



**Europass
Curriculum Vitae**

Personal information

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 Nationality Italian

Work experience

Dates	From May 2015
Occupation or position held	In charge of the Child and Adolescent Psychiatric Unit and Consultant at the Pediatric Neurology Unit
Name and address of employer	Children's Hospital A. Meyer - University of Florence
Dates	From April 2009 onwards
Occupation or position held	Consultant at the Pediatric Neurology Unit
Name and address of employer	Children's Hospital A. Meyer - University of Florence
Dates	June 2008 to March 2009
Occupation or position held	Research fellow
Name and address of employer	Aston University, Birmingham, UK
Dates	May 2007 to May 2008
Occupation or position held	Child neurologist and epileptologist
Name and address of employer	Epilepsy Unit of Child Neurology and Psychiatry Division of the University Hospital of Cagliari
Dates	April 2006 to May 2007
Occupation or position held	Attending Physician in the staff directed by Professor Renzo Guerrini
Name and address of employer	Epilepsy, Neurophysiology and Neurogenetic Unit of the IRCCS Stella Maris Foundation
Dates	July 2005 to March 2006
Occupation or position held	Consultant neurologist and epileptologist and research activity at the Stella Maris Foundation
Name and address of employer	Children's Hospital A. Meyer - University of Florence, Stella Maris Foundation, Pisa
Dates	February 2005 to April 2006
Occupation or position held	Research Fellow in the project on the genetics of epilepsy and cortical malformations coordinated by Prof. Renzo Guerrini
Name and address of employer	October 28, 1999
	Graduated in Medicine and Surgery discussing a thesis on the use of botulinic toxin in cerebral palsy.
	University of Cagliari, Italy
	October 28, 2004
	Specialized in Child Neurology and Psychiatry discussing a thesis on a Sardinian family with Temporal lobe epilepsy due to a new missense mutation.
	University of Cagliari, Italy

Education and training																																									
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Personal skills and competences																																									
Mother tongue(s)	Italian																																								
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(*) Common European Framework of Reference for Languages																																									
Additional information																																									
List of main publications and awards:																																									
1.Pisano T, Marini C, Aridon P, Parrini E, Cianchetti C, Casari G, Pruna D, Guerrini R. Clinical and genetic study of a new large italian family with autosomal dominant nocturnal frontal lobe epilepsy. <i>Epilepsia</i> 44,suppl 8:130,2003 2.Pisano T, Marini C, Pruna D, Mei D, Moro F, Cianchetti C, Guerrini R. New LGI1 missense mutation in an italian family with lateral temporal lobe epilepsy and febrile seizures. <i>Epilepsia</i> 45,suppl 3:123,2004 3.Pisano T, Marini C, Brovedani P, Brizzolara D, Pruna D, Mei D, Moro F, Cianchetti C, Guerrini R. Abnormal phonological processing in familial lateral temporal lobe epilepsy due to a new LGI1 mutation. <i>Epilepsia</i>46 (1):118-23,2005 4 C. Cianchetti, A. Fratta, T.Pisano, Laura Minafra. Pergolide improvement in neuroleptic-resistant Tourette cases: various mechanisms causing tics.<i>Neurol Sci</i> 26(2):137-9,2005 5. Aridon P, Marini C, Di Resta C, Brilli E, De Fusco M, Politi F, Parrini E, Manfredi I, Pisano T, Pruna D, Curia G, Cianchetti C, Pasqualetti M, Becchetti A, Guerrini R, Casari G. Increased sensitivity of the neuronal nicotinic receptor alpha 2 subunit causes familial epilepsy with nocturnal wandering and ictal fear. <i>Am J Hum Genet.</i> 2006 Aug; 79(2):342-50. 6. F. Moro, T. Pisano, B. Della Bernardina, R. Polli, A. Murgia, L. Toccante, F. Darra, A. Battaglia and R. Guerrini. Periventricular heterotopia in Fragile X syndrome <i>Neurology</i> 2006 Aug 22; 67(4):713-5. 7. G. Cossu, A. Mereu, M. Deriu, M. Melis, A. Molari, G. Melis, L. Minafra, T. Pisano et al. Prevalence study of primary blepharospasm (BSP) in Sardinia, Italy. <i>Mov Disord.</i> 2006 Nov;21(11):2005-8. 8. F. Becherini °, T. Pisano *, M. Castagna °, A. Iannelli ° and R. Guerrini à Progressive hemispheric shrinking in hemimegalencephaly: a possible role for seizure related neuronal loss. <i>Dev Med Child Neurol.</i> 2008: 553-7. 9. T. Pisano. M. Meloni, A. Nucaro, M. Falchi, C. Cianchetti and D. Pruna. Megalencephaly, polymicrogyria and hydrocephalus (MPPH syndrome): a new case with syndactyly. <i>Journal of Child Neurology.</i> 2008: 916-8. 10. M Falchi, G Palmas, T. Pisano et al. Incidence of epilepsy in extremely low-birthweight infants (<1000 g): a population study of central and southern Sardinia. <i>Epilepsia.</i> 2009 Jan;50 Suppl 1:37-40 11. Cianchetti C, Cianchetti ME, Pisano T, Hmaidan Y. Treatment of migraine attacks by compression of temporal superficial arteries using a device.<i>Med Sci Monit.</i> 2009: CR185-8. 12. Cianchetti C, Serici MC, Pisano T, Ledda MG. Compression of superficial temporal arteries by a handmade device: a simple way to block or attenuate migraine attacks in children and adolescents.<i>J Child Neurol.</i> 2010 J:67-70. 13. Nucaro AL, Falchi M, Pisano T, Rossino R, Boscarelli F, Stoico G, Milia A, Montaldo C, Cianchetti C, Pruna D. Ring chromosome 14 mosaicism: an unusual case associated with developmental delay and epilepsy, characterized by genome array-CGH. <i>Am J Med Genet A.</i> 2010:234-6. 14. Capovilla G, Beccaria F, Bianchi A, Canevini MP, Giordano L, Gobbi G, Mastrangelo M, Peruzzi C, Pisano T, Striano P, Veggjotti P, Vignoli A, Pruna D. Ictal EEG patterns in epilepsy with centro-temporal spikes. <i>Brain Dev.</i> 2011:301-9. 15. Nucaro A, Chillotti I, Pisano T, Pruna D, Cianchetti C. Progressive spastic paraplegia as a feature of tetrasomy 18p. <i>Am J Med Genet A.</i> 2010:2173-5. 16. Marini C, Mei D, Parmeggiani L, Norci V, Calado E, Ferrari A, Moreira A, Pisano T, Specchio N, Vigevano F, Battaglia D, Guerrini R. Protocadherin 19 mutations in girls with infantile-onset epilepsy.<i>Neurology.</i> 2010:646-53.																																									

17. Melani F, Mei D, **Pisano T**, Savasta S, Franzoni E, Ferrari AR, Marini C, Guerrini R. CDKL5 gene-related epileptic encephalopathy: electroclinical findings in the first year of life. *Dev Med Child Neurol*. 2011 Apr;53(4):354-60.
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19. Filippi L, Catarzi S, Gozzini E, Fiorini P, Falchi M, **Pisano T**, la Marca G, Donzelli G, Guerrini R. Hypothermia for neonatal hypoxic-ischemic encephalopathy: may an early amplitude-integrated EEG improve the selection of candidates for cooling? *J Matern Fetal Neonatal Med*. 2012:2171-6.
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54. De Maria B, Balestrini S, Mei D, Melani F, Pellacani S, **Pisano T**, Rosati A, Scaturro GM, Giordano L, Cantalupo G, Fontana E, Zammarchi C, Said E, Leuzzi V, Mastrangelo M, Galosi S, Parrini E, Guerrini R. Expanding the genetic and phenotypic spectrum of CHD2-related disease: From early neurodevelopmental disorders to adult-onset epilepsy. *Am J Med Genet A*. 2022 Feb;188(2):522-533. doi: 10.1002/ajmg.a.62548.
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- Young Investigator Award at the 25th International Epilepsy Congress, Lisbon October 2003
- Travel Bursary Award at the 6th European Epilepsy Congress, Vienna May

Firenze, 25 marzo 2022

Dott.ssa Tiziana Pisano