

LIST OF PUBLICATIONS OF THE ITALIAN NETWORK (LAST UPDATE NOVEMBER 2018)

1. J Neurol. 2017 Aug;264(8):1777-1784. "Revisiting mitochondrial ocular myopathies: a study from the ItalianNetwork". Orsucci D, Angelini C, Bertini E, Carelli V, Comi GP, Federico A, Minetti C, Moggio M, Mongini , Santorelli FM, Servidei S, Tonin P, Ardisson A, Bello L, Bruno C, Ienco EC, Diodato D, Filosto M, Lamperti C, Moroni I, Musumeci O, Pegoraro E, Primiano G, Ronchi D, Rubegni A, Salvatore S, Sciacco M, Valentino ML, Vercelli L, Toscano A, Zeviani M, Siciliano G, Mancuso M.
2. Acta Myol. 2017 Mar;36(1):25-27. "Mitochondrial ANT-1 related adPEO leading to cognitive impairment: is there a link?" Costanza Simoncini, Gabriele Siciliano, Gloria Tognoni and Michelangelo Mancuso Department of Clinical and Experimental Medicine, Neurological Clinic, University of Pisa, Italy.
3. International Workshop: Outcome measures and clinical trial readiness in primary mitochondrial myopathies in children and adults. Consensus recommendations. 16–18 November 2016, Rome, Italy. Michelangelo Mancuso a, Robert McFarland b, Thomas Klopstock c, Michio Hirano d on behalf of the consortium on Trial Readiness in Mitochondrial Myopathies
4. Neuromuscul Disord. 2016 Apr-May;26(4-5):272-6. "Mitochondrial neuropathies": A survey from the large cohort of the Italian Network". Mancuso M, Orsucci D, Angelini C, Bertini E, Carelli V, Comi GP, Federico A, Minetti C, Moggio M, Mongini T, Tonin P, Toscano A, Bruno C, Ienco EC, Filosto M, Lamperti C, Diodato D, Moroni I, Musumeci O, Pegoraro E, Spinazzi M, Ahmed N, Sciacco M, Vercelli L, Ardisson A, Zeviani M, Siciliano G.
5. Mol Genet Metab Rep. 2016 Sep 1;9:12-4. "Mitochondrial m.3243A > G mutation and carotid artery dissection." Mancuso M, Montano V, Orsucci D, Peverelli L, Caputi L, Gambaro P, Siciliano G, Lamperti C.
6. J Neurol. 2015 May;262(5):1301-9. "Redefining phenotypes associated with mitochondrial DNA single deletion". Mancuso M, Orsucci D, Angelini C, Bertini E, Carelli V, Comi GP, Donati MA, Federico A, Minetti C, Moggio M, Mongini T, Santorelli FM, Servidei S, Tonin P, Toscano A, Bruno C, Bello L, Caldarazzo Ienco E, Cardaioli E, Catteruccia M, Da Pozzo P, Filosto M, Lamperti C, Moroni I, Musumeci O, Pegoraro E, Ronchi D, Sauchelli D, Scarpelli M, Sciacco M, Valentino ML, Vercelli L, Zeviani M, Siciliano G.

7. *Biochem Biophys Res Commun.* 2015 Mar 13;458(3):601-604. "Early onset cardiomyopathy associated with the mitochondrial tRNA^{Leu}((UUR)) 3271T>C MELAS mutation". Brisca G, Fiorillo C, Nesti C, Trucco F, Derchi M, Andaloro A, Assereto S, Morcaldi G, Pedemonte M, Minetti C, Santorelli FM, Bruno C.
8. *J Neurol.* 2015 Mar;262(3):701-10. "Myo-cardiomyopathy" is commonly associated with the A8344G "MERRF" mutation". Catteruccia M, Sauchelli D, Della Marca G, Primiano G, Cuccagna C, Bernardo D, Leo M, Camporeale A, Sanna T, Cianfoni A, Servidei S.
9. *Curr Mol Med.* 2014;14(8):1069-1078. "Mitochondrial Diseases in Childhood". Ardisson A, Lamantea E, Invernizzi F, Zeviani M, Genitrini S, Moroni I, Uziel G.
10. *Mitochondrion.* 2014 Sep;18:49-57. "Riboflavin responsive mitochondrial myopathy is a new phenotype of dihydrolipoamide dehydrogenase deficiency. The chaperon-like effect of vitamin B2". Carozzo R, Torraco A, Fiermonte G, Martinelli D, Di Nottia M, Rizza T, Voza A, Verrigni D, Diodato D, Parisi G, Maiorana A, Rizzo C, Pierri CL, Zucano S, Piemonte F, Bertini E, Dionisi-Vici C.
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27. *J Neurol Sci*. 2012 Apr 15;315(1-2):146-9. "Optic atrophy plus phenotype due to mutations in the OPA1 gene: two more Italian families." Ranieri M, Del Bo R, Bordoni A, Ronchi D, Colombo I, Riboldi G, Cosi A, Servida M, Magri F, Moggio M, Bresolin N, Comi GP, Corti S.
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