BIOTRANSFORMATION AND DRUG DESIGN

Module designation	Biotransformation and Drug Design
Semester(s) in which the module is taught	2
Person responsible for the module	 Dr. Juni Ekowati, M.Si., Apt (Course Coordinator) Prof. Dr. Achmad Syachrani, MS., Apt. Dr. Nuzul Wahyuning D., M.Si., Apt.
Language	Bahasa Indonesia
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 prerequisites for joining the module	NA

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, struggle,
	and entrepreneurship.
	O3: Able to develop and build logical-critical-systematic-
	creative thinking and scientific conceptions through
	scientific research, design creation, or artworks of science
	and technology that pays attention to and applies
	humanities values through an interdisciplinary or
	multidisciplinary approach in the form of a thesis or other
	equivalent forms.
	LO4: Able to develop a pharmaceutical professional
	performance with analytical acumen in solving
	pharmaceutical problems and managing research in the
	and policies, both inter and inter disciplinary approaches
	and policies, both liner and liner-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.
	LO6: Able to make decisions in the context of solving
	problems related to science and technology development
	based on analytical or experimental studies through
	collaboration with colleagues, colleagues in institutions and
	levels and utilizing research results for the benefit of the
	user and other communities
	LO9. Able to carry out molecular manipulation of
	substances and develop formulations and manufacturing of
	pharmaceutical preparations with active pharmaceutical
	ingredients derived from natural products and synthetic
	compounds through the manufacture of polymorphs,
	nanoparticles, solid dispersions.
	LO13: Able to design drug development both from natural
	products and/or synthetic compounds by considering the
	biological mimicry system.

Content	The Biotransformation and Drug Development course describes advanced aspects of drug transformation in the body, including various biotransformation reactions, Phase I and Phase II, enzymes that play a role in biotransformation, effect of the biotransformation process on the physical, chemical, and pharmacological properties of drugs, factors affecting the biotransformation process, biotransformation of an aromatic compound, active metabolites as candidates for drug development, prodrugs in drug development, synthesis of drugs through biotransformation processes.
Exams and assessment formats	Take-home written assignments
Study and examination requirements	the final grade in the module is composed of 30% discussion, 30% presentation, 30% 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	 Michael A. Lieberman, Alisa Peet. 2015. Marks' Essentials of Medical Biochemistry A Clinical Approach 2 nd ed. Wolters Kluwer, Philadelpia B.L. Goodwin. 2005. Handbook of biotransformation of aromatic compounds. CRC Press. Washington. Kamal Shah, Durgesh N. Chauhan. Nagendra S. Chauhan. Pradeep Mishra. 2020. Recent Advancement in prodrugs. CRC Press. Washington. Vivekkumar K. Reasani & Sanjay B.Bari. 2015. Prodrug Design. Perspectives, Approaches, and application in Medicinal Chemistry. ElsevierTokyo Suzana S. et al.2008. Biotransformation of Mefenamic Acid by Cell Suspension Cultures of Solanum Mamosum. NPC 3(2) pp.113-302.