

Teacher(s)	. SOMEBODY ;Marcelis Xavier (coordinator) ;Pronce Thierry ;
Language :	French
Place of the course	Bruxelles Woluwe
Prerequisites	The student will have learned some fundamentals supporting the activities of the pharmaceutical industry and how it is regulated by imposed requirements.
Main themes	<p>Quality and Quality Assurance requirements are defined for each and every level they appear in pharmaceutical industry, i.e. for rooms and environment classification, equipment and machines, products (intermediate raw materials and finished products), documentation (including procedures, methods, processes descriptions, raw data) and working staff (organisation, responsibilities, job description). There are many situational and practical examples available to illustrate the content.</p> <p>Thèmes abordés :</p> <p>Principes de management pharmaceutique, 10h, VANDERHOFSTADT Jean-Michel (ULiège)</p> <p>Assurance qualité, partim a: Concepts de base et organisation de l'assurance qualité, 14,5h, PRONCE Thierry (UCLouvain)</p> <p>Assurance qualité, partim b : Technologie analytique des procédés et analyse de risques, 10h, MARCELIS Xavier (UCLouvain)</p> <p>Anglais appliqué à l'industrie pharmaceutique, 20h, POUPAERT Jacques, 20h, SERBEST Nevin (UCLouvain)</p> <p>Marketing pharmaceutique, 7,5h, BIERLAIRE Vincent (ULiège)</p>
Learning outcomes	<p><b>At the end of this learning unit, the student is able to :</b></p> <ul style="list-style-type: none"> <li>- Define fundamentals of Quality, total Quality Management, Good Manufacturing Practices</li> <li>- Introduce the student to the goals and concept behind Quality Assurance and Quality Control in the pharmaceutical industry.</li> </ul> <p>1</p> <ul style="list-style-type: none"> <li>- Define the scope of Quality Assurance and establish the « compliance » criteria according to the regulatory requirements.</li> <li>- Introduce the notions of "Process Analytical Technology" (PAT) and "Risk Management" in the frame of the FDA initiative called "Pharmaceutical CGMP's for the 21st Century".</li> </ul>
Evaluation methods	The examination will be orally conducted within the course module
Teaching methods	The learning will be provided through slides (PowerPoint) and practical exercises. A copy of those documents will be delivered to the student.
Content	<p>Part A</p> <p>Introduction to Quality systems in general terms and their specific application into the pharmaceutical field.</p> <p>Quality Assurance - Quality Control definitions.</p> <p>Quality Assurance organisation.</p> <p>Quality Assurance philosophy.</p> <p>Quality Assurance pillars with their implication to different levels.</p> <ul style="list-style-type: none"> <li>-personnel</li> <li>-material</li> <li>-equipment</li> <li>-processes</li> <li>-documentation</li> </ul> <p>Installations and equipment qualification.</p> <p>Internal ' External audit preparation.</p> <p>Suitable behaviour during an audit.</p> <p>Part B</p> <p>Process validation.</p> <p>Presentation of the "Process Analytical Technology" (PAT) concept and of the "Risk Management" approach with practical examples</p>

Other infos	The learning will be provided through slides (PowerPoint) and practical exercises. A copy of those documents will be delivered to the student.
Faculty or entity in charge	FARM

<b>Programmes containing this learning unit (UE)</b>				
Program title	Acronym	Credits	Prerequisite	Learning outcomes
Advanced Master in Industrial Pharmacy	FARI2MC	7		