

Block view of the study programme

Or Th Pr Au Cr

Information

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Collegiality

For the ULg : Vincent BIERLAIRE, Bruno BOULANGER, Raphaël DENOZ, Laurence DENIS, Brigitte EVRARD, Marianne FILLET, Michel FREDERICH, Philippe HUBERT, Roland MARINI, Dominique MARTIN, Bernard PIROTTE, Régis RADERMECKER, Jean-Michel VANDERHOFSTADT, Eric ZIEMONS.

For the ULB : Karim AMIGHI, Sonja BEKEN, Daniel BRASSEUR, Jean-Antoine DE MUYLDER, Patrick DI STEFANO, Pierre DUEZ, François DUFRASNES, Véronique FONTAINE, Ghanem GHANEM, Jean-Michel KAUFFMAN, A. LENAERS, Pierre VAN ANTWERPEN, Francis VANDERBIST, David VERMIJLEN.

For the UCL : Catherine DRUEZ, Laure ELEN, Xavier MARCELIS, François-Xavier MATHY, Jacques POUPAERT, Thierry PRONCE, Joëlle QUETIN-LECLERCQ, Nevin SERBEST, Rita VAN BEVER.

Study programme

Block 1

Compulsory courses

Code	Course Title	Q1	Th	Pr	Au	Cr
PHIN2004-1	<i>Active substances</i>	Q1				6
	- <i>Substances issues de recherches pharmacochimiques, part a</i> - Bernard PIROTTE	10	-	-		
	- <i>Substances issues de recherches pharmacochimiques, part b</i> - Françoise DUFRASNE	5	-	-		
	- <i>Substances issues des biotechnologies</i> - David VERMIJLEN	15	-	-		
	- <i>Substances d'origine naturelle, part a</i> - Joëlle QUETINLECLERCQ	5	-	-		
	- <i>Substances d'origine naturelle, part b</i> - Pierre DUEZ	5	-	-		
	- <i>Produits radiopharmaceutiques</i> - Ghanem GHANEM	10	-	-		
PHIN2008-2	<i>Clinical viewpoints</i>	Q1				6
	- <i>Métabolisme des médicaments et paramètres pharmacocinétiques</i> - FrançoisXavier MATHY	20	-	-		
	- <i>Aspects théoriques et pratiques des études cliniques (y compris les méthodes statistiques appliquées aux études cliniques)</i> - Régis RADERMECKER	15	-	-		
	- <i>Information et pharmacovigilance</i> - Raphaël DENOZ	10	-	-		
PHIN2013-2	<i>Quality assurance and pharmaceutical management</i>					7
	- <i>Principles of pharmaceutical management</i> - JeanMichel VAN DER HOFSTADT	10	-	-		
	- <i>Quality assurance, part a : basic concepts and quality assurance organisation</i> - Thierry PRONCE	20	-	-		
	- <i>Quality assurance, part b : Analytical technology of procedures and risk analysis</i> - Xavier MARCELIS	7,5	-	-		
	- <i>English applied to the pharmaceutical industry</i> - Jacques POUPAERT, Nevin SERBEST	20	-	-		
- <i>Pharmaceutical marketing</i> - Vincent BIERLAIRE	7,5	-	-			
PHIN2033-1	<i>Industrial pharmaceutical technology</i>					8
	- <i>Industrial pharmaceutical microbiology</i> - Véronique FONTAINE	15	-	-		
	- <i>Preformulation and selection of galenical forms</i> - Karim AMIGHI	15	-	-		
	- <i>Industrial production of galenical forms</i> - Brigitte EVRARD	15	-	-		
	- <i>Industrial production of biomolécules</i> - Rita VAN BEVER	15	-	-		
	- <i>Industrial aspects of technological development including packaging</i> - Laurence DENIS	10	-	-		
PHIN2023-1	<i>Drug analysis</i>					7
	- <i>Analytical methods and pharmaceutical and biopharmaceutical control, part a</i> - JeanMichel KAUFFMAN, Pierre VAN ANTWERPEN	10	-	-		
	- <i>Analytical methods and pharmaceutical and biopharmaceutical control,</i>	10	-	-		

	<i>part b - Marianne FILLET</i>			
	- <i>Methods in pharmaceutical and biopharmaceutical analysis, part a : Validation and qualification of equipment</i> - Philippe HUBERT, Roland MARINI DJANG'EING'A	15	-	-
	- <i>Methods in pharmaceutical and biopharmaceutical analysis, part b : Process analytical technology</i> - Philippe HUBERT, Eric ZIEMONS	5	-	-
	- <i>Statistical methods applied to the pharmaceutical industry</i> - Laure ELENS	15	-	-
	- <i>Experimental planning and "quality by design"</i> - Bruno BOULANGER	10	-	-
PHIN2029-2	<i>Regulation and the medical-social environment</i>			8
	- <i>Economic aspects of drug development</i> - Dominique MARTIN	10	-	-
	- <i>Legislation and procedures applied to pharmaceutical industry, part a : legislation</i> - Catherine DRUEZ	10	5	-
	- <i>Legislation and procedures applied to pharmaceutical industry, part b : patents and protection</i> - Patrick DI STEFANO	5	-	-
	- <i>Macroeconomic environment and pharmaco-economics</i> - Pierre DUEZ	10	-	-
	- <i>CTD File (Common Technical Document)</i> - JeanAntoine DE MUYLDER	15	-	-
	- <i>Regulations of preclinical and clinical studies, part a : pharmaceutical toxicological files</i> - Sonia BEKEN	15	-	-
	- <i>Regulations of preclinical and clinical studies, part b : clinical studies</i> - Anne LENAERS	5	-	-
	- <i>Regulations of preclinical and clinical studies, part c : pediatric studies</i> - Daniel BRASSEUR	2,5	-	-
	- <i>Specific regulatory issues, part a : medicine and herbal dietary supplement</i> - Michel FREDERICH	5	-	-
	- <i>Specific regulatory issues, part b : Preformulation and galenical development documentation</i> - Francis VANDERBIST	5	-	-
PHIN2032-1	<i>Visits and seminars organised in the pharmaceutical industry</i> - Françoise DUFRASNE, Marianne FILLET, Rita VAN BEVER - [77,5h Vis.]	-	-	[+] 3
MTFE2000-1	<i>Final work</i> - Brigitte EVRARD, Philippe HUBERT	-	-	15