PERSONAL INFORMATION

Angela Zampella



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Sex Female | Date of birth 13/09/1967 | Nationality Italian

Enterprise	University	EPR
☐ Management Level	□ Full professor	☐ Research Director and 1st level Technologist /
		First Researcher and 2nd level Technologist
☐ Mid-Management Level	☐ Associate Professor	☐ Level III Researcher and Technologist
☐ Employee / worker level	☐ Researcher and Technologist of IV, V, VI and VII	☐ Researcher and Technologist of IV, V, VI and VII
	level / Technical collaborator	level / Technical collaborator

WORK EXPERIENCE

Since December 2018 Head of the Department

Department of Pharmacy, University of Naples "Federico II"

Since October 2020

Board Member of Ceinge -Biotecnologie Avanzate operating in the field of molecular biology and advanced biotechnology applied to Human Health, s an excellence in Italy and abroad for the Research and Diagnostics of genetic diseases (hereditary and acquired).

December 2016- now

Full Professor of Organic Chemistry

Department of Pharmacy, University of Naples "Federico II"

2013- now

Member of the board for PhD programme in Pharmaceutical Science Department of Pharmacy, University of Naples "Federico II"

Associate Professor of Organic Chemistry

November 2002-December 2016

Department of Pharmacy, University of Naples "Federico II"

September 1998- Ottobre 2002

Researcher of Organic Chemistry

Department of Pharmacy, University of Naples "Federico II"

EDUCATION AND TRAINING

1995 PhD in "Sostanze Naturali Farmacologicamente attive"

Department of Pharmacy University of Naples "Federico II"

May-October 1994

Visiting Researcher

Institut de Chimie des Substances Naturelles- CNRS- Gif sur Yvette (France)

PERSONAL SKILLS

Mother tongue

Italian

Other language

English and French (advanced)

Organisational skills and competences

Head of the Department of Pharmacy, University of Naples, Federico II, recognized as Department of Excellence (MUR 2018-2022 and 2023-2027). Angela Zampella is leader of a large research group including permanent staff, PhD and post-docs, working on the isolation from natural sources, design, synthesis and pharmacological profiling of new drugs in metabolic disorders, with the research activity results reported in 7 family patents (see below), one of which licensed worldwide. Notably, one of the discovered molecules has recently entered in Phase I clinical trial (ClinicalTrials.gov Identifier: NCT05203367).

WORK ACTIVITIES

Awards

Selection of the contribution "BAR502, a dual FXR and GPBAR1 agonist, reverses steatosis and fibrosis in rodent model of NASH by modulating autophagic genes", in the International Conferences DDW (Digestive Disease Week) San Diego, May 21-24, 2016.

Editorial activity

Referee for J. Med. Chem, Steroids, Marine Drugs, Org. Lett., Bioorg. Med. Chem., Scientific Reports, Nature Communication, PLos One, Tetrahedron, J. Org. Chem, Journal of Immunology

Grants

- Leader for a Pharma collaborating project (2015 to date) in the development of new drugs in liver fibrosis (about 500.000 €).

- -PI and Coodinator for Federico II of "National Center for Gene Therapy and Drugs based on RNA Technology" PNRR MUR M4C2 Project number: E63C22000940007 (27 M€)
- -Unit coordinator (Federico II) for "Infrastructure for precision medicine in oncology (PREMIO)", [26/11/2018 Current] (2.7 M€).
- -Department of Excellence "Project TRAVEL" 2023-2027 (9,2 M€ MUR)
- -Unit Coordinator (Federico II) of Regione Campania project "Fighting Cancer resistance: Multidisciplinary, integrated Platform for a technological Innovative Approach to Oncotherapies (Campania Oncotherapies)", Project N. B61G18000470007 [01/01/2018 Current] (660.000 €);
- -Department of Excellence "Project Health" 2018-2022 (9,3 M€ MUR + 4 M€ cofound)
- -Head of RU for the project PRIN2017: Bile acids activated receptors and liver metabolism: discovery and development of novel therapeutic targets in the treatment of steato-hepatitis (NASH). Funded by MUR.

Patents

- Sterols in the treatment and/or prevention of sars-cov-2 infection. PCT International patent application (PCT/IB2021/054142, filed 14.05.2021)
- 6-Substituted, 22-cyano hyodeoxycholanic analogues and uses thereof. PCT International patent application (PCT/IB2020/061695, filed 09.12.2020)
- Isoxazoles as FXR receptor agonists and their preparation. PCT International patent application PCT/IB2019/056114, filed 17.07.2019
- Preparation of oxadiazoles as FXR receptor antagonists. PCT International patent application PCT/IB2019/054238, filed 22.05.2019, licensed as IT201800005598A1 (22.11.2019)
- Synthesis of hyodeoxycholic acid derivatives for pharmaceutical use. PCT International patent application PCT/IB2017/053959, filed 30.06.2017), licensed as IT201600068742A1 (01.01.2018)
- Derivati chimici del colano per l'uso nel trattamento e/o nella prevenzione delle malattie mediate dai recettori FXR e TGR5/GP-BAR1. Patent application ITUA20161663A, filed 15.03.2016, licensed as ITUA20161663A1 (15.09.2017)
- Cholane derivatives for use in the treatment and/or prevention of FXR and TGR5/GPBAR1 mediated diseases, PCT International patent application (PCT/EP2015/061802, filed 28.05.2015), licensed as AU2015265893B2; CA2948585A1; CN106661079B; CN110003301A; DK3149019T3; EA032820B1; EA201692316A1; EP3149019A1; EP3149019B1; EP3626725A1; ES2768718T3; HRP20200225T1; HUE048351T2; JP2017516856A; JP6820253B2; KR20170008767A; LT3149019T; MA39881B1; MX2016015724A; PH12016502327A1; PL3149019T3; PT3149019T; RS59910B1; SG10201809362RA; SG11201609403UA; SI3149019T1; US10407462B2; US11117926B2; US2017190731A1; US2019352328A1; WO2015181275A8

ADDITIONAL INFORMATION

Publications

Author and co-author of 165 scientific products; From Scopus: H-index: 41; total citations: 5008.

The most relevant publications

- Zampella A. et al. Combinatorial targeting of G-protein-coupled bile acid receptor 1 and cysteinyl leukotriene receptor 1 reveals a mechanistic role for bile acids and leukotrienes in drug-induced liver injury. Hepatology, 2022
- Zampella A. et al. Atorvastatin protects against liver and vascular damage in a model of diet induced steatohepatitis by resetting FXR and GPBAR1 signaling. 2022, 36(1), e22060
- Zampella A. et al. Structural Basis for Developing Multitarget Compounds Acting on Cysteinyl Leukotriene Receptor 1 and G-Protein-Coupled Bile Acid Receptor 1. Journal of Medicinal Chemistry, 2021, 64(22), pp. 16512-16529
- Zampella A. et al. Bile acids and their receptors in metabolic disorders. Progress in Lipid Research 2021, 82,101094
- Zampella A. et al Identification of cysteinyl-leukotriene-receptor 1 antagonists as ligands for the bile acid receptor GPBAR1. Biochemical Pharmacology 2020, 177, 113987
- Zampella A. et al Bile acids activated receptors regulate innate immunity. Frontiers in Immunology 2018,1853
- Zampella A. et al. The bile acid receptor GPBAR1 regulates the M1/M2 phenotype of intestinal macrophages and activation of gpbar1 rescues mice from murine colitis. Journal of Immunology 2017, 199(2), pp. 718-733

Naples March 20, 2023

Augda Lampella