



Janine Kirby

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Publications (3)

Shepherd 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 Belleruche 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.

Projects (0)

No results found.

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