# Course unit name: CLINICAL PHARMACOKINETICS OF ANTINEOPLASTIC DRUGS

# 1.- General information

Code	303008	Plan		ECTS	3	
Туре	Elective	Course 2021/2022 Periodicity 1st Semeste				
Department	Pharmaceutical Sciences					
Virtual	Platform:	Studium:				
Platform	URL de Acces:	https://studium.usal.es/				

# Faculty

Professor coodinator	Dr. Jonás Samuel Pérez Blanco				
Department	Pharmaceutical Sciences				
Research area	Pharmacy and Pharmaceutical Technology				
Center	Faculty of Pharmacy				
URL Web	diarium.usal.es/jsperez/				
E-mail	jsperez@usal.es	Phone	+34 677584202		

Professor coodinator	Dra. Amparo Sánchez Navarro				
Department	Pharmaceutical Sciences				
Research area	Pharmacy and Pharmaceutical Technology				
Center	Faculty of Pharmacy				
URL Web	https://farmaciaytecnologia.org/				
E-mail	asn@usal.es	Phone	+34 677584152		

Professor	Dra. Marina Holgado Madruga				
Department	Phisiology and Pharmacology				
Research area	Pharmacology				
Center	Faculty of Medicine				
E-mail	mholgado@usal.es Phone 923294500 Ext.:14		923294500 Ext.:1488		

Professor	Maria José García Sánchez				
Department	Pharmaceutical Sciences				
Research area	Pharmacy and Pharmaceutical Technology				
Center	Faculty of Pharmacy				
URL Web	https://farmaciaytecnologia.org/				
E-mail	mjgarcia@usal.es	Phone	+34 677584201		

Professor	José Germán Sánchez Hernández			
Department	Hospital Pharmacy Service			
Research area	Clinical Pharmacokinetics			
Center	University Hospital of Salamanca			
URL Web	https://farmaciaytecnologia.org/			
E-mail	Jgermansanchez@salu dcastillayleon.es	Phone	+34 685552072	

### 2.- The course in the context of the Master's Program

### Treaning Module

Thirth block (out of six) of master program organization.

### General aim of the subject

The objective of this subject is to review and apply the basic concepts of clinical pharmacokinetics (PK) together with "in silico" tools for optimization of the pharmacological treatment of cancer diseases. Clinical PK contributes to the precision medicine in personalising drug treatments according to patient idyosincrasy. In addition, therapeutic drug monitoring (TDM) allows dosage individualization to improve clinical outcomes in terms of efficacy and/or safety.

### Professional specialization

This subject is oriented to researchers and clinicians involved in clinical investigation, drug development and improvement of antineoplastic drugs. These include pharmacists, clinicians, biologists, biotechnologists and other professionals who are integrated in multidisciplinary groups working on optimization of pharmacological treatment of cancer diseases.

### 3.- Previous recommendations

No prior requirements.

### 4.- Aims of the subject

To adquire theoretical and practical knowledge about antineoplastic drugs oriented to the study of its pharmacokinetics (PK) and the main factors responsible for PK variability.

To adquire the ability to apply "in silico" and TDM tools to incorporate PK variability for precision dosage in the clinical practice.

### Specific aims:

- To know the mecanism of action of the main antineoplastic drugs used in the clinical practice
- To understand the population pharmacokinetic (PopPK) modeling and simulation methodology and the factors with a relevant impact on patient exposure to antineoplastic drug (demographics, phisiopathological, genetics, etc.)
- To study the concepts and tools regarding TDM of antineoplastic drugs in the clinical routine
- To learn about phisiological based pharmacokinetic (PBPK) modeling and simulations approach and its application to *in silico* clinical trials
- To achieve a holistic knowledge on the operation in a hospital pharmacy service for development, validation and follow-up of oncological therapies

#### 5.- Contents

### **TOPICS (LECTURES):**

- 1. Mecanism of action of the main anticancer drugs used in the clinical practice
- 2. Clinical pharmacokinetics: basic concepts and application to antineoplastic drugs
- 3. Population pharmacokinetics (PopPK)
- 4. Physiological Based Pharmacokinetic (PBPK) models
- 5. Model-informed precision dosing and follow-up criteria in oncologyc treatments

### **SEMINARS and HANDS-ON:**

- 1. Data handling and Bayesian estimation
- 2. Implementation of population pharmacokinetic models
- 3. Parameters estimation of antineoplastic drugs: case reports
- 4. Aplication of PBPK models to oncology patients
- 5. In silico clinical trials
- 6. Development and validation of the oncology therapy in a hospital pharmacy service

# 6.- Skills to be acquired

### Basic skills

Understanding the usefulness of clinical PK to evaluate factors with a significant impact on the response to pharmacological treatments

Capacity to apply dosage individualization tools in the oncological patient

Ability to use the PopPK models to improve the efficacy and safety of treatments with antineoplastic drugs

### Specific skills

- Interpretation and aplicacion of TDM results to optimize and individualize pharmacological treatments with antineoplastic drugs
- Using pharmacokinetic information to select the dosage regimen with the optimal benefit/risk ratio for antineoplastic drugs
- Ability to perform clinical trial in virtual populations
- Understanding the multidisciplinarity of the clinical team involved in the validation and follow-up of onco-hematolologic therapies

## 7.- Teaching methodology

- Lectures
- Seminars
- Hands-on
- Case reports discussions
- Focused activities: presentation, analysis and proposals related to scientific papers

# 8.- Estimated learning time

		Hours tutored by the teacher		Individual	TOTAL
		Attendance required (hours)	Distance learning (hours)	work (hours)	HOURS
Lectures		11	8	10	29
	- In classroom				
Drasticas	- In laboratory				
Practices	- In computer classroom	6			6
	- Countryside				
	- Visualization classroom				
Seminars		4		2	6
Work presentations and debates		4		7	11
Tutorials		5		2	7
Online activities					
Work preparation		2	5	5	12
Other activities		4			4
	TOTAL	36	13	26	75

### 9.- Materials

#### Books

Individualizing Dosage Regimens of Antineoplastic Agents. In Individualized Drug Therapy for Patients: Basic foundations, Relevant software and clinical applications. Ed. Jelliffe R and Neely M. Elsevier. 281-306, 2017.

<u>A First Course in Pharmacokinetics and Biopharmaceutics</u> by David Bourne: <u>http://www.boomer.org/c/p4/</u>

### Scientific Journals

- Therapeutic Drug Monitoring
- Clinical Pharmacokinetics
- British Journal of Clinical Pharmacology

#### Recommended lecture

- Veal GJ, et al. Pharmacodynamic Therapeutic Drug Monitoring for Cancer:
  Challenges, Advances, and Future Opportunities. Ther Drug Monit;41:142–159. 2019
- Evan J. B & Paul K. L. A unified pharmacokinetic approach to individualized drug dosing. Br J Clin Pharmacol. 73:365-2125. 2011
- Guidi M, Csajka C, Buclin T. Parametric Approaches in Population Pharmacokinetics.
  J Clin Pharmacol. 2020 Oct 26. doi: 10.1002/jcph.1633. Epub ahead of print. PMID: 33103774. https://doi.org/10.1002/jcph.1633
- Darwich, A. S., Polasek, T. M., Aronson, J. et al. (2021). Model-Informed Precision Dosing: Background, Requirements, Validation, Implementation, and Forward Trajectory of Individualizing Drug Therapy. Annual Review of Pharmacology and Toxicology, 61(1), 225–245. <a href="https://doi.org/10.1146/annurev-pharmtox-033020-113257">https://doi.org/10.1146/annurev-pharmtox-033020-113257</a>

# 10.- Assessment

### Assessments on the performance of the student

Continous follow-up of the capacities, abilities and knowledge adquired by the students. Student participation will be highly appreciated and positively taking into account.

Attendence and active participation in lectures and seminars (hands-on).

Comments and proposals to the case-studies

Scientific accuracy of the commentaries and answers to the questions arise

Presentation and debate of a scientific paper

## Recommendations

Active participation in the proposed activities

Debate about the multidisciplinarity of the oncological treatments.