UCLouvain

wfari2105

2025

Affaires réglementaires et environnement médico-social

The version you're consulting is not final. This course description may change. The final version will be published on 1st June.

8.00 credits	72.0 h	Q1

This learning unit is not open to incoming exchange students!

Teacher(s)	. SOMEBODY ;Druez Catherine ;
Language :	French
Place of the course	Bruxelles Woluwe
Learning outcomes	
Evaluation methods	The module exam could be placed outside the official June session. Written exam including questions proposed by each speaker. In the event of exceptional circumstances, examination arrangements may be reviewed.
Content	GMP (good manufacturing practices) and International Standards in the engineering of pharmaceutical and related plants. Impact on plant construction, premises, personnel and production equipment. Circuit analysis. Harmonization of pharmaceutical legislation within the European Union. International technical harmonization (ICH: International Conference of Harmonisation). Drug marketing authorization (AMM) procedures. Intellectual property and patents.
Faculty or entity in charge	FARM

Programmes containing this learning unit (UE)

Program title	Acronym	Credits	Prerequisite	Learning outcomes
Advanced Master in Industrial Pharmacy	FARI2MC	8		