



Postgraduate Certificate

Drug Research and Development in Nursing

» Modality: online

» Duration: 12 weeks

» Certificate: TECH Global University

» Credits: 12 ECTS

» Schedule: at your own pace

» Exams: online

Website: www.techtitute.com/us/nursing/postgraduate-certificate/drug-research-development-nursing

Index

 $\begin{array}{ccc} \textbf{O1} & \textbf{O2} \\ \underline{\textbf{Introduction}} & \underline{\textbf{Objectives}} \\ & & \underline{\textbf{p. 4}} \end{array}$

03 04 05
Course Management Structure and Content Methodology

p. 12 p. 16 p. 22

06 Certificate

p. 30





tech 06 | Introduction

With the development of this Postgraduate Certificate in Drug Research and Development in Nursing, students will delve into the study of preclinical drug research, that is, from the time a molecule with therapeutic activity is discovered until it is marketed.

Within the field of research, professionals must also have statistical notions that allow them to carry out the Clinical Trials as accurately as possible. The use of statistics in Clinical Trials allows the researcher to reach reasonable and accurate conclusions from the data collected, and to sound out decisions when certainties are scarce.

In addition, a very important part of the Drug Research and Development process is to know how to communicate the new discoveries, which will allow further research in this field and promote its use in a generalized way, achieving the consequent benefit in patients.

Another point in favor of this program is that it is offered totally online, free of rigid schedules and the need to attend a physical center. In this way, it will be the students themselves who decide from where to study and at what time, eliminating any physical barrier. In this way, the students only need to have a computer or mobile device with an internet connection to access all the material available on the virtual campus.

This Postgraduate Certificate in Drug Research and Development in Nursing contains the most complete and up-to-date scientific program on the market. The most important features include:

- Practical cases presented by experts in Drug Research and Development
- The graphic, schematic, and practical contents with which they are created, provide scientific and practical information on the disciplines that are essential for professional practice
- News on Drug Research and Development
- Practical exercises where the self-assessment process can be carried out to improve learning
- Its special emphasis on innovative methodologies in Drug Research and Development
- Theoretical lessons, questions to the expert, debate forums on controversial topics, and individual reflection work
- Content that is accessible from any fixed or portable device with an internet connection



Prepare with us in Research and Development of Medicines and specialize until you achieve excellence in this field"



This Postgraduate Certificate is the best investment you can make in the selection of a refresher program for two reasons: in addition to updating your knowledge in Drug Research and Development, you will obtain a degree endorsed by TECH"

The teaching staff includes professionals from the health sector, who bring their experience to this educational program, as well as renowned specialists from leading societies and prestigious universities.

Theultimerdia content, developed with the latest educational technology, will provide the professional with situated and contextual learning, i.e., a simulated environment that will provide immersive education programmed to learn in real situations.

This program is designed around Problem-Based Learning, whereby the professional must try to solve the different professional practice situations that arise throughout the program. For this purpose, the professional will be assisted by an innovative interactive video system developed by renowned and experienced experts in the field of Drug Research and Development.

Do not hesitate to take this educational program with us. You will find the best teaching material with virtual lessons.

This 100% online Postgraduate Certificate will allow you to balance your studies with your professional work while increasing your knowledge in this field.





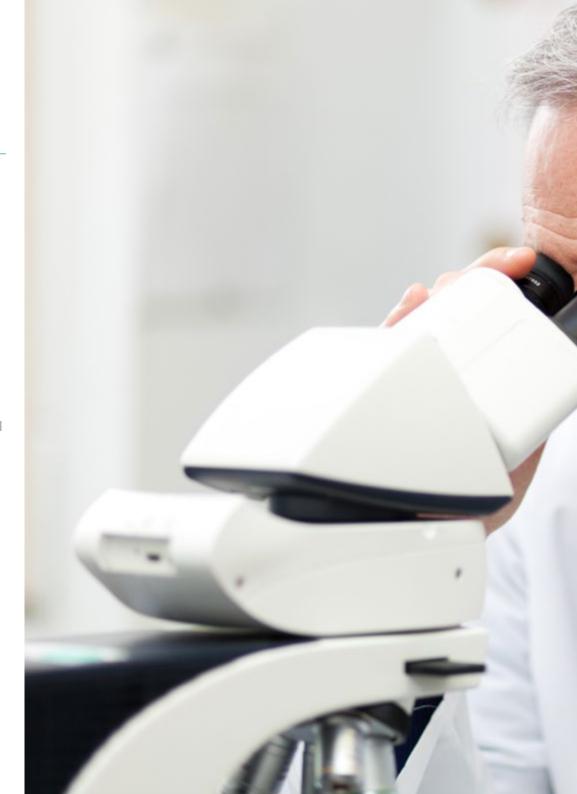


tech 10 | Objectives



General Objectives

- Establish the phases involved in the development of a new drug
- Analyze the steps prior to the development of a clinical trial (preclinical research)
- Examine how a drug is introduced into the market after the clinical trial has been conducted
- Develop knowledge that provides a basis or opportunity for originality in the development and/or application of ideas, often in a research context
- Apply the acquired knowledge and resolution skills in the development of protocols
- Structure statistical methods and techniques
- Communicate and transmit statistical results through the preparation of different types of reports, using terminology specific to the fields of application
- Compile, identify and select sources of public biomedical incapacitation, from international agencies and scientific organizations, on the study and dynamics of populations
- Analyze the scientific method and work on skills in the handling of sources of incapacitation, bibliography, elaboration of protocols and other aspects considered necessary for the design, execution and critical assessment
- Demonstrate logical thinking and structured reasoning in determining the appropriate statistical technique







Specific Objectives

Module 1. Research and Development of Medicines

- Explain the pharmacokinetic processes that a drug undergoes in the organism
- Identify the legislation that regulates each of the steps in the development and authorization of a medication
- Define the specific regulation of some drugs (biosimilars, advanced therapies).
- Define the use in special situations and their types
- Examine the process of financing a drug
- Specify strategies for the dissemination of research results
- Present how to read scientific incapacitation critically
- Compile sources of drug incapacitation and their types

Module 2. Biostatistics

- Identify and incorporate in the advanced mathematical model, which represents the experimental situation, those random factors involved in a high-level biosanitary study
- Design, collect and clean a data set for subsequent statistical analysis
- $\bullet\,$ Identify the appropriate method for determining the sample size
- Distinguish between different types of studies and choose the most appropriate type of design according to the research objective
- Communicate and transmit statistical results correctly, through the preparation of reports
- Acquire an ethical and social commitment





tech 14 | Course Management

Management



Dr. Gallego Lago, Vicente

- Military pharmacist at HMC Gómez Ulla
- Doctoral studies with the qualification of Outstanding
- Honors Degree in Pharmacy from the Complutense University of Madrid
- Resident Internal Pharmacist Examination (F.I.R) obtaining the No. 1 in this selective test
- Resident Internal Pharmacist (F.I.R) of the Pharmacy Service of the "12 de Octubre Hospital

Professors

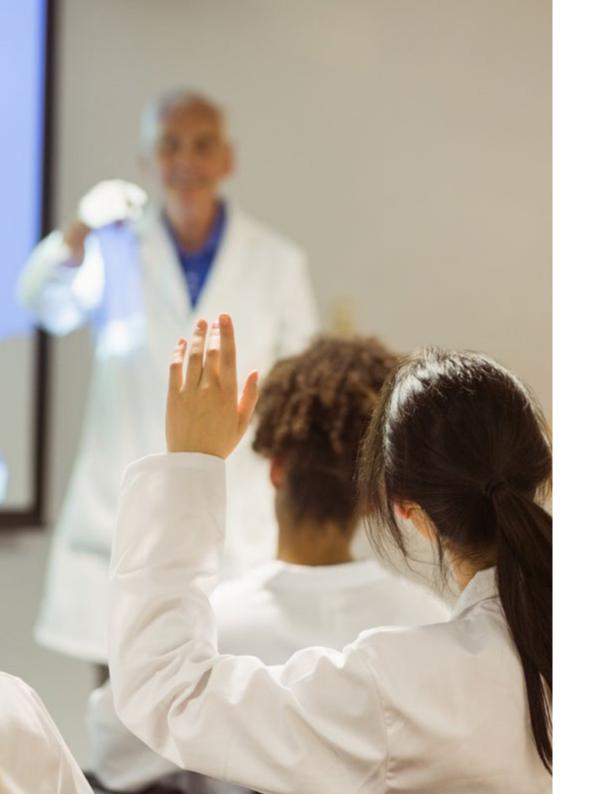
Ms. Valtueña Murillo, Andrea

- Technician in Quality, Regulation and Pharmacovigilance in Cantabria Labs
- Master in Pharmaceutical and Parapharmaceutical Industry in CESIF
- Degree in Pharmacy at Complutense University of Madrid

Ms. Martín-Arriscado Arroba, Cristina

- Biostatistics at the Research and Scientific Support Unit of the 12 de Octubre University Hospital (i+12) and the Clinical Research Units and Clinical Trials Platform (SCReN)
- Member of the Drug Research Ethics Committee of the 12 de Octubre University Hospital







A path to achieve education and professional growth that will propel you towards a greater level of competitiveness in the employment market"





tech 18 | Structure and Content

Module 1. Research and Development of Medicines

- 1.1. Development of New Drugs
 - 1.1.1. Introduction
 - 1.1.2. Development Phases of New Drugs
 - 1.1.3. Discovery Phase
 - 1.1.4. Preclinical Phase
 - 1.1.5. Clinical Phase
 - 1.1.6. Approval and Registration
- 1.2. Discovery of an Active Substance
 - 1.2.1. Pharmacology
 - 1.2.2. Seeding Trials
 - 1.2.3. Pharmacological Intervention
- 1.3. Pharmacokinetics
 - 1.3.1. Methods of Analysis
 - 1.3.2. Absorption
 - 1.3.3. Distribution
 - 1.3.4. Metabolism
 - 1.3.5. Excretion
- 1.4. Toxicology
 - 1.4.1. Single Dose Toxicity
 - 1.4.2. Repeated Dose Toxicity
 - 1.4.3. Toxicokinetics
 - 1.4.4. Carcinogenicity
 - 1.4.5. Genotoxicity
 - 1.4.6. Reproductive Toxicity
 - 1.4.7. Tolerance
 - 1.4.8. Dependency
- 1.5. Regulation of Drugs for Human Use
 - 1.5.1. Introduction
 - 1.5.2. Authorization Procedures
 - 1.5.3. How a Drug is Evaluated: Authorization Dossier
 - 1.5.4. Technical Data Sheet, Package Leaflet and EPAR
 - 1.5.5. Conclusions





Structure and Content | 19 tech

- 1.6. Pharmacovigilance
 - 1.6.1. Pharmacovigilance in Development
 - 1.6.2. Pharmacovigilance in Marketing Authorization
 - 1.6.3. Post-Authorization Pharmacovigilance
- 1.7. Uses in Special Situations
 - 1.7.1. Introduction
 - 1.7.2. Regulations BORRAR
 - 1.7.3. Examples:
- 1.8. From Authorization to Commercialization
 - 1.8.1. Introduction
 - 1.8.2. Drug Financing
 - 1.8.3. Therapeutic Positioning Reports
- 1.9. Special Forms of Regulation
 - 1.9.1. Advanced Therapies
 - 1.9.2. Accelerated Approval
 - 1.9.3. Biosimilars
 - 1.9.4. Conditional Approval
 - 1.9.5. Orphan Drugs
- 1.10. Dissemination of Research
 - 1.10.1. Scientific Article
 - 1.10.2. Types of Scientific Articles
 - 1.10.3. Quality of Research Checklist
 - 1.10.4. Drug Incapacitation Sources

Module 2. Biostatistics

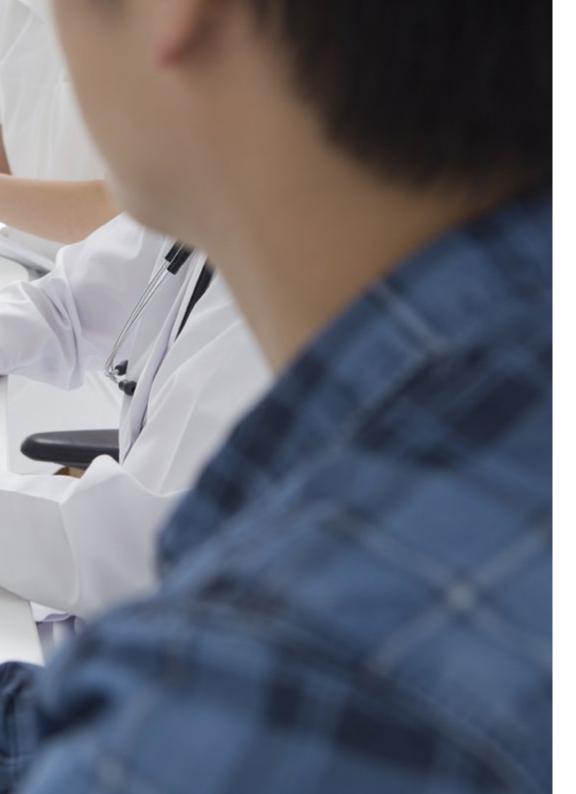
- 2.1. Study Design
 - 2.1.1. Research Question
 - 2.1.2. Population to Analyze
 - 2.1.3. Classification
 - 2.1.3.1. Comparison between Groups
 - 2.1.3.2. Maintenance of the Described Conditions
 - 2.1.3.3. Assignment to Treatment Group
 - 2.1.3.4. Degree of Masking
 - 2.1.3.5. Modality of Intervention
 - 2.1.3.6. Centers Involved

tech 20 | Structure and Content

2.2. Types of Randomized Clinical Trials: Validity	and Biases
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- 2.2.1. Types of Clinical Trials
 - 2.2.1.1. Superiority Study
 - 2.2.1.2. Equivalence or Bioequivalence Study
 - 2.2.1.3. Non-Inferiority Study
- 2.2.2. Analysis and Validity of Results
 - 2.2.2.1. Internal Validity
 - 2.2.2.2. External Validity
- 2.2.3. Biases
 - 2.2.3.1. Selection
 - 2.2.3.2. Measurement
 - 2.2.3.3. Confusion
- 2.3. Sample Size Protocol Deviations
 - 2.3.1. Parameters Used
 - 2.3.2. Protocol Justification
 - 2.3.3. Protocol Deviations
- 2.4. Methodology
 - 2.4.1. Missing Data Handling
 - 2.4.2. Statistical Methods
 - 2.4.2.1. Description of Data
 - 2.4.2.2. Survival
 - 2.4.2.3. Logistic Regression
 - 2.4.2.4. Mixed Models
 - 2.4.2.5. Sensitivity Analysis
 - 2.4.2.6. Multiplicity Analysis
- 2.5. When Does the Statistician Become Part of the Project
 - 2.5.1. Statistician Role
 - 2.5.2. Points of the Protocol to be Reviewed and Described by the Statistician
 - 2.5.2.1. Study Design
 - 2.5.2.2. The Primary and Secondary Objectives of the Study
 - 2.5.2.3. Sample Size Calculation
 - 2.5.2.4. Variables
 - 2.5.2.5. Statistical Justification
 - 2.5.2.6. Material and Methods used to Study the Objectives of the Study





Structure and Content | 21 tech

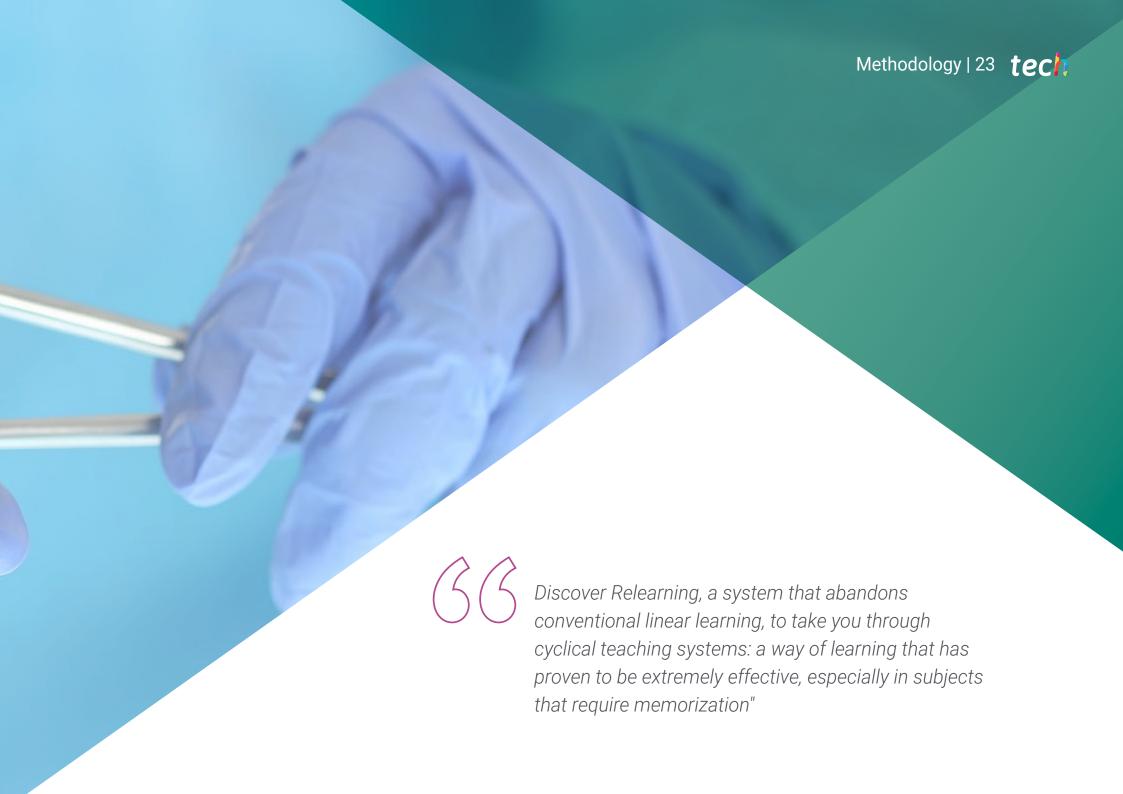
- 2.6. CRD Design
 - 2.6.1. Incapacitation Collection: Dictionary of Variables
 - 2.6.2. Variables and Data Entry
 - 2.6.3. Database Security, Testing and Debugging
- 2.7. Statistical Analysis Plan
 - 2.7.1. What is a Statistical Analysis Plan?
 - 2.7.2. When to Perform a Statistical Analysis Plan
 - 2.7.3. Statistical Analysis Plan Parts
- 2.8. Intermediate Analysis
 - 2.8.1. Reasons for an Early Stopping of a Clinical Trial
 - 2.8.2. Implications of Early Termination of a Clinical Trial
 - 2.8.3. Statistical Designs
- 2.9. Final Analysis
 - 2.9.1. Final Report Criteria
 - 2.9.2. Plan Deviations
 - 2.9.3. Guidelines for the Elaboration of the Final Report of a Clinical Trial
- 2.10. Statistical Review of a Protocol
 - 2.10.1. Checklist
 - 2.10.2. Frequent Errors in the Review of a Protocol





This academic program offers students a different way of learning. Our methodology uses a cyclical learning approach: **Relearning.**

This teaching system is used, for example, in the most prestigious medical schools in the world, and major publications such as the **New England Journal of Medicine** have considered it to be one of the most effective.



tech 24 | Methodology

At TECH Nursing School we use the Case Method

In a given situation, what should a professional do? Throughout the program, students will face multiple simulated clinical cases, based on real patients, in which they will have to do research, establish hypotheses, and ultimately resolve the situation. There is an abundance of scientific evidence on the effectiveness of the method. Nurses learn better, faster, and more sustainably over time.

With TECH, nurses can experience a learning methodology that is shaking the foundations of traditional universities around the world.



According to Dr. Gérvas, the clinical case is the annotated presentation of a patient, or group of patients, which becomes a "case", an example or model that illustrates some peculiar clinical component, either because of its teaching power or because of its uniqueness or rarity. It is essential that the case is based on current professional life, in an attempt to recreate the real conditions in professional nursing practice.



Did you know that this method was developed in 1912, at Harvard, for law students? The case method consisted of presenting students with real-life, complex situations for them to make decisions and justify their decisions on how to solve them. In 1924, Harvard adopted it as a standard teaching method"

The effectiveness of the method is justified by four fundamental achievements:

- Nurses who follow this method not only grasp concepts, but also develop their mental capacity, by evaluating real situations and applying their knowledge.
- 2. The learning process has a clear focus on practical skills that allow the nursing professional to better integrate knowledge acquisition into the hospital setting or primary care.
- 3. Ideas and concepts are understood more efficiently, given that the example situations are based on real-life.
- 4. Students like to feel that the effort they put into their studies is worthwhile. This then translates into a greater interest in learning and more time dedicated to working on the course.





Relearning Methodology

At TECH we enhance the case method with the best 100% online teaching methodology available: Relearning.

This university is the first in the world to combine case studies with a 100% online learning system based on repetition combining a minimum of 8 different elements in each lesson, which is a real revolution compared to the simple study and analysis of cases.

The nurse will learn through real cases and by solving complex situations in simulated learning environments.

These simulations are developed using state-of-the-art software to facilitate immersive learning.



Methodology | 27 tech

At the forefront of world teaching, the Relearning method has managed to improve the overall satisfaction levels of professionals who complete their studies, with respect to the quality indicators of the best online university (Columbia University).

With this methodology we have trained more than 175,000 nurses with unprecedented success in all specialities regardless of practical workload. Our pedagogical methodology is developed in a highly competitive environment, with a university student body with a strong socioeconomic profile and an average age of 43.5 years old.

Relearning will allow you to learn with less effort and better performance, involving you more in your specialization, developing a critical mindset, defending arguments, and contrasting opinions: a direct equation to success.

In our program, learning is not a linear process, but rather a spiral (learn, unlearn, forget, and re-learn). Therefore, we combine each of these elements concentrically.

The overall score obtained by TECH's learning system is 8.01, according to the highest international standards.

This program offers the best educational material, prepared with professionals in mind:



Study Material

All teaching material is produced by the specialists who teach the course, specifically for the course, so that the teaching content is really specific and precise.

These contents are then applied to the audiovisual format, to create the TECH online working method. All this, with the latest techniques that offer high quality pieces in each and every one of the materials that are made available to the student.



Nursing Techniques and Procedures on Video

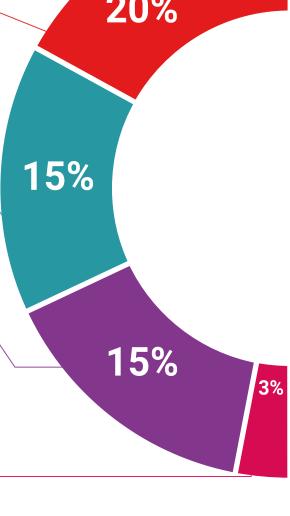
We introduce you to the latest techniques, to the latest educational advances, to the forefront of current medical techniques. All of this in direct contact with students and explained in detail so as to aid their assimilation and understanding. And best of all, you can watch them as many times as you want.



Interactive Summaries

The TECH team presents the contents attractively and dynamically in multimedia lessons that include audio, videos, images, diagrams, and concept maps in order to reinforce knowledge.

This exclusive educational system for presenting multimedia content was awarded by Microsoft as a "European Success Story".





Additional Reading

Recent articles, consensus documents and international guidelines, among others. In TECH's virtual library, students will have access to everything they need to complete their course.



Testing & Retesting

We periodically evaluate and re-evaluate students' knowledge throughout the program, through assessment and self-assessment activities and exercises, so that they can see how they are achieving their goals.



Classes

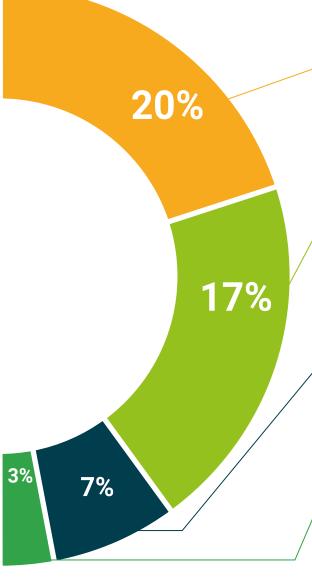
There is scientific evidence suggesting that observing third-party experts can be useful.

Learning from an Expert strengthens knowledge and memory, and generates confidence in future difficult decisions.

Quick Action Guides

TECH offers the most relevant contents of the course in the form of worksheets or quick action guides. A synthetic, practical, and effective way to help students progress in their learning.









tech 32 | Certificate

This program will allow you to obtain your **Postgraduate Certificate in Drug Research** and **Development in Nursing** endorsed by **TECH Global University**, the world's largest online university.

TECH Global University is an official European University publicly recognized by the Government of Andorra (*official bulletin*). Andorra is part of the European Higher Education Area (EHEA) since 2003. The EHEA is an initiative promoted by the European Union that aims to organize the international training framework and harmonize the higher education systems of the member countries of this space. The project promotes common values, the implementation of collaborative tools and strengthening its quality assurance mechanisms to enhance collaboration and mobility among students, researchers and academics.

This **TECH Global University** title is a European program of continuing education and professional updating that guarantees the acquisition of competencies in its area of knowledge, providing a high curricular value to the student who completes the program.

Title: Postgraduate Certificate in Drug Research and Development in Nursing

Modality: online

Duration: 12 weeks

Accreditation: 12 ECTS



Mr./Ms. ______, with identification document ______ has successfully passed and obtained the title of:

Postgraduate Certificate in Drug Research and Development in Nursing

This is a program of 360 hours of duration equivalent to 12 ECTS, with a start date of dd/mm/yyyy and an end date of dd/mm/yyyy.

TECH Global University is a university officially recognized by the Government of Andorra on the 31st of January of 2024, which belongs to the European Higher Education Area (EHEA).

In Andorra la Vella, on the 28th of February of 2024





Postgraduate Certificate Drug Research and Development in Nursing

- » Modality: online
- » Duration: 12 weeks
- » Certificate: TECH Global University
- » Credits: 12 ECTS
- » Schedule: at your own pace
- » Exams: online

