White matter abnormalities on MRI in neuroacanthocytosis

Neuroacanthocytosis denotes a group of uncommon heterogenous neurodegenerative disorders associated with acanthocytosis in the absence of any lipid abnormality. A variety of modes of inheritance have been proposed (X linked and autosomal recessive are clearly described, but a recent report of dominantly inherited chorea acanthocytosis appears to be caused by Huntington’s disease-like type 2 expansions in the junctophilin-3 gene) and mutations in two genes have been identified, the XK gene (in the X linked McLeod phenotype) and the CHAC gene (9q21; autosomal recessive). A wide variety of clinical features including chorea, orofacioclingual dyskinesia, dysphagia, dysarthria, peripheral neuropathy, myopathy, seizures, and dementia has been described in these disorders.

Case reports

Case 1

This patient was briefly described as case 19 in the report of Danek et al. He was a 61 year old white male who had been well until 3 years previously, when he took early retirement from teaching owing to ‘disillusionment’. He subsequently developed a progressive dementing illness, associated with facial tics, grunting noises, dysarthria, and chorea over the subsequent 3 years. There was no family history of neurodegenerative disease. He first presented to a neurologist having had an isolated generalised tonic–clonic seizure. On examination, he had a frontal dementia (Mini Mental State Examination (MMSE) score of 27/30) with evidence of self neglect and choreiform movements in all four limbs, and a prominent facial tic. He had little insight into his current illness. All tendon reflexes were absent. Investigation demonstrated numerous acanthocytes on blood films. Creatine kinase was raised at 1125 IU/l. Kell antigens were only weakly positive, which conformed to the McLeod phenotype. DNA analysis for Huntington’s disease was negative, but a R133X mutation in exon 2 of the XK gene was found. All other investigations were negative (full blood count, copper studies, lipid studies, protein electrophoresis, vasculitic screen (antinuclear antibody, anti-neutrophil cytoplasmic antibody, double stranded DNA antibodies) syphilis serology, neutrophil cytoplasmic antibody, double stranded DNA antibodies), anti-cardiolipin antibodies, HIV serology, Epstein–Barr virus serology, and analysis of the CHAC gene. MRI head scan on May 12, 2002 showed widespread areas of increased signal within the white matter of both cerebral hemispheres, especially within the lentiform nucleus bilaterally, but also within the thalamus, cerebral peduncles, andpons, and involving the corpus callosum (white arrow, fig 1B).

Case 2

This 56 year old Italian male developed chorea at the age of 42 years, and subsequently neuropsychological problems. The clinical aspects of this case have been reported previously. Numerous acanthocytes were seen on blood films, with weak Kell antigen. Analysis of the XK gene identified a R133X mutation. An MRI scan of the head showed mild increased signal within white matter temporally and parietally. T2, proton density and fibre linked FLAIR MRI sequencing (fig 1C) in the posterior periventricular white matter.

Case 3

A 32 year old Indian male, born of consanguineous parents, who presented with progressive disinhibition, altered personality and chorea over a 2 year period. His clinical details have not been reported previously. His family had noticed intermittent unusual head movements in which he would appear to be looking around the room into empty spaces while conversing. Although these movements were involuntary, he was able to stop them temporarily if asked to do so. His personality had become more volatile with emotional outbursts and frequent loss of temper. On examination, his MMSE was 27/30. There were continuous choreiform movements of head and neck, and of all four limbs. He was able to interrupt these temporarily if asked to do so. His speech was slightly dysarthric but there was no involuntary tongue protrusion or evidence of self mutilatory behaviour affecting the tongue or lips. The remainder of his neurological examination was normal. Numerous acanthocytes were seen on blood films (fig 1D). Kell serology was normal, with exclusion of the McLeod phenotype. All other investigations including Huntington’s mutational analysis, CSF, and white cell enzyme analysis were negative. Analysis of the CHAC locus is ongoing, but no mutations were identified in the XK gene. MRI head scan (fig 1E,F) demonstrated abnormally high signal in the periventricular white matter bilaterally, with involvement of the corpus callosum and cerebellar atrophy, but without contrast enhancement.

Discussion

Both computed tomography and MRI have been reported to show caudate and more generalised cerebral atrophy in neuroacanthocytosis. Although increased signal on T2 weighted MRI in the caudate and putamen has been noted previously, the increased signal throughout the cerebral hemispheres (including the corpus callosum in cases 1 and 3) reported here has not been reported previously. Extensive investigation for alternative causes of white matter abnormalities (vasculitic screen, and analysis

Figure 1 Axial T2 weighted (A) and sagittal (B) MRI from case 1, showing numerous areas of signal increase within the white matter, and involving the corpus callosum (arrow). (C) Axial T2 FLAIR MRI from case 2 showing mild signal increase within the white matter in the posterior periventricular area. (D) Blood film from case 3 showing numerous acanthocytes (arrow). Axial T2 weighted (E) and sagittal (F) MRI from case 3 showing similar, but less marked, white matter abnormalities to case 1, involving the corpus callosum (arrow).
of CSF, very long chain fatty acids, mitochondrial white cell enzymes and plasma lysosomal enzymes) was negative and there was no history of hypertension. In view of the ages of cases 1 and 2 when these patients were initially assessed, not much weight had been given to their MRI appearances; it was in the assessment of case 3 (a normotensive young male who was being investigated for a possible leukodystrophy) that the significance of both his abnormal blood film and his MRI prompted us to review the previous two cases.

Until we have a better understanding of the functional basis of these rare neurogenetic disorders, it is difficult to speculate as to the mechanism via which such abnormalities appear. Although the appearances reported in these cases are not specific, they widen the spectrum of MRI abnormalities that have been reported in neuroacanthocytosis. Thus, clinicians need to be particularly aware of the possibility of neuroacanthocytosis in any patient presenting with unexplained chorea, as the MRI appearances are so variable.

Acknowledgements
We are grateful to the Dr J A Spillane and the late Professor S Bunney for their evaluation and referral of case 1.

D J Nicholl, I Sutton
Department of Neurology, Queen Elizabeth Hospital, Birmingham, UK

M T Dotti
Department of Neurological and Behavioral Sciences, Università di Siena, Siena, Italy

S G Supple
Institute of Haematology, Royal Prince Alfred Hospital, Camperdown, Sydney, Australia

A Danek
Neurologische Klinik, Ludwig-Maximilians Universität, Munich, Germany

M Lawden
Department of Neurology, Leicester Royal Infirmary, Leicester, UK

Correspondence to: Dr D J Nicholl, Department of Neurology, Queen Elizabeth Hospital, Birmingham B15 2TH, UK; d.j.nicholl@bham.ac.uk
doi: 10.1136/jnnp.2003.026781
Competing interests: none declared

References

Disruption of facial affect processing in word deafness
Word deafness (also known as auditory agnosia for speech, or as auditory verbal agnosia) is a rare neurobehavioral syndrome characterised by an inability to understand spoken language in spite of intact hearing, speaking, reading, writing, and ability to identify non-speech sounds. The lesions associated with this condition tend to be bilateral and symmetrical in nature, and include cortical-subcortical tissue of the anterior part of the superior temporal gyrus. However, Heschl’s gyrus is not always damaged completely in the left hemisphere. Moreover, there have been documented cases of word deafness caused by unilateral left hemisphere cortical and subcortical lesions. Although these lesions are anatomically different, they represent an effective partial hemispheric disconnection.

Hemispheric disconnection has been associated with unusual disruptions of emotional processing. Bowers and Heilman reported a patient with a lesion of the deep white matter of the right occipito-temporo-parietal region. This patient could name famous faces and discriminate affectively neutral faces, but could not name facial emotions or select emotional faces reflecting a named emotion. Bowers and Heilman hypothesised a visual-verbal disconnection resulting in an anaemia for affective faces. More recently, Bowers, Bauer, and Heilman further articulated this idea, suggesting that this patient’s performance resulted from a disconnection between a hypothesised non-verbal affect lexicon in the right hemisphere and the verbal lexicon in left hemisphere, which normally communicate via the deep white matter pathways damaged in their patient.

The documented association between hemispheric disconnection and anoma for facial emotion raises the possibility that similar deficits in emotion processing may be observed in word deafness.

Case report
WD1 was a 45 year old man who had suffered a left posterior temporal lobe hemisphere CVA two years previously. MRI had demonstrated an acute lesion of the left temporal lobe and a chronic lesion of the right temporal lobe. His new stroke produced an initial Wernicke’s aphasia. On the whole, his speech was functional.

- He was able to differentiate and accurately recognise a range of environmental sounds, although he had trouble with high pitched sounds. His recognition was fast and accurate.
- He had no apraxia or other motor problems, and he was able to communicate by gestures.

Overall, the results of his neuropsychological evaluation were within normal limits. His specific deficits were consistent with those seen in word deafness.

Emotion processing
We administered a modified version of the Florida Affect Battery (FAB), including both facial and vocal prosody subtests, in an attempt to determine whether word deafness was associated with a disruption in the processing of affective prosody. The FAB consists of 10 subtests that evaluate emotion processing by different modalities: visual (facial expression), auditory (prosody), and visual/auditory cross-modal. WD1’s performance was compared with that of 20 healthy adult controls. The test was modified, in that all instructions and emotion labels were presented in written form rather than orally.

WD1 performed at chance level on the prosody tasks, regardless of their affective content. This may have been related to a premorbid occupational sensory hearing loss. The possibility that his word deafness also contributed to his poor performance cannot be ruled out. However, the relative influence of word deafness cannot be determined in the absence of control subjects with impaired hearing.

WD1 was able to complete the visual subtests of the FAB, and his ability to discriminate facial identity and facial affect was within normal limits (table 1). His ability to match a stimulus facial expression with one from a target array was also within normal limits. However, he was moderately impaired relative to controls in his ability to match a printed affective name to facial expressions. He was also severely impaired in his ability to select the correct affective face from an array of faces when presented with a printed emoticon label—that is, happy, sad, angry, frightened, neutral—despite intact reading and ability to discriminate affective facial expressions.

Discussion
WD1’s pattern of performance on the FAB was identical to that of Bowers’ and Heilman’s patient, and consistent with a visual-verbal disconnection. This finding...
raises the possibility that a very specific disturbance of visual affect processing is a component of the word deafness syndrome. However, many neurocognitive syndromes lack a unitary functional basis and instead are artefacts of the behavioural geography of the brain. That is probably so with the affective processing disturbance observed in this case. The documentation of intact naming of affect in another word deafness case would answer this question definitively. At the same time, the functional auditory deficits and characteristic neuroanatomy of word deafness raise intriguing questions about the status of auditory emotion processing in word deafness, in view of this patient’s preserved ability to identify non-speech sounds.

K W Greve, M T Heinly
Department of Psychology, University of New Orleans, LA, USA

K W Greve, C L Joffe, K J Bianchini
Jefferson Neurobehavioral Group, University of New Orleans, LA, USA

K J Bianchini
Department of Psychology, University of New Orleans, LA, USA

Correspondence to: Dr K W Greve; kgreve@uno.edu
doi: 10.1136/jnnp.2003.021790

References

A case of acute urinary retention caused by periaqueductal grey lesion

Diseases of the central nervous system often cause disturbances in micturition. These diseases include lesions in the spinal cord, pons, cerebellum, hypothalamus, basal ganglia, and cerebrum. Of these regions, the dorsomedial pontine tegmentum (pontine micturition centre, PMC), frontal lobe, and sacral spinal cord are considered important in controlling micturition. Recent studies in healthy humans using positron emission tomography (PET) have shown a significant increase in blood flow in the PMC and midbrain periaqueductal grey (PAG) during micturition and urine storage. Thus, in addition to the PMC, the PAG may play an important role in micturition control. However, to our knowledge, there is no clinical report that identifies the role of the PAG in micturition. Here we report a case of acute urinary retention caused by a small lesion in the PAG. A favourable response to steroid therapy resulted in the normalisation of micturition.

Case report

A 31 year old man had sudden voiding difficulty resulting in urinary retention and was referred to a neurologist. Although no particular abnormalities were observed except for an abnormal signal intensity on T2-weighted magnetic resonance image (T2-WI) showing hyperintensity in the periaqueductal grey (PAG) where a significant increase in blood flow has been observed on positron emission tomography during micturition and urine storage in healthy humans. (B) T2-WI showing a reduction in the intensity of the PAG lesion after steroid therapy.

Figure 1 (A) T2-weighted magnetic resonance image (T2-WI) showing hyperintensity in the periaqueductal grey (PAG) where a significant increase in blood flow has been observed on positron emission tomography during micturition and urine storage in healthy humans. (B) T2-WI showing a reduction in the intensity of the PAG lesion after steroid therapy.

A filling cystometrogram revealed an atomic bladder with diminished bladder sensation. There was no overflow incontinence.

Function, motor and sensory systems, and autonomic nervous system were intact with preserved anal reflex, penile erection, and ejaculation. Nerve conduction studies on all four extremities and thermography of the upper extremities were normal.

A filling cystometrogram revealed an atomic bladder with diminished bladder sensation. There was no overflow incontinence.

Laboratory tests and analysis of the cerebrospinal fluid were all within normal reference ranges including immunological examinations. However, MRI of the brain showed a small abnormal signal in the right dorsal part of the PAG that was hypointense on T1-weighted image (WI) and hyperintense on T2-WI and fluid-attenuated inversion recovery (FLAIR) (fig 1A). The lesion was not enhanced with contrast material. No other abnormalities were found on the MRI.

Although we were unable to establish a diagnosis despite the thorough work up, we considered the PAG lesion to be responsible for his urinary symptoms and a disease originating from an immunologic abnormality such as vasculitis, was suspected based on the MRI findings and the favourable response to the steroid therapy. Therefore, 1 g methylprednisolone was given intravenously for three days (steroid pulse therapy), followed by 60 mg oral prednisolone for two weeks which was then tapered at a rate of 10 mg/week. After the steroid therapy was initiated, the patient’s symptoms and the PAG lesion on subsequent MRI of the brain improved and he was able to void (fig 1B). However, the inability to void recurred, and a second course of pulsed steroid therapy was given. Day by day his symptoms improved again and resolved completely.

Comment

The patient reported here presented with acute urinary retention and diminished bladder sensation. The only abnormality detected by imaging, laboratory, and electrophysiological studies was a small PAG lesion. Therefore, we concluded that the PAG lesion was responsible for his symptoms. Unfortunately, we could not establish a diagnosis. However, on the basis of the favourable response to steroid therapy he was suspected to have a disease caused by some immunological abnormality.

Blok et al reported that in human PET studies the right dorsomedial pontine tegmentum and the PAG were significantly activated during micturition. In addition, the results of various studies also suggest that the PAG, especially the right dorsal part, plays a critical role in the control of micturition, possibly as the relay centre from the spinal cord to the PMC. The PAG lesion in the present case was located at the site identified in PET studies to be significantly activated during micturition. Although more similar cases are needed to establish a true relation, our findings in the present case provide direct clinical evidence of the role of the PAG in integrating the micturition reflex in humans.

H Yaguchi, H Soma, Y Miyazaki, J Tashiro, I Yabe, S Kikuchi, H Sasaki
Department of Neurology, Hokkaido University Graduate School of Medicine, Sapporo, Japan

H Kakizaki
Department of Urology, Hokkaido University Graduate School of Medicine, Sapporo, Japan

www.jnnp.com
proximal or truncal muscles remains obscure

the placement of larger lesions carries

this procedure for proximal tremors because

continues to surround the advisability of using

rated our earlier findings.

Kim demonstrated that stereotactic surgical ablation of

tremor synergistically abolishes Holmes’ tremor

stimulation and GPi pallidotomy

Combination of thalamic Vim

References

Knotschild et al., who demonstrated

thalamic nucleus ventrointermedial
(Vim) markedly improved Holmes’ tremor

in a patient with midbrain tumour, corroborated our earlier findings. In this patient, Vim thalamotomy alleviated tremor in both the distal and proximal segments of the upper extremity. However, controversy continues to surround the advisability of using thalamic Vim stimulation for proximal tremors because the placement of larger lesions carries increased risks and the somatotopy of the proximal or truncal muscles remains obscure in these segments.

Here we present a patient with a pontine haemorrhage in whom the combination of thalamic Vim deep brain stimulation (DBS) and globus pallidus internus (GPi) pallidotomy abolished Holmes’ tremor.

This 53 year old right-handed man with a history of essential hypertension suddenly developed right hemiparesis and cerebellar ataxia in February 2000. He was admitted to a hospital where radiological examinations showed a left upper brainstem haemorrhage (fig 1A). His neurological state gradually improved. However, in October 2001 a coarse, slowly progressive tremor arose in his right upper extremity. It was severely disabling and he could not use his right arm. He was admitted to our hospital in December 2001.

On admission, he was alert and oriented. His speech was mildly dysarthric and slurred. There was palatal tremor. Mild hemiparesis with increased stretch reflexes and Babinski sign were noted on the right side. There were mild deficits of position, vibratory sense, and superficial sensation of light touch and pain in his right upper and lower extremities. Dysmetria was more pronounced on the right. Because of severe truncal and gait ataxia, he could not remain upright without support; he was unable to walk even with assistance. There was coarse and severe tremor in the right upper extremity. It persisted at rest and its amplitude increased during maintenance of a fixed posture and intentional voluntary movements. It rendered the right arm useless and prevented him from feeding and caring for himself. He was exhausted because of the severe tremor that persisted throughout his waking hours.

Surface electromyograms showed rhythmic grouping discharges of 3.6 Hz in the right forearm muscles. His preoperative score on the Tremor Rating Scale (TRS) for his right upper extremity (Part A, score 5) was 11. Magnetic resonance imaging (MRI) study (December 2001) showed a haemosiderin ring around the lesion in the left pontine tegmentum (fig 1B). On T2-weighted images, a high signal lesion was seen in the left inferior olive, as consistent with the hypertrophic olivary degeneration (fig 1C).

As sequential pharmacotherapy using clonazepam (3×0.5 mg/day) and benserazide/levodopa (3×25/100 mg per day) was only slightly effective, he was referred for surgery. Prior informed consent was obtained from the patient and his family.

In January 2002, a quadripolar DBS electrode (Model 3387; Medtronic Inc., Minneapolis, MN, USA) was implanted in the left thalamic Vim nucleus with the aid of MRI, third ventriculography, and microelectrode guidance, as previously described. The optimal target was determined to be 7 mm posterior and 14.5 mm lateral to the midpoint of the AC–PC line, and 1 mm dorsal to the floor of the third ventricle. After creating a test lesion (42°C, 60 sec), a permanent anatomical lesion was made by heating the electrode tip to 72°C for up to 70 sec. The electrode was moved in 2 mm increments in the medial, lateral, and dorsal directions, and the lesioning process was repeated to increase the overall size of the lesion (fig 1F). GPi pallidotomy completely abolished his proximal tremor. However, it produced only a small effect on his distal tremor and discontinuation of Vim stimulation resulted in its reappearance at almost the preoperative level. Without stimulation, theTRS score for his upper extremity tremor (Part A, score 5) was 9. The combination of Vim stimulation and GPi pallidotomy had synergistic effects in abolishing Holmes’ tremor in our patient. The therapeutic benefits remain unchanged at the time of writing and theTRS score for his upper extremity tremor (Part A, score 5) is 0. His palatal tremor did not respond to Vim stimulation and pallidotomy and remains unresolved.

Figure 1

(A) Computed tomography (CT) scan showing a haematoma in the pontine tegmentum. (B, C) Axial views of T2-weighted magnetic resonance images at chronic stage (22 months after onset) demonstrating a haemosiderin ring around the lesion in the pontine tegmentum (B, arrow) and a high signal intensity area in the left inferior olivary nucleus indicating hypertrophic olivary degeneration (C). (D, E) Location of the electrode superimposed on the frontal (D) and lateral (E) view of a selective third ventriculography. The target point is indicated by the asterisk. (F) CT scan demonstrating the coagulative lesion made by the left GPi pallidotomy (arrow). The CT scan was carried out 10 days after pallidotomy. AC, anterior commissure; PC, posterior commissure; ML, midline.


Stereotactic Vim surgery, either thalamotomy or thalamic stimulation, is a mainstay in the surgical treatment of parkinsonian or essential tremor. Its efficacy in tremor suppression is superior to that of pallidotomy in parkinsonian patients. However, as evidenced by our case, it does not always provide satisfactory results in patients with Holmes-tremor, particularly with respect to their proximal tremors. The basal ganglia outflow pathway from the GPi exerts a direct influence on not only the thalamus but also the brainstem motor centres such as the pedunculopontine nucleus related to the mesencephalic tegmental field that controls the axial and proximal appendicular musculature via the descending reticulospinal tract. Therefore, unlike thalamic surgery, which interrupts the thalamocortical output that controls distal appendicular musculature via descending corticospinal and corticobulbar tracts, GPi pallidal surgery influences the control of otherwise inaccessible axial and proximal muscles. This may be the reason why GPi pallidotomy produced a marked alleviation of the proximal tremor in our patient. Due to the limited efficacy of thalamic Vim surgery on proximal tremors, the use of other or additional surgeries with greater effects—for example, pallidotomy or subthalamic area stimulation—should be considered for the treatment of Holmes’ tremor.

S Goto, K Yamada
Department of Neurosurgery, Graduate School of Medical Sciences, Kumamoto University, Kumamoto, Japan

Correspondence to: S Goto, MD, PhD, Department of Neurosurgery, Graduate School of Medical Sciences, Kumamoto University, Kumamoto 860-8556, Japan; sgoto@kaiju.med.kumamoto-u.ac.jp
doi: 10.1136/jnnp.2003.023077

References


No association of the mitochondrial DNA A12308G polymorphism with increased risk of stroke in patients with the A3243G mutation

There is a striking phenotypic variability among patients with the A3243G (rRNA(CU(C)) mutation of mitochondrial DNA (mtDNA), the most common heteroplasmic mtDNA mutation. It is responsible for ~80% of cases of MELAS (mitochondrial encephalomyopathy with lactic acidosis and stroke-like episodes) and is also associated with several other phenotypes including maternally inherited diabetes and deafness (MIDD) and chronic progressive external ophthalmoplegia (CPEO). Only 50% of patients carrying the A3243G mutation have stroke-like episodes and the reason for this clinical variability remains poorly understood. Although the percentage level of the A3243G mutation in clinically relevant tissues appears to be important, this relationship is far from clear. High percentage levels of the A3243G mutation in muscle are associated with stroke-like episodes, but approximately one in five patients harbouring >80% A3243G in muscle remain stroke free, suggesting that additional environmental and genetic factors may influence the phenotypic expression of this mutation.

One possibility is that background mtDNA sequence variation influences phenotype. There is a well-recognised association between the mtDNA genetic background (or haplogroup) and the risk of developing visual failure in another mtDNA disorder, Leber’s hereditary optic neuropathy, and a similar mechanism may influence the incidence of stroke-like episodes in patients harbouring the A3243G mutation. Intrafamilial clustering of clinical phenotypes in A3243G patients would indirectly support a role for the mtDNA background, though our own clinical experience suggests that there is significant clinical variability between families.

Pulkes et al. have previously reported an increased risk of stroke associated with the presence of a homoplasmic, polymorphic (A12308G) variant in 48 patients with the A3243G mutation. The A12308G polymorphism, which is also found in the second mitochondrial tRNA gene encoding leucine (tRNALeu(CUN)), occurs with a frequency of 21% in a population of European origin and defines the mtDNA super-haplogroup U/K together with two other polymorphisms (A1467G and G1372A). As haplogroup U has also been reported to be a risk factor for sporadic occipital stroke in patients with migraine, these observations could have profound implications for understanding of mitochondrial genotype and its relationship to the clinical phenotype. Here we report on the investigation of the A12308G polymorphism in a larger group of well-characterised, unrelated A3243G index cases.

Methods

We carried out a large, multicentre study to investigate the A12308G polymorphism in a group of 107 unrelated family index cases harbouring the A3243G mutation. The patients (>95% Caucasian) were from England, Germany, USA, Australia, and Finland. All patients were referred to a neurology clinic, where stroke-like episodes were diagnosed clinically by experienced neurologists based upon a characteristic clinical history and brain imaging with computed tomography (CT) or magnetic resonance imaging (MRI); in all cases, a molecular diagnosis of the A3243G mutation was made at a centre with expertise in the investigation of patients with mtDNA disorders.

To investigate the A12308G and G1372A polymorphisms, a 249 bp fragment spanning this mtDNA region was polymerase chain reaction (PCR)-amplified using a forward primer (5’ CATTTGAAATCTGACACAGGGCTT3’ ; nt 12164–12189) and a reverse primer (5’ GTTAAACGGGGTGTAAGGATG3’ ; nt 12412–12390). Amplified products were purified and sequenced using BigDye terminator cycle sequencing chemistry on an ABI 377 automated DNA sequence (Applied Biosystems, Warrington, UK).

Results

The A12308G polymorphism was present in 32 of the 107 patients, while 56 had a history of stroke-like episodes. Nine of the 56 patients with a history of a stroke and 23 of 51 patients without a stroke were harbouring the A12308G polymorphism. Every patient with the A12308G polymorphism also harboured the G1372A variant, indicating that they belong to the same mtDNA super-haplogroup U/K. As shown in fig 1, our study alone revealed an apparent negative association between stroke-like episodes and the A12308G polymorphism, an observation in direct contrast to the positive association found by Pulkes et al. Meta-analysis of all available data however, including the present study (n = 107) and the published study of Pulkes et al. (n = 48), revealed that 16 of the 77 patients with a history of a stroke and 25 of 78 patients without stroke harbouring the A12308G polymorphism. This did not show a statistically significant association between the A12308G polymorphism and stroke-like episodes (x^2 = 2.53, p = 0.112).

Discussion

The aim of our study was to examine whether a previously described association between the A12308G polymorphism and an increased risk of stroke in patients with the ‘mitochondrial mutation’ was reflected in a larger study group. In agreement with previous reports, 52% of our patients experienced stroke-like episodes and 30% harboured the A12308G polymorphism, confirming that our cohort of

![Figure 1](http://jnnp.bmj.com/ on May 12, 2022 by guest. Protected by copyright.)
A12308G polymorphism and stroke-like episodes

The clinical diversity associated with the A3243G mutation clearly involves multiple factors. We have previously shown a correlation between clinical phenotype and mutation load in muscle. Age may well be a contributing factor, although there was a tendency for patients with stroke-like episodes in our group to be younger than those without. This argues against age as a risk factor for stroke-like episodes, as seen in common stroke.

Importantly our findings serve to highlight the difficulty of performing association studies on small numbers of patients. This is particularly difficult for mitochondrial genetic association studies because of the high variability of the mitochondrial genome. Understanding the phenotypic differences between patients with specific, pathogenic mtDNA mutations will ultimately involve studies of large cohorts of patients, unless we are able to gain clues from experimental studies that may highlight factors involved in the altered expression or segregation of mtDNA mutations.

Acknowledgements

The authors thank Geoff Taylor (University of Newcastle upon Tyne) for help with the sequencing. This study was supported by a fellowship from the European Neurological Society (MD), the Wellcome Trust (PFC, DMT and RWT) and the Wellcome Trust (JPC, IWT and RWT). DRT is supported by the Muscular Dystrophy Association and the Australian NHMRC.

M Deschauer, P F Chinnery, A M Schaefer, D M Turnbull, R W Taylor
School of Neurology, Neurobiology and Psychiatry, The Medical School, University of Newcastle upon Tyne, UK

M Deschauer, S Zierz
Department of Neurology, Martin-Luther-University Halle-Wittenberg, Halle/Saale, Germany

S Shanske, S DiMauro
Department of Neurology, Columbia University College of Physicians and Surgeons, New York, USA

K Majamaa
Department of Neurology, University of Oulu, Finland

E Wilichowski
Department of Paediatrics and Neuropaediatrics, Georg-August University Göttingen, Germany

D R Thorburn
Murdoch Childrens Research Institute, Royal Children's Hospital, Melbourne, Australia

Correspondence to: Dr R W Taylor, School of Neurology, Neurobiology and Psychiatry, The Medical School, Framlington Place, University of Newcastle upon Tyne, Newcastle upon Tyne NE2 4HH, UK, r.w.taylor@ncl.ac.uk: doi: 10.1136/jnnp.2003.026278

Table 1

<table>
<thead>
<tr>
<th>Symptom or problem</th>
<th>Recorded in the hospital records (n = 92)</th>
<th>Elicited from patients and relatives at home interviews (n = 92)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Weakness</td>
<td>55 (60)</td>
<td>51 (55)</td>
</tr>
<tr>
<td>Headache</td>
<td>49 (53)</td>
<td>48 (52)</td>
</tr>
<tr>
<td>Epilepsy</td>
<td>35 (38)</td>
<td>44 (48)</td>
</tr>
<tr>
<td>Sensory loss</td>
<td>32 (35)</td>
<td>37 (40)</td>
</tr>
<tr>
<td>Cognitive loss</td>
<td>30 (33)</td>
<td>42 (46)</td>
</tr>
<tr>
<td>Dysphasia</td>
<td>29 (32)</td>
<td>23 (25)</td>
</tr>
<tr>
<td>Personality change</td>
<td>14 (15)</td>
<td>28 (30)</td>
</tr>
<tr>
<td>Fatigue</td>
<td>13 (14)</td>
<td>44 (48)</td>
</tr>
</tbody>
</table>

Values are n (%).

References


Early symptoms of brain tumours

Malignant cerebral glioma is the most common adult primary brain tumour but surprisingly few studies report how patients with early symptoms present in primary or secondary care. A retrospective audit in south east Scotland found considerable variation in the referral of patients with primary brain tumours: only one quarter of 439 patients were initially referred directly to specialist centres. This must relate in part to the way in which symptoms develop and the difficulty of distinguishing them from more common but sinister problems. For example, a large case record review of initial symptoms experienced by 653 glioma patients presenting to the National Hospital for Neurology and Neurosurgery, Queen Square, London, between 1955 and 1975 found a relatively low prevalence of neurological problems such as epilepsy (38%), headache (35%), mental change (17%), and hemiparesis (10%); by the time of diagnosis the prevalences were 54%, 71%, 52%, and 43%, respectively. Few studies focus on the accounts of patients and relatives. One qualitative interview study of 28 Swedish patients suggested that relatives noticed general changes including cognitive and personality change and took the initiative in seeking help more often than the patients themselves. The recently published last diaries of the politician and historian the late Alan Clark provoked us to reconsider the significance of early symptoms from the perspective of patients and their close relatives. Clark provides us with a moving account of the gradual onset of symptoms from a glioma—fatigue, problems with thinking and concentration, and intermittent headache over nine months. He also describes vividly the anxiety of knowing something was wrong but without any explanation, before his tumour was diagnosed.

During a study of quality of life already described, we had opportunity to visit glioma patients at home after diagnosis, to listen to their accounts, and to question relatives separately. Here we report data on 92 patients (table 1), suggesting a differing development of symptoms and problems from that described in their medical records, and a distinctly similar picture, in some, to that described by Alan Clark. Interviews tended to elicit histories of more subtle problems such as fatigue and cognitive and personality change almost as often as the neurological problems typically associated with brain tumours. Of the 48 patients with headache only two had developed no other symptoms by the time of diagnosis.

Our sample is limited to patients who were well enough for radiotherapy and to receive home visits after diagnosis. It therefore excludes those most disabled and confused at diagnosis and treated with steroids alone. The data only cover problems that had developed before diagnosis. We did not have access to primary care records to explore how symptoms were presented to general practitioners, but 41% (38 of 92) were referred to a neurologist. Of the 64 patients whom we questioned on the topic, 19% (12) were critical of the initial management by their general practitioner and 28% of 88 relatives thought there had been significant delay by the health care system as a whole. This issue remained salient for many, even after the patient had died. Of 56 whom we saw as part of a study after bereavement, one third (17) spontaneously mentioned concerns they continued to have about delay in diagnosis and the effect this might have had on quality of life or survival. The problems they identified ranged across primary, secondary, and tertiary care and included their perception that referrals had not been made quickly enough or that waiting for appointments and imaging had been excessive.

The lack of data on the development of symptoms means that current national criteria for urgent referral rely on data from patients presenting to specialist centres rather than on the predictive power of symptoms in the population attending primary and secondary care. The data elicited here confirm the earlier suggestion by McKeran and Thomas that the significance of headache may lie in its association with altered patterns of behaviour and disability. Although retrospective accounts cannot be used to define predictive factors for earlier diagnosis, they do suggest some implications for future research and practice. First, more detailed study of patients’ and relatives’...
experience might help further define the subacute presentation of cognitive and personality change and their relation to other complaints. Second, the predictive power of neurological symptoms presented to general practitioners could be explored using existing large primary care research datasets. Third, relatives of patients referred urgently should be asked to attend with them to clarify aspects of the history that the patient may be unaware of. Beginning to discuss openly the difficulty of earlier diagnosis may help families come to terms with this lasting aspect of their concern. This might also help repair unnecessary rifts in relations with general practitioners, who are best placed to provide local support and palliative care these patients so often need.

Acknowledgements
We thank Sue Hall and Maureen O’Connor for help collecting the data and patients and relatives for agreeing to be interviewed. Data collection was supported by the Cancer Research Campaign, grant number CP 1017, and writing up by the BMA TP Grayson Award for Health Education in Cancer. The views expressed do not represent those of either organisation.

E Davies
Department of Palliative Care and Policy, Guy’s, King’s and St Thomas’ School of Medicine, Weston Education Centre, Cottambe Road, Denmark Hill, London SE5 9RJ, UK
C Clarke
National Hospital for Neurology and Neurosurgery, University College Hospitals, Queen Square, London WC1N 3BG, UK

Correspondence to: Dr Elizabeth Davies; elizabethdavies@doctors.org.uk
doi: 10.1136/jnnp.2003.033308
Competing interests: none declared

References
5 Davies E, Clarke C, Hopkins A. Malignant cerebral glioma. II. Perspectives of patients and relatives on the value of radiotherapy. BJH 1996;313:1512–16.

Five year follow up of a patient with spinal and bulbar muscular atrophy treated with leuprorelin
Spinal and bulbar muscular atrophy (SBMA; MIM 313200) is an X linked late onset motor neurone disease characterised by slowly progressive proximal and bulbar muscle weakness, muscle atrophy, postural hand tremor, gynecomastia, and endocrine disturbances that include signs of partial androgen suppression. 

Case report
A 75 year old male noticed bilateral finger tremor at age 57. At age 63, he noticed weakness in his arms. He was admitted to our hospital in December 1991, when he was 64 years old. On initial examination, he had bilateral gynecomastia. Neurological examinations revealed facial weakness and lingual atrophy with fasciculations. Mild muscular atrophy was observed in the proximal parts of the upper extremities. Muscle strength was approximately in the range of 3/5 to 4/5 in the proximal parts, and 5/5 in the distal parts of the upper extremities. Fasciculations were observed in upper and lower extremities. Deep tendon reflexes were either lost or markedly diminished. Babinski signs were absent. Laboratory examinations revealed that the serum creatinine kinase (CK) level increased to 803 IU/L (normal range 43–239 IU/L), LH (5.9 IU/L; normal range 1.8–5.2 IU/L) and follicle stimulating hormone (20.5 IU/L; normal range 2.8–9.2 IU/L) levels were elevated. After his informed consent was obtained, high molecular weight genomic DNA was extracted from peripheral leucocytes of the patient according to standard protocols. Genetic analysis of the AR gene was performed and the expansion of a CAG repeat (45 repeats) in exon 1 of the AR gene was identified, leading to a diagnosis of SBMA.

At age 67, he developed weakness in the legs, and noticed difficulty in climbing up stairs or standing up from a chair. Serum CK levels gradually increased to 1717 IU/L at age 70. In January 1998, when he was 71 years old, he was diagnosed as having prostate cancer, and was intramuscularly injected with 3.75 mg of leuprorelin every 28 days, which enhanced the growth of prostate cancer. These drugs eventually inhibited LH production, which in turn inhibits production of testosterone and dihydrotestosterone (DHT), which enhances the growth of prostate cancer cells. One month after the start of treatment, he noticed that his gait disturbance was rapidly exacerbated; however, the gait disturbance returned to the level before the start of treatment by April 1998. After the episode of transient exacerbation, his muscle weakness and atrophy exhibited no apparent deterioration to date. On the contrary, an improved muscle strength was recorded in the neck flexor, biceps brachii, and quadriceps femorius muscles. Furthermore, serum CK levels gradually decreased from 1717 IU/L to 834 IU/L after the leuprorelin treatment. After the treatment, the serum CK levels were 100 IU/L (0.1 IU/L). Testosterone (<0.1 IU/L; normal range 1.2–8.0 IU/L) were decreased by the leuprorelin injections.

Discussion
The experience of a 5 year follow up of this patient treated with leuprorelin is highly indicative of the following. Firstly, leuprorelin treatment induced a transient deterioration of the motor function in humans, as demonstrated in a transgenic mouse model of SBMA. Secondly, after the initial transient deterioration, long term stabilisation of the motor function was obtained. Finally, leuprorelin treatment was effective even when the treatment was started in the advanced stage of the disease, although the patient’s muscle weakness and atrophy have not completely disappeared. These findings provide grounds for the proposal made by Katsumo et al that leuprorelin is a promising candidate for the treatment of SBMA.

At least nine neurodegenerative diseases are known to be caused by expanded CAG repeats. SBMA is unique among these diseases because the disease protein, AR, has a specific ligand, testosterone. It has been demonstrated that the nuclear translocation of ARs is solely dependent on testosterone. Recently, a transgenic mouse model carrying a polyglutamine stretch of the androgen receptor (AR) gene encoding a polyglutamine stretch.1

Recently, Katsumo et al reported that leuprorelin, a luteinising hormone releasing hormone (LHRH) agonist that reduces the level of testosterone release from the testis, rescued motor dysfunction and nuclear accumulation of mutant ARs in a male transgenic mouse model of SBMA. This result indicates that ligand dependent nuclear translocation of mutant ARs containing expanded polyglutamine is the main source of the pathogenesis of SBMA, and that leuprorelin suppresses this translocation. We read this report with great interest, because we followed up a patient with SBMA, who has been administered leuprorelin for 5 years to treat his coexisting prostate cancer.
desensitised and stops releasing LH. When that occurs, the testes stop releasing testosterone. During the period of the initial stimulation, more LH is released, consequently there is a surge in the secretion of testosterone and DHT from the testes (so called “androgen surge”). It is reasonable to assume that administration of leuprorelin causes a transient exaggeration of the above finding in patients with SBMA due to the transient androgen surge.

In conclusion, we report the beneficial effect of leuprorelin on SBMA. Our current experience warrants further investigations to determine whether leuprorelin may be of benefit for the treatment of SBMA in humans.

T Shimohata, T Kimura, M Nishizawa
Department of Neurology, Brain Research Institute, Niigata University, Niigata, Japan

O Onodera
Department of Molecular Neuroscience, Resource Branch for Brain Disease Research, Center for Bioresource-based Researches, Brain Research Institute, Niigata University, Niigata, Japan

S Tsuji
Department of Neurology, University of Tokyo, Tokyo, Japan

Correspondence to: T Shimohata, Department of Neurology, Brain Research Institute, Niigata University, 1-757 Asahimachi, Niigata 951-8585, Japan; t-shimo@bri.niigata-u.ac.jp
doi: 10.1136/jnnp.2003.030064
Received 9 October 2003
In revised form 9 December 2003
Accepted 14 December 2003
Competing interests: none

References

Cessation of migraine following central retinal vein occlusion

Case of retinal vein occlusion with migraine

A case of retinal vein occlusion with migraine has been described since 1882.1 An interesting case of central retinal vein occlusion (CRVO) which coincided with complete cessation of longstanding, severe migraines is reported.

Case report

A 44 year old lady developed classic migraine at the age of 11 years. Her symptoms comprised a visual aura of flashing lights followed by severe headache (not localised to one side), photophobia, and nausea, which generally lasted for two days. There were no identifiable triggers. Her management consisted of sumatriptan, which she took on experiencing the visual aura. This considerably reduced the severity of her headache and usually limited the duration of her symptoms to one day. The migraines occurred frequently and randomly with the longest migraine-free period being one month.

One morning, she awoke with visual loss in her right eye. She assumed this to be the visual aura of a migraine (although it was atypical since there was no photopsia or subsequent headache), and took her normal dose of sumatriptan. The visual loss occurred before taking the medication. The visual defect fragmented into black patches followed by gradual visual improvement over the next few days. She then consulted her general practitioner who referred her to the eye department.

At presentation to the eye department one week after the initial visual loss, her visual acuity was 6/9 right and 6/4 left. There was a right relative afferent pupillary defect. Funduscopy revealed retinal haemorrhages in all four quadrants with a swollen optic disc. A diagnosis of non-isaemic CRVO was made. She was advised to take aspirin 75 mg daily.

On follow up, her visual acuity continued to improve with resolution of the retinal haemorrhages and the disc oedema. The following investigations were normal: full blood count, erythrocyte sedimentation rate, electrolytes, fasting glucose, fasting cholesterol, and plasma protein electrophoresis. General medical examination was normal. She is a non-smoker with no family history of cardiovascular disease. At the 18 month follow up her visual acuity was 6/5 right and 6/4 left. There was no relative afferent pupillary defect. The fundal appearance returned to normal.

Follow up to date is two years and she has not experienced a single migraine since developing the CRVO. There have been no other factors to account for the cessation of her migraines during this period.

Discussion

There have been numerous reports of retinal vaso-occlusion and migraine in the context of “complicated migraine”1. We have presented an interesting patient who instead experienced complete cessation of migraine in association with the development of a CRVO. In the natural history of migraine there is a gradual reduction in severity and frequency of attacks with age.2 The abrupt cessation of migraine following development of a CRVO suggests a causal relationship. She had no risk factors for a retinal vascular event.

It has been proposed that prophylactic use of platelet antagonists, such as aspirin, may reduce the occurrence of migraine. Serotonin is released locally in cerebral tissue shortly before the onset of a migraine attack. Since platelets contain all of the plasma serotonin platelet function has been implicated as a factor in migraine.1 The role of serotonin in migraine is complex. To the best of our knowledge there is no report of platelet antagonists causing complete cessation of migraine. It seems unlikely that aspirin was solely responsible for the cessation of migraine in our patient, however this remains a possibility.

The pathophysiology of migraine is complex but involves neuronal events linked to alterations in the calibre of intracerebral blood vessels. During a migraine aura cerebral blood flow decreases. The subsequent hypertrophy leads to headache by activation of fibres originating in the trigeminal ganglion. These trigeminovascular afferents reside primarily within the ophthalmic division of the trigeminal nerve.3 The retinal vasculature is very similar to the cerebral vasculature both in structure and response to vasoactive substances.4 This probably accounts for cases of “complicated migraine” leading to retinal vein occlusion.

We postulate that an initial neuronal event occurred in our patient that resulted in a functional alteration in her trigeminovascular system leading to the complete cessation of migraine. This neuronal event also produced a temporary decrease in central retinal artery perfusion and the subsequent development of a CRVO.5 This case therefore demonstrates the potential for intracerebral events to influence the retinal vasculature.

S P Meredith, D K Newman
Department of Ophthalmology, Addenbrooke’s Hospital, Cambridge, UK
Correspondence to: S P Meredith, spmeredith@doctors.org.uk
doi: 10.1136/jnnp.2003.024869
Competing interests: none declared

References