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2. Weissbach A, Wittke C, Kasten M, Klein C. 'Atypical'Parkinson's disease–genetic. In: International review of neurobiology. Vol 149. Elsevier; 2019:207-235.

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4. Weissbach A, Klein C. Hereditary dystonia and parkinsonism: two sides of the same coin? Brain. 2014;137:2402-4.
5. Weissbach A, Bäumer T, Rosales R et al. Neurophysiological fingerprints of X-linked dystonia-parkinsonism – a model basal ganglia disease; Mov. Disord. 2015;30:873-5.
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7. Weissbach A, Bäumer T, Pramstaller P, et al. Abnormal premotor-motor interaction in heterozygous Parkin- and Pink1 mutation carriers. Clin. Neurophysiol. 2016;128:275-280
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2. Vulinovic F, Lohmann K, Rakovic A, Schmidt A, Capetian P, Weissbach A et al. Unraveling cellular phenotypes of novel TorsinA/TOR1A mutations. Hum Mutat. 2014;35:1114-22.
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