# **COURSE DETAIL**

### **ISSUES IN DRUG DESIGN AND DEVELOPMENT**

## **Country**

New Zealand

#### **Host Institution**

University of Auckland

## Program(s)

University of Auckland

#### **UCEAP Course Level**

**Upper Division** 

## **UCEAP Subject Area(s)**

Chemistry

### **UCEAP Course Number**

189

### **UCEAP Course Suffix**

#### **UCEAP Official Title**

ISSUES IN DRUG DESIGN AND DEVELOPMENT

## **UCEAP Transcript Title**

DRUG DESIGN & DEVEL

# **UCEAP Quarter Units**

6.00

### **UCEAP Semester Units**

4.00

### **Course Description**

Intellectual property and patent law in the pharmaceutical industry. An overview of the legal and regulatory framework for drug design and development. Clinical trials: formulation of a drug; phase I, phase II and phase III protocols. An introduction to the principles involved in the Codes of Good Manufacturing Practice and Good Laboratory Practice (quality control and quality assurance procedures) as applied to the manufacture of drug products and the quantification of drugs and metabolites in biological fluids. Examples of drug development. Case studies of selected drugs from design to release.

### Language(s) of Instruction

English

### **Host Institution Course Number**

**CHEM 392** 

### **Host Institution Course Title**

ISSUES IN DRUG DESIGN AND DEVELOPMENT

# **Host Institution Campus**

Auckland

# **Host Institution Faculty**

**Host Institution Degree** 

# **Host Institution Department**

Chemistry

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