

**PNRR Missione 4, Componente 2, Investimento 1.4 “Potenziamento strutture di ricerca e creazione di "campioni nazionali di R&S" su alcune Key Enabling Technologies”**  
*Iniziativa finanziata dall'Unione europea — NextGenerationEU.*

**National Center for Gene Therapy and Drugs based on RNA Technology**  
**Sviluppo di terapia genica e farmaci con tecnologia a RNA**

Codice progetto MUR: **CN00000041** – CUP UNINA: **E63C22000940007**

**Doctorate of National Interest**  
**RNA THERAPEUTICS AND GENE THERAPY**

**TITLE OF THE RESEARCH PROJECT**

Multi-omics approaches based on MS techniques for drug target efficacy, safety and mechanism elucidation.

**SELECT ONE OF THE FOLLOWING RESEARCH AREA:**

- ☒ **Mechanisms of Diseases and Drug Target Identification**
- ☐ **Design and Delivery of New Gene Therapy and RNA-Based Medicines**
- ☐ **Validation and Safety In Preclinical and Clinical Studies**

**LOCATION OF THE RESEARCH ACTIVITY (INSTITUTION/DEPARTMENT):**

University of Salerno, Department of Pharmacy

**TUTOR:** Prof. Pietro Campiglia, Co-TUTOR Prof Eduardo Maria Sommella

**PROPOSED RESEARCH ACTIVITIES (max 300 words):**

Reduce issues during preclinical trials is crucial to evaluate the in vivo efficacy and toxicity of any drug candidate early in the drug discovery process. The lack of translational efficacy stems from inadequate knowledge of the drug's mechanism of action. The employment of multi-omics approaches, eg the combination of metabolomics, lipidomics and proteomics, early in drug development is particularly useful, as they can individuate metabolite, lipid or protein biomarkers or signatures for both efficacy and side effects. Additionally, these studies provide a comprehensive understanding of drug action and potential toxicity mechanisms. The objective of this project is to integrate multiple omics approaches



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using state-of-the-art hyphenated mass spectrometry techniques. The project will develop and apply novel integrated omics methods to cellular models, organoids, and biofluids and tissues from in vitro and ex vivo preclinical studies. This approach aims to elucidate the molecular effects of novel drugs and their formulations, uncover new molecular insights, facilitate target deconvolution, and highlight key pathways to better understand pathophysiology and drug mechanisms.