# CONTENTS

<table>
<thead>
<tr>
<th>OBJECTIVES OF THE COURSE</th>
<th>.................................................................</th>
<th>1</th>
</tr>
</thead>
<tbody>
<tr>
<td>COURSE CO-ORDINATOR and LECTURERS</td>
<td>.................................................................................</td>
<td>1</td>
</tr>
<tr>
<td>COURSE STRUCTURE and TEACHING STRATEGIES</td>
<td>.................................................................................</td>
<td>2</td>
</tr>
<tr>
<td>APPROACH TO LEARNING AND TEACHING</td>
<td>.................................................................................</td>
<td>2</td>
</tr>
<tr>
<td>TEXTBOOKS AND OTHER RESOURCES</td>
<td>.................................................................................</td>
<td>2</td>
</tr>
<tr>
<td>STUDENT LEARNING OUTCOMES</td>
<td>.................................................................................</td>
<td>3</td>
</tr>
<tr>
<td>ASSESSMENT PROCEDURES</td>
<td>.................................................................................</td>
<td>3</td>
</tr>
<tr>
<td>COURSE EVALUATION AND DEVELOPMENT</td>
<td>.................................................................................</td>
<td>4</td>
</tr>
<tr>
<td>GENERAL INFORMATION</td>
<td>.................................................................................</td>
<td>5</td>
</tr>
<tr>
<td>Official Communication</td>
<td>.........................................................................</td>
<td>5</td>
</tr>
<tr>
<td>Attendance Requirements</td>
<td>.........................................................................</td>
<td>5</td>
</tr>
<tr>
<td>Practical Classes</td>
<td>.........................................................................</td>
<td>6</td>
</tr>
<tr>
<td>Handwriting</td>
<td>.........................................................................</td>
<td>6</td>
</tr>
<tr>
<td>Special Consideration</td>
<td>.........................................................................</td>
<td>6</td>
</tr>
<tr>
<td>Student Support Services</td>
<td>.........................................................................</td>
<td>6</td>
</tr>
<tr>
<td>Appeal Procedures</td>
<td>.........................................................................</td>
<td>6</td>
</tr>
<tr>
<td>Academic Integrity and Plagiarism</td>
<td>.........................................................................</td>
<td>6</td>
</tr>
<tr>
<td>LECTURE and PRACTICAL OUTLINES</td>
<td>..................................................................................</td>
<td>7</td>
</tr>
<tr>
<td>TIMETABLE</td>
<td>..................................................................................</td>
<td>11</td>
</tr>
<tr>
<td>ASSESSMENT TASKS</td>
<td>..................................................................................</td>
<td>12</td>
</tr>
<tr>
<td>Formative Assessment</td>
<td>..................................................................................</td>
<td>12</td>
</tr>
<tr>
<td>Laboratory Notebook</td>
<td>..................................................................................</td>
<td>12</td>
</tr>
<tr>
<td>Marking Criteria – Laboratory Notebook</td>
<td>.........................................................................</td>
<td>15</td>
</tr>
<tr>
<td>Molecular Techniques Collaborative Learning Session</td>
<td>.........................................................................</td>
<td>16</td>
</tr>
<tr>
<td>Marking Criteria – Molecular Techniques Wiki</td>
<td>.........................................................................</td>
<td>19</td>
</tr>
<tr>
<td>Marking Criteria – Molecular Techniques Learning Activity</td>
<td>.........................................................................</td>
<td>20</td>
</tr>
</tbody>
</table>
PHAR3102 Course Information

Molecular Pharmacology (PHAR3102) is a 3rd year Science Course worth six units of credit (6 UOC). The course is usually undertaken as part of a major in Pharmacology in a Bachelor of Science or Bachelor of Medical Sciences and as part of the Medicinal Chemistry Program. This course builds on the information you have already gained in Introductory Pharmacology and Toxicology (PHAR2011).

OBJECTIVES OF THE COURSE

This course will examine the molecular basis of drug action and explore how cutting edge biotechnology and biomedical research can advance pharmacological knowledge, increasing our understanding of how drugs work. The following areas will be studied in detail; genetic variability in drug action, protein structure-activity relationships, receptor-ligand interactions, signal transduction, biochemical and molecular aspects of G-protein coupled receptors and their signalling mechanisms. Research and analytical skills will be developed in practical classes.

COURSE CO-ORDINATOR and LECTURERS

Course Coordinator:
Dr Angela Finch
Rm 326 Wallace Wurth Building East ph: 9385 1325

Co-coordinator
Dr Lu Liu
Rm 325 Wallace Wurth Building East ph: 9385 8762

Students wishing to see the course coordinators should make an appointment via email as our offices are not readily accessible. We will organize to meet you in a convenient location elsewhere in the building.

Lecturers in this course:
Dr Angela Finch a.finch@unsw.edu.au
Dr Nicole Jones n.jones@unsw.edu.au
Dr Trevor Lewis t.lewis@unsw.edu.au
Dr Lu Liu lu.liu@unsw.edu.au
Dr Nicola Smith n.smith@victorchang.edu.au

Please read this manual/outline in conjunction with the following pages on the School of Medical Sciences website:
- Advice for Students
- Learning Resources

(or see "STUDENTS" tab at medicalsciences.med.unsw.edu.au)
COURSE STRUCTURE and TEACHING STRATEGIES

Learning activities occur on the following days and times:

- Lectures: Thursday 10-11 am and Friday 11-12pm
- Collaborative Learning Session: Wednesday 4-5pm or 5-6pm*
- Practicals: Monday 10 am -1 pm or 2-5pm*

* Once enrolled in one of the two sessions, students cannot change.

Students are expected to attend all scheduled activities for their full duration (2 hours of lectures per week and up to 4 hours of practical and collaborative learning sessions per week). Students are reminded that UNSW recommends that a 6 units-of-credit course should involve about 150 hours of study and learning activities. The formal learning activities are approximately 72 hours throughout the semester and students are expected (and strongly recommended) to do at least the same number of hours of additional study.

Lectures will provide you with the concepts and theory essential for an understanding of molecular pharmacology. To assist in the development of research and analytical skills practical classes and collaborative learning sessions will be held. These classes allow students to engage in a more interactive form of learning than is possible in the lectures. The skills you will learn in practical classes are relevant to your development as professional scientists.

APPROACH TO LEARNING AND TEACHING

The learning and teaching philosophy underpinning this course is centred on student learning and aims to create an environment which interests and challenges students. The teaching is designed to be engaging and relevant in order to prepare students for future careers.

Although the primary source of information for this course is the lecture material, effective learning can be enhanced through self-directed use of other resources such as textbooks and Web based sources. Your practical classes will be directly related to the lectures and it is essential to prepare for practical classes before attendance. It is up to you to ensure you perform well in each part of the course; preparing for classes; completing assignments; studying for exams and seeking assistance to clarify your understanding.

TEXTBOOKS AND OTHER RESOURCES

Due to the cutting edge nature of this course and the rapid advances made in the field of Molecular Pharmacology, a single primary text which adequately covers the content of this course has not been identified. Therefore each lecturer will provide you with additional resources to supplement their lecture material. These resources will take the form of text books, journal articles or web-based resources. If available, links to the electronic form of these resources will be put on the course Moodle page.

Three textbooks have been identified that together cover the majority of the course content. These texts are also available as online resources from the UNSW library

“Pharmacology in drug discovery: understanding drug response” by T. P. Kenakin, will be used as a reference text for lectures given in weeks 4, 7-12.

“Molecular Pharmacology: From DNA to Drug Design” by Dickenson, Freeman, Lloyd Mills, Thode, & Sivasubramaniam, will be used as a reference text for lectures in weeks 1, 2, 5-11.

“General and Molecular Pharmacology: Principles of Drug Action” Edited by Francesco Clementi and Guido Fumagalli, will be used as an additional reference text throughout the course.
STUDENT LEARNING OUTCOMES

PHAR3102 will develop those attributes that the Faculty of Science has identified as important for a Science Graduate to attain. These include; skills, qualities, understanding and attitudes that promote lifelong learning that students should acquire during their university experience.

**Graduate Attributes**
- A. Research, inquiry and analytical thinking abilities
- B. The capability and motivation for intellectual development
- C. Ethical, social and professional understanding
- D. Effective communication
- E. Teamwork, collaborative and management skills
- F. Information Literacy – the skills to locate, evaluate and use relevant information.

On completion of this course students should:
1. be able to describe the genomic regulation of drug action
2. be able to discuss the molecular pharmacology of receptors, channels and enzymes
3. have gained a knowledge of molecular biology techniques used in pharmacology
4. be able to accurately record experimental data and draw conclusions from experimental data
5. be able to demonstrate their ability to work in teams and communicate scientific information effectively

ASSESSMENT PROCEDURES

- Progress exam (40 min duration) 15%
- Laboratory notebook 10%
- Molecular techniques wiki and learning activity 10%
- End of session examination (2 hours duration) 60%
- Formative Assessment 5%

The **practicals** are provided to support lecture material and practise analytical skills. The practical classes and collaborative learning sessions help you to develop graduate attributes A, C, D & E. At the completion of the practical course you will be required to submit your laboratory notebook covering all of the practical sessions.

In the collaborative learning sessions students will work in teams to research a technique used in molecular pharmacology. They will build a *wiki* and facilitate a *learning activity* in the collaborative learning session. This assessment task will allow you to develop your research, information literacy, communication and time management skills, as well as allowing you to demonstrate your ability to work in a team and collaborate successfully (Graduate Attributes A, D, E &F).

A penalty will apply for late submissions of assessment tasks (10% per day).

The **progress examination** will be held in the lecture slot on **Thursday the 21st of April**. This exam will give you feedback on how you are succeeding in the course. The **progress examination** and **end of session examination** will test not only your knowledge of the molecular pharmacology of receptors, channels and enzymes, and molecular techniques used in pharmacology but also your ability to apply the knowledge you have acquired from
multiple lectures, collaborative learning sessions and practicals to molecular pharmacology scenarios. The examinations may be in the format of MCQ, short or long answer questions. The questions will be based on the material covered in the lectures, practical classes and collaborative learning sessions. Material covered prior to the progress exam may be again examined in the final exam. The examinations will address graduate attributes A and B. The end of session examination will be held during the official examination period.

COURSE EVALUATION AND DEVELOPMENT

Each year feedback is sought from students about the courses offered in the Department of Pharmacology and continual improvements are made based on this feedback. The Course and Teaching Evaluation and Improvement (CATEI) Process of UNSW is the way in which student feedback is evaluated and significant changes to the course will be communicated to subsequent cohorts of students. Also a staff-student liaison group will be set up and students will be invited to become class representatives to seek feedback from their colleagues and meet with academic staff to discuss any issues that arise. Based on feedback given in these meetings changes will be implemented during the course and for future years.

Based on the feedback received; in 2009 and 2010: questions were provided to help focus the reading of journal articles for collaborative learning sessions, the proportion of total marks for the final examination was reduced, marks to encourage participation in collaborative learning sessions were given, smaller practical classes and reduction in the length of each experiment to ensure it can be completed within a three hour practical class were implemented; In 2011: the journal club questions are referenced back to the lectures to a greater extent. Dr Finch has worked with Dr Kenakin to develop a textbook that covers some parts of the course; in 2012: formative quizzes have been added to provide more continual feedback and a new textbook will be trialled. In 2013: the practical manual was revised. In 2014: the order of the topics covered in the collaborative learning sessions has been changed to better match with the lecture content; in 2015: the unannounced ‘spot quizzes’ are now timetabled quizzes additional information is provided for each wiki topic to help focus the wiki to the most relevant information.
GENERAL INFORMATION

The Department of Pharmacology is part of the School of Medical Sciences and is within the Faculty of Medicine. It is located in the Wallace Wurth building. General inquiries can be made at the BABS.SOMS.BEES (B.S.B.) Student Office, located on the Ground Floor Room G27, of the Biosciences Building. Office hours are 9.00 am - 5:00pm.

Professor Margaret Morris is Head of Department and appointments to meet with her may be made via email (m.morris@unsw.edu.au).

Departmental Vacation Scholarships: The School of Medical Sciences supports several summer vacation scholarships each year to enable good students to undertake short research projects within the school. For further details contact the Administrative Officer.

There is an honours program conducted by the School. The Honours program is coordinated by Dr Thomas Fath (t.fath@unsw.edu.au), Ph: 9385 8495. Any students considering an Honours year should discuss the requirements with the coordinator.

Honours Administrator: Vicky Sawatt (v.sawatt@unsw.edu.au) Ph: 9385 8195.

Postgraduate degrees

The Department of Pharmacology offers students the opportunity to enter into the following graduate programs:

Course Work Masters: Masters in Drug Development. For more information contact Dr Orin Chisholm (o.chisholm@unsw.edu.au)

Research Masters: In Pharmacology. For more information contact the post-graduate coordinator Dr Pascal Carrive (p.carrive@unsw.edu.au)

Doctorate (Ph.D): In Pharmacology. For more information contact the post-graduate coordinator Dr Pascal Carrive (p.carrive@unsw.edu.au)

Enrolment and administrative help

Ms Carmen Robinson and Ms Justine Maguire-Scarvelli are available to help with problems with enrolment and scheduling, and should be the first point of contact for administrative problems. They can be found in the BSB Student Office, Room G27, Ground floor of the BioSciences Building. Ph: 9385 2464,

Email: Carmen.Robinson@unsw.edu.au; j.maguire-scarvelli@unsw.edu.au

Official Communication

All communicate will be via your official UNSW email please see Advice for Students-Official Communication for more details.

Attendance Requirements

For details on the Policy on Class Attendance and Absence see Advice for Students and the Policy on Class Attendance and Absence.

Guidelines on extra-curricular activities affecting attendance can be found on the School of Medical sciences Website. http://medicalsciences.med.unsw.edu.au/sites/default/files/Extra-curricularActivitiesSOMS.pdf

Attendance at practical classes is compulsory, and must be recorded in the class roll at the start of each class. Arrival more than 15 minutes after the start of the class will be recorded as non-attendance. It is your responsibility to ensure that the demonstrator records your
attendance and no discussions will be entered into after the completion of the class. Satisfactory completion of the work set for each class is essential. It should be noted that non-attendance for other than documented medical or other serious reasons, or unsatisfactory performance, for more than 1 practical class during the session may result in an additional practical assessment exam or ineligibility to pass the course. Students who miss practical classes due to illness or for other reasons must submit a copy of medical certificates or other documentation to the course coordinator.

**Practical Classes**

The practical class is an opportunity for students to develop graduate attribute C by behaving in an ethical, socially responsible and professional manner within the practical class.

Students must take due care with biological and hazardous material and make sure all equipment is left clean and functional. In the interests of safety, special attention should be paid to any precautionary measures recommended in the notes. If any accidents or incidents occur they should be reported immediately to the demonstrator in charge of the class who will record the incident and recommend what further action is required.

For more details see [Advice for Students-Practical Classes](#)

**Handwriting**

Please see [Student Advice-handwriting](#).

**Special Consideration**

Please see [UNSW-Special Consideration](#) and [Student Advice-Special Consideration](#)

The supplementary exams for the School of Medical Sciences in Semester 1, 2016 will be held on the 12th, 13th and 14th July, 2016.

If you unavoidably miss the progress exam in PHAR3102, you must lodge an application with UNSW Student Central for special consideration. If your request for consideration is granted an alternative assessment will be organised which may take the form of a supplementary exam or increased weighting of the final exam.

**Student Support Services**

Details of the available student support services can be found at [Student Advice-Student support services](#).

**Appeal Procedures**

Details can be found at [Student-Advice-Reviews and Appeals](#)

**Academic Integrity and Plagiarism**

The [UNSW Student Code](#) outlines the standard of conduct expected of students with respect to their academic integrity and plagiarism. More details of what constitutes plagiarism can be found [here](#)
LECTURE and PRACTICAL OUTLINES

The course timetable is appended at the end of these notes

The course is divided into 4 main themes covering the molecular basis of drug action.
- Genomic Regulation of Drug Actions
- Molecular Pharmacology of Receptors, Channels and Enzymes
- Signal Transduction and Modulation
- Receptor Theory

Genomic Regulation of Drug Actions

Pharmacogenetics and Pharmacogenomics
The concept of pharmacogenetics and pharmacogenomics will be covered in these lectures. The types of genetic mutations: single nucleotide polymorphisms; tandem repeat polymorphisms; gene insertion and deletion; gene duplications; alternative splicing and their effects on drug targets will be explored. The influence of genetic background on drug efficacy and the use of pharmacogenomics to individualise therapy will also be covered.

Pharmacogenetics: Practical
In this practical we will investigate the role polymorphisms of cytochrome P450 2D6 play in inter-individual differences in drug metabolism and their contribution to either a lack of efficacy or adverse side effects of drugs. To identify these polymorphisms, we will be using publically accessible databases, other computer based techniques and data obtained from PCR.

The Regulation of Gene Transcription
This lecture will briefly discuss the process of gene transcription and go on to examine transcription factors in more detail – including their different structures and roles in biological functions such as development, responses to environmental stimuli (e.g. heat or low oxygen), and gene transcription. A number of examples of therapeutic agents that can act by modulating gene transcription – including hormones acting at nuclear receptors, will be discussed.

Pharmacological Regulation of Gene Expression: Practical
Drug induced changes of gene expression of transcription factors will be examined. Techniques learnt include tissue/cell culture, RNA isolation, RNA quantification, RT-PCR and gel electrophoresis. Primer design will also be introduced.

Molecular Pharmacology of Enzymes, Channels and Receptors

Protein Structure and Receptor-Ligand Interactions
In this lecture we will review the non-covalent interactions underlying the molecular recognition between a protein (‘receptor’) and a small molecule (‘ligand’). We will focus on the relationships between affinity and specificity, enthalpy and entropy and these non-covalent interactions. We will also review the basics of protein structure, including the structure and properties of amino acid side chains and the four levels of protein architecture, primary through to quaternary structure. We will focus on the importance of non-covalent interactions in protein structure.

How Enzymes Work
This lecture will cover general principles of how enzymes work including; catalytic transition state, binding energy and reaction specificity, equilibrium constant and enzyme kinetics (the Michaelis-Menten equation).
Drug Modulation of Enzyme Function
This lecture will cover the basic principles of how drugs modulate enzyme activity including competitive, non-competitive and uncompetitive inhibition. A brief overview of analytical techniques used in enzyme activity characterisation will also be given.

Voltage-gated ion channels
This lecture introduces the families of voltage-gated channels and identifies the key organs where they have an important role (eg. nervous system, heart, skeletal muscle). The main structural domains of the voltage-gated ion channels with respect to the crystal structures that are available and how they relate to the experimental evidence for function will be discussed. We will relate these structural domains to common drug actions. Introduces the idea of ion channels as 'enzymes' where the product is the conduction of ions across the cells membrane.

Ligand-gated ion channels
This lecture introduces the families of ligand-gated ion channels. The main structural domains with respect to the crystal structures available and how they relate to the experimental evidence for function will be discussed. We will discuss some key actions of ligand-gated ion channels (eg. neurotransmission), how they are altered by common drug actions, and the kinetic mechanism of the actions.

Catalytic Receptors
This lecture introduces the catalytic receptors, covering the five main types of catalytic receptors. The main structural features of catalytic receptors and how they relate to function of these receptors will be examined. The signalling pathways of each main type of catalytic receptors and how signalling is regulated will also be discussed.

G-Protein Coupled Receptors (GPCRs)
This lecture will provide an introduction to the six G-protein coupled receptors (GPCR) families. It will explore the structural similarities and diversity between these families. Representative members from each family will be examined in more detail. The role of receptor dimerisation and receptor activity modifying proteins (RAMPs) in producing different receptor phenotypes will also be covered.

GPCRs: Role of Structural Motifs in Binding, Activation and Regulation of Signalling
This lecture will take a more detail look at the key structural regions of GPCRs and their role in receptor activation and regulation. The structural regions examined will include the N-terminus, extracellular loops, specific transmembrane helices, the DRY motif, the NPXXY motif, intracellular loops and tails.

Signal Transduction and Modulation

Second Messengers
This lecture will review the types of second messenger molecules. Several examples of second messenger systems, including: the phosphoinositol, Ca²⁺, cAMP, cGMP and arachidonic acid systems will be covered. The main signal transduction pathways used by GPCRs and catalytic receptors will be introduced in this lecture and the role of second messengers as drug targets will be explored.

Guanine Nucleotide-Binding Proteins (G-proteins)
This lecture will review the members of the G-protein superfamily. It will explore the structural characteristics of this family and the mechanisms of G-protein activation and regulation, including the GTPase and GTP switch. G-protein dependent signalling pathways will be covered. The role of G-proteins in disease will be discussed.
Regulation of GPCR Signalling
Mechanisms by which receptor desensitisation occurs, including internalisation, phosphorylation, binding of β-arrestins, and degradation will be covered. The role of homologous and heterologous in receptor regulation will be explored. Some of the key enzymes involved in modulation of signalling include; second messenger dependant kinases, G-protein receptor kinases, regulators of G protein signaling (RGS) proteins and guanine nucleotide exchange factors (GEFs) that facilitate GDP dissociation, GTPase activating proteins (GAPs) that stimulate GTP hydrolysis and guanine dissociation inhibitors (GDIs). The function and regulation of these enzymes will be covered.

Receptor Internalisation & Alternative Signalling Pathways
Ligand mediated receptor endocytosis and the regulation of this process will be discussed. The role of RAB and ARF proteins and other small GTP-binding proteins that control trafficking and the role of ubiquitylation in the process will be covered. The classification of desensitisation of GPCRs into Class A and Class B and the role of internalisation in non-G-protein mediated receptor signalling will also be discussed.

Receptor Signalling: Practical
This practical will examine the ability of the β2 adrenergic receptor to activate extracellular signal-regulated kinase (ERK) via G-protein dependent and independent pathways.

Advanced Pharmacodynamics
The pharmacological concept of potency, efficacy, pD₂, pA₂ will be reviewed. The calculation of pD₂ and pA₂ values from concentration-response curves of agonists and antagonists will be covered. Factors affecting pharmacodynamic variability and the role of this variability in drug efficacy and toxicity will be discussed.

Determining Antagonist Potency: Practical
In this computer-aided practical the antagonist potency of mepyramine against histamine-induced contractile responses of guinea-pig ileum will be determined. EC₅₀ values will be obtained from concentration-response curves generated by semi-log paper and Prism. The antagonist pA₂ value will be calculated using the Arunklakshana & Schild method.

Constitutively Active Receptors and Inverse Agonists:
The concept of constitutively active receptors will be discussed in this lecture. Examples of wide-type receptors, naturally occurring receptor mutants, receptor variants created by site-directed mutagenesis, showing constitutive activity will be covered. The concept of inverse agonism and its discovery through molecular pharmacology techniques will be discussed. Examples of inverse agonists will be given and their potential as therapeutics will be discussed.

Allosteric Modulators:
This lecture will cover the principles of receptor allosteric modulation. The concepts of allosteric versus orthosteric binding sites will be explored. The allosteric mechanisms in activation of enzymes, ligand-gated channels and GPCRs will be covered. The role of allosteric sites as novel drug targets will be explored.

Signalling-Bias,
Signalling-bias (also called; ligand-directed signalling, functional selectivity, agonist-directed trafficking, biased agonism, or protean agonism) describes the observation that different ligands acting on the same receptor cause different patterns of response. These observations have led to a change in the concept of the receptor as either off or on but rather existing in a spectrum of conformational states each of which gives rise to a different
signalling outcome. This lecture will explore these concepts and the influence they have on drug development.

**Receptor Theory**
Since the 1920's models have been developed to assist in the understanding of the complex events that occur upon ligand binding to receptor. The first simple model was the two-state model, however with advances in pharmacology (such as molecular pharmacology) this model could no longer explain the results obtained, this lead to the development of more complex models; including the Occupational and Operational models of agonist action followed by the Ternary complex model and the Cubic ternary complex model. These lectures will discuss the development of these models and examine specific examples of experimental results which support the receptor states described by these models. The Induction versus Conformational Selection hypotheses of ligand action will also be covered.

**New concepts in Pharmacology**
In this lecture we will discuss the emerging concepts in receptor pharmacology that are currently being published and debated at scientific meetings, including heteromers, bitopic ligands and signalling texture.
# TIMETABLE

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<tr>
<th>Wk</th>
<th></th>
<th>Practical Monday 10-1 or 2-5pm Wallace Wurth 120</th>
<th>Collaborative Learning Wed 4-5 or 5-6 Colombo LG01</th>
<th>Lecture Thu 10-11 CLB3</th>
<th>Lecture Fri 11-12 CLB2</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>29/2</td>
<td>Introduction to Molecular Pharmacology and Assessment Tasks (AF)</td>
<td></td>
<td>Pharmacogenetics/genomics I (L Liu)</td>
<td>Pharmacogenetics/genomics II (L Liu)</td>
</tr>
<tr>
<td>2</td>
<td>7/3</td>
<td>Pharmacogenetics (AF)</td>
<td>Microarray/ siRNA</td>
<td>Nuclear Receptors/Transcription Factors (N Jones)</td>
<td>Protein Structure &amp; Receptor-Ligand Interactions (R Griffith)</td>
</tr>
<tr>
<td>3</td>
<td>14/3</td>
<td>Pharmacological Regulation of Gene Expression (Part A) (LL)</td>
<td>Journal Club</td>
<td>Enzymes (R Griffith)</td>
<td>Enzyme Modulation (R Griffith)</td>
</tr>
<tr>
<td>4</td>
<td>21/3</td>
<td>Pharmacological Regulation of Gene Expression (Part B) (LL)</td>
<td>Genetically Engineered Animals/ Reporter Gene Assays</td>
<td>Channels (T Lewis)</td>
<td>Good Friday Holiday</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<td></td>
<td><strong>Mid-semester Break</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>4/4</td>
<td>Pharmacological Regulation of Gene Expression (Part C) (LL)</td>
<td>Journal Club</td>
<td>Channels (T Lewis)</td>
<td>Catalytic Receptors (A Finch)</td>
</tr>
<tr>
<td>6</td>
<td>11/4</td>
<td>Pharmacological Regulation of Gene Expression (Part D) (LL)</td>
<td>Protein Crystallography/Receptor binding assays</td>
<td>GPCRs: Introduction to Families A, B, C &amp; F (A Finch)</td>
<td>GPCRs: Role of Structural Motifs in Binding, Activation and Regulation of Signalling (A Finch)</td>
</tr>
<tr>
<td>7</td>
<td>18/4</td>
<td>Receptor Signalling (Part A) (AF)</td>
<td>Journal Club</td>
<td><strong>Progress Exam</strong></td>
<td>Second Messengers (L Liu)</td>
</tr>
<tr>
<td>8</td>
<td>25/4</td>
<td>ANZAC Day Holiday</td>
<td>Careers in Pharmacology Symposium</td>
<td>G-proteins (L Liu)</td>
<td>Regulation of GPCR Signalling (L Liu)</td>
</tr>
<tr>
<td>9</td>
<td>2/5</td>
<td>Receptor Signalling (Part B) (AF)</td>
<td>Bioluminescence Resonance Energy Transfer/ Confocal Microscopy</td>
<td>Receptor Internalisation &amp; Alternative Signalling Pathways (A Finch)</td>
<td>Allosteric Modulators (A Finch)</td>
</tr>
<tr>
<td>10</td>
<td>9/5</td>
<td>Receptor Signalling (Part C) (AF)</td>
<td>Journal Club</td>
<td>Constitutive Active Receptors (L Liu)</td>
<td>Advanced Pharmacodynamics (L Liu)</td>
</tr>
<tr>
<td>11</td>
<td>16/5</td>
<td>Receptor Signalling (Part D) (AF)</td>
<td>Calcium signalling/ cAMP Assays</td>
<td>Signalling Bias (A Finch)</td>
<td>Receptor Theory I (A Finch)</td>
</tr>
<tr>
<td>13</td>
<td>30/5</td>
<td>Advanced Pharmacodynamics (LL)</td>
<td></td>
<td></td>
<td>Exam Revision</td>
</tr>
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ASSESSMENT TASKS

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<tr>
<th>Task</th>
<th>Due Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Molecular Techniques Wiki (draft) /feedback meeting</td>
<td>No later than one week before due date</td>
</tr>
<tr>
<td>Molecular Techniques Wiki</td>
<td>9 am the Friday prior to the week of scheduled presentation</td>
</tr>
<tr>
<td>Learning Activity</td>
<td>During assigned collaborative learning session</td>
</tr>
<tr>
<td>Open-book Quizzes</td>
<td>In practical class in weeks 4, 6, 9 &amp; 12</td>
</tr>
<tr>
<td>Progress Exam</td>
<td>Thursday, 21st April, 10 am</td>
</tr>
<tr>
<td>Laboratory Notebook</td>
<td>Monday 23rd May, 10am</td>
</tr>
<tr>
<td>Final Examination</td>
<td>Official exam period</td>
</tr>
</tbody>
</table>

Formative Assessment

The goal of formative assessment is to provide ongoing feedback that you can use to improve your learning. Formative assessment tasks help students identify their strengths and weaknesses and therefore the areas they should focus on.

Journal Club

The journal article and the questions to be answered will be posted the week before the collaborative learning session. You need to come to the collaborative learning session with the questions answered. You will be given credit for attempting to answer the questions and participating in the class discussion.

Learning Activity Participation

You will get credit if you participate meaningfully in the planned activity. You should come to class prepared to discuss any of the information found in the wiki.

Quizzes.

Four times in the semester, during a practical class, a 15 minute open-book quiz will be given. This quiz will be a mixture of MCQ and short answer questions. You will be able to use your lecture notes (slides and your handwritten notes) and other notes you have made from textbooks and other additional reading suggested by your lecturers.

Laboratory Notebook

You will be required to keep a laboratory notebook for all practical classes in this course. Keeping a laboratory notebook is an important skill for every scientist to develop. Laboratory notebooks are a complete record of all the procedures carried out and data collected for each experiment. Enough detail needs to be recorded so as someone could reproduce your experiment at a later date. A laboratory notebook is a legal document and as such certain conventions and procedures must be followed.
The requirements for laboratory notebooks are given below:

- The lab notebook must be a bound book and not a spiral notebook or loose sheets in a folder.
- The first 2 pages should be reserved for a table of contents.
- Each page is numbered.
- All entries are to be made in blue or black ink.
- Never remove any pages from your laboratory notebook.
- All information should be recorded directly into your notebook. Write down in detail what you do for each step of the experiment and record your results as you obtain them. Graph, films, printouts etc should be stuck directly into your notebook. Do not record data or calculations on scraps of paper and later copy them into your book. Whilst your laboratory notebook should be legible and as neat as possible it is more important that you record all the information.
- If you make a mistake strike it through with a single line and initial and date the correction. Do not use correction fluid or scribble out the mistake. The mistake should still be readable as sometimes you will realize that the entry was not a mistake after all and will want to be able to read it.
- Do not skip pages in your laboratory notebook (for example to allow space for data that will be collected later) instead make reference such as "continued from page 2" or "continued on page 5". You can start a new a new experiment on a new page however any blank space on the preceding page must have an X put through it.
- Each entry in the laboratory notebook should begin with the date, the title and a brief description of the aims of the experiment.
- At the end of each day your laboratory notebook should be signed and dated by a witness (i.e. a demonstrator).

The following information should be recorded in your laboratory notebook for each experiment:

To be completed before you come to class:

1. A brief and informative title.
2. The aims of the experiment. This section should clearly and concisely describe the purpose of the experiment (two or three sentences maximum) in your own words, explaining what the question is that you are asking (your hypothesis) and the methods you are using to answer your question. N.B. This only needs to be done once for multi-week experiments.
3. Preparation questions for each practical done (keep this brief).

To be completed during the class:

4. A concise step by step description of what you actually do when performing the experiment (this should be in point form).

The protocol should contain enough information to allow another researcher to repeat what you've done. If you reference the lab manual, you need to be specific and record which pages you are referring to. Also, if you did the experiment differently to what is written in the lab manual, you need to indicate this. Key information such as centrifuge speeds, temperatures, master mix recipes, incubation times and concentrations, along with notes on major deviations from the manual/protocol needed to be recorded.
Extraneous information like telling the reader you were labelling tubes should not be included. It is OK to note what is in each tube and its label, but you don’t need to tell us you wrote a label on the tubes.

5. Record any calculations you have done
6. Any observations (e.g. gel has a bubble, not all the sample loaded, current not stable).
7. All data collected with correct units, titles and labelled axes on graphs, and titles and labels for all drawings, films and printouts.

To be completed at the end of the experiment:

8. A summary of your findings and conclusions (two or three sentences maximum, this can be in point form).

How do your findings relate to your original aims/hypothesis? If the results were not as expected, suggest possible reasons for this. How do your findings relate to other studies in the scientific literature? N.B. This only needs to be done once for multi-week experiments.

Laboratory Notebooks are due before 10am, Monday 23rd May. Laboratory notebooks are to be submitted at the BABS.SOMS.BEES (B.S.B.) Student Office, G27 Biosciences Building. You will need to staple or use sticky-tape to attach the Assessment Cover Sheet to the front cover of your book.
Marking Criteria – Laboratory Notebook

<table>
<thead>
<tr>
<th>Criteria</th>
<th>Excellent</th>
<th>Very Good</th>
<th>Good</th>
<th>Needs Improvement</th>
<th>Unacceptable</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>&gt;8.5</td>
<td>8.4-7.5</td>
<td>7.4-6.5</td>
<td>6.4-5.0</td>
<td>&lt;5.0</td>
</tr>
<tr>
<td><strong>Organisation</strong></td>
<td>Each experiment has a title and date and appears in the table of contents. Errors are neatly struck through, no correction fluid used. All entries in pen. Each page numbered. Each day is initialled.</td>
<td>Each experiment has a title and date and appears in the table of contents. Most errors are neatly struck through. All entries in pen. Each page numbered. Each day is initialled.</td>
<td>All but one experiment has a title and date and appears in the table of contents. Some errors are not neatly struck through. Some entries not in pen. Each page numbered. Each day is initialled.</td>
<td>Not all experiments have a title and date and don't appear in the table of contents. Not all errors are neatly struck through. Not all entries are in pen or each page numbered. One day not initialled.</td>
<td>Experiments lack a title and date and don't appear in the table of contents. Errors are not neatly struck through. Entries not in pen. Page numbers lacking. More than one day is not initialled.</td>
</tr>
</tbody>
</table>
|                                | \[
\frac{______/10 \times 0.5}{______/10 \times 0.5}
\] | | | | |
| **Safety**                     | All safety summaries attached and signed. | 8-9 safety summaries attached and signed. | 6-7 safety summaries attached and signed. | 6-5 safety summaries attached and signed | >50% of safety summaries not attached and/or signed. |
|                                | \[
\frac{______/10 \times 0.5}{______/10 \times 0.5}
\] | | | | |
| **Preparation**                | All questions answered correctly. | All questions answered with minor errors. | Questions answered, with some errors. | Not all questions answered, some answered with errors. | Majority of questions not answered or incorrect. |
|                                | \[
\frac{______/10 \times 1}{______/10 \times 1}
\] | | | | |
| **Aims**                       | The aims of each experiment or the questions to be answered is clearly identified and stated. | The aims of each experiment or the questions to be answered are stated. | The aims of most experiments or the questions to be answered are identified but more detail needed. | The aims of some experiments or the questions to be answered are partially identified, however more details are needed. | The majority of the aims of each experiment or the questions to be answered are not given. |
|                                | \[
\frac{______/10 \times 2}{______/10 \times 2}
\] | | | | |
| **Experimental protocol**      | The protocol for each experiment is succinctly described with all key information included. All calculations are correct. | The protocol for each experiment is described with all key information included. Only minor errors in calculations. | The protocol for each experiment is described however, key information missing in some experiments. A few errors in calculations. | Describes how the experiments were done, but key information missing. Errors in calculations. | Does not accurately describe how the experiments were done. Calculations are incorrect or missing. |
|                                | \[
\frac{______/10 \times 2}{______/10 \times 2}
\] | | | | |
| **Data**                       | All data collected are presented. The data are clearly labelled. Correct units are given. | All data collected are presented. The data are clearly labelled with minor omissions. Minor errors in units given. | Data collected are presented and clearly labelled but with minor omissions. Errors in units given. | Some data not presented or not labelled. Errors in units. | Most of the data collected not presented. The data are not labelled. Correct units not used. |
|                                | \[
\frac{______/10 \times 2}{______/10 \times 2}
\] | | | | |
| **Conclusions**                | Clear and concise conclusions given. Conclusions are valid and supported by appropriate reference(s). | Clear conclusions are given. Conclusions are mostly valid and supported by appropriate reference(s). | Conclusions are given. Minor errors in conclusions. Appropriate references needed. | Conclusions are given for most experiments. Some conclusions are not valid and or not supported by appropriate reference(s). | Conclusions do not accurately describe the results of the experiments or are not presented for the majority of experiments. |
|                                | \[
\frac{______/10 \times 2}{______/10 \times 2}
\] | | | | |

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PHAR3102 Course Outline

15
Molecular Techniques Collaborative Learning Session

Overview
Publishing a scientific paper is the primary way that a scientist contributes new knowledge or methods to the scientific community. Collectively, these journal articles chronicle advances in science and technology. Without this foundation of work—whether a seminal contribution or a simple finding—future experiments would have no context and scientific research could not progress. Thus, understanding the literature is vital to understanding scientific research.

To understand scientific literature, however, it is important to know what tools and techniques scientists use to ask and answer questions. Over the course of the semester, you will explore several molecular biology techniques that are essential to understanding the scientific papers publish in the area of molecular pharmacology. Each student group will design and lead an exploration of a laboratory technique that is found in the papers to be discussed in each "journal club" collaborative learning session.

Instructions for the in-class activity

Part 1: Make a wiki about the technique
To help your classmates prepare for the activity you will design, your group will write a wiki about the technique that contains the following information:

- How the technique works. For example, describe the materials needed or provide an annotated diagram of the important steps involved, or the molecular process that occur during each step of the technique.

- How the technique is used. Include a description of what type of information this technique provides and what types of questions can be answered using the technique.

- Two Hypotheses that could be tested using this technique and two hypotheses that could not be tested using this technique. Include an explanation of why they can or cannot be tested using this technique.

- Benefits and limitations of the technique.

- References. List any resources that your group used to develop the fact sheet or in-class activity, and highlight other important resources where your classmates can find more information.

The Wiki should not be longer than 1000 words. Each member of the group must write and/or edit a minimum of 100 words. Contributions of less than this will result in a grade of 0%. Each member must log onto Moodle themselves; if you use someone else’s log-on, or work collaboratively outside Moodle your contribution will not be recognised.

The course coordinators are available at any point in the planning process to answer questions about the technique or this project.

Each group must organise a meeting with the Course coordinator for the week prior to their presentation to review their wiki content and learning activity. It is recommended that these meetings are booked as earlier as possible as your preferred time may not be available.
Researching your assigned technique

Places to look for general information:
- Textbooks
- Internet: Beware that not all sites will be accurate! Good information can usually be found at university websites, textbook publishers and scientific company websites.

Places to look for specific information:
- The library has many books covering molecular biology techniques
- Scientific literature (hint: limit your search to reviews or methods journals)
- Journal Websites such as: Nature Methods, Molecular Pharmacology, Trends in Pharmaceutical Sciences, Cell, Science Signaling.

Part 2: Design and lead an in-class activity

Each group will work together to design and lead an in-class activity that will:
1. Actively engage your classmates involved in learning and understanding the technique
2. Help your classmates determine how well they understand the technique

The entire activity should take no more than 15 minutes. After the activity, plan to spend 5 minutes to tie everything together and answer any questions your classmates might have. Altogether, you will have 20 minutes of collaborative learning session time, so use it wisely.

Some ideas for activities
- **Interpret data from an experiment** that employs the technique. Develop a series of key questions that will encourage discussion about what conclusions can be drawn from those data.

- **Sequence the important steps in the technique.** Diagram each step on a separate sheet of paper. Have groups of students describe what is happening at each step, arrange the diagrams into the correct order, and explain why they ordered the steps in that way.

- **Act out the important steps of the technique.** Provide materials for your classmates to serve as critical molecules, reactions, or other “players” in technique, then have each person describe what their role is, and have the entire class act out the technique. Discuss key points afterwards.

- **Solve a scenario where the technique is done or used incorrectly** (e.g. steps missing or out of order, or incorrect conclusions drawn from an experiment). Have your classmates work together to determine the correct order of the steps, propose more appropriate conclusions (and justify their answers), or answer questions about the scenario.

- **Compare and contrast.** Have classmates compare and contrast your technique with another related technique. Give your classmates a set of scientific questions and have each group decide which technique to use for each and explain their decisions.
TIMELINE (what is due when):

Draft wiki at least one week before your wiki is due (1%)
Organise a time to meet with the course coordinator that suits the majority of the group. Complete the draft of the wiki at least 24h before the meeting. The coordinator will review your draft of the wiki and a description of your activity (and any accompanying visual aids, worksheets etc.) and feedback will be given in the meeting with the coordinator where suggestions for improvement will be given.

Final wiki on the Friday before you present (4%)
The coordinator will open your wiki for viewing by the whole course at 9am Friday so that your classmates can study it before the collaborative learning session.

Learning activity during collaborative learning session (5%)
- Introduce the activity, give instructions
- Guide classmates in completing the activity
- Wrap-up; tie everything together and answer questions
- Do not give a “lecture” on the topic as the rest of the class will have covered that prior to the class by reading your wiki

What to do on the weeks another group will be presenting:
Study the wiki to familiarise yourself with the technique so that you can participate meaningfully in the planned activity. You should come to class prepared to discuss any of the information found in the wiki and how the technique is used in the paper for discussion in the collaborative learning session.

Adapted from a teaching unit developed by Amy Hubert and Bridget Jacques-Fricke University of Wisconsin–Madison, Madison, WI 53706 USA.
# Marking Criteria – Molecular Techniques Wiki

<table>
<thead>
<tr>
<th>Criteria</th>
<th>Excellent (&gt;8.5)</th>
<th>Very Good (8.4-7.5)</th>
<th>Good (7.4-6.5)</th>
<th>Needs Improvement (6.4-5.0)</th>
<th>Unacceptable (&lt;5.0)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Completeness</strong></td>
<td><strong>_____/10 x 15</strong></td>
<td><strong>_____/10 x 15</strong></td>
<td><strong>_____/10 x 15</strong></td>
<td><strong>_____/10 x 15</strong></td>
<td><strong>_____/10 x 15</strong></td>
</tr>
<tr>
<td>The wiki contains all assigned elements and all details necessary to fully explain the technique. Your wiki can stand alone as a comprehensive source of information on the technique.</td>
<td>The wiki contains all assigned elements, but lacks a few minor details. Your wiki provides readers with most of what they should know about the technique but could be more comprehensive.</td>
<td>The wiki contains all assigned elements, but lacks some details. Your wiki provides readers with most of what they should know about the technique but lacks a few minor details.</td>
<td>The wiki contains all assigned elements, but lacks important details. Your wiki provides a general overview, but readers would need to look elsewhere to fill in the gaps.</td>
<td>One or more assigned elements are missing or incomplete. Your wiki lacks key information needed to give the reader a basic understanding of the technique.</td>
<td></td>
</tr>
<tr>
<td><strong>Accuracy</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
</tr>
<tr>
<td>All information included in the wiki is accurate. The reader can confidently rely on the wiki as a source of information about the technique.</td>
<td>The information is accurate except for a few minor errors. The wiki is useful as a resource but may mislead the readers on a few small details.</td>
<td>The information is accurate except for a few minor errors. The wiki is useful as a resource but may mislead the readers with some details.</td>
<td>The information contains several errors. While no significant errors are made, the wiki contains enough errors to detract from its usefulness as a source of information about the technique.</td>
<td>A significant error is made that causes confusion. The reader cannot depend on your work as a reliable source of information about the technique.</td>
<td></td>
</tr>
<tr>
<td><strong>Clarity</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
</tr>
<tr>
<td>The wiki is well written and easy to read. All terms are clearly defined and topics are fully explained. Your writing allows readers to easily understand the meaning of all points presented.</td>
<td>The majority of the wiki is well written and easy to read, but a few minor terms or details are unclear. Your writing allows readers to understand the meaning of all points presented.</td>
<td>The majority of the wiki is well written and easy to read, but some terms or details are unclear. Your writing requires readers to infer your meaning regarding a few details.</td>
<td>Some parts are unclear or poorly written. The lack of clarity in your writing is distracting to readers and causes them to question your meaning, but they can still draw appropriate conclusions with some effort.</td>
<td>Major portions or key details of the wiki are unclear or poorly written. Your writing is unclear enough to cause the reader to misinterpret your meaning, leading to confusion about the technique.</td>
<td></td>
</tr>
<tr>
<td><strong>Creativity</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
<td><strong>_____/10 x 1.5</strong></td>
</tr>
<tr>
<td>The wiki makes optimal use of visual aids or other creative elements (pictures, drawings, flow charts, figures, etc.) to illustrate key points of the technique. Your creativity greatly enhances your wiki as a learning tool and provides additional means for the reader to gain understanding about the technique beyond what is stated in the text.</td>
<td>The wiki makes use of visual aids or other creative elements to illustrate key points of the technique. Your creativity enhances your wiki as a learning tool and allows the reader to gain greater understanding about the technique.</td>
<td>The wiki includes visual aids or other creative elements to illustrate the technique, but they are not original or inclusive of details specific to the technique. The readers gain something from the tool but may have trouble visualizing or fully understanding parts of the technique.</td>
<td>The wiki includes pictures or diagrams, but they do not contribute to the readers’ understanding of the technique.</td>
<td>Visual aids or creative elements are not used. The reader does not gain a full sense of the technique from your information sheet.</td>
<td></td>
</tr>
<tr>
<td><strong>First draft</strong></td>
<td><strong>_____/10 x 2</strong></td>
<td><strong>_____/10 x 2</strong></td>
<td><strong>_____/10 x 2</strong></td>
<td><strong>_____/10 x 2</strong></td>
<td><strong>_____/10 x 2</strong></td>
</tr>
<tr>
<td>The first draft was complete. No coordinator contribution needed to bring wiki up to minimal required standard. Minor corrections needed.</td>
<td>The first draft was complete. Minor coordinator contribution needed to bring wiki up to minimal required standard. Some corrections required.</td>
<td>The first draft was complete. Some coordinator contribution needed to bring wiki up to minimal required standard. Significant corrections required.</td>
<td>The first draft was incomplete. Significant coordinator contribution needed to bring wiki up to minimal required standard. Major corrections required.</td>
<td>Large amount of coordinator contribution needed to bring wiki up to minimal required standard. Extensive corrections required.</td>
<td></td>
</tr>
<tr>
<td><strong>Evidence of Collaboration Writing Process</strong></td>
<td><strong>_____/10 x 1</strong></td>
<td><strong>_____/10 x 1</strong></td>
<td><strong>_____/10 x 1</strong></td>
<td><strong>_____/10 x 1</strong></td>
<td><strong>_____/10 x 1</strong></td>
</tr>
<tr>
<td>Revision history indicates substantial group collaboration. Extensive discussion, drafting, editing, and revision were evident throughout the collaboration.</td>
<td>Revision history indicates group collaboration. Discussion, drafting, editing, and revision were evident throughout the collaboration.</td>
<td>Revision history shows some evidence of group collaboration. Some evidence of drafting, editing, writing, and revising in the final product.</td>
<td>Revision history indicates little group collaboration. Very little evidence of a drafting, editing, writing, and revision process.</td>
<td>The revision history indicates no group collaboration. No evidence of a drafting, editing, writing, and revision process.</td>
<td></td>
</tr>
<tr>
<td><strong>Individual contribution to the wiki</strong></td>
<td><strong>_____/10 x 1</strong></td>
<td><strong>_____/10 x 1</strong></td>
<td><strong>_____/10 x 1</strong></td>
<td><strong>_____/10 x 1</strong></td>
<td><strong>_____/10 x 1</strong></td>
</tr>
<tr>
<td>Contributes extensively to the researching, writing, and editing. Shows appropriate wiki etiquette when editing and respects the work of others.</td>
<td>Contributes to the researching, writing, and editing. Shows appropriate wiki etiquette when editing and respects the work of others.</td>
<td>Contributes to the researching, writing or editing. Displays appropriate wiki etiquette most of the time and generally respects the work of others.</td>
<td>Provides minimal contribution to the researching, writing and editing. Displays appropriate wiki etiquette some of the time but often fails to respect the work of others.</td>
<td>Does not contribute to the researching, writing and editing (&lt;100 words).</td>
<td></td>
</tr>
</tbody>
</table>

Adapted from a teaching unit developed by Amy Hubert and Bridget Jacques-Fricke University of Wisconsin–Madison, Madison, WI 53706 USA.
## Marking Criteria – Molecular Techniques Learning Activity

<table>
<thead>
<tr>
<th>Criteria</th>
<th>Excellent (&gt;8.5)</th>
<th>Very Good (6.4-7.5)</th>
<th>Good (7.4-6.5)</th>
<th>Needs Improvement (6.4-50)</th>
<th>Unacceptable (&lt;5.0)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Choice of content</td>
<td>The activity addresses key or difficult aspects of the technique. Your classmates will leave with a deeper understanding or reinforced knowledge of the technique.</td>
<td>The activity addresses an important aspect of the technique. Your classmates will learn about the technique but would benefit more from a more in depth choice of content.</td>
<td>The activity addresses an important aspect of the technique, but not the most important or difficult aspects. Your classmates will learn something about the technique but would benefit more from a different choice of content.</td>
<td>The activity addresses a minor or easily understood aspect of the technique. This will not help further their understanding of the technique beyond what they could easily grasp on their own.</td>
<td>The activity does not address any important details of the technique or its uses. Your classmates will not learn from it.</td>
</tr>
<tr>
<td>Knowledge of the technique</td>
<td>The presenters serve as experts on the technique. The content of the activity is clear and accurate, and the group is able to provide thorough and accurate answers to all reasonable questions raised by the class. Your classmates can depend on you to teach them all they need to know about the technique.</td>
<td>The presenters are knowledgeable about the technique. The content of the activity is clear and accurate. The group is able to accurately answer most reasonable questions raised by the class without assistance. You are able to give your classmates a sound knowledge of the technique.</td>
<td>The presenters have some knowledge of the technique. The content of the activity is clear and mostly accurate. The group is able to answer most reasonable questions raised by the class but requires assistance from the instructor in some cases. You are able to teach your classmates most of what they need to know about the technique.</td>
<td>The presenters know a little more about the technique than their classmates. Some content of the activity is unclear or inaccurate, or the group cannot answer basic questions raised by their classmates about the technique. Your classmates cannot depend on you to teach them much about the technique.</td>
<td>The presenters do not understand the technique themselves. Much of the content of the activity is inaccurate or vague, or the group cannot answer basic questions about the technique. Your lack of preparation causes confusion for your classmates.</td>
</tr>
<tr>
<td>Activity design</td>
<td>The chosen activity engages all members of the class in learning about the technique. The activity is well-designed, creative, and generates useful discussion. Your classmates will gain something from the activity beyond what they would gain from simply reading the information sheet.</td>
<td>The chosen activity engages all members of the class in learning about the technique and generates good discussion. Your classmates will learn from the activity, but would benefit from a more creative format.</td>
<td>The chosen activity engages most members of the class in learning about the technique and generates some discussion but lacks originality. Your classmates will learn from the activity, but would benefit from a more engaging format or better planned activity.</td>
<td>The chosen activity engages the class in learning about the technique but is not well-designed or does not generate useful discussion. Poor planning reduces the effectiveness of the activity in helping your classmates learn about the technique.</td>
<td>The chosen activity does not engage the class in learning about the technique. Your classmates could learn the information just as well by reading the information sheet.</td>
</tr>
<tr>
<td>Group participation (group graded as a whole)</td>
<td>All members of the presenting group work together to lead the activity and make sure it runs smoothly and all members of the group are willing and able to answer questions about the technique. Everyone contributes equally and works as a team.</td>
<td>All members of the presenting group participate in leading the activity and are willing and able to answer questions about the technique, but all do not contribute equally.</td>
<td>All members of the presenting group participate in leading the activity, however, the members of the group work independently rather than as a team.</td>
<td>Some members of the presenting group show little effort to participate in leading the activity or are not prepared to answer questions about the technique.</td>
<td>The presenting group does not work together to lead the activity and answer questions. Some members do not participate, or a lack of preparation and communication leads to confusion.</td>
</tr>
</tbody>
</table>

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