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| Dr Angela Lucia Berardinelli, Curriculum sketch |
| **Career profile (Education and Employment)** |
| Education and training   * July 1982 : High School : Maturità Classica (60/60) liceo Classico B. Zucchi, Monza(MI) * October 10th 1988 MD, graduated in Medicine cum laude at University of Milan * July 12th 1993 Speclisation in Child Neuorlogy and Psychiatry,full marksUniversity of Pavia * 2ndJanuary 1989- 1993: residency Child Neurolgy and Psychiatry, University of * Pavia, with special interest in neuromuscular disorders, * July 1993, March-June 1994, March-;June 2001: visiting fellow Hammersmith * Hospital , London (Prof. Dubowitz, Prof. Muntoni) * 1st Summer School of Myology, Institute de Myologye, Hopital Salpetrière, Paris * (Prof. Fardeau) * Perodic stages in Neuromuscular Laboratory, Padova (Prof. Angelini) and Verona (Neuropathology Clinic) * Visiting fellow Hammersmith Hospital , London (Prof. Dubowitz, Prof. Muntoni)   Emlpoyment and research experience   * Employment: since February1st 1994 permanent position at the Child Neurology and Psychiatry Department, C. Mondino Foundation University of Pavia, Pavia : diagnosis and clinical management of neuromuscular disorders in childhood, |

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| Total Publications |  | | | | |
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| Journal Articles | Reviews | Book Chapters | Books | \*Conference associated publications | Other |
| 66 | # | 2 | # | 1 | 10 |
| 1. **Selected pubblications**   1: Chieffo D, Brogna C, Berardinelli A, D'Angelo G, Mallardi M, D'Amico A,  Alfieri P, Mercuri E, Pane M. Early Neurodevelopmental Findings Predict School  Age Cognitive Abilities in Duchenne Muscular Dystrophy: A Longitudinal Study.  PLoS One. 2015 Aug 14;10(8):e0133214. doi: 10.1371/journal.pone.0133214.  eCollection 2015. PubMed PMID: 26275215; PubMed Central PMCID: PMC4537199.  2: Pane M, Fanelli L, Mazzone ES, Olivieri G, D'Amico A, Messina S, Scutifero M,  Battini R, Petillo R, Frosini S, Sivo S, Vita GL, Bruno C, Mongini T, Pegoraro E,  De Sanctis R, Gardani A, Berardinelli A, Lanzillotta V, Carlesi A, Viggiano E,  Cavallaro F, Sframeli M, Bello L, Barp A, Bianco F, Bonfiglio S, Rolle E, Palermo  C, D'Angelo G, Pini A, Iotti E, Gorni K, Baranello G, Bertini E, Politano L,  Sormani MP, Mercuri E. Benefits of glucocorticoids in non-ambulant boys/men with  Duchenne muscular dystrophy: A multicentric longitudinal study using the  Performance of Upper Limb test. Neuromuscul Disord. 2015 Oct;25(10):749-53. doi:  10.1016/j.nmd.2015.07.009. Epub 2015 Jul 17. PubMed PMID: 26248957.  3: Fattori F, Maggi L, Bruno C, Cassandrini D, Codemo V, Catteruccia M, Tasca G,  Berardinelli A, Magri F, Pane M, Rubegni A, Santoro L, Ruggiero L, Fiorini P,  Pini A, Mongini T, Messina S, Brisca G, Colombo I, Astrea G, Fiorillo C, Bragato  C, Moroni I, Pegoraro E, D'Apice MR, Alfei E, Mora M, Morandi L, Donati A, Evilä  A, Vihola A, Udd B, Bernansconi P, Mercuri E, Santorelli FM, Bertini E, D'Amico  A. Centronuclear myopathies: genotype-phenotype correlation and frequency of  defined genetic forms in an Italian cohort. J Neurol. 2015 Jul;262(7):1728-40.  doi: 10.1007/s00415-015-7757-9. Epub 2015 May 10. PubMed PMID: 25957634.  4: Rubboli G, Veggiotti P, Pini A, Berardinelli A, Cantalupo G, Bertini E,  Tiziano FD, D'Amico A, Piazza E, Abiusi E, Fiori S, Pasini E, Darra F, Gobbi G,  Michelucci R. Spinal muscular atrophy associated with progressive myoclonic  epilepsy: A rare condition caused by mutations in ASAH1. Epilepsia. 2015  May;56(5):692-8. doi: 10.1111/epi.12977. Epub 2015 Apr 3. PubMed PMID: 25847462.  5: Ripolone M, Ronchi D, Violano R, Vallejo D, Fagiolari G, Barca E, Lucchini V,  Colombo I, Villa L, Berardinelli A, Balottin U, Morandi L, Mora M, Bordoni A,  Fortunato F, Corti S, Parisi D, Toscano A, Sciacco M, DiMauro S, Comi GP, Moggio  M. Impaired Muscle Mitochondrial Biogenesis and Myogenesis in Spinal Muscular  Atrophy. JAMA Neurol. 2015 Jun;72(6):666-75. doi: 10.1001/jamaneurol.2015.0178.  PubMed PMID: 25844556.  6: Graziano A, Bianco F, D'Amico A, Moroni I, Messina S, Bruno C, Pegoraro E,  Mora M, Astrea G, Magri F, Comi GP, Berardinelli A, Moggio M, Morandi L, Pini A,  Petillo R, Tasca G, Monforte M, Minetti C, Mongini T, Ricci E, Gorni K, Battini  R, Villanova M, Politano L, Gualandi F, Ferlini A, Muntoni F, Santorelli FM,  Bertini E, Pane M, Mercuri E. Prevalence of congenital muscular dystrophy in  Italy: a population study. Neurology. 2015 Mar 3;84(9):904-11. doi:  10.1212/WNL.0000000000001303. Epub 2015 Feb 4. PubMed PMID: 25653289; PubMed  Central PMCID: PMC4351663.  7: Pane M, Mazzone ES, Sivo S, Fanelli L, De Sanctis R, D'Amico A, Messina S,  Battini R, Bianco F, Scutifero M, Petillo R, Frosini S, Scalise R, Vita GL, Bruno  C, Pedemonte M, Mongini T, Pegoraro E, Brustia F, Gardani A, Berardinelli A,  Lanzillotta V, Viggiano E, Cavallaro F, Sframeli M, Bello L, Barp A, Busato F,  Bonfiglio S, Rolle E, Colia G, Bonetti A, Palermo C, Graziano A, D'Angelo G, Pini  A, Corlatti A, Gorni K, Baranello G, Antonaci L, Bertini E, Politano L, Mercuri  E. The 6 minute walk test and performance of upper limb in ambulant duchenne  muscular dystrophy boys. PLoS Curr. 2014 Oct 7;6. pii:  ecurrents.md.a93d9904d57dcb08936f2ea89bca6fe6. doi:  10.1371/currents.md.a93d9904d57dcb08936f2ea89bca6fe6. PubMed PMID: 25642376;  PubMed Central PMCID: PMC4208936.  8: Pane M, Mazzone ES, Sivo S, Sormani MP, Messina S, D Amico A, Carlesi A, Vita  G, Fanelli L, Berardinelli A, Torrente Y, Lanzillotta V, Viggiano E, D Ambrosio  P, Cavallaro F, Frosini S, Barp A, Bonfiglio S, Scalise R, De Sanctis R, Rolle E,  Graziano A, Magri F, Palermo C, Rossi F, Donati MA, Sacchini M, Arnoldi MT,  Baranello G, Mongini T, Pini A, Battini R, Pegoraro E, Previtali S, Bruno C,  Politano L, Comi GP, Bertini E, Mercuri E. Long term natural history data in  ambulant boys with Duchenne muscular dystrophy: 36-month changes. PLoS One. 2014  Oct 1;9(10):e108205. doi: 10.1371/journal.pone.0108205. eCollection 2014. Erratum  in: PLoS One. 2015;10(3):e0121882. PubMed PMID: 25271887; PubMed Central PMCID:  PMC4182715.  9: Esposito S, Bruno C, Berardinelli A, Filosto M, Mongini T, Morandi L, Musumeci  O, Pegoraro E, Siciliano G, Tonin P, Marrosu G, Minetti C, Servida M, Fiorillo C,  Conforti G, Scapolan S, Ansaldi F, Vianello A, Castaldi S, Principi N, Toscano A,  Moggio M. Vaccination recommendations for patients with neuromuscular disease.  Vaccine. 2014 Oct 14;32(45):5893-900. doi: 10.1016/j.vaccine.2014.09.003. Epub  2014 Sep 16. PubMed PMID: 25223270.  10: Ricci G, Scionti I, Sera F, Govi M, D'Amico R, Frambolli I, Mele F, Filosto  M, Vercelli L, Ruggiero L, Berardinelli A, Angelini C, Antonini G, Bucci E, Cao  M, Daolio J, Di Muzio A, Di Leo R, Galluzzi G, Iannaccone E, Maggi L, Maruotti V,  Moggio M, Mongini T, Morandi L, Nikolic A, Pastorello E, Ricci E, Rodolico C,  Santoro L, Servida M, Siciliano G, Tomelleri G, Tupler R. Large scale  genotype-phenotype analyses indicate that novel prognostic tools are required for  families with facioscapulohumeral muscular dystrophy. Brain. 2013 Nov;136(Pt  11):3408-17. doi: 10.1093/brain/awt226. Epub 2013 Sep 11. PubMed PMID: 24030947;  PubMed Central PMCID: PMC3808686. | | | | | |