COURSE SYLLABUS AND COURSE REQUIREMENTS ACADEMIC YEAR 2024-2025 SEMESTER I.

Course title	Quality Assurance and Device Certification
Course Code	MSM610AN-EA-00
Hours/Week: le/pr/lab	2/1/0
Credits	4
Degree Programme	biomedical Engineering MSc
Study Mode	
Requirements	
Teaching Period	fall
Prerequisites	
Department(s)	
Course Director	
Teaching Staff	_

COURSE DESCRIPTION

A short description of the course (max. 10 sentences). Neptun: Instruction/Subjects/Subject Details/Basic data/Subject description

•••

SYLLABUS

Neptun: Instruction/Subjects/Subject Details/Syllabus

1. GOALS AND OBJECTIVES

 ${\it Goals, student learning outcome.}$

Neptun: Instruction/Subjects/Subject Details/Syllabus/Goal of Instruction

•••

2. COURSE CONTENT

Neptun: Instruction/Subjects/Subject Details/Syllabus/Subject content

TOPICS

LECTURE	1.	History of Quality Assurance
	2.	Quality assurance and quality control topic
	3.	ISO 13485
	4.	ISO 13485
PRACTICE	1.	Target group of MDs and the needs of costumers
	2.	Quality control of MD production
	3.	Quality assurance of MD; documentation
	4.	Audit of MD production

DETAILED SYLLABUS AND COURSE SCHEDULE

ACADEMIC HOLIDAYS INCLUDED

LECTURE

LECTO				
week	Торіс	Compulsory reading; page number (from to)	Required tasks (assignments, tests, etc.)	Completion date, due date
1.	Introduction to quality assurance. Quality control and quality assurance in historical ages.			
2.	Introduction to quality assurance. Quality control and quality assurance in historical ages.			
3.	Quality standards. Lean quality standards, six sigma, lean six sigma in health care			
4.	ISO 13485 – 1. Scope, 2. Normative references, 3. Terms and definitaions, 4. Quality management system (general and documentational requirements)			
5.	ISO 14385 5 Management responsibility 5.1 Management commitment 5.2 Customer focus 5.3 Quality policy 5.4 Planning 5.5 Responsibility, authority and communication 5.6 Management review			
6.	Resource management			
<i>7.</i>	ISO 13485			
8.	ISO 13485			
9.	FALL BREAK			
10.	Medical Device regulation in the USA			
11.	Validation of medical devices			
12.	Measurements			
13.	Audits – internal audit			

week	Торіс	Compulsory reading; page number (from to)	Required tasks (assignments, tests, etc.)	Completion date, due date
1.	Target group of Medical devices – measurements of demands and needs			
2.	Quality assurance of MD production – Quality policy; chapters of the manual			
3.	Planning the design of MD according to consumers demand			
4.	Teams and workers – division of labor and responsibilities			
5.	Planning the production of MD – raw materials – suppliers; production – equipments			
6.	Planning the production of MD – transport of MD to consumers			

7.	Quality control of production of MD – flow		
	charts		
8.	Quality control of production – hazard		
	analysis; critical points		
9.	Documentations of suppliers		
10.	Documentations of production		
11.	Documentations of transport		
12.	Consumer complaints – root cause analysis		
	(backward and forward)		
13.	Validation of medical device		

3. ASSESSMENT AND EVALUATION

(Neptun: Instruction/Subjects/Subject Details/Syllabus/Examination and Evaluation System)

ATTENDANCE

In accordance with the Code of Studies and Examinations of the University of Pécs, Article 45 (2) and Annex 9. (Article 3) a student may be refused a grade or qualification in the given full-time course if the number of class absences exceeds 30% of the contact hours stipulated in the course description.

Method for monitoring attendance (e.g.: attendance sheet / online test/ register, etc.)

attendance sheet, continuous preparation and presentation of reports

ASSESSMENT

Cells of the appropriate type of requirement is to be filled out (course-units resulting in mid-term grade or examination). Cells of the other type can be deleted.

Course resulting in mid-term grade (PTE TVSz 40§(3))

Mid-term assessments, performance evaluation and their ratio in the final grade (The samples in the table to be deleted.)

Туре	Assessment	Ratio in the final grade
Preparation of practical material based on the lecture	140points	100%
material		

Opportunity and procedure for re-takes (PTE TVSz 47§(4))

The specific regulations for improving grades and resitting tests must be read and applied according to the general Code of Studies and Examinations. E.g.: all tests and assessment tasks can be repeated/improved at least once every semester, and the tests and home assignments can be repeated/improved at least once in the first two weeks of the examination period.

Preparation of full documentation

Grade calculation as a percentage

Course grade	Performance in %
excellent (5)	85 %
good (4)	70 % 85 %
satisfactory (3)	55 % 70 %
pass (2)	40 % 55 %
fail (1)	below 40 %

The lower limit given at each grade belongs to that grade.

Course-unit with final examination

Mid-term assessments, performance evaluation and their weighting as a pre-requisite for taking the final exam (The samples in the table to be deleted.)

Туре	Assessment	Weighting as a proportion of the
		proportion of the

		pre-requisite for taking the exam
1. e.g: Test 1	eg. max 20 points	eg. 20 %
2. e.g.: Test 2	eg. max 30 points	eg. 30 %
3. e.g.: home assignment (project documentation)	eg. max 30 points	eg. 30 %
4	eg. max 15 points	eg. 20 %

Requirements for the end-of-semester signature

(Eg.: mid-term assessment of 40%)

Re-takes for the end-of-semester signature (PTE TVSz 50§(2))

The specific regulations for grade betterment and re-take must be read and applied according to the general Code of Studies and Examinations. E.g.: all the tests and the records to be submitted can be repeated/improved each at least once every semester, and the tests and home assignments can be repeated/improved at least once in the first two weeks of the examination period.

Type of examination (written, oral):

The exam is successful if the result is minimum ... %. (The minimum cannot exceed 40%.)

Calculation of the grade (TVSz 47§ (3))

Calculation of the final grade based on aggregate performance in percentage.

Course grade	Performance in %
excellent (5)	85 %
good (4)	70 % 85 %
satisfactory (3)	55 % 70 %
pass (2)	40 % 55 %
fail (1)	below 40 %

The lower limit given at each grade belongs to that grade.

4. Specified literature

In order of relevance. (In Neptun ES: Instruction/Subject/Subject details/Syllabus/Literature)

COMPULSORY READING AND AVAILABILITY

- [1.] **REGULATION (EU) 2017/745 ON MEDICAL DEVICES (MDR)**, FULLY APPLICABLE FROM 26 MAY 2021; HTTPS://EURLEX.EUROPA.EU/LEGAL-CONTENT/EN/TXT/?URI=CELEX:02017R0745-20200424
- [2.] **REGULATION (EU) 2017/746 ON IN VITRO DIAGNOSTIC MEDICAL DEVICES (IVDR)**, FULLY APPLICABLE FROM 26 MAY 2022. https://eur-lex.europa.eu/legal-content/EN/TXT/?uri=CELEX:02017R0746-20170505 https://www.iso.org/
- [3.] ISO 13485 HTTPS://WWW.ISO.ORG/OBP/UI#ISO:STD:ISO:13485:ED-3:V1:EN
- [4.] PETER D. MAUCH: Quality Management Theory and Application. https://pqmonline.com/assets/files/lib/books/mouch.pdf
 RECOMMENDED LITERATURE AND AVAILABILITY
- [5.] Juran's Quality Handbook. https://gmpua.com/QM/Book/quality%20handbook.pdf
- [6.] Mohamed Zairi (2002) Beyond TQM implementation: the new paradigm of TQM sustainability, Total Quality Management, 13:8, 1161-1172, DOI:10.1080/09544120200000011
- [7.] O. M. Ikumapayi, E. T. Akinlabi, F. M. Mwema et al., Six sigma versus lean manufacturing An overview, Materials Today: Proceedings,

https://doi.org/10.1016/j.matpr.2020.02.986

[8.] Improta, G, Balato, G, Romano, M, et al. Improving performances of the knee replacement surgery process by applying DMAIC principles. *J Eval Clin Pract*. 2017; 23: 1401–1407. https://doi.org/10.1111/jep.12810