# UC 9121 06F Page 1 of 2

## CBM003 ADD/CHANGE FORM

☐ Undergraduate Council	or Graduate/Professional Studies Council
New Course ☐ Course Change	New Course Course Change
Core Category: NONE Effective Fall 2007	Effective Fall
Core Category. 1101112 Entetry of an 2007	J. Zinetti va Tan
1. Department: ET College: TECH	RECEIVED OCT 1 3 2006
2. Person Submitting Form: Rupa Iyer Telephon	ne: <u>713-743-4076</u>
<ul> <li>Course Information on New/Revised course:</li> <li>Instructional Area / Course Number / Long of BTEC / 3100 / Intrumentation And Measure</li> </ul>	
<ul> <li>Instructional Area / Course Number / Short ( BTEC / 3100 / INSTRUMENT &amp; MEASURED / 3100 / INSTRUMENT &amp; MEASURED / 3100 / 3</li></ul>	· · · · · · · · · · · · · · · · · · ·
• SCH: <u>1.00</u> Level: <u>JR</u> CIP Code: <u>261201</u> (	0022 Lect Hrs: 0 Lab Hrs: 3
4. Justification for adding/changing course: To pr	ovide for new discipline areas
	fered as a special topics course?  Yes No
Content ID: Start Date (yyyy3):	
6. Is this course offered for undergraduate credit or	nly? 🛛 Yes 🔲 No
<ul> <li>Authorized Degree Program(s): BS, Biotechnolo</li> <li>Does this course affect major/minor requirem</li> <li>Does this course affect major/minor requirem</li> <li>Are special fees attached to this course?</li> <li>Can the course be repeated for credit?</li> </ul>	nents in the College/Department?
8. Grade Option: <u>Letter (A, B, C)</u> Instruct	tion Type: <u>laboratory</u>
9. If this form involves a change to an existing course the course inventory: Instructional Area / Cours	rse, please obtain the following information from se Number / Long Course Title
• Start Date (yyyy3): Content I.D.: _	
10. Proposed Catalog Description:  Cr: (0-3). Prerequisites: BIOL 3301. Descript techniques and instrumentation used in the mode	
1. Dean's Signature:	Date: 60/12/06
Print/Type Name: Fred D. Lewallen	· <del>v maa.</del>

## UC 9121 06F Page 2 of 2

# University of Houston Proposed Course Outline for BTEC 3100, Instrumentation and Measurement laboratory

Course Objectives: Students who successfully complete this course will be able to:

- Make calculations and prepare stock laboratory solutions..
- Purify and maintain bacterial strains on agar plates.
- Culture pure microbial strains in liquid media.
- Clone DNA sequences into bacterial plasmids
- Transform bacteria with recombinant DNA plasmids
- Purify recombinant plasmid DNA from bacteria.
- Perform agarose gel electrophoresis to separate DNA fragments.
- Analyze and map recombinant DNA fragments.
- Set up PCR reactions and operate a thermal cycler.
- Analyze PCR reaction products and calculate genetic frequencies.
- Analyze DNA sequence information.
- Perform protein assays
- Determine protein structure
- Analyze proteins by SDS polyacrylamide gel electrophoresis..
- Maintain a laboratory notebook according to biotechnology industry standards.

#### **Course Outline**

#### 1. Analyze Clones

- a. Verification of clones
- b. Isolation of Plasmid DNA
- c. Restriction Cleavage
- d. Restriction Activity

#### 2. Structure of OPH Gene

- a. Sequencing and Sequence analysis
- b. Homology searches of genomic databases
- c. Homology alignments and identification of conserved residues

#### 3. Expression of OPH

#### 4. Enzyme Assay of Organophosphorus Hydrolase

- a. A qualitative assay
- b. A quantitative assay
- c. Discuss factors affecting enzyme function and production
- d. Broad spectrum activity: a survey of activity against a variety of OP pesticides

### 5. Characteristics of Organophosphorus Hydrolase

- a. Introduction to protein structure
- b. Visualization using PyMol
- c. SDS PAGE (general protein stain) vs. native PAGE (OPH specific activity stained)