Learning Outcomes Department of Chemistry and Biochemistry

Biochemistry Major (typical electives) Pharmacological Chemistry Major

This document outlines overarching learning objectives for undergraduate students majoring in Biochemistry or Pharmacological Chemistry in the Department of Chemistry and Biochemistry at UC San Diego. The table below lists a specific goal with corresponding broad objectives and notes the courses in which the goal is introduced to students (initial assessment) as well as with one or two later courses wherein students may be further assessed. Note that the last of the general goals, M, does not appear as a column, because it refers to an objective that applies more or less equally to all goals and can be assessed only after students graduate.

Broad Goals or Learning Objectives:

- A. Have firm foundations in the fundamentals and applications of current chemical theories for the physical world.
- B. Use molecular understanding in fields that are based upon chemistry: biology, environmental science, and engineering.
- C. Be skilled in problems solving, critical thinking, and analytical reasoning.
- D. Know the proper procedures and regulations for safe handling and use of chemicals and follow the proper procedures and regulations for safety when using chemicals.
- E. Design, carry out, record, and analyze the results of chemical experiments.
- F. Use a broad variety of modern instrumentation and classical techniques in the course of experimentation.
- G. Interpret and evaluate results critically. Identify and quantify uncertainties in measurements and limitations in methodologies.
- H. Use modern library searching and retrieval methods to obtain information about a topic, chemical, chemical technique, or an issue relating to chemistry, going beyond textbooks and common handbooks.
- I. Communicate results of work to chemists and non-chemists, including respect for the tradition of careful citation of prior contributions, both orally and in effective writing.
- J. Collaborate effectively as part of a team to solve problems, debate different points of view, and interact productively with a diverse group of team members.
- K. Understand the ethical, historic, philosophical, and environmental dimensions of problems and issues facing chemists.
- L. Be able to identify and solve chemical problems and explore new areas of research.
- M. Find gainful employment in industry or government, be accepted at graduate or professional schools, or find employment in school systems as instructors or administrators.

Specific Goals:

Objective		A	В	\mathbf{C}	D	\mathbf{E}	\mathbf{F}	G	H	T	J	K	L	Where
Objective	Taught	1.					-	•	4.1	_		-11		Assessed
Meet the objectives of introductory calculus as	Math													Many
specified by the Department of Mathematics	20A/B/	✓		✓										
	C/D													
Meet the objectives of elementary physics as	Physics													105A
specified by the Department of Physics	2A/B/D	✓		✓										
	2CL													
Recognize elemental symbols and place the more	6A	✓	√											All
common elements on a Periodic Chart		Ľ	•											
Use a Periodic Chart to predict elemental and atomic	6A													6C
properties, such as electronegativity, size, state of		✓												120A
matter, likely reaction partners														
Count molecules in units of moles and write balanced	6A	✓		✓										All
chemical reactions in terms of mole numbers		لنا		Ť										
Recognize a limiting reagent, calculate amounts of	6A	✓		/										6BL
reaction product and yield														
Recognize the differences among materials that are	6A	✓	✓											120A
metallic, ionic, or covalently bonded			•											
Use molecular orbital theory to explain differences	6A	✓												140A
among second row diatomic molecules		لنا												
Appreciate the role of nonbonding interactions, in	6A	✓	√											6C
particular with respect to solubilities			•											
Use quantum mechanical descriptions for electronic	120A													120B
orbitals and molecular symmetry principles to		✓												124
describe chemical bonding														
Use Lewis Diagrams to predict molecular	6A	✓												140A
connectivity														
Use valence shell repulsion theory to predict shapes	6A	✓												120A
of symmetric molecules														140A
Sketch 1s, 2s and 2p atomic orbitals and combine	6A	✓												140A
them to interpret sp3, sp2 and sp hybrid orbitals.	140A													156
Sketch molecular orbitals (bonding and antibonding)	140A													140B
for any 2-carbon molecule, with peripheral atoms,														156
showing the mathematical signs of the lobes and		✓												
approximate relative energies. Sketch pi molecular														
orbitals of conjugated systems. Sketch the structures														
of carbocations, carbanions and radicals.	1404	1												1.40D
Understand bond formation and bond energies, and	140A	✓												140B
predict which bonds are weak and which are strong.	140A													156
Extend valence shell repulsion theory to treat strain	140A	✓												140B
II	(0													156
Use a simplified crystal field theory to rationalize	6C	✓												120A
structure and reactivity of transition metal complexes and their colors when dissolved in water		*												
	120B													124
Use ligand field theory and other quantum methods	120 D	✓												223
to predict the molecular structures of transition metal		•												223
complexes and extend this to organometallics Solve the Schroedinger Equation for a 1-d harmonic	126													230
oscillator to derive eigenvalues and eigenfunctions.	133	✓		/										230
Note the equal-spaced energy levels	133			•										
Solve the Schroedinger equation for a 1-d square	126													230
well and for a rigid rotor, noting that energy levels	133	✓		✓										230
well and for a rigid rotor, noting that energy levels				•						1				ı

	0												
Solve the Schroedinger Equation for a Coulomb	126												230
potential, noting that energy levels are spaced more	133	✓		✓									
closely at high energies													
Explain energies and transitions for simple atoms at	126												230
an intermediate level	133	✓		✓									
Develop a proper quantum interpretation of bonding	126								_				230
for simple molecules	133												230
	6A								-	-			6BL
Use and be able to interconvert among the several	θA			✓									
ways of denoting solutions concentrations													6C
Use the four colligative properties to calculate	6A												6BL
concentrations or molar masses, depending on known		✓		✓									100A
information.													
State the 4 great laws of thermodynamics and explain	6B	✓	_										127
why they are considered great		V	√										131
Distinguish state functions from such non-quantities	6B								_				127
as heat and work	OB	✓											131
	127								\dashv	-			230
Manipulate partial derivatives of state quantities		✓		✓									230
using relations such as the Maxwell relations	131	$\vdash \vdash$							4			$\ \cdot \ $	
Calculate the idealized maximum efficiency of a	127		,	,									
heat engine or a refrigerator as deduced from a	131		✓	√									
reversible Carnot cycle													
Calculate the maximum efficiency of a less-than-	127		✓	<				Ţ					
ideal reversible cycle, such as those of Otto or Diesel	131		٧	٧									
Identify the fallacy in the creationists' erroneous	6B												127
assertion that evolution is inconsistent wit the Second		✓	✓										131
Law													151
Explain why it is that reactions that heat their	6B								-	-			127
	ОБ												
surroundings are likely to be spontaneous and why it		✓											131
is that even some that cool their surroundings can be													
spontaneous.													
Use Hess's Law to combine thermal energies for	6B												127
chemical reactions when one combines consecutive		✓	✓	✓									131
atomic combinations													
Use tables of free energies to compute equilibrium	6B		,	,									127
constants		✓	✓	√									131
Evaluate equilibrium constants from information	6B								_				127
about concentrations or partial pressures; or use	OD												131
		✓		./									131
equilibrium constants to deduce concentrations or		V		٧									
partial pressures at equilibrium, given some initial													
condition													
Distinguish strong and weak acids and bases	6B	✓	✓										140A
Convert between the pH scale and concentrations of	6B		✓	./				T	T				6BL
protons or proton acceptors in aqueous solution			•	•									100A
Carry out titrations to determine the pH of an	6B							1	T				100A
unknown aqueous solution to acceptable accuracy	6BL			✓	✓	✓	√	✓					
and precision.	(DL												
Generalize the concept of a titration to any chemical	114A	\vdash							+	+		⊢∦	100BL
			✓	✓									
or biochemical measurement	112A	$\vdash \vdash$				-			4		\perp		112B
Desire and prepare a pH buffer of required pH and	6B		✓		√	✓		✓					112A
ionic strength	100A												
Compare and contrast Arrhenius, Bronsted, and	6B	✓											140A
Lewis acids													
Write balanced equations for oxidation-reduction	6C		,	,				T	Ī				100A
reactions, including the participation of solvent water		✓	✓	√									120A
Use redox tables to predict the spontaneous direction	6C							<u> </u>	1			\Box	120A
for reactivity in redox reactions, and have some		✓	✓										12011
intuitive notions even without a table of potentials			•					-					
					i	I	1			1		1 1	

Calculate the reversible emf expected for an arbitrary	6C												127
redox reaction, using tables, for any combination of				✓									132
concentrations of solutes and pressures of gasses													
Deduce reaction rate laws and rate constants from	6C	✓		✓									127
initial rate data													132
Transform data from measurements of kinetic	6C			,									127
processes to produce a linear plot and deduce				✓									132
reaction order and rate constants from such plots	-60												105A
Explain the role of catalysts in a reaction and give	6C	✓											140B
some examples	-60												1.10.0
Distinguish addition polymers from condensation	6C	✓	✓										140C
polymers and give examples of each.	4444												44.5
Recognize and use Michaelis-Menten kinetic scheme	114A	✓	✓										116
And the single transfer to the second of	127												
Apply the principles of gravimetry to determine the	6BL			✓	✓	✓	✓	✓					
amount of analyte in an unknown sample	(DI												
Titrate a weak acid with a strong base to determine	6BL			✓	✓	✓	✓	✓					
the molar mass, pKa, and identity of the acid	6BL												
Determine the specific heat of a metal, the heat of fusion of water, and the heat of neutralization of an	ODL			✓	./	/	./						
acid-base reaction via coffee-cup calorimetry				•	•	•	•						
Use oxidation-reduction titration to determine the	6BL												
oxalate content in the iron oxalate complex	ODL			✓	✓	✓	✓	✓					
Synthesize an iron (III) oxalate complex	6BL				✓	_	√						
Use spectrophotometry to determine the iron content	6BL				·	_	•						
in the iron oxalate complex	ODL			✓	✓	✓	✓	✓					
Understand and follow a semimicro qualitative	6BL												
analysis scheme to characterize a mixture of common	ODL			1	✓	✓	√						
metal ions													
Investigate the atomic emission spectra of various	6BL												
elements						~	V						
Maintain a clearly written lab notebook as a	6BL												
permanent record of experimental results						~			✓				
Write a simple report in standard format emulating	6BL									√			100A
publication in a science journal										•			All labs
Demonstrate skill using a computer spreadsheet	100A						✓						105A
Demonstrate proficiency with computer graphing	100A						✓			✓			105A
Characterize reaction kinetics in a laboratory	100A				✓	✓	✓			✓			105A
Measure chemical equilibria in solution	100A			✓	✓	✓	✓	✓		✓			
Use electrochemical techniques and ion selective	100A			√	/			_					
electrodes to determine ion concentrations				•	•	•	•	•		v			
Use column chromatography to separate components	100A			√	./	~	✓	✓	√	✓			100B
of a mixture				٧	٧	٧	٧	٧	٧	٧			Others
Use gas chromatography to separate mixtures, using	100BL												
several different detection strategies, including mass	143B		✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	
spectrometry	143C												
Use high performance liquid chromatography to	100A		<	\	/	✓	100B						
separate mixtures	100BL				Ĺ								
Demonstrate proficiency in statistical analysis and	105A											_	100BL
error estimation beyond what was learned in the	100BL	✓	✓			✓		✓		✓		✓	105B
lower labs	1.40.1												1.407
Draw conformations of alkanes and cycloalkanes	140A												140B
(Newman projections, wedge/dotted-line structures).													156
Graph the relation between conformation and		✓		✓									
potential energy for these molecules. Predict													
preferred conformations, including those of													
substituted cyclohexanes. Calculate the ratio of													

Explain how conformations around bonds translate into global shape changes and dictate the overall structure of big molecules, emphasizing relevancy for biological structures. Recognize strain in various conformations and predict effect on stability and as a driving force for reactivity and rearrangements. Define and recognize stereoisomer, enantiomer, diastereomer, conformation, configuration, meso, epimer, resolution. Recognize inversion, retention and racemization. All these for any molecule. Sketch a molecule with a chiral center so as to show unambiguously the configuration using both Fischer projection and perspective drawing. Determine the configuration (R or S) of any chiral center from a perspective drawing. Calculate "specific rotation" from the experimental optical rotation and concentration. Determine the configuration (E or Z) of any double bond. Describe the formation and relative stabilities of carbocations as related to hyperconjugation. Convert IUPAC names of simple molecules to chemical structures. Analyze inter and intramolecular forces and estimate 140A 140B 140B 140A 140B 140A 140B 140A 140B 140A 140B 140A 140B
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contributor. 156 Analyze inter and intramolecular forces and estimate 140A 143
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a stubility, malting point and bailing point Describe
solubility, melting point and boiling point. Describe
the molecular events occurring during the processes
of dissolving, melting and boiling.
Explain the unique role of water as a solvent. 6A
Use the unique solvation properties of water to 140A 140A 140
predict or retrodict organic molecular structure with 140C
emphasis on molecules of biochemical interest
Estimate relative acidities and basicities of organic 140A 140
compounds based on estimation of the stabilities of 140B
their conjugate base and acid. Calculate the pH of a
solution of a weak acid or base from the analytical
concentration and Ka. Calculate the proportions of
protonated and non-protonated species at a given pH.
protonated and non-protonated species at a given pri.
Legate reactive cites within a malegula and decree
Locate reactive sites within a molecule and draw 140A 140B
correct electron-pushing arrows for reactions based 140B \(\)
on electronic properties and structure instead of rote
memorization of mechanisms.
Explain by words and equations the factors affecting 140A 140
the rate of a chemical reaction, including 140B 156
temperature. Analyze kinetic data and determine the
order of a reaction. Validate reaction mechanisms by
comparison with kinetic data.
Distinguish between kinetic and thermodynamic 140B 140
products of reactions. Explain reasons for obtaining 154 \checkmark 154
products of reactions. English reasons for octaming
one product rather than the other.
Use the concepts of delocalization and resonance for 140A 140

	1.405	11 1	1			1			-1		— г	1.400
Draw conclusions about a reaction mechanism from	140B											140C
the stereochemistry of the products. Given a	140C	✓		✓								154
proposed mechanism for a reaction, predict the	154											156
stereochemistry.												
Define and recognize regioselective, stereoselective	140A											140B
and stereospecific reactions. Describe resolution of a				/	✓	/						156
racemic mixture by converting it to a diastereomeric				٧	'	•						
mixture.												
Understand and explain the importance of chiral	140A								1			140B
recognition in biological systems.	140C		✓									114B
Distinguish nucleophiles from electrophiles and list	140A	+							+	-		140C
	II .											
examples of each. Write chemical equations to	140B											154
describe the currently accepted mechanism(s) for	155											156
major reactions: radical, SN1, SN2, E1, E2,												
electrophilic addition, electrophilic substitution,		✓										
conjugate addition, addition-elimination, pericyclic.												
Explain how each mechanism is deduced from												
experimental kinetic data and stereochemistry of the												
products. Be able to specify structures and energetics												ļ ļ
of intermediates in multistep reactions.												
Describe how the terms oxidation and reduction are	140A											143A
used in organic chemistry.		✓			Ī							1.2.2
Recognize and predict rearrangements of	140A	+							+			140B
carbocations.	156	✓										154
		╂							+	-	-	
Identify all major functional groups and the reactivity	140A											140C
of each.	140B	✓										152
		1										155
Identify the functional groups prominent in reactions	140C		√									114B
that biomolecules undergo.												
Conduct a retrosynthetic analysis of a given	140B											140C
compound and outline the forward steps and reagents	140C				✓	✓	✓	✓				155
that are required	152											
Be aware of the pervasiveness of organic substances	140A											140B
in the environment.	149B		✓								✓	140C
Identify and discuss some of the common polymers	140C								1			114A
and macromolecules, at a level more soils than in C.	1400		✓								✓	114D
UV-VIS: Use the terms chromophore, molar	140B	-							+			143D
	140 D						./					
absorptivity, wavelength at maximum, transition, pi							•					158
pi*, n - pi*.		4							_			
UV-VIS: Use UV-VIS data to calculate	6BL											143D
concentrations and assist in determining chemical	100A			√		✓	✓					158
structure.	140B											
UV-VIS: Explain the effect of conjugation on the	140B											143A
absorption wavelength by sketching the molecular												
orbitals and relative energies.												
IR: Describe the molecular transitions responsible for	140B	1						-	1	+		143A
the infrared absorption.	עטיי	✓			Ī							143A 143B
IR: Use the characteristic absorption frequencies	143A	+							+		+ +	143B
	II			./	Ī			√				ll .
(data provided) of functional groups to assist in	143C			✓	Ī	*		v				146
determining the structure of an unknown compound.	1.10=	\bot				 			_	4	\sqcup	1.10=
IR: Predict how electronic and structural factors	140B											143B
affect the infrared absorption of functional groups,	143A	✓		✓	Ī		✓					140C
particularly carbonyls.					L			l				158
IR: Explain the connection between infrared	140A		√									140C
absorption and the "greenhouse effect".	149A		•								*	173
NMR: Magnetic resonance of protons and carbon:	140B			,					1			143B
Identify the number of non-equivalent protons and				✓	Ī		✓	✓				140C
recently the number of non equivalent protons and	Ш				L	li .						1 700

	1	1			1							1	1.50
carbons in a given molecule based on symmetry.													158
Assign peaks of an NMR spectrum to likely chemical													
environments. Identify the relative numbers of													
protons of an unknown using integration. Identify the													
presence of neighboring protons from splitting													
patterns and coupling constants. Use NMR spectrum													
to elucidate the structure of an unknown compound.	1.400	-											1.105
NMR: Predict the NMR spectrum from a structure	140B		✓			✓	✓						143B
(number of peaks, multiplicity and chemical shift).	1.400												143C
NMR: Use the proton decoupled ¹³ C NMR spectrum	140B		_			_	_						143B
to assist in the determination of the structure of an			✓			✓	✓						143D
unknown compound.	1.40D	-											158
NMR: Distinguish solvent and reference NMR	140B						✓						143B
signals from that of the sample	1.40D	-											143D
NMR: Understand and explain conformational	140B	✓				✓	✓						143C
averaging in NMR spectra.	1.42.4	-											158
Characterize reaction products by spectroscopic	143A	✓			✓	✓							143C
methods, as available.	143B	-											143D
Document data and observation accurately.	6BL				✓	✓							All labs
X I G	143A												1001
Lab: Carry out a task with a proficient and confident	6BL			✓	✓								100A
manner while working alone	143A	-											1000
Lab: Work as a member of team in an efficient	105A			✓	✓	✓	✓	✓	✓	✓	✓		100BL
manner toward a common goal	143D												
Lab: Maintain safe practices for oneself and others	6BL	-		✓	✓								All labs
Lab: Minimize waste and dispose of waste legally	6BL			✓	✓								All labs
and correctly	143A	-											
Lab: Relate laboratory procedures, whether synthetic	105A	✓			✓	✓							All labs
or analytical, to underlying theory	143A												1.420
Lab: Demonstrate and use subsequently:	143A				_	_							143B
Recrystallization, extraction, evaporation, TLC,				~	V	v							
column chromatography, distillation.	1.40.4	-											122
Lab: Set up and use apparatus to carry out a variety	143A			✓	✓	✓							123
of reaction types	143B	-											143D
Lab: Demonstrate when and how to reduce hazards	6BL												123
by using hoods, glove boxes, or oxygen-free	143A			•	•								143B 143D
techniques	(DI	-											
Lab: Document procedures and results completely,	6BL				_				/	_			All labs
accurately, and with complete honesty in notebooks	100A				•				•	•			
kept to professional standards	143A	-											A 11 1 . 1
Operate a variety of laboratory instruments and	100A				✓	_	_						All labs
apparatus for synthesis and for analysis, with explicit	105A			•	•	•	•						
direction or, eventually, following written manuals.	143A												105 4
Analyze experimental data, using proper statistical	6BL						√	./	./				105A
methods and construction of graphs that re effective	100A				*		*	*	•				All labs
in communicating results to others	100A	-			\vdash								100DI
Distinguish precision and accuracy. Distinguish													100BL Others
systematic from random error and blatant mistakes.	105A				✓		✓		✓	✓			Others
Identify these in reports and present quantitative													
limits on error when it is possible to do so.		-											
Counch and natriava abamical information from	105 4												1.42D
Search and retrieve chemical information from	105A			✓	✓			✓					143D Others
various databases.	143C	-			\vdash								Others
Read, analyze and critically evaluate journal papers	105A							✓			✓		143D Others
in various subfields of chemistry Write scientific reports in a concise, organized and	Et al. 6BL	-			\vdash								Others 105A
effective style	100A								✓		✓		105A 100BL
effective style	100A												TUUDL

	143A/B						-				143C/D
Report scientific findings and inferences in oral	105A						-				100BL
presentations in clear and organized fashion, using	103/1							√	١,	/	143D
visual tools, mostly PowerPoint® computer methods											1130
Recognize relation between molecular structure and	140A							+			154
reactivity	11011	✓		✓							131
Explain the theory of origin of life	114A	✓		√				+	١,	/	
Describe the difference between eukaryotic and	114A										
prokaryotic cells	117/1	✓	✓								
Recognize the 20 amino acids and explain the	114A										
differences in their chemical properties.	11711	✓									
Explain and sketch the periodic arrangements of	114A										
secondary structures within a protein fold	117/1	✓									
Understand the packing of secondary structure units	114A										
to form a tertiary fold	114/1	✓									
Identify the packing of tertiary folds to form specific	114A	1					-	+			
quaternary structures	114/1	✓									
Use analysis of hydrophobic interactions and	114A	1					-	+			
properties of water and how they influence protein	114/1	✓									
folding in solution and in membranes											
Distinguish and explain negative and positive	114A	1				+	-	+			
cooperativity and allosteric interactions	114/1	✓									
Describe the organization of the membranes and the	114A	1				+	-	+			
influence of specific chain properties on the fluidity	114/1	✓									
of the membrane											
Know that specific classes of proteins called	114A	1					-	+			
enzymes are catalysts of chemical reactions	114A	✓									
Recognize and use Michaelis-Menten kinetic scheme	114A			√			-				
Review the properties of buffers and concept of pH	114A 114A			•			-				
and explain how solution pH can influence protein	114A	✓		✓							
stability and enzyme kinetics		*		•							
Distinguish competitive, non-competitive, and	114A						-	+	-		
uncompetitive inhibitors affect observed rates of	114/1	✓		✓							
reactions											
Explain how inhibitors can be used as drugs.	114A		√								
Describe the structure and properties of DNA in	114A	1				+	-	+			
terms that take advantage of insights provided by	114/1	✓									
organic chemistry.											
Understand the difference between anabolic and	114B	✓				+	-	+			
catabolic processes in metabolism	מדוו										
Use knowledge from organic chemistry reaction	114B	✓									
mechanisms to follow metabolic pathways	1170			✓							
Understand experimental approaches to tracing	114B	✓									
metabolic pathways	1170			✓							
Be able to describe anabolic and catabolic proceses	114B	✓						+		+	
are coupled with energetics from hydrolysis of ATP	עדוו										
Understand oxidation and reduction and electron	114B	✓				+	-	+	+		
transfer reactions in biological systems	עדוו										
Understand that reaction coordinate diagrams are	114B	✓					+	+	+		
useful for thermodynamics of coupling of anabolic	עדוו										
and catabolic processes in metabolism											
Be able to trace through the Calvin cycle	114B	✓				-	-	+	+	1	
Follow the fate of precursors and radioactive labels	114B					+	-	+	+		-
in the metabolic reactions	עדוו	✓		✓							
Relate glycogen metabolism to diseases.	114B	✓	√			\dashv	\dashv	+	١,	+	
Use knowledge of thermodynamics to describe	114B		•			+	+	+	+	+	
transport through membranes	עדנו	✓									
transport unough memoranes	II	1		<u> </u>						1	1

Use knowledge of kinetics to describe transport	114B	✓											
through membranes													
Identify the enzymes and molecules involved in the citric acid cycle	114B	✓											
Understand the mechanism of common metabolic diseases	114B	✓	✓								✓		
	114A/C	✓	√										
Describe the central dogma		∨	· ·										
Know the reaction in photosynthesis	114C	•	~										
Know the properties of nucleic acid structure and	114C	✓											
how that forces the conformation of the DNA													
Know the genetic code	114C	✓	✓								✓		
Know the concepts of translation and transcription	114C	✓									✓		
Know how recombinant DNA technology works	114C	✓											
Know the difference between RNA and DNA	114C	✓											
Understand the mechanism of DNA repair and their	114C												
relationship to diseases		✓	✓								✓		
Know the structure of viral particle and their	114C												
mechanism of infection	1140	✓	✓								✓		
	114C												
Understand the concepts of gene expression and	114C	✓									✓		i l
genomic organization	1121	 											
Develop the skills to purify proteins from tissues and	112A				√	✓	√						
recombinant sources													
Develop the skills to analyze purity of proteins	112A				✓	✓	✓	✓					
Develop the skills to analyze how purity and specific	112A				✓	./		✓					
activity are coupled					•	•		V					
Develop the skills to identify posttranslational	112A					_							
modification of proteins					✓	~	√		✓	✓			
Develop the skills to conduct accurate kinetic	112A												
analyses	11211			✓	√	√	✓	✓		✓			
Develop the skills to run polyacrylamide gel	112A												
electrophoresis and isoelectric focusing	112A				✓	✓	✓						
	1104												
Develop the skills to identify tissue specific	112A				✓	✓	✓						
difference in isoenzyme contents													
Develop the skills to subclone DNA fragment into	112B				✓	✓	√						
plasmid vectors													
Develop the skills to use restriction enzymes to cut	112B				/	/	/						
DNAs					•	•	•						
Develop the skills to run agarose gel electrophoresis	112B				✓	✓	✓						
Develop the skills to isolate DNA fragment from	112B				_								
agarose gels					•	•	•						
Develop the skills to transform hors organism for	112B				,								
protein expression and drug resistance					✓	✓	√			√			
Understand how stability and thermodynamics are	113												
related to driving protein folding	110	✓		✓									
Describe the theory of funneled landscapes and	113	1											
evolution in driving efficient folding	113	✓											i l
	112	\vdash								\vdash			
Be able to calculate the difference in thermodynaic	113	✓		✓									
stability of wild-type and mutant proteins	112	1											
Be able to calculate the populations of foleded and	113	✓		✓									
unfolded proteins from thermodynamic parameters													ļ
Be able to calculate the entropy and enathalpy of	113	✓											
ligand binding		Ĺ											
Be able to read and use journal scientific literature	113								✓		✓	✓	
Make detailed analysis of enzyme catalysis from	116											/	
primary (recent) literature.		✓							✓			√	
Be able to find strength and weaknesses in particular	116												
approach to analyzing enzyme mechanisms		✓		✓					✓			√	
approuch to unaryzing onzyme mechanisms	11	II				1			I		 <u> </u>		

For Pharmacology: delete the previous 8 goals and add the following 4 goals:													
Complete an introduction to pharmacy and pharmacology as professions	92		✓								✓		
Understand, explain, and interpret the interaction between chemo-biological agents (drugs) and the human physiology and global nature	118	~	✓	✓	✓			✓		√	✓		
Complete one year of study in biology with some laboratory experience, meeting the goals specified by the Division of Biology	various	~									✓		
Complete one quarter of economics, meeting the goals specified by the Department of Economics	Econ 1 or 3										✓		
For both majors:													
Independent research is encouraged but not required	199				✓	✓		✓	✓	✓		✓	