Module	Pharmaceutical Sciences Technologies
Code	MLS_S23
Degree Program	Master of Science in Life Sciences (MSLS)
Cluster	Bio/Pharma
Specialization	Applied Biosciences
ECTS Credits	4
Workload	120 h: Contact 56 lessons = 42 h; Self-study 78 h
Module Coordinator	Name Origène Nyanguile
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Lecturers	PhD. Origène Nyanguile, HES-SO Valais / Wallis
	PharmD. Philippe Buthier
	Guest lecturers
Entry Requirements	Bachelor of Science in Life Technologies (orientation Biotechnology or Analytical Chemistry) or in a related course of study (Bachelor level).
Learning Outcomes and Competences	After completing the module, the student will have an end to end overview of Drug Products manufacturing processes from development stages to manufacturing activities in Pharmaceutical Industry.
	The aim of this master is to be as close as possible to the industrial practices and to give to the student an overview of activities performed in pharmaceutical Industry. Most of the lectures will be given by experts who are currently working in different key positions inside Pharmaceutical Industries.
	 The student will be able: To study the process of bringing a new pharmaceutical drug to the market, from Drug Substance discovery to Drug Product manufacturing To understand the basic of pharmaceutical manufacturing technologies, excipients and equipment used To understand the organization within the pharmaceutical industry To understand the working methodologies used by the pharmaceutical Industry

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Module Content	Drug discovery, hit to lead optimization,
	Drug Product definition (different dosage forms),
	Pharmaceutical Packaging Overview,
	 Pharmaceutical solid dosages manufacturing processes and control strategy description,
	 Pharmaceutical liquid and semi-liquid dosages manufacturing processes and controls description
	Pharmaceutical Product Development and continuous improvements
	Pharmaceutical Product Manufacturing Activities, visit of Drug Products manufacturing site and discussion with key people of the organization
	Main Excipients used in Pharmaceutical Industry, DCI, trade names, suppliers, description and regular utilization.
	 Case studies on one Drug Product from development stage to industrial technical transfer.
Teaching / Learning	Lectures, seminar-style work, practical work, and exercise.
Methods	Active participation in the module is requested
	The reports and presentations related to practical work or for seminars must be validated to gain access to the module examination.
	Final examination (written): 100 % of the final grade
	Reassessment: written exam
Bibliography	The Practice of Medicinal Chemistry: Camille Georges Wermuth
	Handbook of Pharmaceutical Excipients sixth edition: Raymond C. Rowe, Paul J. Sheskey, Siân C. Owen, American Pharmacists Association
	Handbook of Pharmaceutical Granulation Technology: Dilip M. Parikh
	Pharmacie galénique : formulation et technologie pharmaceutique : Prof Werhlé
	Phi 41 : groupe IMT
	ICH guidelines, European Pharmacopoeia.
	SUPAC Guidelines from FDA,
	European Pharmacopoeia,
	United States Pharmacopeia.
Language	English
Comments	

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