Absence epilepsy and the CHD2 gene: an adolescent male with moderate intellectual disability, short-lasting psychoses, and an interstitial deletion in 15q26.1–q26.2

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Abstract: Deletions of the 15q26 region encompassing the chromodomain helicase DNA binding domain 2 (CHD2) gene have been associated with intellectual disability, behavioral problems, and several types of epilepsy. Including the cases mentioned in ECARUCA (European cytogeneticists association register of unbalanced chromosome aberrations) and DECIPHER (database of genomic variation and phenotype in humans using ensemble resources), so far, a total of 13 intellectually disabled patients with a genetically proven deletion of the CHD2 gene are described, of whom eleven had a history of severe forms of epilepsy starting from a young age. In this article, a moderately intellectually disabled 15-year-old male with a 15q26.1–q26.2 interstitial deletion is reported, who was referred for analysis of two recent short-lasting psychotic episodes that were nonresponsive to antipsychotic treatment and recurrent disinhibited behaviors since early infancy. Careful interdisciplinary assessment revealed that the psychotic phenomena originated from a previously unrecognized absence epilepsy. Treatment with valproic acid was started which resulted in full remission of psychotic symptoms, and consequently, substantial improvement of behavior. It was concluded that in case of (rare) developmental disorders with genetically proven etiology, a detailed inventory of anamnestic data and description of symptomatology over time may elucidate epilepsy-related psychopathology for which a specific treatment regimen is needed.

Keywords: 15q26.1 interstitial deletion, CHD2 gene, intellectual disability, epilepsy, psychotic disorder, neuropsychology

Introduction

Microarray analysis has been shown to be of crucial importance for the understanding of the genetic etiology of intellectual disability (ID). With this technique, numerous (de novo) copy number variations have been discovered leading to several circumscribed microdeletion syndromes.1–2 Especially in patients in whom neuropsychiatric disorders or congenital anomalies are present, in addition to ID, extensive genetic analysis is warranted.

Partial deletions of the long arm of chromosome 15, particularly in the 15q26 region, have been typically associated with somatic anomalies, especially congenital diaphragmatic hernia.3 In addition, ID, epilepsy, and behavioral problems are frequently mentioned.4–6 With respect to the 15q26.1 subregion, the chromodomain helicase DNA binding domain 2 (CHD2) gene (OMIM: 602119), originally characterized by Woodage et al,7 has been repeatedly reported to play a pivotal role in cerebrocortical development.8 Both microdeletions encompassing this gene and de novo loss of
function mutations lead to a syndrome characterized by ID and epileptic encephalopathy with generalized seizures.9,10

So far, a total of nine patients with ID (seven females and two males) has been reported with a genetically proven deletion of the CHD2 gene. Of these patients, eight had a history of severe forms of epilepsy starting from the young age.4–6,11,12

Here, a 15-year-old male with a 15q26.1–q26.2 interstitial deletion is described, who was referred to the specialized outpatient department for psychopathology and genetics because of short-lasting psychotic episodes and problematic management of disinhibited behaviors for which over many years symptomatic treatment with antipsychotics (ie, haloperidol) was ineffective.

Clinical description

The patient is a 15-year-old male of Chinese origin, who was adopted at the age of 18 months in a malnourished condition from an orphanage by his Dutch foster parents. Shortly after birth, in the People’s Republic of China, he underwent the first surgical correction for bilateral cheilognathopala-toschisis followed by several reconstructive surgeries in the Netherlands up to the age of 5 years. Upon arrival, he was referred to a pediatrician of a university hospital for evaluation of developmental delay as well as bizarre autistic and disinhibited behaviors. At examination, apart from moderate auditory impairment secondary to recurrent upper airway infections, mild motor delay was noticed for which physical therapy was applied. Full laboratory investigation according to the Dutch adoption protocol, including screening for metabolic disorders, disclosed no abnormalities. Electroencephalogram (EEG) registration revealed focal frontocentral abnormalities that were difficult to interpret because of muscular artifacts.

At the age of 2 years, a symptomatic treatment with a low dose of haloperidol was started because of persisting challenging behavior. Subsequently, he was extensively examined by a child neurologist at the age of 5 years. Weight, length, and head circumference were 17.4 kg (−1 standard deviation [SD]), 110 cm (0 SD), and 46 cm (−2 SD), respectively. No abnormalities were found apart from motor overactivity. His attention was undisturbed, whereas communication was adequate with the use of a hearing aid. Since his parents frequently noticed brief moments with a blank staring and unresponsiveness to his surroundings, EEG recording was performed, which showed similar frontal irregularities without, however, specific epileptic configurations. A diagnosis of aspecific psychomotor retardation was made. Six months later, due to aggressive, self-injurious, and autistic behaviors, reevaluation by a pediatrician was performed, who advised a more congruent and structured behavioral approach.

During subsequent years, various interventions with antipsychotics for behavioral control were implemented, which, however, frequently induced unwanted side effects such as dyskinesias, akathisia, and facial edema. In this period, intelligence testing was performed twice with an interval of 3 years showing a total intelligence quotient (IQ) of 55 and a developmental age of ~3.6 years (calendar age: 8 years). At the age of 12 years, microarray analysis (Agilent 180K oligoarray, amadid 023363 [Agilent Technologies, Santa Clara, CA, USA]) demonstrated an 1.8 Mb interstitial deletion in 15q26.1–q26.2, encompassing the CDH2 and RGM4 genes that were considered to be etiologically involved (karyotype: arr[hg19]15q26.1q26.2(93,408,532-95,188,862)x1; Figure 1). Magnetic resonance imaging scan of the brain showed no abnormalities.

In the next 18 months, two short-lasting episodes occurred characterized by marked global regression, anxieties, hallucinatory experiences (visual and auditory), as well as
of psychopathology by means of the psychopathology instrument for mentally retarded adults pointed specifically at chaotic, anxious, and generally inadaptive behaviors (total score: 32; maximum: 56).^{17}

Based on the aforementioned information, a preliminary diagnosis of epilepsy-related recurrent brief psychosis was made and it was advised to perform EEG recording. The latter revealed that some epileptic paroxysms left frontotemporal and sporadic polyspike–wave complexes. For this reason, the antiepileptic compound valproic acid was prescribed, whereas symptomatic treatment for behavioral control with a low dose of haloperidol (1.6 mg daily [gtt]; plasma concentration: 1 μg/L) was continued. This antiepileptic intervention (600 mg valproic acid; plasma concentration: 85 mg/L) resulted in a full remission of psychotic symptoms and a gradual normalization of behavior to the premorbid level that, apart from incidentally occurring odd beliefs, was paralleled by stabilization of mood and marked improvement of social adaptation.

**Discussion**

In the moderately intellectually disabled patient presented here, psychotic phenomena appeared to be the result of a previously unrecognized absence epilepsy. Treatment with valproic acid induced a substantial improvement of his behavior while keeping free from further psychotic episodes. As stressed by several authors, epilepsy-related psychoses, both post- and interictal, are still a clinical challenge to recognize and treat appropriately.^{18,19}

The 15q26.1 microdeletion, encompassing among others the CHD2 gene, is in general associated with febrile seizures starting at early age followed by various epileptic features, myoclonic as well as tonic–clonic and absence seizures, and not typically clear dysmorphisms.^{9,10} In the current patient, absence epilepsy was only present. Out of a total of nine published cases, this type of epilepsy is mentioned in two, whereas in one, no epilepsy was present (Table 1). As can be inferred, in addition to the published cases, four

**Table 1** Reported patients with a 15q26 microdeletion encompassing the CHD2 gene, intellectual disability, and challenging behaviors

<table>
<thead>
<tr>
<th>Study</th>
<th>Patient number</th>
<th>Sex/age (years)</th>
<th>Deletion size</th>
<th>Dysmorphisms</th>
<th>Epilepsy</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>Veredice et al</td>
<td>1</td>
<td>F/2.6</td>
<td>−5 Mb</td>
<td>+</td>
<td>+^{b}</td>
<td>Congenital hypothyroidism</td>
</tr>
<tr>
<td>Dhamija et al</td>
<td>1</td>
<td>F/9</td>
<td>−731 kb</td>
<td>+</td>
<td>+^{b}</td>
<td>Gate ataxia</td>
</tr>
<tr>
<td>Capelli et al</td>
<td>1</td>
<td>F/6</td>
<td>−500 kb</td>
<td>+</td>
<td>+</td>
<td>Also deletion of RGMA</td>
</tr>
<tr>
<td>Courage et al</td>
<td>1</td>
<td>M/25</td>
<td>−415 kb</td>
<td>+</td>
<td>+</td>
<td>Also deletion of RGMA</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>M/21</td>
<td>−106 kb</td>
<td>+</td>
<td>+</td>
<td>Also deletion of RGMA</td>
</tr>
<tr>
<td>Chénier et al</td>
<td>1</td>
<td>F/11</td>
<td>−191 kb</td>
<td>+</td>
<td>+</td>
<td>Auditory neuropathy</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>F/9</td>
<td>−210 kb</td>
<td>+</td>
<td>+</td>
<td></td>
</tr>
</tbody>
</table>

(Continued)
patients are registered in international databases of whom one (DECIPHER 304608) was not diagnosed with epilepsy. Finally, two patients were detected in large copy-number variation studies, but no clinical information about their phenotype or presence of epilepsy is provided.

In eight patients, the 15q26.1 deletion also encompassed the RGMA gene (OMIM: 607362) that is assumed to be involved in immune functionality but not regarded to be of clinical significance for ID or epilepsy.  

### Conclusion

This case demonstrates the vital importance of detailed anamnesis and careful description of the symptom–cognition–behavior complex over time, both to ensure early detection of epilepsy-related psychopathology, especially in people with ID, and to avoid diagnostic misinterpretation and inappropriate treatment.

### Disclosure

The authors report no conflicts of interests in this work.

### References


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