

Chiara Fiorillo, MD, PhD

born in Naples 11/09/1975

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Education and Training

- 1994-1999 School of Medicine and Surgery, University of Naples, Federico II.
- 1999-2004 Resident Registrar in Neurology at University of Naples, Federico II.
- 2004-2008 PhD in Neuroscience
- 2006-2007 Research Visiting Fellow at Institute of Human Genetic, New Castle University, UK.
- 2008-2009 Research Visiting Fellow (Honorary Contract) at Neuropathology Lab, RJAH Hospital, Oswestry, UK

Current position

From 2018 RTDA University of Genoa, Department of Neuroscience (DINOEMI) and Medical Consultant at Paediatric Neurology, IRCCS Gaslini Children Hospital, Genoa, Italy

Previous employments

- 2014-2018 Clinical researcher at University of Genoa, Department of Neuroscience (DINOEMI) and Gaslini Children Hospital, Genoa, Italy.
- 2010-2014 Clinical Researcher at Unit of Molecular Medicine and Neuromuscular Disorders, IRCCS Stella Maris. Pisa
- 2009-2010 Research fellowship from Telethon Foundation as curator of DMD and BMD registry within Treat NMD project.

Teaching

Seminars and tutoring small group at the School of Medicine and Surgery, Course of Paediatrics, from year 2015

Course of Paediatric Neurology, CL in Scienze Infermieristiche from year 2019

Tutor of PhD and graduate students.

Fundings and Grants as PI (past 5 yrs)

PTC Therapeutics International Ltd., Private Grants: Next generation sequencing for detection of point mutations in DMD

AFM French Telethon 2018 #22341 Defining molecular and functional consequences of titin mutations in human muscle progenitors from affected patients.

Bando AIFA 2017 Ricerca Indipendente. TRS-2018-00001625 TREAT-LMNA- "Deflazacort TREATment in LMNA related congenital muscular dystrophy: study of clinical effectiveness and search for reliable markers

Italian Ministry of Health. Ricerca Finalizzata 2019. Progetto di rete: New models for patient management in the Emergency Department: looking for improvement of efficiency and quality

Telethon UILDM 2019 Clinical Project GUP19002C Trial readiness and endpoint assessment in congenital and childhood Myotonic Dystrophy

Telethon UILDM 2021 Clinical Project GUP21006 Characterization of the phenotypic diversity in DupEx2 Duchenne Muscular Dystrophy and identification of predictive/prognostic markers.

Publication/Editorial activity

Author of 106 articles in International Peer-reviewed Journals

Member of editorial staff for the following journals: Acta Myologica and Frontiers in Paediatric Neurology

Reviewer for the following journals: Neuroscience Letters, European Journal of Paediatric Neurology, BMC Medical Genetics, Acta Neuropathologica, Neurological Science, Muscle and Nerve, DNA and Cell Biology, Human Mutation.

*Indici citazioni Tutte Dal 2017

Citazioni 1635 1277

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**fonte Google Scholar*

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