Study and examination requirements	the final grade in the module is composed of 45% mid term exam, 45% final exam , 10% in-class participation and soft- skills assessment. Students must have a final grade of 70% or higher to pass		
Reading list	 Whalen K, Pharmacology (Lippincott's Illustrated Reviews Series), 7th Edition, Lippincott Williams and Wilkins, 2019 Katzung B, Masters S, Trevor A, Basic and Clinical Pharmacology, LANGE Basic Science, 15th edition. Lange, 2021 Brenner GM and Stevens CW, Pharmacology, Second Edition, Saunders, 2006 Brunton L, Chabner B, and Knollman B, Goodman And Gilman's The Pharmacological Basis of Therapeutics, 13th edition, McGraw Hill Profesional, 2017 Vauquelin G and Mentzer BV, G Protein-coupled Receptors: Molecular Pharmacology, A John Willey and Sons, 2008 Offermanns S and Rosenthal W, Encyclopedia of Molecular Pharmacology, 3rd edition, Springer, 2021 		

PHARMACOMETRICS

Module designation	Pharmacometrics		
Semester(s) in which the module is taught	2		
Person responsible for the module	 Chrismawan Ardianto, S. Farm., Apt., M.Sc., Ph.D.(Course Coordinator) Prof. Junaidi Khotib, S.Si., Apt., M.Kes., Ph.D Prof. Dr. apt. Aty Widyawaruyanti, MSi. 		
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:
Module objectives/intended learning outcomes	Students are: 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. 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 research communities at both national and international levels and utilizing research results for the benefit of the user and other communities. LO11: Able to develop systems for evaluating the bioavailability of drugs in the body, pharmaceutical products circulation permits, and their in-vitro and in-vivo evaluations with specific delivery systems with appropriate analytical
Content	methods. The Pharmacometrics course provides an introduction to pharmacometrics (definition, scope, objectives, and benefits of pharmacometrics), the development of diseased-animal models, the validation of diseased-animal models including models of infectious disease (malaria, viruses induced and bacterial-induced), models of degenerative diseases (cancer, diabetes, hypertension, atherosclerosis, kidney failure, and aging), models of other diseases (chronic pain, inflammation, and drug dependency), drug compound potency and effectiveness testing both in vitro and in vivo, calculation of drug compounds potency and effectivity, pharmacokinetic and pharmacodynamic relationship (based on circumstances, physiology, mechanism of action, and allometric scaling), pharmacometrics models in the development of pharmacometrics models in the development of drug compounds, computer programs and their use in pharmacometrics.
Exams and assessment formats	Mid term exam and final exam
Study and examination requirements	the final grade in the module is composed of 45% mid term exam, 45% final exam , 10% in-class participation and soft- skills assessment. Students must have a final grade of 70% or higher to pass

Reading list	1. 2. 3. 4. 5.	Brutton L, et al, Goodman and Gilman's the Pharmacological Basis of Therapeutics, 13th Edition, McGraw-Hill Education 2017 Katzung B, Masters S, Trevor A, Basic And Clinical Pharmacology, Lange Basic Science, 14th edition. Lange, 2017 Vela JM., Maldonado R, Hamon M., In Vivo Models For Drug Discovery, Wiley VCH Press, 2014 Brenner GM and Stevens CW, Pharmacology, 5th Edition, Saunders, 2017 International Conference on Harmonization Good Clinical Practice (ICH GCP-E6) Integrated addedum,
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