

# Author Index Volume 1 (2014)

The issue number is given in front of the pagination

- Aartsma-Rus, A., *Dystrophin Analysis in Clinical Trials* (1) 41–53  
Aartsma-Rus, A., see van den Bergen, J.C. (1) 91–94  
Abicht, A., see Chaouch, A. (1) 75–90  
Ando, Y., see Yamashita, S. (1) 111–115
- Bartsakoulia, M., see Pyle, A. (1) 55–63  
Beeson, D., see Chaouch, A. (1) 75–90  
Blakely, E.L., see Pyle, A. (1) 55–63  
Boczonadi, V., see Pyle, A. (1) 55–63  
Bronze-da-Rocha, E., see Oliveira, J. (2) 169–179  
Bushby, K., see Chaouch, A. (1) 75–90
- Carr, B., see Montes, J. (2) 151–161  
Chaouch, A., V. Porcelli, D. Cox, S. Edvardson, P. Scarcia, A. De Grassi, C.L. Pierri, J. Cossins, S.H. Laval, H. Griffin, J.S. Müller, T. Evangelista, A. Töpf, A. Abicht, A. Huebner, M. von der Hagen, K. Bushby, V. Straub, R. Horvath, O. Elpeleg, J. Palace, J. Senderek, D. Beeson, L. Palmieri and H. Lochmüller, *Mutations in the Mitochondrial Citrate Carrier SLC25A1 are Associated with Impaired Neuromuscular Transmission* (1) 75–90  
Chinnery, P.F. see Pyle, A. (1) 55–63  
Chinnery, P.F., see Sommerville, E.W. (2) 119–133  
Cobben, N.A.M., see van den Bergen, J.C. (1) 99–109  
Cossins, J., see Chaouch, A. (1) 75–90  
Cox, D., see Chaouch, A. (1) 75–90
- de Coo, I.F.M., see van den Bergen, J.C. (1) 99–109  
De Grassi, A., see Chaouch, A. (1) 75–90  
de Groot, I.J.M., see van den Bergen, J.C. (1) 99–109  
De Vivo, D.C., see Montes, J. (2) 151–161  
Donlin-Smith, C.M., see Statland, J. (2) 181–190  
Douroudis, K., see Pyle, A. (1) 55–63  
Dowling, P., A. Holland and K. Ohlendieck, *Mass Spectrometry-Based Identification of Muscle-Associated and Muscle-Derived Proteomic Biomarkers of Dystrophinopathies* (1) 15–40  
Duff, J., see Pyle, A. (1) 55–63  
Dunaway, S., see Montes, J. (2) 151–161
- Edvardson, S., see Chaouch, A. (1) 75–90  
Eglon, G., see Pyle, A. (1) 55–63  
Elpeleg, O., see Chaouch, A. (1) 75–90  
Evangelista, T., see Chaouch, A. (1) 75–90
- Farooq, F., see MacKenzie, D. (1) 65–74  
Fineza, I., see Oliveira, J. (2) 169–179  
Fock, J.M., see van den Bergen, J.C. (1) 99–109  
Fook-Chong, S., see Gosavi, T.D. (2) 163–168  
Fujimoto, A., see Yamashita, S. (1) 111–115
- Gagnon, C., see Gallais, B. (1) 95–98  
Galban-Horcajo, F., see Miller, J.A.L. (2) 191–195  
Gallais, B., C. Gagnon, J. Mathieu, L. Richer, S. Jean and L. Laberge, *Cognitive Deficits Associated with Sleep Apnea in Myotonic Dystrophy Type 1* (1) 95–98  
Garber, C.E., see Montes, J. (2) 151–161  
Ginjaar, H.B., see van den Bergen, J.C. (1) 99–109  
Ginjaar, H.B., see van den Bergen, J.C. (1) 91–94  
Gomez-Duran, A., see Pyle, A. (1) 55–63  
Gonçalves, A., see Oliveira, J. (2) 169–179  
Gorman, G.S., see Sommerville, E.W. (2) 119–133  
Gosavi, T.D., G.Z. Huang, S. Fook-Chong and Y.L. Lo, *Predictive Value of Electrophysiology for Presence of Thymic Pathology in Myasthenia Gravis* (2) 163–168  
Griffin, H., see Chaouch, A. (1) 75–90  
Griffin, H., see Pyle, A. (1) 55–63
- Herczegfalvi, A., see Pyle, A. (1) 55–63  
Hirahara, T., see Yamashita, S. (1) 111–115  
Hirano, T., see Yamashita, S. (1) 111–115  
Holland, A., see Dowling, P. (1) 15–40  
Horemans, A.M.C., see van den Bergen, J.C. (1) 99–109  
Horvath, R., see Chaouch, A. (1) 75–90  
Horvath, R., see Pyle, A. (1) 55–63  
Huang, G.Z., see Gosavi, T.D. (2) 163–168  
Huebner, A., see Chaouch, A. (1) 75–90
- Jaros, E., see Miller, J.A.L. (2) 191–195  
Jean, S., see Gallais, B. (1) 95–98

- Kamil-Rosenberg, S., see Montes, J. (2) 151–161  
 Kampelmacher, M.J., see van den Bergen, J.C. (1) 99–109  
 Karcagi, V., see Pyle, A. (1) 55–63  
 Kaufmann, P., What got us here will not get us there (2) 117–118  
 Kramer, S.S., see Montes, J. (2) 151–161
- Laberge, L., see Gallais, B. (1) 95–98  
 Lamar, K.-M. and E.M. McNally, Genetic Modifiers for Neuromuscular Diseases (1) 3–13  
 Laval, S.H., see Chaouch, A. (1) 75–90  
 Lek, M. and D. MacArthur, The Challenge of Next Generation Sequencing in the Context of Neuromuscular Diseases (2) 135–149  
 Lo, Y.L., see Gosavi, T.D. (2) 163–168  
 Lochmüller, H., see Chaouch, A. (1) 75–90  
 Lochmüller, H., see Pyle, A. (1) 55–63
- MacArthur, D., see Lek, M. (2) 135–149  
 MacKenzie, A., see MacKenzie, D. (1) 65–74  
 MacKenzie, D., F. Shamim, K. Mongeon, A. Trivedi, A. MacKenzie and F. Farooq, Human Growth Hormone Increases *SMN* Expression and Survival in Severe Spinal Muscular Atrophy Mouse Model (1) 65–74  
 Maeda, Y., see Yamashita, S. (1) 111–115  
 Man, P.Y.W., see Pyle, A. (1) 55–63  
 Mathieu, J., see Gallais, B. (1) 95–98  
 McNally, E.M., see Lamar, K.-M. (1) 3–13  
 McNally, E.M., see Wyatt, E.J. (2) 197–206  
 Miller, J.A.L., A. Spyropoulos, E. Jaros, F. Galban-Horcajo, R.G. Whittaker and H.J. Willison, Anti-GQ1b ganglioside positive Miller Fisher syndrome – evidence of paranodal pathology on nerve biopsy (2) 191–195  
 Mongeon, K., see MacKenzie, D. (1) 65–74  
 Montes, J., C.E. Garber, S.S. Kramer, M.J. Montgomery, S. Dunaway, S. Kamil-Rosenberg, B. Carr, N.E. Strauss, D. Sproule and D.C. De Vivo, A Randomized, Controlled Clinical Trial of Exercise in Patients with Spinal Muscular Atrophy: Methods and Baseline Characteristics (2) 151–161  
 Montgomery, M.J., see Montes, J. (2) 151–161  
 Moore, D., see Pyle, A. (1) 55–63  
 Mori, A., see Yamashita, S. (1) 111–115  
 Mori, Y., see Yamashita, S. (1) 111–115  
 Müller, J.S., see Chaouch, A. (1) 75–90
- Niks, E.H., see van den Bergen, J.C. (1) 99–109  
 Niks, E.H., see van den Bergen, J.C. (1) 91–94
- Ohlendieck, K., see Dowling, P. (1) 15–40  
 Oliveira, J., A. Gonçalves, M.E. Oliveira, I. Fineza, R.C.M. Pavanello, M. Vainzof, E. Bronze-da-Rocha, R. Santos and M. Sousa, Reviewing Large LAMA2 Deletions and Duplications in Congenital Muscular Dystrophy Patients (2) 169–179  
 Oliveira, M.E., see Oliveira, J. (2) 169–179
- Palace, J., see Chaouch, A. (1) 75–90  
 Palmieri, L., see Chaouch, A. (1) 75–90  
 Pangalila, R., see van den Bergen, J.C. (1) 99–109  
 Pavanello, R.C.M., see Oliveira, J. (2) 169–179  
 Pierri, C.L., see Chaouch, A. (1) 75–90  
 Porcelli, V., see Chaouch, A. (1) 75–90  
 Pyle, A., V. Ramesh, M. Bartsakoulia, V. Boczonadi, A. Gomez-Duran, A. Herczegfalvi, E.L. Blakely, T. Smertenko, J. Duff, G. Eglon, D. Moore, P.Y.W. Man, K. Douroudis, M. Santibanez-Koref, H. Griffin, H. Lochmüller, V. Karcagi, R.W. Taylor, P.F. Chinnery and R. Horvath, Behr's Syndrome is Typically Associated with Disturbed Mitochondrial Translation and Mutations in the *C12orf65* Gene (1) 55–63
- Ramesh, V., see Pyle, A. (1) 55–63  
 Richer, L., see Gallais, B. (1) 95–98  
 Rijlaarsdam, M.E.B., see van den Bergen, J.C. (1) 99–109
- Santibanez-Koref, M., see Pyle, A. (1) 55–63  
 Santos, R., see Oliveira, J. (2) 169–179  
 Scarcia, P., see Chaouch, A. (1) 75–90  
 Senderek, J., see Chaouch, A. (1) 75–90  
 Shamim, F., see MacKenzie, D. (1) 65–74  
 Smertenko, T., see Pyle, A. (1) 55–63  
 Sommerville, E.W., P.F. Chinnery, G.S. Gorman and R.W. Taylor, Adult-onset Mendelian PEO Associated with Mitochondrial Disease (2) 119–133  
 Sousa, M., see Oliveira, J. (2) 169–179  
 Sproule, D., see Montes, J. (2) 151–161  
 Spyropoulos, A., see Miller, J.A.L. (2) 191–195  
 Statland, J., C.M. Donlin-Smith, S.J. Tapscott, S.M. van der Maarel and R. Tawil, Multiplex Screen

- of Serum Biomarkers in Facioscapulohumeral Muscular Dystrophy (2) 181–190
- Straathof, C.S.M., see van den Bergen, J.C. (1) 99–109
- Straub, V., see Chaouch, A. (1) 75–90
- Strauss, N.E., see Montes, J. (2) 151–161
- Sweeney, H.L., see Wyatt, E.J. (2) 197–206
- Tapscott, S.J., see Statland, J. (2) 181–190
- Tawil, R., see Statland, J. (2) 181–190
- Taylor, R.W., see Pyle, A. (1) 55–63
- Taylor, R.W., see Sommerville, E.W. (2) 119–133
- Töpf, A., see Chaouch, A. (1) 75–90
- Trivedi, A., see MacKenzie, D. (1) 65–74
- Uchino, M., see Yamashita, S. (1) 111–115
- Vainzof, M., see Oliveira, J. (2) 169–179
- van den Bergen, J.C., H.B. Ginjaar, A.J. van Essen, R. Pangalila, I.J.M. de Groot, P.J. Wijkstra, M.P. Zijnen, N.A.M. Cobben, M.J. Kampelmacher, B.H.A. Wokke, I.F.M. de Coo, J.M. Fock, A.M.C. Horemans, M. van Tol, E. Vroom, M.E.B. Rijlaarsdam, C.S.M. Straathof, E.H. Niks and J.J.G.M. Verschuuren, Forty-Five Years of Duchenne Muscular Dystrophy in The Netherlands (1) 99–109
- van den Bergen, J.C., H.B. Ginjaar, E.H. Niks, A. Aartsma-Rus and J.J.G.M. Verschuuren, Prolonged Ambulation in Duchenne Patients with a Mutation Amenable to Exon 44 Skipping (1) 91–94
- van der Maarel, S.M., see Statland, J. (2) 181–190
- van Essen, A.J., see van den Bergen, J.C. (1) 99–109
- van Tol, M., see van den Bergen, J.C. (1) 99–109
- Verschuuren, J.J.G.M., see van den Bergen, J.C. (1) 99–109
- Verschuuren, J.J.G.M., see van den Bergen, J.C. (1) 91–94
- von der Hagen, M., see Chaouch, A. (1) 75–90
- Vroom, E., see van den Bergen, J.C. (1) 99–109
- Whittaker, R.G., see Miller, J.A.L. (2) 191–195
- Wijkstra, P.J., see van den Bergen, J.C. (1) 99–109
- Willison, H.J., see Miller, J.A.L. (2) 191–195
- Wokke, B.H.A., see van den Bergen, J.C. (1) 99–109
- Wyatt, E.J., H.L. Sweeney and E.M. McNally, New Directions in Biology and Disease of Skeletal Muscle 2014 (2) 197–206
- Yamashita, S., A. Fujimoto, Y. Mori, T. Hirahara, A. Mori, T. Hirano, Y. Maeda, M. Uchino and Y. Ando, Coexistence of Amyotrophic Lateral Sclerosis and Myasthenia Gravis (1) 111–115
- Zijnen, M.P., see van den Bergen, J.C. (1) 99–109