Block view of the study programme

Information

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Collegiality

Coordinator : Philippe HUBERT
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

Bloc 1 du programme de l'année

Compulsory courses

			Or	Th	Pr	Au	Cr
PHIN2004-1	*Active substances*		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*						**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*						**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*						**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*						**7**
	- Analytical methods and pharmaceutical and biopharmaceutical control,			10	-	-	

	part a - JeanMichel KAUFFMAN, Pierre VAN ANTWERPEN - *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*				**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.]	-	-	[+]	**3**
MTFE2000-1	*Final work* - Françoise DUFRASNE, Brigitte EVRARD, Rita VAN BEVER	-	-	-	**15**