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Case study

CLN6 disease caused by the same mutation originating in Pakistan has varying pathology

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ABSTRACT

The neuronal ceroid lipofuscinoses (NCLs), the most common neurodegenerative diseases in children, are characterised by storage of autofluorescent material that has a characteristic ultrastructure. We report two families with variant late infantile NCL, both originating from Pakistan. Probands from both families were homozygous for the same mutation (c.316dupC) but had variable pathology to that currently thought to be typical for CLN6 disease, late infantile variant. The observed pathology of one proband resembled condensed fingerprints, previously described in late infantile CLN7 and CLN8 diseases, and pathology from the second proband was thought to be absent even after repeated skin biopsy, but observed after review. This mutation is the most common NCL mutation in families originating from Pakistan and could be prioritised for testing. Finally, this report contains the first prenatal diagnosis for late infantile CLN6 disease, initially made on the basis of EM and now confirmed by mutation analysis.

1. Introduction

The neuronal ceroid lipofuscinoses (NCLs) are the most common neurodegenerative diseases in children, characterised by storage of autofluorescent material that has a characteristic ultrastructure.[1] Mutations in seven known genes (*CLN2/TPP1*, *CLN5*, *CLN6*, *CLN7/MFSD8*, *CLN8*, and *CLN1/PPT1* or *CLN10/CTSD*) cause NCL disease with onset in late infancy.[2,3] CLN1,

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CLN2 and CLN10 diseases can be diagnosed by enzyme assay, the remainder can be distinguished only by sequencing to identify or exclude a disease-causing variation. However, the ultrastructural pathology of the storage material is commonly used to guide the order of sequential sequencing.[4]

We report two families, both originating from Pakistan, in which the probands were homozygous for the same mutation but had variable pathology to that currently thought to be typical for CLN6 disease, late infantile variant.

2. Case reports

2.1. *Case 1 (UCL468)*

The proband was born to parents who were first cousins originating from Pakistan. Normal development was observed until his first seizure just before the age of 3 years after which he became unsteady on his feet, and developed myoclonic epilepsy. There was an 18 month history of nystagmus, ataxia and wide based gait, rough skin with increased pigment and a decelerating head circumference. He presented with bilateral optic atrophy and ERG deterioration by age 4. An EEG showed general bursts of sharp and slow waves and brain MRI displayed deficient inferior vermis and large cysterna magna at 4 years; by 5 years the patient had deteriorated, with abnormal myelination and ex-vacuo dilatation of ventricles thereby suggesting an atrophic process. By this age he was blind, in a wheelchair and unable to perform any skills. Later he became bed-ridden, with gastrostomy, and was stiff and uncomfortable when moved. Hearing was retained. He died aged 11 years. Examination of blood films stained with May-Grundwald giemsa showed no evidence of lymphocyte vacuolation. Rectal biopsy and blood samples were examined comprehensively by light microscopy and subsequently, ultrastructural analysis by transmission electron microscopy (EM). Submucosal ganglion cells showed prominent storage granules and stained positively with periodic acid Schiff's reaction, Luxol fast blue, Sudan black. The storage material exhibited strong autofluorescence when examined by ultraviolet light. The smooth muscle cells of the muscularis layer also had autofluorescent inclusions and stained positively for acid phosphatase. A muscle biopsy showed large active acid phosphatase positive lysosomes. Ultrastructural examination confirmed the presence of sparse storage deposits in lymphocytes. The storage material was membrane-bound as compact lipopigments with fingerprint profiles and amorphous material, confirming NCL disease (Fig. 1(a) and (b)). Variant late infantile NCL was diagnosed at age 5. *CLN7* and *CLN8* genes were excluded as the cause of disease by sequencing.

Following diagnosis of NCL but prior to genetic confirmation, there was a pregnancy terminated at 14 weeks due to the presence of inclusions in chorionic villus sample (CVS) (Fig. 1(c) and (d)), with inclusions confirmed in the aborted foetus (Fig. 1(e)). These were amorphous and of low density. A subsequent pregnancy for which CVS was normal resulted in a healthy sister.

DNA from the proband was exome sequenced and whole genome genotyped. Genomic DNA was prepared according to Illumina's TruSeq Sample preparation and Exome Enrichment protocols (Illumina, CA). Captured DNA was sequenced on an Illumina HiSeq 2000 using 2×100 bp paired-end reads. Resulting sequences were aligned to the human genome reference (hg19) and variants called using GATK.[5] Variants were annotated according to their presence in publicly available databases (1000Genomes and dbSNP build 135). The initial filtering process excluded all synonymous and heterozygous variants present in the proband. There were 57 homozygous variants, of which one was a pathogenic mutation in *CLN6* (NM_017882: c.316dupC: p.Arg106ProfsX26). Both parents were carriers, and DNA from the aborted foetus was homozygous for this same change.

2.2. *Case 2 (UC593)*

The proband is the only child of first degree consanguineous parents from Rawalpindi in northern Pakistan. Early development was normal. His speech and walking began to deteriorate when 3 years old. When 4 years he developed shaking of both legs which progressed to myoclonic jerks within a year, and he experienced his first generalised tonic clonic seizure at age 6, his current age. He now has continuous wringing hand movements, can no longer bear weight and has lost eating and drinking skills. MRI showed cerebral and cerebellar atrophy with ex-vacuo dilation of lateral ventricles. The periventricular white matter and posterior limb of internal capsules along with deep cerebellar white matter showed high T2 signal with hazy appearances. The optic nerve and chiasm were thinner than normal. Flash electroretinograms were absent bilaterally with giant flash visual evoked responses. This was further confirmed with abnormal somatosensory evoked potentials (giant cortical response) and present C reflexes. Giant flash visual responses were evoked at slow flash rate of 1.2 Oz-Fz.

NCL was suspected. However, ultrastructural examination of a skin biopsy twice failed to reveal any inclusions. EM of buffy coat was not requested. Nevertheless, DNA was submitted for sequencing of genes that typically cause NCL with onset in late infancy and a known pathogenic mutation (c.316dupC: p.Arg106ProfsX26) was found in homozygous form in *CLN6*. Subsequent ultrastructural analysis re-examination of the second skin biopsy by transmission electron microscopy showed distinct storage inclusions in sweat gland epithelium, endothelial cells and smooth muscle cells (Fig. 2). The storage was of a mixed type with curvilinear-like and fingerprint profiles.

2.3. *Ethical standards*

All work was approved by UCL Research Ethics Committee. The parents of the probands gave informed consent prior to inclusion in this study and to its publication.

3. Discussion

These two cases demonstrate important considerations regarding pathology in CLN6 disease, and perhaps NCL in general, since both are atypical. In family UCL468 the pathology is more condensed than previously reported for late infantile CLN6 disease, but resembles that reported for CLN7