Cycle view of the study programme

Information

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Collegiality

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ULg : Vincent BIERLAIRE, Bruno BOULANGER, Raphaël DENOOZ, Dominique MARTIN, Laurence DENIS, Brigitte EVRARD, Marianne FILLET, Bernard PIROTTE, Eric ROZET, André SCHEEN, Monique TITS, Jean-Michel VANDERHOFSTADT, Eric ZIEMONS.
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.
Pour l'UCL : Catherine DRUEZ, Bernadette GOVAERTS, Philippe LEVEQUE, Xavier MARCELIS, Jacques POUPAERT, Joëlle QUETIN-LECLERCQ, Nevin SERBEST, Jean SCOUVART, Rita VAN BEVER, Roger VERBEECK, N...

Study programme

Compulsory courses (B1 : 60Cr)

		Bl	Or	Th	Pr	Au	Cr
PHIN2004-1	*Active substances*	B1	Q1				**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					**6**
	- *Métabolisme des médicaments et paramètres pharmacocinétiques* - Roger VERBEEK		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					**7**
	- *Principles of pharmaceutical management* - JeanMichel VAN DER HOFSTADT		10	-	-		
	- *Quality assurance, part a : basic concepts and quality assurance organisation* - Jean SCOUVART		20	-	-		
	- *Quality assurance, part b : Analytical technology of procedures 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					**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* - Philippe LEVEQUE, Rita VAN BEVER		15	-	-		
	- *Industrial aspects of technological development including packaging* - Laurence DENIS		10	-	-		
PHIN2023-1	*Drug analysis*	B1					**7**
	- *Analytical methods and pharmaceutical and biopharmaceutical control, part a* - JeanMichel KAUFFMAN, Pierre VAN ANTWERPEN		10	-	-		

	- *Analytical methods and pharmaceutical and biopharmaceutical control, part b* - Marianne FILLET		10	-	-		
	- *Methods in pharmaceutical and biopharmaceutical analysis, part a : Validation and qualification of equipment* - Philippe HUBERT, Roland MARINI DJANG'EING'A		15	-	-		
	- *Methods in pharmaceutical and biopharmaceutical analysis, part b : Process analytical technology* - Philippe HUBERT, Eric ZIEMONS		5	-	-		
	- *Statistical methods applied to the pharmaceutical industry* - Bernadette GOVAERTS		15	-	-		
	- *Experimental planning and "quality by design"* - Bruno BOULANGER		10	-	-		
PHIN2029-2	*Regulation and the medical-social environment*	B1					**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* - Pierre DUEZ		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* - A 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, Brigitte EVRARD, Rita VAN BEVER - [77,5h Vis.]	B1		-	-	[+]	**3**
MTFE2000-1	*Final work* - Françoise DUFRASNE, Brigitte EVRARD, Rita VAN BEVER	B1		-	-	-	**15**