

# BB1110 Gene Technology and Molecular Biology 7.0 credits

Genteknik och molekylärbiologi

This is a translation of the Swedish, legally binding, course syllabus.

If the course is discontinued, students may request to be examined during the following two academic years

## Establishment

Course syllabus for BB1110 valid from Autumn 2011

## Grading scale

A, B, C, D, E, FX, F

#### **Education cycle**

First cycle

## Main field of study

Biotechnology, Technology

## Specific prerequisites

Completed upper secondary education including documented proficiency in English corresponding to English A. For students who received or will receive their final school grades after 31 December 2009, there is an additional entry requirement for mathematics as follows: documented proficiency in mathematics corresponding to Mathematics A. Specific requirements in mathematics, physics and chemistry are corresponding to Mathematics E, Physics B and Chemistry A.

#### Language of instruction

The language of instruction is specified in the course offering information in the course catalogue.

#### Intended learning outcomes

Breakthroughs in molecular biology and biotechnology have the last decades paved the way for completely new strategies that hold promise to solve real-world problems; some are related to the diagnosis and treatment of disease, others to the use of genetically modified organisms for detoxification of the environment or production of biofuels, while still others deal with the engineering of proteins to adapt them for specific applications.

The course's main goal is to provide a good insight into the principles and methods on which modern (molecular) biotechnology is based as well as an understanding of their inherent possibilities and limitations to address and solve modern day problems.

After passing the course, the student should be able to:

- describe the function of commonly used enzymes within the field of molecular biotechnology
- from a given problem, design a suitable PCR-setup/strategy; for example, how to clone a certain gene, and explain the function of all necessary components
- explain the principle behind different DNA-sequencing methods and discuss their possible strengths and weaknesses
- give examples of different physical and genetic strategies for modification/manipulation of gene expression and describe which consequences this will have at a cellular level
- describe different mutagenesis, screening, and selection methods that are used within protein engineering and suggest strategies for how these techniques can be applied in order to solve/address a given issue
- from a given issue or problem, choose an appropriate combination of host-vector system and describe its specific advantages and disadvantages in relation to other conceivable combinations. The student should also be able to describe/explain the function of the different vector component/elements
- describe the principles behind modern gene technology-based therapeutics such as modern vaccines and gene therapy, and give examples on some of the advantages/disadvantages and possible limitations compared with traditional treatments
- give examples of methods for transcriptome and proteome analysis and explain the underlying principles
- present and evaluate a laboratory exercise in a written report

#### **Course contents**

The course focuses on the most important gene technology principles/methods and thereto-relevant concepts in molecular biology will be reviewed. An in-depth look at prokaryotic and eukaryotic gene expression and recombinant protein production and optimization will be central. In addition, several important applications of molecular biotechnology will be presented and discussed. Some of the topics covered:

- transcription/translation regulation
- recombinant DNA (enzymes, vectors and host cells)
- PCR techniques
- DNA sequencing
- mutagenesis, gene libraries
- screening and selection methods
- design of recombinant processes (promoters, vectors, host cells, gene fusions etc)
- therapeutic strategies (vaccine technology, gene therapy etc)
- DNA diagnostics
- transgenic organisms
- functional genomics
- laboratory exercise (site directed mutagenesis, screening/selection, DNA sequencing, protein expression and characterization)

#### **Course literature**

Biotechnology: Applying the genetic revolution. 1st ed., 2009

David P. Clark & Nanette J. Pazdernik

Elsevier Academic Press ISBN 978-0-12-175552-2

#### Examination

- LAB1 Laboratory Work, 1.5 credits, grading scale: P, F
- TEN1 Examination, 5.5 credits, grading scale: A, B, C, D, E, FX, F

Based on recommendation from KTH's coordinator for disabilities, the examiner will decide how to adapt an examination for students with documented disability.

The examiner may apply another examination format when re-examining individual students.

LAB1 - Laboratory Work, 1.5 credits, grade scale: P, F

TEN1 - Examination, 5.5 credits, grade scale: A, B, C, D, E, FX, F

#### Other requirements for final grade

A written exam (TEN1; 5,5 credits, grading scale A-F) and labs (LAB1; 1,5 credits, grading scale Pass/Fail).

# Ethical approach

- All members of a group are responsible for the group's work.
- In any assessment, every student shall honestly disclose any help received and sources used.
- In an oral assessment, every student shall be able to present and answer questions about the entire assignment and solution.