Pearls & Oy-sters: Family history of Huntington disease disguised a case of dentatorubral-pallidoluysian atrophy

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Pearls

- Additional neurologic signs such as mild cerebellar ataxia and uncommon EEG and MRI findings in patients with generalized chorea and a positive family history should prompt clinicians to also consider Huntington disease–like syndromes.

Oy-sters

- A self-reported positive family history for Huntington disease may disguise dentatorubral-pallidoluysian atrophy, an autosomal dominantly inherited trinucleotide repeat expansion disorder that is very rare outside of Asian countries.
- Mild to moderate cerebellar ataxia may be overlooked in the presence of dominating generalized chorea.

Case report

A 42-year-old woman of Caucasian ancestry presented with an episode of acute paranoid psychosis, an 8-month history of slowly progressive generalized choreoathetosis, gait and speech disturbance, as well as cognitive impairment. Her choreoathetosis remarkably improved due to olanzapine, and subsequently, cerebellar signs became obvious upon neurologic examination (video, <http://links.lww.com/WNL/A45>). Cranial MRI revealed moderate atrophy of the vermal part of the cerebellum but not of the caudate nucleus. EEG showed runs of frontal intermittent high-amplitude delta activity. The patient reported a positive family history for Huntington disease (HD) (figure). Genetic testing was negative twice for HD and spinocerebellar ataxias (SCAs) including SCA17, but revealed 59 ± 2 CAG repeats (<48) in the *ATN1* gene confirming the first German case of adult-onset dentatorubral-pallidoluysian atrophy (DRPLA).^{1,2}

Discussion

The differential diagnosis of choreatic movement disorders can be challenging. At first sight, the presence of characteristic clinical signs such as adult-onset progressive generalized choreoathetosis and the combination with neuropsychiatric features may prompt clinicians to order a genetic test for HD.³ This is particularly true for patients where the family history for HD is reported to be positive. Nevertheless, this seemingly obvious diagnosis sometimes may be misleading. We report a case of DRPLA where the correct diagnosis was initially concealed by a positive family history of HD. Further, cerebellar ataxia as an unusual clinical sign for HD was not initially identified and only noticed when choreoathetosis improved following neuroleptic treatment. After our patient was tested negative twice for a repeat expansion in *Huntingtin*, further genetic testing for HD-like syndromes revealed 59 ± 2 CAG repeats in the *ATN1* gene, confirming the correct genetic diagnosis. Retrospectively, the reported HD-positive family members happened to be diagnosed with HD before genetic testing was available—highlighting DRPLA as a differential diagnosis, also in populations where it was not described before.

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Video

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Figure Family history, imaging, and EEG of the index patient

Pedigree of the family (A), MRI (B), and routine EEG (C) of the index patient. Arrow = index patient (*ATN1* mutation); * = affected individuals; (?) = clinical status unknown. MRI reveals moderate atrophy of the vermal part of the cerebellum but not of the caudate nucleus. EEG shows runs of frontal intermittent high-amplitude delta activity.

DRPLA is caused by an autosomal dominantly inherited pathogenic CAG trinucleotide expansion in the *ATN1* gene.^{1,2} The CAG repeat length is normal in the range between 6 and 35, while the full penetrance is reported for ≥ 48 CAG repeats. The size of the expanded *ATN1* CAG repeat and the age at onset show an inverse correlation. Moreover, the clinical phenotype may vary dependent on repeat length and age at onset.^{2,4} Progressive ataxia and cognitive impairment are the most commonly reported features in all patients, while epilepsy and myoclonus are predominantly found in children, and choreoathetosis and dementia in adults.⁴ Progressive myoclonus epilepsy occurs usually in patients with an age at onset lower than 20 years.⁴ In individuals with an age at onset of 20 years and above, predominant features include choreoathetosis, dementia, and psychiatric disturbances such as psychosis.⁵ The total mean age at onset is 31.5 years, ranging from less than 12 months to 72 years.² DRPLA has initially been reported to occur predominantly in Japan with a prevalence of 0.48:100,000, but several families of non-Asian ancestry have been identified, including North and South American, African American, and European populations.² Analyses of the haplotype associated with DRPLA showed similarities between the Japanese and Portuguese haplotype, suggesting a founder effect.²

Brain imaging in DRPLA revealed a distinct atrophy of the brainstem and the cerebellum depending on the CAG repeat length.⁶ This pattern is different from HD, in which the striatum is early and usually predominantly involved.⁷

DRPLA is an important differential diagnosis in individuals with adult-onset progressive ataxia, choreoathetosis, or dementia, even if the patients are of non-Asian ancestry and report a positive family history for other genetically inherited neurodegenerative disorders.

Author contributions

Sinem Tunc: acquisition and interpretation of data, drafting the manuscript. Vera Tadic: acquisition and interpretation of

data, drafting the manuscript. Christine Zühlke: interpretation of data, critical revision of manuscript for intellectual content. Yorck Hellenbroich: interpretation of data, critical revision of manuscript for intellectual content. Norbert Brüggemann: study concept and design, interpretation of data, critical revision of manuscript for intellectual content.

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