

PERSONAL INFORMATION

Andrea MAZZANTI



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Sex M | Date of birth [REDACTED] Nationality Italian

Enterprise	University	EPR
<input type="checkbox"/> Management Level	<input type="checkbox"/> Full professor	<input type="checkbox"/> Research Director and 1st level Technologist / First Researcher and 2nd level Technologist
<input type="checkbox"/> Mid-Management Level	<input type="checkbox"/> Associate Professor	<input type="checkbox"/> Level III Researcher and Technologist
<input type="checkbox"/> Employee / worker level	<input checked="" type="checkbox"/> Researcher and Technologist of IV, V, VI and VII level / Technical collaborator	<input type="checkbox"/> Researcher and Technologist of IV, V, VI and VII level / Technical collaborator

WORK EXPERIENCE

- 12/2018 – Current **Research assistant professor**
 University of Pavia - Department of Molecular Medicine, Pavia, Italy (www.unipv.it)
 Clinical and translational research in cardiology (inherited arrhythmias and cardiomyopathies); mentoring of undergraduate and Ph.D. students; lecturing different undergraduate and graduate courses.
 Business or sector: Cardiology
- 2017 – Current **Co-coordinator of the European Registry on Short QT Syndrome**
 ERN GUARD-HEART: Gateway to Uncommon and Rare Diseases of the Heart
<https://guardheart.ern-net.eu/>
 Management of the patients' registry in an EU funded project for the study and treatment of rare inherited arrhythmogenic conditions.
 Business or sector: Cardiology
- 01/2017 – Current **Visiting scientist**
 Centro Nacional de Investigaciones Cardiovasculares Carlos III, Madrid, Spain (www.cnics.es)
 Monthly visits (4-5 days) to conduct research in the field of gene therapy and experimental electrophysiology.
 Business or sector: Cardiology – Translational Research
- 07/2012 – Current **Clinical cardiologist**
 IRCCS ICS Maugeri, Pavia, Italy (www.icsmaugeri.it)
 Outpatient and ward clinic specialized in inherited arrhythmogenic diseases and cardiomyopathies; Cardiovascular imaging; Clinical and translational research in cardiology.
 Business or sector: Cardiology
- 02/2011 – 06/2012 **Clinical cardiologist and Researcher**
 Health in Code – La Coruña, Spain (www.healthincode.com)
 Outpatient clinic for patients with inherited arrhythmogenic diseases and cardiomyopathies; Development of new screening tools for genetic conditions of the heart; Business development in a highly competitive academic spin-off company.
 Business or sector: Cardiology – Genetic Heart Diseases
- 08/2010 – 01/2011 **Clinical cardiologist**
 The Salam Centre for Cardiac Surgery – NGO Emergency, Khartoum, Sudan (www.emergency.it)
 Head of the cardiology ward; Outpatient clinic and cardiac intensive care unit; Intraoperative transthoracic and transoesophageal echocardiography
 Business or sector: Cardiology – Cardiac Surgery – Emergency Medicine

EDUCATION AND TRAINING

- 2019 – Current **National Scientific Qualification (Professor "Il Fascia") – Settore 06/D1**
 Italian Ministry of Education, University and Research (www.miur.gov.it)
 Field: Cardiovascular Diseases
- 10/2016 – 12/2019 **Doctor of Philosophy in Biomedical Sciences (XXXII cycle)** EQF 8
 University of Pavia, Pavia, Italy (<https://web.unipv.it/>)
 Thesis: Development and Characterization of the First Large Mammal Model of Long QT Syndrome

2015 – 2016	Master in Medical and Genomic Statistics University of Pavia, Pavia, Italy (https://web.unipv.it/) Field(s) of study: Life sciences, mathematics and statistics: Statistics	EQF 8
06/2006 – 06/2010	Specialist in Cardiology (final grade 70/70) University of Turin, Turin, Italy (https://en.unito.it/) Thesis: "Long-Term Follow-Up of Patients with Short QT Syndrome" (<i>J Am Coll Cardiol.</i> 2011; 58:587-95)	EQF 7
10/1998 – 03/2005	Medical Doctor (final grade 110/110 magna cum laude) University of Turin, Turin, Italy (https://en.unito.it/)	EQF 6
1993– 1998	High School Diploma (final grade 60/60) Liceo Classico Vittorio Alfieri, Turin, Italy (http://liceoalfieri.it/)	EQF 4

WORK ACTIVITIES

Honours and Awards	<ul style="list-style-type: none"> -2020: Young Scientific Board – Società Italiana di Medicina e Chirurgia Rigenerativa Polispecialistica Responsabile prof. E. Caradonna – Campobasso, Italy - 2020: Member of PhD Commission – University of La Coruña, Spain - 2014: Member of PhD Commission – University of Murcia, Spain - 2006: Research scholarship on heart failure (A.O.U. Città della Salute e della Scienza di Torino, Turin, Italy)
Editorial Activity	Member of the Editorial Board: Journal of Child and Adolescent Health Member of the Editorial Board: Journal of Cardiovascular Electrophysiology Reviewer: Circulation, Journal of the American College of Cardiology, Circulation Research, Nature Reviews Cardiology
Invited Presentation	More than 50 invitations in the last five years by the leading cardiological societies (ESC, EHRA, HRS, AHA)
Grants	2014 – 2018: Gilead Research Grant Group Member: "Long QT Syndrome type 3: epidemiological profile, risk stratification and personalized management" 2016: Gilead Research Grant Group Member: "Role of Calmodulin kinase II inhibitors in Catecholaminergic Polymorphic Ventricular Tachycardia in knock-in animal models"
Patents	Compositions and methods for the treatment of dominantly inherited catecholaminergic polymorphic ventricular tachycardia. Publication number: WO/2019/193563 Predicting patient response to sodium channel blockers Application number: 16708312

ADDITIONAL INFORMATION

Research Output Summary	<ul style="list-style-type: none"> • Last 10 years: 64 indexed papers, of which 21 as the first author. Total impact factor (IF) 757. Average IF/paper 11.8. Total citations 4,124 (Scopus); H-index 22 (Scopus).
Publications (5 Selected)	<ul style="list-style-type: none"> • Mazzanti, A., Kanthan, A., Monteforte, et al. Novel insight into the natural history of short QT syndrome (2014) <i>Journal of the American College of Cardiology</i>, 63, pp. 1300-1308. Cited 129 times. PMID: 24291113. IF 16.5 • Mazzanti, A., Maragna, R., Faragli, et al. Gene-specific therapy with mexiletine reduces arrhythmic events in patients with long QT syndrome type 3 (2016) <i>Journal of the American College of Cardiology</i>, 67, pp. 1053-1058. Cited 105 times. PMID: 26940925. IF 19.9 • Mazzanti, A., Ng, K., Faragli, A., et al. Arrhythmogenic Right Ventricular Cardiomyopathy: Clinical Course and Predictors of Arrhythmic Risk (2016) <i>Journal of the American College of Cardiology</i>, 68, pp. 2540-2550. Cited 91 times. PMID: 27931611. IF 19.9 • Mazzanti, A., Maragna, R., Vacanti, et al. Interplay Between Genetic Substrate, QTc Duration, and Arrhythmia Risk in Patients with Long QT Syndrome (2018) <i>Journal of the American College of Cardiology</i>, 71, pp. 1663-1671. Cited 66 times. PMID: 29650123. IF 18.6 • Bongianino R, Denegri M, Mazzanti A, et al. Allele-Specific Silencing of Mutant mRNA Rescues Ultrastructural and Arrhythmic Phenotype in Mice Carriers of the R4496C Mutation in the Ryanodine Receptor Gene (RYR2) (2018) <i>Circulation Research</i>, 121, page 525-536. Cited 31 times. PMID: 28620067. IF 15.9

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