Kantonsspital St.Gallen

Janine Kirby

Kontakt

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Publikationen (3)

Shepheard S, Parker M, Cooper-Knock J, Verber N, Tuddenham L, Heath P, Beauchamp N, Place E, Sollars E, Turner M, Malaspina A, Fratta P, Hewamadduma C, Jenkins T, McDermott C, Wang D, Kirby J, Openshaw P, Project MINE Consortium, Project MinE. Value of systematic genetic screening of patients with amyotrophic lateral sclerosis. J Neurol Neurosurg Psychiatry 2021; 92:510-518.

Cooper-Knock J, Kirby J, McDermott C, Hautbergue G, Ince P, PROJECT MINE ALS SEQUENCING CONSORTIUM, Kazoka M, Walsh T, Higginbottom A, Heath P, Wyles M, Niedermoser I, Robins H, Shaw P. Targeted Genetic Screen in Amyotrophic Lateral Sclerosis Reveals Novel Genetic Variants with Synergistic Effect on Clinical Phenotype. Front Mol Neurosci 2017; 10:370.

Shaw C, Shaw P, Robberecht W, Van Damme P, Veldink J, Van den Berg L, Ticozzi N, Taroni F, Gellera C, Silani V, Kirby J, Pall H, Morrison K, Al-Chalabi A, Weale M, Brown R, Landers J, Andersen P, Baas F, Vianney de Jong J, de Belleroche J, Morris A, Asbroek A, Schelhaas H, Scott K, Troakes C, Lee Y, Miller J, Johnson L, Topp S, Vance C, Shatunov A, Newhouse S, Jones A, Gray I, Wright J, Nestor P, Weber M, Sapp P, Lovestone S, Lupton M, Powell J, Rogelj B, Al-Sarraj S, Hortobágyi T, Smith B. The C9ORF72 expansion mutation is a common cause of ALS+/-FTD in Europe and has a single founder. Eur J Hum Genet 2012; 21:102-8.

Projekte (0)

Keine Resultate gefunden.

Kantonsspital St.Gallen Rorschacher Strasse 95 CH-9007 St.Gallen T: +41 71 494 11 11 support.forschung@kssg.ch