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CASE REPORT

Autosomal recessive cerebellar ataxia caused by mutations in the PEX2 gene

Caroline Sevin1*, Sacha Ferdinandusse2, Hans R Waterham2, Ronald J Wanders2, Patrick Aubourg1

Abstract

Objective: To expand the spectrum of genetic causes of autosomal recessive cerebellar ataxia (ARCA).

Case report: Two brothers are described who developed progressive cerebellar ataxia at 3 1/2 and 18 years, respectively. After ruling out known common genetic causes of ARCA, analysis of blood peroxisomal markers strongly suggested a peroxisomal biogenesis disorder. Sequencing of candidate PEX genes revealed a homozygous c.865_866insA mutation in the PEX2 gene leading to a frameshift 17 codons upstream of the stop codon. PEX2 gene mutations usually result in a severe neurological phenotype (Zellweger spectrum disorders).

Conclusions: Genetic screening of PEX2 and other PEX genes involved in peroxisomal biogenesis is warranted in children and adults with ARCA.

Background

Main causes of autosomal recessive cerebellar ataxia (ARCA) include Friedreich ataxia, ataxia telangiectasia and oculomotor apraxia type 1 and 2 [1,2]. Cerebellar ataxia may sometimes be the leading neurologic symptom in a limited number of autosomal recessive metabolic diseases [1,2] but has not been reported as such in inherited peroxisomal diseases due to isolated enzyme deficiency or peroxisomal biogenesis defect [3,4]. Patients with adrenomyeloneuropathy, the adult form of X-adrenoleukodystrophy, or adult Refsum disease (resulting from mutations in PHYH gene coding for the peroxisomal phytanoyl-CoA hydroxylase enzyme, or in PEX7 gene coding for the peroxin 7 receptor protein) often develop cerebellar ataxia but have always associated neurological symptoms (peripheral neuropathy, retinitis pigmentosa, spastic paraplegia) [5,6]. Patients with peroxisomal biogenesis defects (Zellweger spectrum disorders) may also display cerebellar dysfunction among many other neurological symptoms when they have prolonged survival [3,4]. Peroxisomal biogenesis disorders (PBDs) are characterized by the loss of multiple peroxisomal metabolic functions caused by mutation in 13 PEX genes involved in the import of peroxisomal matrix proteins [3,4]. Except for rhizomelic chondrodysplasia punctata due to mutations in the PEX7 gene, all patients have multiple metabolic impairments involving: (1) oxidation of very long chain fatty acids (VLCFA), branched-chain fatty acids (phytanic and pristanic acids, bile acid intermediates) and L-pipecolic acid; (2) and plasmalogen biosynthesis. We report two brothers who developed isolated progressive cerebellar ataxia at 3 1/2 and 18 years. After ruling out common genetic causes of ARCA, screening of peroxisomal metabolites revealed in the youngest brother at 9 years a moderate increase in the plasma levels of phytanic and pristanic acids suggesting PBD. This patient was afterwards found to have a homozygous mutation in the PEX2 gene. His older brother carrying the same PEX2 gene mutation developed cerebellar ataxia at 18 years.

Case presentation

The younger brother (P1) was evaluated for progressive gait disturbance at the age of 9 years. He was the fourth child of non-consanguineous healthy parents. His milestones were normal but mild dysarthria was noticed at 2 1/2 years. Gait disturbance appeared at 3 1/2 years. At 7 years, he walked with a wide-based gait but was able to run, rise from the floor and climb stairs. On neurological examination, the child had moderate truncal ataxia, dysarthria but no tremor and pyramidal signs in the lower limbs. Symptoms worsened at the age of 9 years: he was...
no longer able to run, could walk only 100 meters without support and had difficulties with writing. On the Wechsler Intelligence Scale for Children (WISC-III), his full, verbal and performance IQ were 75, 84 and 79. He had truncal ataxia, moderate cerebellar tremor, mild scanning dysarthria, nystagmus, slow saccades without oculo-motor apraxia and hyporeflexia. His total ataxia score evaluated by the International Cooperative Ataxia Rating Scale (ICARS) was 17/100. He had no other neurological symptoms and his physical examination was normal. Cerebellar ataxia exacerbated slowly up to the age of 14 years but he was still able to walk without support. The ICARS score was 33/100. No other neurological signs were present and cognitive function remained normal.

At the age of 9 years, a brain MRI showed atrophy of the vermis and lateral hemispheres of the cerebellum that worsened at the age of 14 years, but without signs of demyelination or neuronal migration defects in forebrain and cerebellum (Figure 1). Fundoscopic examination, electroretinogram, EEG, electromyography, peripheral nerve conduction, visual, brainstem auditory and somatosensory evoked potentials, and audiogram were normal at 9 and 14 years. Other normal tests included ECG, cardiac ultrasound, liver enzymes, ammonia, lactate, vitamin E, lipid electrophoresis, alpha-foeto protein, plasma and urinary amino acids and urinary organic acids. Molecular studies of frataxin and aprataxin genes ruled out Friedrich ataxia and cerebellar ataxia with oculomotor apraxia 1 (AOA1).

His older brother (P2) had normal neurologic examination at 14 years, but at the age of 18 years he started to develop cerebellar signs with impaired gait, dysmetria, ataxia of the trunk, dysarthria, hyporeflexia and slow saccades with oculomotor apraxia. The ICARS score was 17/100. He had no other neurological symptoms. On the WISC-IV, his full, verbal and performance IQ were 80, 74 and 75 (normal range). Brain MRI showed isolated cerebellar atrophy at 18 years (Figure 1). Electroretinogram, peripheral nerve conduction, visual, brainstem auditory, and somatosensory evoked potentials were normal. These two patients had a 7-year-old brother and two sisters, aged of 22 and 25 years, who had a normal neurologic examination.

After ruling out Friedreich ataxia, AOA1 and common metabolic causes of cerebellar ataxia, peroxisomal parameters were analysed in plasma of patient P1 and his brother and sisters. Plasma VLCFAs were nearly normal in patients P1 and P2, but a moderate increase in the levels of phytanic and pristanic acids was found, prompting us to suspect a peroxisomal disorder (Table 1). Further studies showed a moderate increase of bile acid intermediates (DHCA and THCA) in plasma and normal levels of pipicolic acid in plasma and CSF (Table 1).

The findings of a moderate elevation of pristanic and phytanic acids with normal VLCFAs in plasma from P1 and P2 suggested initially that they might suffer from α-methylacyl-CoA racemase (AMACR) deficiency [7]. This was ruled out by the measurement of R- and S-isomers of DHCA and THCA in plasma and AMACR activity in fibroblasts from P1. The presence of DHCA and THCA traces also excluded the diagnosis of adult Refsum disease. Further metabolic studies performed in fibroblasts [7,8] from P1 showed normal plasmalogen biosynthesis, DHAP-AT activity, oxidation of VLCFAs, and pristanic acid (Table 2). Oxidation of phytanic acid was only slightly abnormal. Taken together, those analyses pointed towards a mild peroxisomal defect, but did not clearly indicate at which level. Since few patients with PEX2, PEX10 and PEX12 gene mutations display a relatively mild PBD phenotype, a complete genomic sequencing of these genes was performed. No mutations were identified in the PEX10 and PEX12 gene, but an apparent homozygous c.865_866insA mutation was found in the PEX2 gene in patients P1 and P2. The mother was heterozygous for the c.865_866insA mutation. Paternal genotype could not be determined because the proband’s father had lost contact with the family. No mutation was found in P1 and P2 asymptomatic siblings, who had normal peroxisomal parameters in plasma and no neurologic symptoms.

**Discussion**

The PEX2, PEX10 and PEX12 genes encode peroxins that are integral peroxisomal membrane proteins with a
cytosolic carboxy-terminal RING finger domain that act as ubiquitin ligases required for the ubiquitination of the PTS1-receptor (PEX5) in the peroxisomal membrane [9].

We do not have any reliable background data on the frequency of the c.865_866insA mutation in the proband’s ethnic background, and this mutation has not been reported in the database that we have consulted (http://www.hgmd.cf.ac.uk; http://www.dbpex.org; http://www.ncbi.nlm.nih.gov/snp). However, because the c.865_866insA mutation introduces a frameshift 17 codons upstream of the stop codon, resulting in a PEX2 protein with an altered C-terminus (p.Ser289LysfsX36), it is very unlikely that it could be a non-pathogenic variation occurring in the normal population. Additionally, homozygosity for this mutation co-segregated undeniably with the disease in the family. It should be noted, however, that homozygosity for this mutation could not be confirmed due to the unavailability of paternal DNA. This leaves the possibility that the paternal mutant allele harbors a different mutation, for example a (partial) deletion of the PEX2 gene, which will not be recognized when in trans with the c.855_866insA allele. In that case the c.855_866insA allele will also appear as homozygous.

The fact that peroxisomes in the patient’s fibroblasts were normally present and contained catalase suggests that the mutated PEX2 is localized correctly in the peroxisomal membrane and still is partly active. How this moderate impairment in peroxisome biogenesis impairs selectively neuronal cells in cerebellum is unknown. It is possible that environmental factors and modifier genes contribute to the different age of onset of clinical symptoms in the two affected brothers.

### Table 1 Analysis of plasma VLCFAs, branched-chain fatty acids and bile acids intermediates

|                      | Controls (5% - 95% range) | Patient 1 | Patient 2 |
|----------------------|---------------------------|-----------|-----------|
| **VLCFAs**           |                           |           |           |
| - C22:0              | 40-119                    | 34        | 36        |
| - C24:0              | 33-84                     | 33        | 38        |
| - C26:0              | 0.45-1.32                 | 0.76      | 0.95      |
| - C24:0/C22:0        | 0.57-0.92                 | 0.98      | 1.07      |
| - C26:0/C22:0        | 0-0.02                    | 0.02      | 0.03      |
| **Branched-chain fatty acids** |                   |           |           |
| - Phytanic acid      | 0.9                      | 23.5      | 10.3      |
| - Pristanic acid     | 0.4                      | 27.9      | 16.7      |
| - Pristanic/Phytanic ratio | 0.05-0.4             | 1.19      | 1.62      |
| **Bile acids intermediates** |                 |           |           |
| - DHCA               | 0-0.02                    | 0.4       | 0.1       |
| - THCA               | 0-0.08                    | 0.1       | 0.2       |

All values are given in μM.

### Table 2 Biochemical abnormalities in fibroblasts from patient 1 and 5 other patients with PEX2 mutations

|                      | Controls (5% - 95% range) | Patient 1 | Other PEX2 deficient patients |
|----------------------|---------------------------|-----------|-------------------------------|
| **Peroxisomal beta-oxidation** |                           |           |                               |
| - C26:0              | 1214-1508                 | 1296      | 186                           |
| - Pristanic acid     | 675-1121                  | 1114      | 2                             |
| **Phytanic acid alpha-oxidation** |                   |           |                               |
| - 8pPE in PE         | 728-814                   | 796       | 58.5                          |
| - 8pPC in PC         | 3.3-5.5                   | 5.8       | 1.1                           |
| **Plasmalogen de novo synthesis** |               |           |                               |
| - %pPE in PE         | 58.2-62.3                 | 83        | 0.7                           |
| - %pPC in PC         | 3.8-5.5                   | 5.8       | 0.5                           |
| **DHAPAT-activity**  | 5.8-12.3                  | 8.3       | 0.7                           |
| **Catalase immunofluorescence** |               |           |                               |
| - Acyl-CoA oxidase (70/50/20 kDa) | +/-             | +/-       | +/-                           |
| - Peroxosomal thiolase (44/41 kDa) | +/+               | +/+       | +/-                           |

Patients 1-4 are described in ref 12. Patient 5 is a newly identified PEX2 patient with severe clinical phenotype. °pmol/(h.mg protein); °nmol/(2 h.mg protein).

**List of abbreviations:** DHAP-AT, dihydroxyacetonephosphate-acyltransferase; DHCA, dihydroxycholestanolic acid; PC, total phosphatidylcholine; pPC, plasmalogen phosphatidylcholine; PE, total phosphatidylethanolamine; pPE, plasmalogen phosphatidylethanolamine; THCA, trihydroxycholestanolic acid; VLCFA, very-long chain fatty acids.
Cerebellar defects (cerebellar hypoplasia, altered folial pattern, abnormal Purkinje cell dendritic arborization) are observed in PEX2 deficient mice [10]. Tissue-selective elimination of peroxisomes in the mouse’s brain or liver suggests that the folial abnormalities may be due mainly to the lack of hepatic peroxisomes, whereas the Purkinje cells defects may imply both brain and liver peroxisomal abnormal metabolism [11]. The patients that we describe seem to have progressive cerebellar atrophy rather than malformativ cerebellar hypoplasia.

The clinical phenotype of these two patients strongly differs from PBD patients with PEX2 gene mutations who may display cerebellar symptoms in addition to other severe neurological signs in the context of Zellweger spectrum disorders [12,13].

While this work was in progress, PBD patients with mild phenotypes including cerebellar ataxia were shown to have mutations in PEX10 gene or PEX16 gene [14,15]. However, these patients displayed additional neurological symptoms including axonal motor neuropathy and decreased vibration sense in PEX10- mutated patients and spastic paraparesia, leukodystrophy peripheral neuropathy and cataracts in patients with PEX16 mutation. In contrast, our patients have a milder phenotype restricted to isolated cerebellar ataxia.

Conclusion
These clinical observations broaden the spectrum of phenotypes associated with PEX gene mutations but, most importantly, indicate that a search for PEX gene defect must be considered as potential cause of ARCA. Systematic screening for such PEX gene defect can easily and reliably be performed by the simple measurement of peroxisomal metabolites in plasma and will probably increase the frequency of PBD patients with mild phenotypes such as ARCA.

Consent
Written consent was obtained from the patient P2 and parents of patient P1 for publication of this case report.

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Authors’ contributions
CS and PA were involved in the clinical evaluation and follow-up of the patient, the data analysis and interpretation, and drafted the manuscript. SF, HRW and RJW carried out the biochemical and molecular genetic studies and the interpretation of the results. SF was involved in the write-up of the manuscript. All authors read and approved the final manuscript.

Competing interests
The authors declare that they have no competing interests.

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