Cycle view of the study programme

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Information

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Collegiality

For ULiege: Vincent BIERLAIRE, Bruno BOULANGER, Raphaël DENOOZ, Laurence DENIS, Brigitte EVRARD, Marianne FILLET, Michel FREDERICH, Philippe HUBERT, Roland MARINI, Dominique MARTIN, Bernard PIROTTE, Régis RADERMECKER, Caroline STEVIGNY, Jean-Michel VANDERHOFSTADT, Eric ZIEMONS.
For ULB: Patrick DI STEFANO, François DUFRASNE, Véronique FONTAINE, Jonathan GOOLE, Anne LENAERS, Hugues MALONNE, Thierry SCHURMANS, Pierre VAN ANTWERPEN, Francis VANDERBIST, Karen VAN MALDEREN, David VERMIJLEN.
For UCL: Catherine DRUEZ, Laure ELENS, Xavier MARCELIS, François-Xavier MATHY, Jacques POUPAERT, Thierry PRONCE, Joëlle QUETIN-LECLERCQ, Nevin SERBEST, Rita VANBEVER.

Compulsory courses (B1 : 60Cr)

PHIN2004-1	*Active substances*	B1	TA				**6**
	- Substances issues de recherches pharmacochimiques, part a - Bernard PIROTTE		10	-	-		
	- Substances issues de recherches pharmacochimiques, part b - Françoise DUFRASNE		5	-	-		
	- Substances issues des biotechnologies - David VERMIJLEN		15	-	-		
	- Substances d'origine naturelle, part a - Joëlle QUETINLECLERCQ		5	-	-		
	- Substances d'origine naturelle, part b - Pierre DUEZ		5	-	-		
	- Produits radiopharmaceutiques - Ghanem GHANEM		10	-	-		
PHIN2008-2	*Clinical viewpoints*	B1	TA				**6**
	- Métabolisme des médicaments et paramètres pharmacocinétiques - FrançoisXavier MATHY		20	-	-		
	- Aspects théoriques et pratiques des études cliniques (y compris les méthodes statistiques appliquées aux études cliniques) - Régis RADERMECKER		15	-	-		
	- Information et pharmacovigilance - Raphaël DENOOZ		10	-	-		
PHIN2013-2	*Quality assurance and pharmaceutical management*	B1	TA				**7**
	- Principles of pharmaceutical management - JeanMichel VAN DER HOFSTADT		10	-	-		
	- Quality assurance, part a : basic concepts and quality assurance organisation - Thierry PRONCE		20	-	-		
	- Quality assurance, part b: Qualification and validation approach and risk analysis - Xavier MARCELIS		7,5	-	-		
	- English applied to the pharmaceutical industry - Jacques POUPAERT, Nevin SERBEST		20	-	-		
	- Pharmaceutical marketing - Vincent BIERLAIRE		7,5	-	-		
PHIN2033-1	*Industrial pharmaceutical technology*	B1	TA				**8**
	- Industrial pharmaceutical microbiology - Véronique FONTAINE		15	-	-		
	- Preformulation and selection of galenical forms - Karim AMIGHI		15	-	-		
	- Industrial production of galenical forms - Brigitte EVRARD		15	-	-		
	- Industrial production of biomolecules - Rita VAN BEVER		15	-	-		
	- Industrial aspects of technological development including packaging - Laurence DENIS		10	-	-		
PHIN2023-1	*Drug analysis*	B1	TA				**7**
	- Methods in pharmaceutical and biopharmaceutical analysis, part a : Detection of (bio)pharmaceutical products - JeanMichel KAUFFMAN, Pierre VAN ANTWERPEN		10	-	-		
	- Methods in pharmaceutical and biopharmaceutical analysis, part b :		10	-	-		

	Preparation and separation of (bio) pharmaceutical products for quality control and pharmacokinetic study - Marianne FILLET						
	- Methods in pharmaceutical and biopharmaceutical analysis, part c : Validation and qualification of equipment - Philippe HUBERT, Roland MARINI DJANG'EING'A		15	-	-		
	- Methods in pharmaceutical and biopharmaceutical analysis, part d : Vibrational spectroscopy - Eric ZIEMONS		5	-	-		
	- Statistical methods applied to the pharmaceutical industry - Laure ELENS		15	-	-		
	- Experimental planning and "quality by design" - Bruno BOULANGER		10	-	-		
PHIN2029-2	*Regulation and the medical-social environment*	B1	TA				**8**
	- Economic aspects of drug development - Dominique MARTIN		10	-	-		
	- Legislation and procedures applied to pharmaceutical industry, part a : legislation - Catherine DRUEZ		10	5	-		
	- Legislation and procedures applied to pharmaceutical industry, part b : patents and protection - Patrick DI STEFANO		5	-	-		
	- Macroeconomic environment and pharmaco-economics - Hugues MALONNE		10	-	-		
	- CTD File (Common Technical Document) - JeanAntoine DE MUYLDER		15	-	-		
	- Regulations of preclinical and clinical studies, part a : pharmaceutical toxicological files - Sonia BEKEN		15	-	-		
	- Regulations of preclinical and clinical studies, part b : clinical studies - Anne LENAERS		5	-	-		
	- Regulations of preclinical and clinical studies, part c : pediatric studies - Daniel BRASSEUR		2,5	-	-		
	- Specific regulatory issues, part a : medicine and herbal dietary supplement - Michel FREDERICH		5	-	-		
	- Specific regulatory issues, part b : Preformulation and galenical development documentation - Francis VANDERBIST		5	-	-		
PHIN2032-1	*Visits and seminars organised in the pharmaceutical industry* - Françoise DUFRASNE, Marianne FILLET, Rita VAN BEVER - [77,5h Vis.]	B1	TA	-	-	[+]	**3**
MTFE2000-1	*Final work* - Philippe HUBERT	B1	TA	-	-	-	**15**