Kantonsspital St.Gallen

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Publications (2)

Czell D, Sapp P, Neuwirth C, Weber M, Andersen P, Brown R. Further analysis of KIFAP3 gene in ALS patients from Switzerland and Sweden. Amyotroph Lateral Scler Frontotemporal Degener 2017; 18:302-304.

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.

Projects (0)

No results found.

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