



PHARMACEUTICAL TECHNOLOGICAL ELEMENTS

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| Enrollment year | 2019/2020 |
| Academic year | 2021/2022 |
| Regulations | DM270 |
| Academic discipline | CHIM/09 (APPLIED TECHNOLOGICAL PHARMACEUTICS) |
| Department | DEPARTMENT OF BIOLOGY AND BIOTECHNOLOGY "LAZZARO SPALLANZANI" |
| Course | BIOTECHNOLOGY |
| Curriculum | PERCORSO COMUNE |
| Year of study | 3° |
| Period | 1st semester (01/10/2021 - 14/01/2022) |
| ECTS | 6 |
| Lesson hours | 48 lesson hours |
| Language | Italian |
| Activity type | WRITTEN TEST |
| Teacher | PERTEGHELLA SARA (titolare) - 6 ECTS |
| Prerequisites | Student must have attended the courses, and acquired the basic knowledge, in biochemistry, general and inorganic chemistry. |
| Learning outcomes | At the end of the course the student will have to know the basic pharmaceutical technology. Students will also have acquired the basic principles necessary to define a pharmaceutical dosage form and to understand the rationale for the formulation of conventional drugs. |
| Course contents | Classification of pharmaceutical dosage forms and administration routes. Principles of biopharmaceutic and pharmacokinetic. Bioavailability and bioequivalence. Conventional solid pharmaceutical dosage forms. Pharmaceutical powders characterization. Grinding and mixing. Capsules and tablets (coating and controls of tablets). Conventional liquid pharmaceutical dosage forms. Solutions and dispersed systems: emulsions and suspensions. Parenteral preparations. Sterilization of |

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| | injectable preparations. Inhalator and pulmonar pharmaceutical dosage forms. pharmaceutical dosage forms for drug controlled release. Site- and time-specific therapeutic systems. Mechanisms for the controll of release rate: reservoir and matrix systems, osmotic pumps. |
| Teaching methods | Frontal lessons |
| Reccomended or required readings | Lessons' slides and books. P. Colombo et al. "Principi di tecnologie farmaceutiche". Casa Editrice Ambrosiana, Milano. A.T. Florence et al. "Physical Pharmacy". Pharmaceutical Press, London. M.E. Aulton "Pharmaceutics: the Science of Dosage Form Design". Churchil Livingstone, New York. |
| Assessment methods | Learning is verified by written exam. The subject of the examination is the contents of the reference texts and the contents of the lectures. |
| Further information | Non contents |
| Sustainable development goals - Agenda 2030 | Not applicable \$lbl legenda sviluppo sostenibile |