Leukoencephalopathy caused by a 17p13.3 microdeletion

INTRODUCTION
Inherited white matter disorders (IWMDs) comprise a group of genetic disorders with onset from childhood to late adulthood in which there is degeneration of central nervous system white matter tracts. They are clinically and genetically heterogeneous. In adults, there may be a wide variety of clinical manifestations including cognitive decline, spasticity and ataxia, and the general course is progressive. Despite advances in imaging, phenotyping and genetic analysis, at least one-third of patient with suspected adult-onset IWMDs are refractory to diagnosis—hindering prognostication, reproductive decision making and the identification of disease-modifying treatments.

In this report, we describe a novel phenotype of extensive, confluent, paucisymptomatic adult-onset leukoencephalopathy in adults caused by a 248 kb microdeletion of 17p13.3, which unlike other similar microdeletion syndromes has no impact on psychomotor development and has not been associated with any syndromic or extraneurological features so far.

Case descriptions
The proband (patient 1) is a woman in her 30s referred with a history of headache. She had no other medical history. Her perinatal and early childhood history was unremarkable, and she met all psychomotor milestones, attended university and excelled at sport. Her neurological examination (including funduscopy) was normal aside from mild essential tremor, and her general examination revealed only mild hypermobility. A brain MRI revealed confluent white matter changes. Routine blood and cerebrospinal fluid white matter investigations were negative, with no evidence of systemic organ

Figure 1  Coronal and axial fluid attenuated inversion recovery (FLAIR) imaging for patient 1 at first presentation (A, B) and after a 5-year interval (C, D) demonstrating extensive deep white matter hyperintensity arranged in concentric band-like distributions (best seen on the coronal plane). Further coronal and axial FLAIR images for the patient 2 (E, F) and patient 3 (G, H) show a similar distribution of white matter changes. More extensive, confluent FLAIR hyperintensity is seen in patient 4 (I, J, K, L), the non-symptomatic father again with sparing of the U-fibres, corpus callosum, external capsule, anterior temporal lobe and infratentorial compartment.
involvement. Renal function was normal. The proband’s younger sister (patient 2) was already known to have similar white matter changes (having had an MRI for head injury), and also suffered from migraine. As a result, a genetic cause was suspected, and the remainder of the family were screened.

The proband’s older sister and father (patients 3 and 4)—neither of whom experienced migraine or other neurological symptoms—were found to also exhibit confluent white matter changes (figure 1). The rest of the family was unaffected clinically and radiologically.

All affected family members completed higher education and held high functioning jobs. There is no history of prenatal or postnatal abnormality, psychomotor disturbance, seizure or facial dysmorphism. Common to all affected individuals was hypermobility, with no other neurological or extraneurological features on examination.

In his mid-60s, patient 4 developed subacute gait problems culminating in repeat imaging that revealed multifocal high-grade glioma. He passed away soon after diagnosis.

Imaging abnormalities

All patients exhibited extensive, confluent, supratentorial deep white matter FLAIR hyperintensities arranged in apparent concentric bands with sparing of the U-fibres, corpus callosum, external capsule and infratentorial region (figure 1). In patients 2 and 4 there was mild posterior-temporal lobe involvement but in patients 1 and 3 the temporal lobe was spared. The changes were static over longitudinal imaging. There was no evidence of atrophy and the cortex was spared. None of the patients exhibited lacunes or diffuse abnormalities and there were no microbleeds on susceptibility weighted imaging. Spinal cord imaging, where available was normal. Magnetic resonance spectroscopy was not performed.

Genetic testing

We initially tested the proband with a targeted 75-gene panel for genetic variants associated with adult-onset white matter disorders, which was negative. All family members then underwent whole exome sequencing which did not identify any segregating likely pathogenic variants. The proband and her parents then underwent trio whole genome sequencing. This revealed heterozygosity for a 248 kb microdeletion on the short arm of chromosome 17 between position 1829683 and 2078186 (online supplemental figure 1). The proband’s siblings underwent targeted chromosomal microarray analysis which confirmed segregation. This deletion has not been found in controls.

DISCUSSION

We describe a novel syndrome of minimally symptomatic adults with florid, static leukoencephalopathy secondary to a 17p13.3 microdeletion. Microdeletion syndromes are not commonly associated with adult-onset IWMDs, and microarray analysis is not routinely performed. Typically, microdeletions that cause white matter abnormalities (1p36, 2q23.1, 6p25, 18q23 and 22q11.2) are detected in childhood and are associated with significant developmental abnormalities. Larger deletions in the 17p13.3 genomic region also cause white matter changes in children. Well-characterised phenotypes associated with these deletions, such as Miller-Dieker syndrome (caused by deletions that include genes YWHAE and PAFAH1B1) and isolated lissencephaly sequence (caused by deletions involving only PAFAH1B1 but not YWHAE) are also typically associated with significant symptoms including dysmorphic features, developmental delay and premature life expectancy. Emrick et al recently described six patients with 17p13.3 microdeletions between the YWHAE and PAFAH1B1 loci (online supplemental figure 1). These patients also presented with white matter abnormalities without significant associated intellectual disability. However, the patients described by Emrick et al are children who presented with dysmorphic facies, behavioural changes or extraneurolological features, unlike the adults described here.

Notably, all deletions in the Emrick et al cohort were larger than the deletion we report, explaining the dysmorphism and other syndromic features in their patients (online supplemental figure 1). Interestingly, the smallest region of overlap between the Emrick et al deletions (Chr17:1770860-2099130) is very similar in size and location to the deletion described here, indicating that this region is responsible for causing a unique static leukoencephalopathy.

There are 10 RefSeq genes within the deletion we report (online supplemental figure 1). RTN4RL1 and SMG6 are the only genes with specific expression in the brain, with the others expressed at low levels. RTN4RL1 regulates axonal and dendritic growth and contributes to normal axon migration, which is of interest given the appearance of concentric hyperintensities in our patients. RTN4RL1 has a high pHaplo score (0.99), indicating a high likelihood of haploinsufficiency. Deletions of its paralog RTN4R cause white matter abnormalities and have been associated with various neurological and psychiatric disorders. SMG6 (pHaplo score 0.99) encodes a component of the telomerase ribonucleoprotein complex, which plays a role in the nonsense-mediated messengerRNA decay pathway.

CONCLUSION

This report contributes to the literature on the genetic basis of white matter disorders. Although microdeletions are not a frequent cause of adult-onset IWMDs, in cases refractory to genetic diagnosis, consideration should be given to whole genome sequencing, chromosomal microarray or both.
Patient consent for publication  Consent obtained directly from patient(s).

Ethics approval  This study was approved by ethics committee under UCL ION code 10/H0721/87. Participants gave informed consent to participate in the study before taking part.

Provenance and peer review  Not commissioned; externally peer reviewed.

Supplemental material  This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

Open access  This is an open access article distributed in accordance with the Creative Commons Attribution 4.0 Unported (CC BY 4.0) license, which permits others to copy, redistribute, remix, transform and build upon this work for any purpose, provided the original work is properly cited, a link to the licence is given, and indication of whether changes were made. See: https://creativecommons.org/licenses/by/4.0/.

© Author(s) (or their employer(s)) 2023. Re-use permitted under CC BY. Published by BMJ.

► Additional supplemental material is published online only. To view, please visit the journal online (http://dx.doi.org/10.1136/jnnp-2023-331986).

To cite  Wade C, Williams T, Labrum R, et al. J Neurol Neurosurg Psychiatry Epub ahead of print: [please include Day Month Year]. doi:10.1136/jnnp-2023-331986

Received 20 June 2023
Accepted 1 August 2023

J Neurol Neurosurg Psychiatry 2023;0:1–3. doi:10.1136/jnnp-2023-331986

ORCID iDs
Charles Wade http://orcid.org/0000-0001-5199-7566

REFERENCES


