

Curriculum Vitae

Accademic Title
MD, PhD, MSc

Current Position and Discipline

Consultant in Pediatrics

Medical Licence / Registration Number and issue date

Register of Physicians and Surgeons of Foggia (registration number: 5742, 18 Sept 2003)

Current Employer

Institution	Azienda Ospedaliero Universitaria Policlinico di Bari
Department	Department of Pediatrics, Unit of Metabolic and Genetic Diseases
Address	Ospedale Pediatrico "Giovanni XXIII", Via Amendola, 207 - Bari

Education *(all relevant major medical qualifications)*

Year	Qualification	Academic Institution
2011	PhD in "Pediatric Sciences"	Università degli Studi di Bari
2010	Certificate Advanced Course in "Infectivity and Pediatric Rheumatology"	Università degli Studi di Firenze
2010	Certificate Advanced Course in "Rare Diseases: Assistance, Training and Research Methodologies"	Università degli Studi di Firenze
2010	Certificate Diploma in "Specialized Pediatrics, Address in Pediatric Endocrinology"	Università degli Studi di Bari
2008	MSc in "Clinical Pediatrics"	University College of London
2007	Specialization in Pediatrics "General and Specialist Pediatrics"	Università degli Studi di Bari
2002	Degree in Medicine and Surgery	Università degli Studi di Foggia

Medical Career History

Dates	Position	Institution and Department
2017-ongoing	Professional Assignment in Neonatal screening and Inherited Metabolic Disorders	Ospedale Pediatrico "Giovanni XXIII", Department of Pediatrics, Unit of Metabolic and Genetic Diseases
2015-ongoing	Senior Pediatrician	Ospedale Pediatrico "Giovanni XXIII", Department of Pediatrics, Unit of Metabolic and Genetic Diseases
2014-2015	Senior Pediatrician	Unit of Inherited Metabolic Diseases, Azienda

		Ospedaliera di Padova
2012-2014	Senior Pediatrician	Ospedale Pediatrico “Giovanni XXIII”, Department of Pediatrics, Unit of Metabolic and Genetic Diseases
2011	Senior Pediatrician	Pediatria, Presidio Ospedaliero “Perrino” dell’ASL di Brindisi

Research experience

Dates	Position	Institution and Department
2018 – ongoing	Principal Investigator	CCD-LMZYMAA1-08 A 24-month Multicenter, Open-label Phase II Trial Investigating the Safety and Efficacy of Repeated velmanase alfa (recombinant human alpha-mannosidase) Treatment in Pediatric Patients below 6 years of age with Alpha-Mannosidosis
2016 – 2019	Principal Investigator	“Long Term Safety Study of Orfadin Treatment in Tyrosinemia type1 Patients in Standard Clinical Care” (OPAL)
2019 – ongoing	Principal Investigator	LOTUS (Long Term Carbaglu® Use in PA and MMA patients) Multicentric Observational Prospective Study on the Long Term Use of Carbaglu® for the Treatment of Propionic and Methylmalonic Acidurias
2019 - ongoing	Principal Investigator	An Observational Disease and Clinical Outcomes Registry of Patients with Lysosomal Acid Lipase (LAL) Deficiency ALX-LALD-501.
2019 - ongoing	Scientific Representative	European network and registry for Homocystinurias and Methylation Defects (E-HOD)

Other scientific experiences

Dates	Institution and Department
March 2018	Invited Advisor to the Pegvaliase for the Treatment of Adult PKU Advisory Board organized by BiMarin Pharmaceutical Inc
Sept 2016 – Jul 2017	Participant to the Nutricia survey ATTITUDE (Analysis of the most relevant and predictive factors influencing adherence to PKU diet)
Jul 2016	Invited Advisor to PKU Advisory Board meeting organized by BiMarin Pharmaceutical Inc
May 2018- March 2019	Invited Advisor to the 1st and 2nd Italian Advisory Board organized By Applied Pharma Research (APR) on Fenyketonuria Dietetic Management
June 2018	Member of the faculty of Orphan Europe meetings entitled: International Young Metabolicians and Central Eastern Europe (CEE) Metabolic Disorders Experts
June 2018	Invited expert for the BioMarin PKU Global Expert Forum

June – December 2019	Tutor for the MPS Multiconnection
June 2019	Phenylketonuria Expert Educator (PEER) meeting
September 2019	Invited Expert for the Meeting “La gestione del paziente con Fenilchetonuria dalla fase pediatrica a quella adulta”
December 2019	Invited Advisor to the ADVISORY BOARD on the use of glicomacropptide in Phenylketonuria organized by Quaris S.r.l.
January 2020	Clinical Representative of the Study Group “Microbiota e Malattie Metaboliche Ereditarie”
April – December 2020	Tutor for the MPS Multiconnection
Sept 2019 – Dec 2020	Member of the Expert Group of the “Italian National Consensus Statement on Pharmacological Treatment of PKU”

Teaching activities

Dates	Institution and Department
June 2019	University Master Hereditary metabolic diseases: social and nutritional problems in pediatric age

Last GCP Certificate date: May 2019

Bari, 20 March 2024