

**C V
(B R E V E)**

NOME: Antonio Orlacchio

LUOGO E DATA DI NASCITA: Perugia, 14/02/1972

CITTADINANZA: Italiana

E-MAIL: antonio.orlacchio@uniroma2.it, a.orlacchio@hsantalucia.it

SITO WEB: http://www.hsantalucia.it/neurogen/index_it.htm

ISTRUZIONE:

2004	Dottorato di Ricerca (Neuroscienze) - Università di Roma “Tor Vergata”
2000	Specializzazione in Neurologia - Università di Perugia
1996	Laurea in Medicina e Chirurgia - Università di Perugia

FORMAZIONE ALL’ESTERO:

1997-2001	Post-Doctoral Fellow, Centre for Research in Neurodegenerative Diseases, University of Toronto, Canada
-----------	---

POSIZIONE ATTUALE:

Professore Aggregato di Neurologia-Ricercatore Confermato (SSD MED/26),
Dipartimento Universitario di Medicina dei Sistemi - Sezione di Neurologia,
Università di Roma ”Tor Vergata”; Dirigente Medico, Responsabile Ambulatorio e
Programma Assistenziale di Neurogenetica, UOC Neurologia, Dipartimento
Assistenziale di Neuroscienze, PTV; Direttore Laboratorio di Neurogenetica, CERC
- IRCCS Santa Lucia, Roma

ATTIVITÀ DI RICERCA:

Neurogenetica

PREMI ED ONORIFICENZE (SELEZIONATI):

2012	Onorificenza Camera di Commercio di Perugia
2010	Premio Internazionale “San Valentino d’Oro”
2003	European Neurological Society Award

PROGETTI DI RICERCA:

Coordinatore Scientifico, 11; Responsabile UO, 21; Collaboratore: 5
Fonti di finanziamento: Ministero della Salute, Fondazione Telethon, Ministero
dell’Università e Ricerca Scientifica, Università di Roma “Tor Vergata”, Rotary
International, Lundbeck Foundation (Danimarca), Nakabayashi Trust for ALS
Research (Giappone), National Institute of Health (USA)

PUBBLICAZIONI:

Lavori *per extenso* revisionati da pari: 68, di cui 65 con *IF*
Primo autore: 24; Autore senior: 2; Autore per corrispondenza: 23
IF totale *ISI 2011*: 300
Capitoli di libro: 6; Abstracts: 137; Altre pubblicazioni: 10

LAVORI PER EXTENO REVISIONATI DA PARI (ULTIMI 5 ANNI):

28. Tarquini F, Tiribuzi R, Crispoltoni L, Porcellati S, Del Pino AM, **Orlacchio A**, Coata G, Arnone S, Torlone E, Cappuccini B, Di Renzo GC, Orlacchio A:
Caspase 3 activation and PARP cleavage in lymphocytes from newborn babies of diabetic mothers with unbalanced glycaemic control.
Cell Biochemistry and Function, in stampa
27. Kawarai T, Pasco PMD, Teleg RA, Kamada M, Sakai W, Shimozono K, Mizuguchi M, Tabuena D, **Orlacchio A**, Izumi Y, Goto S, Lee LV, Kaji R:
Application of long- range polymerase chain reaction in the diagnosis of X-linked dystonia parkinsonism.
Neurogenetics, in stampa
26. Martino S, Montesano S, di Girolamo I, Tiribuzi R, Di Gregorio M, **Orlacchio A**, Datti A, Calabresi P, Sarchielli P, Orlacchio A:
Expression of cathepsins S and D signals a distinctive biochemical trait in CD34+ hematopoietic stem cells of relapsing-remitting multiple sclerosis patients.
Multiple Sclerosis Journal, in stampa
25. Miyashiro A, Sugihara K, Kawarai T, Miyamoto R, Izumi Y, Morino H, Maruyama H, **Orlacchio A**, Kawakami H, Kaji R:
Oromandibular dystonia associated with SCA36.
Movement Disorders 2013, 28(4): 557-559
24. Tiribuzi R, Crispoltoni L, Tartacca F, **Orlacchio A**, Martino S, Palmerini CA, Orlacchio A:
Nitric oxide depletion alters hematopoietic stem cell commitment toward immunogenic dendritic cells.
Biochimica et Biophysica Acta (BBA) - General Subjects 2013, 1830(3): 2830-2838
23. Chiurchiù V, Maccarrone M, **Orlacchio A**:
Compositions and methods for treatment of Parkinson's disease: a patent evaluation of WO2011/102847A1.
Expert Opinion on Therapeutic Patents 2012, 22(2): 181-184
22. Montenegro G, Rebelo AP, Connell J, Allison R, Babalini C, D'Aloia M, Montieri P, Schüle R, Ishiura H, Price J, Strickland A, Gonzalez MA, Baumbach-Reardon L, Deconinck T, Huang J, Bernardi G, Vance JM, Rogers MT, Tsuji S, De Jonghe P, Pericak-Vance MA, Schöls L, **Orlacchio A***, Reid E, Züchner S:
Mutations in the ER-shaping protein reticulon 2 cause the axon-degenerative disorder hereditary spastic paraparesis type 12.
The Journal of Clinical Investigation 2012, 122(2): 538-544
* Co-senior author
21. Mattoli F, Tiribuzi R, D'Angelo F, di Girolamo I, Quattrocelli M, Montesano S, Crispoltoni L, Oikonomou V, Cusella De Angelis MG, Marconi P, **Orlacchio A**, Sampaolesi M, Martino S, Orlacchio A:
Development of a new tool for 3D-modeling for regenerative medicine.
International Journal of Biomedical Imaging 2011, 2011: 236854
20. Tiribuzi R, **Orlacchio A**, Crispoltoni L, Maiotti M, Zampolini M, De Angelis M, Mecocci P, Cecchetti R, Bernardi G, Datti A, Martino S, Orlacchio A:
Lysosomal β-galactosidase and β-hexosaminidase as feature of Alzheimer's disease development in T2DM patients.
Journal of Alzheimer's Disease 2011, 24(4): 785-797

19. **Orlacchio A**, Montieri P, Babalini C, Gaudiello F, Bernardi G, Kawarai T: Late-onset hereditary spastic paraplegia with thin corpus callosum caused by a new *SPG3A* mutation.
Journal of Neurology 2011, 258(7):1361-1363
18. Shimizu H, Oka N, Kawarai T, Taniguchi K, Saji N, Tadano M, Bernardi G, **Orlacchio A**, Kita Y: Late-onset CMT2 associated with a novel missense mutation in the cytoplasmic domain of the MPZ gene.
Clinical Neurology and Neurosurgery 2010, 112(9): 798-800
17. **Orlacchio A**, Bernardi G, Orlacchio A, Martino S: Stem cells and neurological diseases.
Discovery Medicine 2010, 9(49): 546-553. Invited Review
16. **Orlacchio A**, Babalini C, Borreca A, Patrono C, Massa R, Basaran S, Munhoz RP, Rogaeva EA, St George-Hyslop PH, Bernardi G, Kawarai T: SPATAC SIN mutations cause autosomal recessive juvenile amyotrophic lateral sclerosis.
Brain 2010, 133(Pt 2): 591-598
15. **Orlacchio A**, Bernardi G, Orlacchio A, Martino S: Stem cells: an overview of the current status of therapies for central and peripheral nervous system diseases.
Current Medicinal Chemistry 2010, 17(7): 595-608. Invited Review
14. Martino S, di Girolamo I, **Orlacchio A**, Datti A, Orlacchio A: MicroRNA implications across neurodevelopment and neuropathology.
Journal of Biomedicine and Biotechnology 2009, 654346. Invited Review
13. Refenes N, Bolbrinker J, Tagaris G, **Orlacchio A**, Drakoulis N, Kreutz R: Role of the H1 haplotype of microtubule-associated protein tau (*MAPT*) gene in Greek patients with Parkinson's disease.
BMC Neurology 2009, 9: 26
12. **Orlacchio A**, Brusa L, Moschella V, Iani C, Bernardi G, Mercuri NB: Treatment of the symptoms of Huntington's disease: preliminary results comparing Aripiprazole and Tetrabenazine.
Movement Disorders 2009, 24(1): 126-129
* Equal first author
11. Pippucci T, Panza E, Pompili E, Donadio V, Borreca A, Babalini C, Patrono C, Zuntini R, Kawarai T, Bernardi G, Liguori R, Romeo G, Montagna P, **Orlacchio A**, Seri M: Autosomal recessive hereditary spastic paraplegia with thin corpus callosum: a novel mutation in the *SPG11* gene and further evidence for genetic heterogeneity.
European Journal of Neurology 2009, 16(1): 121-126
10. Costanzi E, Persichetti E, Tiribuzi R, Massini C, Bernardi G, **Orlacchio A**, Orlacchio A: Effects of vitamin C on fibroblasts from sporadic Alzheimer's disease patients.
Neurochemical Research 2008, 33(12): 2510-2515
9. **Orlacchio A**, Bernardi G, Orlacchio A, Martino S: Patented therapeutic RNAi strategies for neurodegenerative diseases of the CNS.
Expert Opinion on Therapeutic Patents 2008, 18(10): 1161-1174. Invited Review

8. **Orlacchio A**, Bruce IN, Rahman P, Kawarai T, Bernardi G, St George-Hyslop PH, Gladman DD, Urowitz MB:
The apolipoprotein E2 isoform is associated with accelerated onset of coronary artery disease in systemic lupus erythematosus.
Medical Science Monitor 2008, 14(5): CR233-237
7. **Orlacchio A**, Patrono C, Gaudiello F, Rocchi C, Moschella V, Floris R, Bernardi G, Kawarai T:
Silver syndrome variant of hereditary spastic paraplegia: a locus to 4p and allelism with *SPG4*.
Neurology 2008, 70(21): 1959-1966
6. **Orlacchio A**, Patrono C, Borreca A, Babalini C, Bernardi G, Kawarai T:
Spastic paraplegia in Romania: high prevalence of *SPG4* mutations.
Journal of Neurology, Neurosurgery and Psychiatry 2008, 79(5): 606-607
5. Urbanelli L, Emiliani C, Massini C, Persichetti E, **Orlacchio A**, Pelicci G, Sorbi S, Hasilik A, Bernardi G, Orlacchio A:
Cathepsin D expression is decreased in Alzheimer's disease fibroblasts.
Neurobiology of Aging 2008, 29(1): 12-22
4. **Orlacchio A**, Bernardi G, Orlacchio A, Martino S:
RNA interference as a tool for Alzheimer's disease therapy.
Mini-Reviews in Medicinal Chemistry 2007, 7(11): 1166-1176. Invited Review
3. Matsui M, Kawarai T, Hase Y, Tomimoto H, Iseki K, Rogava EA, **Orlacchio A**, Bernardi G, St George-Hyslop PH, Takahashi R, Matsui M:
A novel mutation in *SPG3A* gene (atlastin) in a Japanese family with hereditary spastic paraplegia.
Journal of Neurology 2007, 254(7): 972-974
2. **Orlacchio A**, Calabresi P, Rum A, Tarzia A, Salvati AM, Kawarai T, Stefani A, Pisani A, Bernardi G, Cianciulli P, Caprari P:
Neuroachantocytosis associated with a defect of the 4.1R membrane protein.
BMC Neurology 2007, 7: 4
1. Pachatz C, Terracciano C, Desiato MT, **Orlacchio A**, Mori F, Rocchi C, Bernardi G, Massa R:
Upper motor neuron involvement in X-linked recessive bulbospinal muscular atrophy.
Clinical Neurophysiology 2007, 118(2): 262-268