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| Course unit English denomination | Nanopharmaceutical and biopharmaceutical: physicochemical/pharmacokinetic correlations  |
| Teacher in charge (if defined)   | SALMASO Stefano, Paolo Caliceti   |
| Teaching Hours                   | 24  |
| Number of ECTS credits allocated | 3   |
| Course period                    | 06-07/2025  |
| Course delivery method           | <input type="checkbox"/> In presence<br><input type="checkbox"/> Remotely<br><input checked="" type="checkbox"/> Blended  |
| Language of instruction          | English   |
| Mandatory attendance             | <input checked="" type="checkbox"/> Yes (75% minimum of presence)<br><input type="checkbox"/> No  |
| Course unit contents             | <p>Introduction into pharmacokinetics and pharmacodynamics, PK and PD of small MW drugs.</p> <p>Pharmacokinetics (PK) and pharmacodynamics (PD) terminology, Physiological basis for drug (small MW compound) distribution and elimination, Major PK parameters: clearance, volume of distribution, elimination half-life, Major properties of compartmental and physiologically-based PK (PBPK) models, PK-PD correlations (sigmoid E max model).</p> <p>PK and PD of nano-drug delivery systems (DDSs). Targeted drug delivery for enhancement of drug effectiveness and safety, Major pathways of nano-DDS disposition following systemic administration. Effect of the nano-DDS formulation properties (size, charge, composition, targeting residues) on their systemic disposition and accumulation in solid tumors. Analytical issues: quantification of nano-DDS-encapsulated vs. free drug in the systemic circulation and at the site of action (solid tumor). Problems with limited drug/DDS permeability into the 'deep' parts of the solid tumor (i.e., cells that are distant from the capillaries). Modeling analysis of rate-limiting steps of nano-DDSs systemic and intratumoral disposition. Strategies to modulate the nano-DDSs disposition for enhancing their therapeutic effectiveness</p> <p>(PK) and pharmacodynamic (PD) properties of biopharmaceuticals. Immunogenicity and PK/PD of biopharmaceuticals. Target-Mediated Disposition of biopharmaceuticals. PK and PK-PD modeling of biopharmaceuticals and its use in pre-clinical and clinical drug development.</p> |
| Learning goals                   | <p>Knowledge: general concepts on PK/PD correlation for diverse biotechnological drugs and colloidal carriers of drugs</p> <p>Skills: Ability to foresee the PK and biodistribution and thus PD of drugs and drug carriers based on rational physico-chemical and physiological concepts.</p> <p>Competencies: Integration of the skills with the capacity of proposing projects related to drug delivery and advanced formulation of drugs with critical features</p>  |
| Teaching methods                 | Frontal teaching  |



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Course on  
transversal,  
interdisciplinary,  
transdisciplinary  
skills

Yes  
 No

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Available for PhD  
students from other  
courses

Yes  
 No

Students external to the PhD Course admitted upon evaluation of the CV by the teachers

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Prerequisites  
(not mandatory)

No specific prerequisites.

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Examination  
methods

Students will be group tasked to propose and design some rational formulation and delivery approaches for difficult to administer drugs based on the knowledge and contents provided during the course. The exam will be oral with in depth discussion.

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Study material

Slides/articles provided by the teacher

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Additional  
information  
(not mandatory)

max 3750 caratteri

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