



Tim M Strom

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Publications (3)

Weishaupt J, Weis J, Dorst J, Volk A, Borck G, Sperfeld A, de Carvalho M, Klopstock T, Sendtner M, Otto M, Schuster J, Andersen P, Ludolph A, Strom T, Meitinger T, Freischmidt A, Danzer K, Braak H, Del Tredici K, Jablonka S, Kubisch C, German ALS network MND-NET, Ruf W, Weydt P, Grosskreutz J, Meyer T, Petri S, Grehl T, Müller K, Yilmaz R, Neuwirth C, Weber M, Zeller D, Hübers A, Günther K, Knehr A, Jordan B, Schrank B, Claeys K, Pinto S, Brenner D. Hot-spot KIF5A mutations cause familial ALS. *Brain* 2018

Brenner D, Danzer K, Volk A, Meitinger T, Strom T, Otto M, Kassubek J, Ludolph A, Andersen P, Wahlqvist M, Borck G, Müller K, Wieland T, Weydt P, Böhm S, Lule D, Hübers A, Neuwirth C, Weber M, Weishaupt J. NEK1 mutations in familial amyotrophic lateral sclerosis. *Brain* 2016

Horn D, Illig T, Bezzina C, Franke A, Spranger S, Villavicencio-Lorini P, Seifert W, Rosenfeld J, Klopocki E, Rappold G, Wohlleber E, Riess O, Bonin M, Kapeller J, Rivera-Brugués N, Moog U, Lorenz-Depiereux B, Eck S, Hempel M, Wagenstaller J, Gawthrop A, Monaco A, Strom T. Identification of FOXP1 deletions in three unrelated patients with mental retardation and significant speech and language deficits. *Hum Mutat* 2010; 31:E1851-60.

Projects (0)

No results found.

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