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Publications (2)

Nordin A, Burkhardt C, Neuwirth C, Holmøy T, Morita M, Tysnes O, Benatar M, Wu J, Lange D, Bisgård C, Asgari N, Tarvainen I, Brännström T, Weber M, Schweikert K, Grehl T, Akimoto C, Wuolikainen A, Alstermark H, Forsberg K, Baumann P, Pinto S, de Carvalho M, Hübers A, Nordin F, Ludolph A, Weishaupt J, Meyer T, Andersen P. Sequence variations in C9orf72 downstream of the hexanucleotide repeat region and its effect on repeat-primed PCR interpretation: a large multinational screening study. *Amyotroph Lateral Scler Frontotemporal Degener* 2016; 18:256–264.

Czell D, Andersen P, Neuwirth C, Morita M, Weber M. Progressive aphasia as the presenting symptom in a patient with amyotrophic lateral sclerosis with a novel mutation in the OPTN gene. *Amyotroph Lateral Scler Frontotemporal Degener* 2013; 14:138–40.

Projects (0)

No results found.

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