

## Curriculum Vitae

### Personal Information

**ARIANNA MANINI**

Address

Telephone number

E-mail

Nationality

Date of birth

Sex

### Position

Doctor of Medicine (M.D.)

### Professional experience

November, 2019 - Present

**Residency School in Neurology at the Department of Neurological Sciences, University of Milan, Italy**

Department of Biomedical and Clinical Sciences "Luigi Sacco", University of Milan, Milan, Italy

Currently attending the second year

February, 2019

**Habilitation to the profession of Medical Doctor – State Exam**

University of Milan, Milan, Italy

February, 2019

**Member of the Medicine Board of Milan – "Ordine dei Medici, Chirurghi ed Odontoiatri di Milano" (n.46239)**

March, 2017 – October, 2019

**Clinical Neurology Internship**

Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico, Department of Pathophysiology and Transplantation (DEPT), Dino Ferrari Centre, Neuroscience Section, University of Milan, Milan, Italy

Clinical activity in Neurology Inpatient and Outpatient Clinic, focused on Movement Disorders and Neurodegenerative Diseases. Actively taking part in clinical rounds, patient diagnosis, treatment and management, clinical cases discussion, Magnetic Resonance Imaging interpretation and practical skills.

March, 2017 – October, 2019

**Research Internship**

Laboratory of Biochemistry and Genetics, Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico, University of Milan, Milan, Italy

Active participation to basic research projects, focused on genetics of Movement Disorders and other Neurodegenerative Diseases.

January, 2017 – February, 2017

**Clinical Infectious Diseases Internship**

III Infectious Diseases Unit, Department of Biomedical and Clinical Sciences "Luigi Sacco", University of Milan, Milan, Italy

October, 2016 – December, 2016

**Clinical Oncology Internship**

Division of early Drug Development for Innovative Therapy, European Institute of Oncology (IEO), IRCCS, Milan, Italy

July, 2016 – August, 2016

**Medical Elective Project - "International Volunteer HQ"**

Pokhara – Nepal

April, 2016 – June, 2016	<b>Clinical Oncology Internship</b> Department of Medical Oncology, Fondazione IRCCS Istituto Nazionale dei Tumori, University of Milan, Milan, Italy
<b>Education and training</b>	
July, 2018	<b>Master's Degree in Medicine and Surgery</b> <ul style="list-style-type: none"> <li>• University of Milan, Milan, Italy</li> <li>• Final score: 110/110, <i>magna cum laude and honorable mention</i></li> <li>• Title of dissertation: "The role of <i>LRP10</i> mutations in Parkinson's Disease and Dementia with Lewy Bodies"</li> </ul>
July, 2012	<b>Scientific High School Graduation</b> <ul style="list-style-type: none"> <li>• Liceo Scientifico Statale Elio Vittorini, Milan, Italy</li> <li>• Final score: 100/100</li> </ul>
<b>Work experience</b>	
July, 2018 – Present	<b>Family Doctor Stand-in</b> Work experience as Family Doctor Stand-in in several Family Medicine Clinics in Milan
March, 2017 – Present	<b>Research Activity</b> Laboratory of Biochemistry and Genetics, Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico, University of Milan, Milan, Italy  Active participation to basic research projects, focused on genetics of Movement Disorders ( <i>Parkinson's disease, Dementia with Lewy Bodies, Spinocerebellar Ataxias</i> ) and other Neurodegenerative Diseases. Experience of laboratory activity in the field of molecular genetics (Next Generation Sequencing; Sanger Sequencing; DNA extraction; RT-PCR). Currently working on the following basic research projects: <ul style="list-style-type: none"> <li>- Genetic screening of <i>LRP10</i> mutations in a cohort of patients affected by Parkinson's disease and Dementia with Lewy Bodies.</li> <li>- Whole Exome Sequencing and Next Generation Sequencing panel approach of patients affected by Neurodegenerative and Neuromuscular Diseases.</li> </ul>
March, 2017 – Present	<b>Clinical Research Activity</b> Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico, Department of Pathophysiology and Transplantation (DEPT), Dino Ferrari Centre, Neuroscience Section, University of Milan, Milan, Italy  Stroke and Dementia Lab, Department of Biomedical and Clinical Sciences "Luigi Sacco", University of Milan, Milan, Italy  Participation to clinical trials, involvement in clinical evaluation of patients, patient's data collection and analysis, creation of databases ( <i>ischemic and hemorrhagic stroke, dementia, Parkinson's disease</i> ), literature review. Currently working on the following clinical projects: <ul style="list-style-type: none"> <li>- Neuropsychological evaluation in the acute phase and during follow-up of patients with ischemic and hemorrhagic stroke.</li> <li>- Detection of atrial fibrillation using implantable loop recorders in patients with cryptogenic ischemic stroke.</li> <li>- Phenotypical and genetic characterization of patients with known and novel <i>NOTCH3</i> (CADASIL) mutations.</li> </ul>
<b>Honours and awards</b>	
January 23 <sup>th</sup> -25 <sup>th</sup> , 2019	<b>Travel grant for the 7<sup>th</sup> Winter Seminar on Dementia and Neurodegenerative Disorders of the SiNdem4Juniors, Bressanone, Italy</b>

May 23<sup>rd</sup>-26<sup>th</sup>, 2020 **Runner-up presentation in Basic Neurology at the European Academy of Neurology (EAN) Tournament for Neurologists in Training (virtual)**

Title: "The role of *LRP10* mutations in Parkinson's Disease and Dementia with Lewy Bodies"

Authors: A. Manini, L. Straniero, E. Monfrini, M. Vizziello, G. Franco, S. Duga, A. Di Fonzo

November 28<sup>th</sup>-30<sup>th</sup>, 2020 **Grant for the 51<sup>th</sup> National Congress of the Italian Neurological Society (SIN) (virtual)**

June 19<sup>th</sup>-22<sup>th</sup>, 2021 **Travel grant for the 7<sup>th</sup> Congress of the European Academy of Neurology (EAN) (virtual)**

## Conferences

January 25<sup>th</sup>, 2019 **Oral presentation at the 7<sup>th</sup> Winter Seminar on Dementia and Neurodegenerative Disorders of SIndem4Juniors, Bressanone, Italy**

Title: "The role of *LRP10* mutations in Parkinson's disease and Dementia with Lewy Bodies"

Authors: A. Manini, E. Monfrini, M. Vizziello, F. Arienti, G. Lazzeri, G. Franco, A. Di Fonzo

October 13<sup>th</sup>, 2019 **Oral presentation at the 50<sup>th</sup> Congress of the Italian Neurological Society (SIN), Bologna, Italy**

Title: "Identification of two novel *MFN2* mutations in CMT2A patients"

Authors: E. Abati, A. Manini, R. Del Bo, S. Salani, F. Rizzo, G. Citterio, P. Mandich, M.T. Bassi, N. Bresolin, G.P. Comi, S. Corti

May 23<sup>rd</sup>, 2020 **Oral presentation at the 6<sup>th</sup> Congress of the European Academy of Neurology (EAN) (virtual)**

Title: "The role of *LRP10* mutations in Parkinson's disease and Dementia with Lewy Bodies"

Authors: A. Manini, L. Straniero, E. Monfrini, M. Vizziello, G. Franco, S. Duga, A. Di Fonzo

November 29<sup>th</sup>, 2020 **Oral presentation at the 51<sup>th</sup> National Congress of the Italian Society of Neurology (SIN) (virtual)**

Title: "The impact of lockdown during SARS-CoV-2 outbreak on behavioral and psychological symptoms of dementia"

Authors: A. Manini, M. Brambilla, L. Maggiore, S. Pomati, L. Pantoni

## Poster presentations

November 15<sup>th</sup>, 2018 **Selected poster presentation at the Fresco International Workshop on synaptic plasticity and advances in Parkinson's Disease, Florence, Italy**

Title: "The role of *LRP10* mutations in Parkinson's disease and Dementia with Lewy Bodies"

Authors: A. Manini, E. Monfrini, M. Vizziello, F. Arienti, G. Lazzeri, G. Franco, A. Di Fonzo

September 23<sup>th</sup>, 2019 **Selected poster presentation at the International Congress of Parkinson's Disease and Movement Disorders, Nice, France**

Title: "The role of *LRP10* mutations in Parkinson's disease and Dementia with Lewy Bodies"

Authors: A. Manini, E. Monfrini, M. Vizziello, F. Arienti, G. Lazzeri, G. Franco, A. Di Fonzo

September 23<sup>th</sup>, 2019 **Selected poster presentation at the International Congress of Parkinson's Disease and Movement Disorders, Nice, France**

Title: "A novel *TGM6* heterozygous mutation in a patient with cerebellar ataxia"

Authors: A. Manini, T. Bocci, E. Monfrini, M. Vizziello, G. Franco, A. Di Fonzo, G.P. Comi

September 24<sup>th</sup>, 2019

**Selected poster presentation at the International Congress of Parkinson's Disease and Movement Disorders, Nice, France**

Title: "Study of the haplotypic context as a modulator of the expressivity of *GBA* gene mutations in Parkinson disease and Gaucher disease"

Authors: M. Vizziello, E. Monfrini, I. Trezzi, G. Franco, M.D. Cappellini, E. Cassinerio, P. Tocco, F. Carubbi, F. Nascimbeni, E.M. Valente, S. Petrucci, F. Arienti, G. Lazzeri, **A. Manini**, G. Bitetto, A. Di Fonzo

May 25<sup>th</sup>, 2020

**Selected poster presentation at the 6<sup>th</sup> Congress of the European Academy of Neurology (EAN) (virtual)**

Title: "A rare p.R342W *TGM6* (SCA35) mutation in a patient with late-onset cerebellar ataxia"

Authors: **A. Manini**, T. Bocci, E. Monfrini, M. Vizziello, G. Franco, A. Di Fonzo, G.P. Comi

June, 2021

**Selected ePoster presentation at the 7<sup>th</sup> Congress of the European Academy of Neurology (EAN) (virtual)**

Title: "Implantable loop recorder to detect Atrial Fibrillation in Cryptogenic Stroke: a real-world experience"

Authors: **A. Manini**, G. Scopelliti, F. Mele, I. Cova, P. Bertora, S. Rosa, M. Schiavone, M. Viecca, G. Forleo, L. Pantoni

#### **Publications on Peer-Reviewed Journals**

**A. Manini**, T. Bocci, A. Migazzi, E. Monfrini, D. Ronchi, G. Franco, A. De Rosa, F. Sartucci, A. Priori, N. Bresolin, G.P. Comi, S. Corti, M. Basso, A. Di Fonzo, "A case report of late-onset cerebellar ataxia associated with a rare p.R342W *TGM6* (SCA35) mutation", *BMC Neurology*, 2020 Nov 7, 20(1):408, doi: 10.1186/s12883-020-01964-1, PMID: 33160304, PMCID: PMC7648302

**A. Manini**, M. Brambilla, L. Maggiore, S. Pomati, L. Pantoni, "The impact of lockdown during SARS-CoV-2 outbreak on behavioral and psychological symptoms of dementia", *Neurological Sciences*, 2021 Mar, 42(3):825-833, doi: 10.1007/s10072-020-05035-8, PMID: 33442845, PMCID: PMC7806279

**A. Manini**, L. Pantoni, "CADASIL from bench to bedside: disease models and novel therapeutic approaches", *Molecular Neurobiology*, 2021 Jan 19, doi: 10.1007/s12035-021-02282-4, PMID: 33464533

#### **Membership of Scientific Societies**

European Academy of Neurology (EAN)

RRFS member of the EAN Scientific Neurogenetics Panel

Società Italiana di Neurologia (SIN)

#### **Professional skills**

Mother tongue(s)

**Italian**

Other languages

**English**

Comprehension		Speaking		Writing
Listening	Reading	Spoken interaction	Spoken production	

C1	C2	C1	C1	C2
SLAM (Servizio Linguistico dell'Ateneo di Milano) – CEFR level: C1, awarded in date 18/09/2017				

Levels: A1 and A2: Basic user – B1 and B2: Independent user – C1 and C2: Proficient user.  
Common European Framework of Reference for Languages – Self-assessment grid.

## Research fields of interest

Genetics of movement disorders (i.e., Parkinson's disease, Dementia with Lewy Bodies, Spinocerebellar Ataxias).  
 Genetics of small vessel diseases (i.e., CADASIL, Fabry Disease).  
 Genetics of neurodegenerative and neuromuscular diseases (i.e., Amyotrophic lateral sclerosis, Congenital myopathies, Congenital neuropathies, Mitochondriopathies).  
 Dementia and vascular cognitive impairment: epidemiology, clinical aspects, diagnosis, neuroimaging.  
 Ischemic and hemorrhagic stroke: diagnosis, treatment of acute phase, primary and secondary prevention, atrial fibrillation, neuroimaging.

## Job-related skills

Experience of clinical management of patients affected by neurological diseases (i.e., Parkinson' disease, dementia, multiple sclerosis, epilepsy, headache).  
 Specific experience of clinical management of patients affected by movement disorders (i.e., Parkinson' disease, Multiple System Atrophy, Progressive Supranuclear Palsy, Dementia with Lewy Bodies, Spinocerebellar Ataxias, dystonia).  
 Specific experience of clinical management of patients affected by neurodegenerative and neuromuscular diseases (i.e., Amyotrophic lateral sclerosis, Congenital myopathies, Congenital neuropathies, Mitochondriopathies).  
 Specific experience of clinical management of patients affected by vascular cognitive impairment and small vessel diseases (i.e., CADASIL, Fabry Disease).  
 Experience of clinical management of the acute phase of ischemic and hemorrhagic stroke (Stroke Unit).  
 Experience of laboratory activity in the field of molecular genetics (Next Generation Sequencing; Sanger Sequencing; DNA extraction; RT-PCR).

Le dichiarazioni contenute nel curriculum formativo e professionale allegato sono veritiere e sono rese ai sensi del D.P.R. 445/00 e di essere a conoscenza delle sanzioni penali di cui all'art.76 del D.P.R. medesimo in caso di false dichiarazioni.

**Firma**

firmato Arianna Manini