DRUG INTERACTIONS

Module designation	Drug Interactions
Semester(s) in which the	2
module is taught	
Person responsible for the	1. Prof. Junaidi Khotib, S.Si.,Apt.,M.Kes.,Ph.D (Course
module	Coordinator)
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	NA
prerequisites for joining the	
module	

	Other law to a way
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. LO3: 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 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. LO6: Able to analytical or experimental studies through collaboration with colleagues, colleagues in institutions and research communities. LO7: Able to analyze natural materials to obtain active ingredients and/or pharmaceutical excipients with due observance of nature conservation. LO8: Able to carry out drug designs through the synthesis of bioactive compounds based on the structure-activity relationship. LO9: Able to carry out molecular
	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. LO7: Able to analyze natural materials to obtain active ingredients and/or pharmaceutical excipients with due observance of nature conservation. LO8: Able to carry out drug designs through the synthesis of bioactive compounds based on the structure-activity relationship. 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. 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 methods. LO13: Able to design drug development both from natural products and/or synthetic compounds by considering the biological mimicry system.
	LO14: Able to build drug management systems from active pharmaceutical ingredients to finished products that are ready for therapeutic uses.

Content	The Drug Interactions course presents: types of drug interactions, the concept of drug interaction mechanisms, prediction of drug interactions, prevention and resolution of drug interactions, drug interactions in the clinic
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	 David S Tatro, 2011. Drug Interaction Facts 2009: TheAuthority on Drug Interactions, Lippincot Williams & Wilkins, Informa Healthcare, New York Ashraf Mozayani D Lionel Raymon, 2012, Handbook of Drug Interactions, A Clinical and Forensic Guide, Humana Press, New York Beverly J. McCabe, Eric H. Frankel, Jonathan J., Wolfe, 2003, Handbook of Food-Interactions, CRC Press, London Karen Baxter, 2010. Stockley's Drug Interactions, Pharmaceutical Press, London Williamson E, Driver Sand Baxter K, 2009. Stockley's Herbal Medicines Interactions, Pharmaceutical Press London Manjunnah P. Pai, Jennifer J. Kiser, Paul O. Gubbins, Keith A. Rodyold (eds.) 2018. Drug Interactions in Infectious Diseases: Mechanisms and Models of Drug Interactions [4th Ed.] Clarke, Simon; Collignon, Ursula;
	 Clarke, Sinon, Congnon, Orsula, Karalliedde, Janaka; Karalliedde, Lakshman 2016. Adverse drug interactions: a handbook for prescribers [Second edition] CRC Press