



# HL2014 Safe Medical Devices

## 7.5 credits

**Säkra medicintekniska produkter**

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 HL2014 valid from Autumn 2019

### **Grading scale**

P, F

### **Education cycle**

Second cycle

### **Main field of study**

Medical Engineering

### **Language of instruction**

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

### **Intended learning outcomes**

The main objective with this course is to give the student substantial understanding about the regulatory framework for medical devices and how personal protection and intended product performance can be assured by the medical device industry and the health care sector.

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

- Describe, explain and apply in practical use the regulatory demands for medical devices.
- Describe the difference between regulatory demands in different countries.
- Explain the interaction between authorities, regulatory bodies, standardization organizations and industry when placing a medical device on the market.
- Define quality and explain different methods for assuring quality in an organization or for products or services.
- Enlarge upon the essential role of risk analysis and quality assurance for the medical device industry.
- Explain and discuss how standardization development enhances the work in the medical device industry and the health care sector.

## Course contents

Regulatory aspects: Legal prerequisites, safety, responsibilities, directives dealing with medical devices, standardization, clinical trials as a tool for demonstrating safety and efficacy, harmonization, certification and testing, product classes for devices and how to place a new product on the market. Quality: Quality as a working tool, total quality management (TQM), quality system. Safety and risk analysis. Development of quality system in industries/organizations, providing services/products for assurance that safe and efficient product reach the market.

## Specific prerequisites

At least two years of science studies at university level . Basic knowledge in medicine and medical engineering in accordance with the course HL1007 Medical engineering, basic course and and knowledge in English corresponding to English B/English 6.

## Course literature

Course literature see literature list. Reference literature and internet sites will be informed on the course website before course start

## Examination

- RED1 - Examination, 7.5 credits, grading scale: P, 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.

Approved written examination and approved project work

## **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.