p>	<p>Period 1: -----</p> <p>Period 6: -----</p>	<p>Period 1: -----</p> <p>Period 6: -----</p>

	MONDAY May 14 Day 4	TUESDAY May 15 Day 5	WEDNESDAY May 16 Day 6	THURSDAY May 17 Day 1	FRIDAY May 18 Day 2
M O L E B I O	<p><u>Lab #24:</u> Creating a Library of Bacteriophage λ and pBLU</p> <p>preparation of duplicate minipreps (PART I)</p>	<p><u>Lab #24:</u> Creating a Library of Bacteriophage λ and pBLU</p> <p>preparation of duplicate minipreps (PART II)</p>	<p><u>Lab #24:</u> Creating a Library of Bacteriophage λ and pBLU</p> <p>set up and run restriction digest of purified plasmid DNA with <i>Bam</i>HI / <i>Hind</i>III</p> <p>prepare 0.8% agarose</p>	<p><u>Lab #24:</u> Creating a Library of Bacteriophage λ and pBLU</p> <p>separate purified plasmid DNA fragments by electrophoresis</p>	<p><u>Lab #24:</u> Creating a Library of Bacteriophage λ and pBLU</p> <p>analyze fragments</p> <p>catalog library!</p>
H W	Review pages: (486)	Review pages: (487 - 490)	Review pages: (490 - 491)		
	<p>Period 1: -----</p> <p>Period 6: -----</p>	<p>Period 1: -----</p> <p>Period 6: -----</p>	<p>Period 1: -----</p> <p>Period 6: -----</p>	<p>Period 1: -----</p> <p>Period 6: -----</p>	<p>Period 1: -----</p> <p>Period 6: Devin</p>

	MONDAY May 21 Day 3	TUESDAY May 22 Day 4	WEDNESDAY May 23 Day 5	THURSDAY May 24 Day 6	FRIDAY May 25 Day X
M O L E B I O	<p><u>Lab #24:</u> Creating a Library of Bacteriophage λ and pBLU</p> <p>(CATCH UP DAY!)</p>	<p><u>Lab #24:</u> Creating a Library of Bacteriophage λ and pBLU</p> <p>(CATCH UP DAY!)</p>	<p>Making Media</p> <p>lots of LB/amp plates & nutrient agar plates</p> <p>streak starters for transformation of <i>E. coli</i> MM294 with PM1/PM2</p>	<p>Dr. Tsien Lecture on GFP</p> <p>streak starter plates</p>	
H W			Read Lab #25	Read Lab #25	

	MONDAY May 28	TUESDAY May 29 Day 1	WEDNESDAY May 30 Day 2	THURSDAY May 31 Day 3	FRIDAY June 1 Day 4
M O L E B I O		<p><u>Lab #25:</u></p> <p>Fluorescent <i>E. coli</i> Art Show!</p> <p>transformation of <i>E. coli</i> MM294 with 'fluorescent' plasmids</p>	<p><u>Lab #25:</u></p> <p>Fluorescent <i>E. coli</i> Art Show!</p> <p>preparation of fluorescent lawns</p>	<p><u>Lab #25:</u></p> <p>Fluorescent <i>E. coli</i> Art Show!</p> <p>painting with bacteria</p>	<p><u>Lab #25:</u></p> <p>Fluorescent <i>E. coli</i> Art Show!</p>
H W		Read Lab #25	Answer Lab #25 Questions		

	MONDAY June 4 Day 5	TUESDAY June 5 Day 6	WEDNESDAY June 6 Day 1	THURSDAY June 7 Day 2	FRIDAY June 8 Day 3
M O L E B I O	SENIOR DAY OF OPTIONAL ATTENDANCE DUE TO A CRITICAL MASS OF APATHY	<u>Lab #26:</u> Swabbing the School... 	<u>Lab #26:</u> Swabbing the School... 	<u>Lab #26:</u> Swabbing the School... 	SENIOR PICNIC
H W					