GENOMIC AND PROTEOMIC ANALYSIS

Module designation	Genomic and Proteomic Analysis
Semester(s) in which the	2
module is taught	
Person responsible for the	1. Prof. Dr. apt. Djoko Agus Purwanto, M.Si (Course
module	Coordinator)
	2 Prof Dr ant Sudiarwo MS
	2. Prof. Dr. apt. Sukardiman, MS
	S. FIOL DL. apl. Sukalullian, WS
Relation to curriculum	Compulsory / elective / specialisation
Teaching methods	lecture discussion assignment
Workload (incl contact	(Estimated) Total workload:
hours self-study hours)	Contact hours (structured activities): 90 67 hours
	Private study including independent learning activites: 90 67
	hours
Credit points	2 SCU / 6 ECTS
Required and recommended	NA
prerequisites for joining the	
module	
Module objectives/intended	Students are:
learning outcomes	LO1: Able to realize excellence based on religious morals
	(excellence with morality), able to work together, and
	show a responsible attitude to work in their field of
	expertise independently
	LO2: Able to internalize the spirit of independence,
	struggie, and entrepreneursnip
	LO4. Able to develop a pharmaceutical professional
	pharmaceutical problems and managing research in the
	pharmaceutical field related to national and global systems
	and policies, both inter and inter-disciplinary approaches.
	LO5: Able to access and review information through an
	Information and Communication Technology (ICT) system,
	decide on a specific subject of study, maintain the feasibility
	of implementing research designs, conduct research,
	analyze data, conclude research results comprehensively,
	and create strategic issues based on the study that reflect
	the latest updates in the field of pharmaceutical sciences,
	and communicate them in the media and scientific forums
	at the national and international level through an
	interdisciplinary or multidisciplinary approach in the form of
	a thesis or other equivalent forms.
	LUTT: ADIE to develop systems for evaluating the
	bioavailability of drugs in the body, pharmaceutical products
	with specific delivery systems with appropriate analytical
	methods
	I O15: Able to plan and organize concepts and procedures
	for quality assurance and recommendations on
	pharmaceutical products, which include drugs, cosmetics.
	foods, and beverages as products and therapeutic goods.
	pnarmaceutical products, which include drugs, cosmetics, foods, and beverages as products and therapeutic goods.

Content	The Genomic and Proteomics Analysis course provide learning topics about the nature and structure of DNA, PCR and electrophoresis analysis, application of PCR and electrophoresis analysis results, gene regulation, Lac operon theory, and tryptophan metabolism, immunohistochemistry, gene expression analysis for cancer research, and gene expression analysis for diabetes research.
Exams and assessment formats	Final exam or take-home written assignments
Study and examination requirements	The final grade in the module is composed of 25% presentation 65% take-home assignments, 10% in-class participation and soft-skills assessment. Students must have a final grade of 70% or higher to pass
Reading list	 Siswandono, ed., 2016. Kimia Medisinal I, Edisi Kedua. Sura¬baya: Airlangga University Press. Siswandono, 2014. Pengembangan Obat Baru. Sura¬baya: Airlangga University Press. Choudhuri, S., 2014. Bioinformatics for Beginners; Genes, Genomes, Molecular Evolution, Databases and Analytical Tools, Amsterdam: Elsevier Inc. Roy, K., 2019. Multi-Target Drug Design Using Chem- Bioinformatic Approaches, New York: Springer Science+Business Media. Cavasotto, C.N., 2016. In Silico Drug Discovery and Design; Theory, Methods, Challenges, and Applications, Boca Raton: Taylor & Francis Group, LLC. Merz, K.M., Ringe, D., Reynolds, C.H., 2010. Drug Design, Structure- and Ligand-Based Approaches, Cambridge: Cambridge University Press