

### Publicazioni ultimi 3 anni

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- 6: Roux T, Barbier M, Papin M, Davoine CS, Sayah S, Coarelli G, Charles P, Marelli C, Parodi L, Tranchant C, Goizet C, Klebe S, Lohmann E, Van Maldergem L, van Broeckhoven C, Coutelier M, Tesson C, Stevanin G, Duyckaerts C, Brice A, Durr A; SPATAX network. Correction: Clinical, neuropathological, and genetic characterization of STUB1 variants in cerebellar ataxias: a frequent cause of predominant cognitive impairment. *Genet Med*. 2020 Dec 22. doi: 10.1038/s41436-020-01064-y. Epub ahead of print. Erratum for: *Genet Med*. 2020 Nov;22(11):1851-1862. PMID: 33353973.
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